EUP OHAMR Joint transnational call 2027



The Project has received funding from the European Union's Horizon Europe research and Innovation programme under Grant Agreement No101217154

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This second EUP OHAMR joint transnational call for projects will explore the emergence and spread of AMR. The call "JTC 2027 - Understand, Model and Stop AMR Transmission", involves 37 funding organisations from 27 different countries with an estimated total call budget of over 30 million Euro. The call is co-funded by the European Union.

1- Call summary

Call title	JTC 2027 - Understand, Model and Stop AMR Transmission
EUP OHAMR focus area	Prevent the emergence and spread of AMR
EUP OHAMR thematic areas	Transmission and evolution; Surveillance; Interventions for prevention and mitigation
Type of Call	Call for research and innovation projects
Initial grant period	Up to 3 years
Call opens	18 November 2026
Call closes	1st February 2027, 13h CET (pre-proposals)
Estimated Call Budget	30.9 Mio €
Link to the submission platform	https://ptoutline.eu/app/OHAMR_JTC-2027
Link to the partner search tool	https://www2.ncn.gov.pl/partners/ohamr-2027/
Link to the pre-proposal template	https://ohamr.eu/app/uploads/2026/11/EUP-OHAMR-JTC2027_Pre-proposal-form.docx
Contact Joint Call Secretariat	EUP-OHAMR@DLR.de

Please note:

- Only transnational research and innovation projects will be supported.

- An online partner search tool is available.

- Applications must be submitted to the EUP OHAMR submission platform and follow the EUP OHAMR pre-proposal/full proposal template.

- Applications to EUP OHAMR joint transnational calls can require the submission of additional information on national funding platforms.

- All applicants must have fulfilled both joint and national requirements for an application to be eligible.

- Applicants are strongly advised to confirm their funding eligibility with their national/regional funding organisation before submission.

- For updates to the call text, please see the History of change section at the bottom of the Call text.

2- Introduction

Antimicrobial resistance (AMR) is a global health challenge that affects human and animal health, food security and the environment. The European Partnership on One Health AMR (EUP OHAMR) is one of the key partnerships that has been identified by the European Commission within the framework of the Horizon Europe funding programme to support Research and Innovation (R&I) to respond to the challenges of AMR.

The EUP OHAMR was launched in June 2025 for a period of ten (10) years. The programme brings together 55 organisations (research and innovation funding organisations, key AMR actors, and stakeholders) from 30 countries in EU and beyond to address AMR challenges across sectors and One Health domains. To ensure its mission, the EUP OHAMR will deploy a wide range of activities organised under four programmes: (i) the R&I Funding programme, to provide a financial support to AMR R&I; (ii) the Capacity Strengthening Programme, to support training, networking and collaboration across disciplines, One Health sectors, professional domains (academia and industry including small and medium enterprises, SMEs), countries and career stages; (iii) the Data Exploitation Programme, to facilitate and promote access, sharing and (re) use of data and research infrastructures; and (iv) the Impact Programme for knowledge mobilization, to facilitate uptake of research results into products, practice and policy for maximum societal impact.

The R&I Funding programme, which includes the launch of annual Joint Transnational Calls (JTCs), will be articulated around the three following focus areas: (i) prevent the emergence and spread of AMR, (ii) strengthen appropriate use of antimicrobials and infection prevention and control and (iii) provide innovative and cost-effective treatment options. Each of the three focus areas covers various aspects of AMR related to therapeutics, diagnostics, surveillance, transmission and evolution, as well as interventions.

For the second EUP OHAMR JTC 37 funding organisations from 27 countries will join their forces.

Resources:

EUP OHAMR Strategic Research and Innovation Agenda: https://ohamr.eu/about/sria/

3- Aim of the Call

This second call will address focus area 1, namely, “Prevent the emergence and spread of AMR”. Current measures to decrease and control the emergence and spread of AMR and recent technical advances are still insufficient to reduce the burden of AMR. More knowledge on transmission dynamics is needed. In the coming years, novel and/or improved measures need to be developed to control AMR, based on a better understanding of the mechanisms driving emergence, evolution, selection, persistence, and transmission of AMR, in and between different One Health (OH) sectors.

4- Call topics

Research and innovation proposals submitted under the EUP OHAMR 2027 Joint Transnational Call must address one of the following topics:

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Topic 1

Model AMR transmission dynamics to predict the risk of AMR emergence and spread to support decision making of public authorities

Proposals addressing this topic should develop AMR transmission models at the global scale to predict the emergence and spread of antibiotic and/or antifungal resistance (within One-Health settings or between different One-Health settings) to support the public authorities in taking evidence-based decisions to curb AMR transmission and spread. The models proposed should consider the contribution of the anthropogenic activities (such as agriculture, travels, trade, large gatherings, migration, war, etc) and/or societal properties (e.g. behavioural aspects of the population, population distribution) and climate change (heatwave, floods, etc) in order to identify points of intervention. Documented and coordinated collection, use and/or re-use of existing data and quality assessments are expected.

Topic 2

Understand and characterize the role of host commensal micro-organisms (including fungi) in AMR dynamics and develop solutions aiming at selective decolonizing of resistant organisms in the host (human and animals) to prevent transmission

Commensal micro-organisms in humans and animals can act as reservoirs of resistance genes, which can be transferred to pathogens. The purpose of this topic is to better understand the role of commensal organisms in the spread of AMR, and propose solutions to prevent AMR transmission.

Topic 3

Develop or improve tools and methods to reduce resistant bacteria, fungi, and resistance genes in environmental hotspots, and investigate the impact of these tools and methods on AMR transmission

Hotspots are where resistant bacteria, fungi, and resistant genes are present in large quantity for example: wastewater treatment plants, hospital pipelines, effluent of antibiotic and antifungal manufacturing production sites and manure. Proposals addressing this topic should identify, develop, improve and/or test innovative solutions facilitating the decontamination of the environmental hotspots (including, but not limited to the sites mentioned above) and demonstrate the impact on AMR transmission and on human health (excluding infection prevention and control measures). Proposals addressing this topic must consider at least two One Health settings (Human, Environment).

Requirements for all call topics

- Proposals should consider how the expected results will impact the risk of resistance in other One Health settings, and how the results could be extended to additional One Health settings.
- Proposals should explain their feasibility by outlining realistic objectives, practical methodologies, and achievable timelines (e.g.: workplan, risk identification, allocation of resources, capacity of the consortium to conduct the work, business plan if the proposal envisages commercial component, etc).
- Proposals must clearly demonstrate the potential health, social and/or economic impact(s) of the expected results. In particular, the proposals should explain how the uptake of the expected project results by the society/end-users/next actors will be facilitated (Pathway to impact).

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- Proposals should clearly demonstrate the unique contribution of each partner (i.e. expertise, resources). In addition, the proposals should demonstrate the added value of transnational collaboration: sharing of expertise and resources (models, databases...), harmonization of data, access to innovative technologies, etc.
- Participation of the private sector (start-up, SMEs, industry) in the consortium is encouraged for all call topics, if appropriate (and in accordance to national/regional funding eligibility rules).
- Proposals submitted to this call should not present overlap with proposals by the same consortium submitted to/granted by other Horizon Europe or EU4Health calls or with projects previously funded by JPIAMR. In their full proposal, applicants will be asked to identify potential related projects and describe how their proposal does not overlap with, but possibly complements, a related EU project (including projects funded by other European Partnerships, projects submitted to external calls, as well as research activities funded internally).
- Clinical trials cannot be funded in this call.

5- Participating countries and funding organisations

5.1- Financial commitment

The following participating funding organisations have agreed to fund this call for transnational research projects:

Country	Funding Organisation (Acronym)	National/ Regional Financial Commitment	Expected number of funded proposals
Australia	National Health and Medical Research Council (NHMRC)	4 000 000 AUD Ca. 2 456 000 €	8
Austria	Fonds zur Förderung der Wissenschaftlichen Forschung (FWF)	1 400 000 €	3-4
Belgium	Fonds de la Recherche Scientifique (FNRS)	300 000 €	1
Belgium	Fonds Wetenschappelijk Onderzoek-Vlaanderen (FWO)	700 000 €	2-3
Belgium	Service Public de Wallonie (SPW)	1 000 000 €	2
Canada	Canadian Institutes of Health Research (CIHR)	600 000 CAD ca. 372 000 €	1
Czech Republic	Ministerstvo Zdravotnictvi Ceske Republiky (MZCR)/ Agentura pro zdravotnicky vyzkum (AZVCR)	500 000 €	2
Denmark	Innovationsfonden, Innovation Fund Denmark (IFD)	1 600 000 €	3-5
Estonia	Regionaal- ja Põllumajandusministeerium (REM)	100 000 €	1
France	Agence Nationale de la Recherche (ANR)	1 800 000 €	6-8
Germany	Bundesministerium für Forschung, Technologie und Raumfahrt (BMFTR)/ Deutsches Zentrum für Luft- und Raumfahrt Projektträger (DLR, without own financial contribution)	3 700 000 €	10
Hungary	Nemzeti Kutatási, Fejlesztési Innovációs Hivatal (NKFIH)	300 000 €	1-2
Ireland	Department of Agriculture, Food and the Marine (DAFM)	200 000 €	1
Ireland	Taighde Éireann - Research Ireland (TÉ-RI)	600 000 €	1-2

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Ireland	The Health Research Board (HRB)	530 000€	1-2
Israel	Ministry Of Health - Chief Scientist Office (CSO-MOH)	360 000 €	2
Italy	Fondazione Regionale per la Ricerca Biomedica (FRRB)	1 500 000 €	3-4
Italy	Ministero della Salute (MOH-IT)	1 000 000 €	2
Latvia	Latvijas Zinatnes Padome (LZP)	600 000 €	2
Lithuania	Lietuvos Mokslo Taryba (LMT)	300 000 €	1-2
Malta	Xjenza Malta (XM)	500 000 €	1-2
Moldova	National Agency for Research and Development (NARD)	100 000 €	1-2
Netherlands	Zorgonderzoek Nederland (ZonMw) & Nederlandse Organisatie voor Wetenschappelijk Onderzoek (NWO)	2 500 000 €	6
Norway	The Research Council of Norway (RCN)	1 340 000 €	2 -3
Poland	Narodowe Centrum Nauki (NCN)	1 000 000 €	3-4
Portugal	Fundacao para a Ciencia e a Tecnologia (FCT)	400 000 €	2-4
Slovakia	Centrum Vedecko Technickych Informacii Slovenskej Republiky (CVTI SR)	300 000 €	1-2
Spain	Instituto Aragones de Ciencias de la Salud (IACS)	150 000 €	1-2
Spain	Agencia Estatal de Investigación (AEI)	800 000 €	4-5
Spain	Instituto de Salud Carlos III (ISCIII)	500 000 €	2-3
Sweden	Vetenskapsrådet, Swedish Research Council (SRC)	15 000 000 SEK ca. 1 385 000 €	3-5
Switzerland	Swiss National Science Foundation (SNSF)	1 364 285 CHF ca. 1 460 338 €	4-5
Türkiye	Turkiye Bilimsel ve Teknolojik Arastirma Kurumu (TUBITAK)	500 000 €	3-4
United-Kingdom	Medical Research Council - United Kingdom Research and Innovation (MRC - UKRI)	1 000 000 £ ca. 1 150 000 €	3-4
United Kingdom	Department of Health and Social Care (DHSC)	500 000 £ ca. 570 000 €	1-2

The funding will be provided by the national/regional research funding organisations participating in the call. Each of them will fund their national/regional applicants, according to their own eligibility rules. To be eligible for funding, applicants must comply with the regulations and scientific remits of their own funding organisations as detailed in the National and Regional Requirements or specific regulations of their corresponding funding partner organisation. Applicants are strongly advised to confirm their funding eligibility with their national/regional funding organisation before submission.

Please note:

- The exchange rates for funders outside the Euro-zone are listed in the National and Regional Requirements.

- For Canada, topic 3 only relevant when environmental work is explicitly connected to human health outcomes.

- For Czech Republic, the Call is administered by the Czech Health Research Council (AZV CR) at the national level but the Ministry of Health of the Czech Republic (MZ CR) is the funding organisation.

5.2- Eligible Call topics and One Health settings

Country	Funding Organisation (Acronym)	Eligible Topic(s)	Eligible One Health settings
Australia	National Health and Medical Research Council (NHMRC)	1, 2, 3	Human Health
Austria	Fonds zur Förderung der Wissenschaftlichen Forschung (FWF)	1, 2	Human Health, Animal Health, Environment, Plant Health
Belgium	Fonds de la Recherche Scientifique (FNRS)	1, 2, 3	Human Health, Animal Health, Environment, Plant Health
Belgium	Fonds Wetenschappelijk Onderzoek-Vlaanderen (FWO)	1, 2, 3	Human Health, Animal Health, Environment, Plant Health
Belgium	Service Public de Wallonie (SPW)	2 and 3	Human Health, Animal Health, Environment, Plant Health
Canada	Canadian Institutes of Health Research (CIHR)	1, 2, 3	Human Health
Czech Republic	Ministerstvo Zdravotnictvi Ceske Republiky (MZCR)/ Agentura pro zdravotnicky vyzkum (AZVCR)	1, 2, 3	Human Health
Denmark	Innovationsfonden, Innovation Fund Denmark (IFD)	1, 2, 3	Human Health, Animal Health, Environment, Plant Health
Estonia	Regionaal- ja Põllumajandusministeerium (REM)	1	Human Health, Animal Health, Environment, Plant Health
France	Agence Nationale de la Recherche (ANR)	1, 2, 3	Human Health, Animal Health, Environment, Plant Health
Germany	Bundesministerium für Forschung, Technologie und Raumfahrt (BMFTR)/ Deutsches Zentrum für Luft- und Raumfahrt Projektträger (DLR)	1, 2, 3	Human Health, Animal Health, Environment, Plant Health
Hungary	Nemzeti Kutatási, Fejlesztési Innovációs Hivatal (NKFIH)	1, 2, 3	Human Health, Animal Health, Environment, Plant Health
Ireland	Department of Agriculture, Food and the Marine (DAFM)	2	Animal Health, Environment, Plant Health
Ireland	Taighde Éireann - Research Ireland (TÉ-RI)	1, 2, 3	Human Health, Animal Health, Environment, Plant Health

Ireland	The Health Research Board (HRB)	1, 3	All One Health settings (Human, Animal, Environment, Plant) related to Human Health.
Israel	Ministry Of Health - Chief Scientist Office (CSO-MOH)	1, 2, 3	Human health Animal health Environmental health (Only with relation to human health).
Italy	Fondazione Regionale per la Ricerca Biomedica (FRRB)	1, 2, 3	Human Health
Italy	Ministero della Salute (MOH-IT)	1, 2, 3	Human Health, Animal Health, Environment, Plant Health
Latvia	Latvijas Zinatnes Padome (LZP)	1, 2, 3	Human Health, Animal Health, Environment, Plant Health
Lithuania	Lietuvos Mokslo Taryba (LMT)	1, 2, 3	Human Health, Animal Health, Environment, Plant Health
Malta	Xjenza Malta (XM)	1, 2, 3	Human Health, Animal Health, Environment, Plant Health
Moldova	National Agency for Research and Development (NARD)	1, 2, 3	Human Health, Animal Health, Environment, Plant Health
Netherlands	Zorgonderzoek Nederland (ZonMw) & Nederlandse Organisatie voor Wetenschappelijk Onderzoek (NWO)	1, 3	All One Health settings (Human Health, Animal Health, Plants, Environment)
Norway	The Research Council of Norway (RCN)	1, 2, 3	All one health settings (human, animal, plant, environment, social sciences) related to human health.
Poland	Narodowe Centrum Nauki (NCN)	1, 2, 3	Human Health, Animal Health, Environment, Plant Health
Portugal	Fundacao para a Ciencia e a Tecnologia (FCT)	1, 2, 3	Human Health, Animal Health, Environment, Plant Health
Slovakia	Centrum Vedecko Technickych Informacii Slovenskej Republiky (CVTI SR)	1, 2, 3	Human Health, Animal Health, Environment, Plant Health
Spain	Instituto Aragones de Ciencias de la Salud (IACS)	1, 2, 3	Human Health, Animal Health, Environment
Spain	Agencia Estatal de Investigación (AEI)	1, 2, 3	Human Health, Animal Health,

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			Environment, Plant Health
Spain	Instituto de Salud Carlos III (ISCIII)	1, 2, 3	Human Health (eligible for funding). Animal Health (ISCIII can only fund research groups working on AH related to zoonotic diseases). Environment (eligible only if the research has a direct connection to human health)
Sweden	Vetenskapsrådet, Swedish Research Council (SRC)	1, 2, 3	Human Health, Animal Health, Environment, Plant Health
Switzerland	Swiss National Science Foundation (SNSF)	1, 2, 3	Human Health, Animal Health, Environment, Plant Health
Türkiye	Turkiye Bilimsel ve Teknolojik Arastirma Kurumu (TUBITAK)	1, 2, 3	Human Health Animal Health Environment, Plant Health
United-Kingdom	Medical Research Council - United Kingdom Research and Innovation (MRC - UKRI)	1	Human Health
United-Kingdom	Department of Health and Social Care (DHSC)	1	Human Health

6- General Conditions for application

- The call will only support transnational research and innovation projects. You can find more information on the rules for composing your consortium in the section *"Composition of the transnational consortium"*.
- The call will follow a two-step evaluation process (submission of a pre-proposal; successful consortia will be invited to submit a full proposal).
- Pre-proposals and full proposals must be written in English, and must follow the format and the guidelines provided in the pre-/full proposal templates. Pre-proposals or full proposals that do not follow the template guidelines (i.e. number of pages of the different sections, number of CVs and letters of intent) will be rejected without further review. The link to download the pre-proposal template can be found below. The full proposal template will be sent to the project coordinators invited to the next evaluation stage.
- Pre-proposals and full proposals must be submitted by the project coordinator on the EUP OHAMR on-line submission platform. No other means of submission (i.e. post or e-mail) will be accepted. Please note that some funding organisations might request an additional mandatory submission on their own national/regional platform (see national rules and requirements). Pre-/full proposals submitted on a national/regional platform but not on the EUP OHAMR submission platform will be rejected without further review.

- In absence of other information, the call will follow the timeline indicated in the timetable presented in the next section. Applicants must respect the submission deadlines indicated. Submission of pre-proposals or full proposals after the submission timeline will not be accepted. Applicants must consider enough time to complete the information on-line. In particular, please consider that collecting some information, such as the Participant Identification Code (PIC) of your research organisation, may require external procedures. Applicants should also ensure that they will have a proper internet connection at the time of submission.
- All questions related to the general eligibility rules and general evaluation process should be addressed to the EUP OHAMR Joint Call Secretariat (JCS 2027) (EUP-OHAMR@DLR.de).
- All questions related to the national/regional eligibility rules and national/regional eligibility costs should be addressed to the national/regional funding organisations.
- The only official communication line between the project consortia and the JCS is through the project coordinator. Throughout the application procedure the JCS will only contact the project coordinators by email, who must forward all information to other partners of their consortia. This includes evaluation results.

Resources:
- Pre-proposal template: https://ohamr.eu/app/uploads/2027/01/EUP-OHAMR-JTC-2027-Pre-proposal-template_260114.docx
- EUP OHAMR on-line submission platform: https://ptoutline.eu/app/OHAMR_JTC-2027
- Participant Identification Code (PIC) of your research organisation: https://ec.europa.eu/info/funding-tenders/opportunities/portal/screen/how-to-participate/participant-register

7- Call Timeline

October/November 2026	Call Pre-announcement
Wednesday 18th November 2026	Publication of the Call
Wednesday 25th November 2026	Webinar Info-day
Monday 1st February 2027	**Submission deadline for pre-proposals**
Wednesday May 5th 2027	Communication, by the JCS, of the results of the pre-proposal evaluation (invitation for full proposal; by e-mail, no publication on the EUP OHAMR website nor national websites)
Wednesday 23rd June 2027	**Submission deadline for full proposals**
Thursday 4th November 2027	Final funding recommendation announced to applicants (e-mail sent by the JCS)

November 2027	Publication of the results on the EUP OHAMR Website during the AMR Awareness Week
December 2027- April 2028	Start of the projects

8- Composition of the transnational consortium

To be eligible, the submitted pre-/full proposals must respect the following eligibility rules regarding the composition of their consortium. Consortia not respecting the following eligibility rules will be rejected without further review.

8.1- Composition

- The consortium must include a **minimum of three (3) eligible partners asking for funding from three (3) different eligible countries** (including **at least two amongst EU Member States or Associated Countries**).
- The consortium can include a **maximum of six (6)** project partners (**including non-funded partners**). The maximum number of partners can be increased **to seven (7)** if the consortium includes:
    - at least one partner from an under-represented country or
    - at least one partner where the Principal Investigator meets the definition of an Early-Career Researcher or
    - a start-up, SME, or an Industry.
- For the purpose of this call, the under-represented countries are
    - Czech Republic, Hungary, Latvia, Lithuania, Malta, Moldova, Poland, and Slovakia.
- A consortium cannot include more than two (2) partners requesting funding from the same funding organisation.
- A consortium cannot include, as funded partners as well as non-funded partners, any legal entities established in Russia, Belarus, or in non-government-controlled territories of Ukraine, or any legal entities established outside Russia but whose proprietary rights are directly or indirectly owned for more than 50% by a legal person, entity or body established in Russia. This list is not extensive and may evolve. Please consult the general annexes of the 2023-2025 Horizon Europe funding programme for more information on other restrictive measures.
- A principal investigator can apply for max. two proposals, one as partner and one as coordinator (and in accordance to national/regional funding eligibility rules).
- At both the pre- and full proposal stages, all partners, including non-funded partners, must submit a signed letter of intent along with their pre-/full proposal (the templates to be used for the letters of intent are included in the pre-/full proposal templates). In the absence of these letters, the proposal will be declared ineligible. At the pre-proposal stage, the letter of intent must be signed by the project investigators representing the different partner institutions. At the full proposal stage, the letter of intent should also be signed by a person having the legal mandate to represent the partner institutions.

Please note:
- For the purpose of this call, an Early-Career Researcher (ECR) is a PhD holder, up to 8 years after the year of PhD award, holding a position at a recognized institution. The 8-year period may be extended to allow for career breaks including documented: parental leave, positions of trust in trade union organizations and student organizations, mandatory military or civil service, illness (own illness or care for close family members), medical internships or medical fellowship (applies to clinically active professionals). The last two categories may involve periods of up to 24 months each. Please note that the EUP OHAMR definition of an ECR may differ from the national/regional definition. Please contact the JCS and your national/regional funding organisation to confirm your eligibility.
- For UK partners, the limit of two applies to both UK funding organisations, meaning **that the consortium cannot include more than 2 UK partners funded either by MRC-UKRI or DHSC**.

Resources:
- The general annexes of the 2023-2025 Horizon Europe funding programme: https://ec.europa.eu/info/funding-tenders/opportunities/docs/2021-2027/horizon/wp-call/2023-2024/wp-13-general-annexes_horizon-2023-2024_en.pdf
- Check-list for applicants: https://ohamr.eu/calls/call-2026-new-treatments-to-tackle-amr/jtc-2026-checklist-for-applicants/https://ohamr.eu/calls/joint-transnational-call-2027/jtc2027-checklist-for-applicants/

8.2- Modification of the composition of your consortium - Widening Process

Composition of the consortium should **not** be modified between the pre- and the full proposal except for widening (see below), in case of force majeure/unforeseen event (e.g. change of professional affiliation, lab relocation, prolonged absence of the Principal Investigator, etc.), or upon recommendation of the Peer Review Panel or request of the Call Steering Committee. In any case, changes in the composition of the consortium must be approved by the funders of the respective proposal **ahead** of the submission of the full proposal. In particular, the project coordinator should send an e-mail to the JCS, and to the respective funding organisations, at least 15 calendar days before the full proposal submission deadline to confirm the eligibility of the proposed modifications.

- **Widening**: Consortia which are invited to the second stage of the call (full proposal) will be able to increase their initial size by adding one (1) new partner eligible for funding by an under-subscribed funding organisation. The total number of partners in the consortium must not exceed eight (8). An under-subscribed funding organisation is understood as a funding organisation that, based on the outcome of the pre-proposal stage and budget planning by the funders, is expected to underspend the full amount of funds it has committed to the call. The list of such organisations will be determined and documented by the funders. Project coordinators will be notified of this option in their invitation letter to submit a full proposal. The list of under-subscribed funding organisations will be included in the full proposal

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template. The widening process promotes inclusiveness, ensure global participation, relevance and impact of the submitted projects in and outside Europe, as well as maximisation of the use of the committed resources.

9- Applicants and National Rules

9.1- Partners

Joint research proposals may be submitted by partners belonging to the following categories (according to national/regional regulations, certain categories may not be eligible for funding by a specific funding organisation, or some categories may be mandatory, please see national rules and requirements):

- Academia (public and private universities, other higher education institutions or research institutes).
- Clinical/public health sector (hospitals/public health and/or other health care settings and health organisations, including primary health care).
- Enterprises (private companies of all sizes involved in research and innovation).
- Operational stakeholders – e.g. patient advocacy organisations, municipalities and local governments, local/national NGOs. Operational stakeholders should be able to provide useful knowledge to the consortium, ensure the consortium's research is useful and translatable to their (or other) organizational contexts, and/or influence decision making or create change within their organisations. Operational stakeholders should be engaged in the research process from conception of the study to dissemination.

Each partner institution will be represented by **one** Principal Investigator who will be a researcher scientifically responsible for the implementation of tasks assigned.

To avoid conflicts of interest, employees of the EUP OHAMR beneficiaries are not allowed to apply to this call, with the exception of the staff working in AGES, ICARS, ISCIII, IACS, TUBITAK and DLR. AGES and ICARS have been excluded from the call preparation. Firewall measures (such as organisation of the work in separate divisions) have been put in place internally in ISCIII, IACS, TUBITAK and DLR to guarantee an equal treatment of the applicants and applications. In particular, the persons working on the call implementation will ensure that no confidential information will be shared with their research community.

Resources: EUP OHAMR Beneficiaries: https://ohamr.eu/about/partners/

9.2- Non-funded partners

- Project partners not eligible for funding (e.g. from countries not participating in the call or not fundable according to national/regional regulations of the funding partner organisations) may be involved in projects if they bring their own funding (in-kind).
- The budget of non-funded partners must be included in the proposal (in-kind Budget) and shall not exceed 30% of the requested total transnational project budget.
- The number of non-funded project partners in a consortium must not exceed the number of funded partners.

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- A project partner not receiving funding cannot be the coordinator of a proposal.
- Non-funded partners have the same responsibilities as funded partners, i.e. they must accept all EUP OHAMR rules and guidelines as set out in this Call Text.

9.3- National Rules

Please consult the National Rules section for more information on the national/regional regulations.

10- Evaluation

The evaluation process will comply with the principles of transparency, equal treatment and non-discrimination.

10.1- Eligibility check and evaluation procedure

Formal check and evaluation of pre-proposals

The JCS will check all proposals to ensure that they meet the call's formal criteria, i.e. date of submission; number and category of participating countries; inclusion of all necessary information in English; appropriate limits on length; signature of the letters of intent, number of proposals per investigator. In parallel, the JCS will forward the proposals to the national/regional funding organisations, which will perform a control for compliance with national/regional regulations. Each proposal passing both eligibility controls will be evaluated independently by two to three reviewers for a first evaluation (see evaluation criteria below). Potential conflicts of interests of the evaluators will be taken into consideration during the allocation of the proposals. The reviewers will perform the evaluation of the pre-proposals and complete a written evaluation form with scores and comments for each evaluation criterion. The pre-proposal written evaluation will result in 3 ranking lists, one ranking list for each topic. The ranking lists are based on the average score received by each pre-proposal.

The pre-proposals that have been considered in scope by the reviewers and that obtained an average score >13 will be automatically invited to submit a full proposal. The pre-proposals that were not recommended for the next stage by at least two of the reviewers and that obtained an average score <10 with no strong score discrepancy (StDev<2) will be automatically rejected. The following pre-proposals will be discussed during a Peer Review Panel (PRP) meeting:

- the pre-proposals that obtained an average score ≥10 and ≤ 13;
- the pre-proposals that obtained an average score <10 with a strong score discrepancy (StDev>2) and that were recommended for the next stage by at least 2 out of the 3 reviewers;
- the pre-proposals that obtained a score > 13 but that were considered out of scope by at least one of the reviewers.

As a result of the PRP meeting, a consensus score will be identified for those pre-proposals. To avoid conflicts of interest during the PRP meeting, evaluators with a conflict related to a specific proposal (i.e. co-publication with one of the applicants during the last 5 years, current collaboration with one of the applicants, same research centre as one of the applicants, personal or professional links with one of the applicants that may compromises the evaluator impartiality, involvement in the preparation of the proposal) will not participate in the discussion of that proposal. The Call Steering Committee (CSC) will meet to decide which of those proposals will be invited to submit a full proposal based on the

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reviewers' scores and to ensure a reasonable balance of requested and available national/regional budgets. Pre-proposals which do not pass this evaluation will not be invited for the full proposal stage. The consortia will receive a consensus report without scores written by one of the experts.

Formal check and evaluation of full proposals

The JCS will check the full proposals to ensure that they meet the call's formal criteria and have not changed substantially from the respective pre-proposals (e.g. composition of the consortium, the objectives of the project or the requested budget). In parallel, the JCS will forward the proposals to the national/regional funding organisations, which will perform a check for compliance with national/regional regulations. Each full proposal passing both checks will be allocated to three reviewers taking the potential conflicts of interest into consideration. The reviewers will perform the evaluation of the full proposal and complete a written evaluation form with scores and comments for each criterion (see evaluation criteria below). During a second PRP meeting, the reviewers will discuss all proposals and produce 3 ranking lists of proposals recommended for funding (one ranking list for each call topic). To avoid conflicts of interest, evaluators with a conflict related to a specific proposal will not participate in the discussion of that proposal and will leave the meeting. The final summary review report prepared by the evaluators will be sent to the respective project coordinators.

Ethics and legal requirements

Please note that at the full proposal stage, applicants will be required to complete a self-assessment checklist for ethics and to provide details on safety, animal studies, genetically modified organisms and microorganisms, environmental hazards and waste handling, data management, statistical methods, ethics and legal issues. Applicants should anticipate this requirement and ensure that they have consulted with relevant experts to verify the feasibility of the project, and that the proposal can be completed within the defined budget and within the prescribed time window.

Full proposals recommended for funding by the PRP and selected for funding by the board of funders (CSC) will undergo an ethics review by an Ethics Panel. Ethics experts will remotely check the selected proposal for their compliance with ethical norms and regulations. A meeting will also be organised for a discussion between the various ethics experts. If necessary, the ethics experts may ask the consortium for clarifications. The ethics experts may highlight some vigilance points that need to be monitored during the implementation of the funded project. Only those proposals approved by both the scientific evaluation and ethics assessment (complying with all central Horizon Europe and regional/national ethical requirements) will be recommended for funding.

Decision

The board of funders (CSC) will take their funding decision, based on the ranking lists established by the PRP, the available funding and the ethics panel recommendations. The JCS will send **by e-mail** the funding recommendation to the project coordinator, who is then responsible to communicate this information to the respective project partners.

Redress Procedure

Applicants can appeal against the evaluation outcome if they suspect a breach in the implementation of the evaluation and selection procedures. This redress procedure only covers the procedural aspects of the evaluation.

A mere disagreement with peer reviewers or panel members' comments are not grounds for an appeal. The redress procedure will not call into question the scientific or technical judgement of appropriately qualified experts.

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The applicants shall submit their appeal against the evaluation outcome to the JCS via e-mail (EUP-OHAMR@DLR.de) up to 7 calendar days (168 hours) after the date of the notification of evaluation outcome sent by the JCS at the end of each step (evaluation of the pre- or full proposal).

For an appeal to be admissible the following conditions must be met:

- The appeal must be submitted by the project coordinator of the proposal to which the appeal relates

- Only one appeal per proposal can be submitted after each step

- The appeal must contain the following minimum information: the name of the call for proposals, the proposal acronym, the title of the proposal, a description of the alleged shortcomings of the evaluation procedure.

The appeal must demonstrate a procedural irregularity, factual or manifest errors in the evaluation process, misuse of powers, or a conflict of interest. Appeals that do not meet the above conditions, or do not deal with the evaluation of a specific proposal or express mere disagreement with the result or the reasoning of the evaluation will be judged as not suitable for redress.

Upon receipt of an appeal, an acknowledgement of receipt will be sent by the JCS as soon as the e-mail is read. The acknowledgement shall report the redress process and the anticipated date by which a decision on the appeal will be communicated to the appellant. All appeals received by the 7 calendar days deadline will be processed together by a designated redress committee and the decision will be communicated to the appellant within 14 calendar days from the deadline for submitting the appeals.

Questions related to the national/regional eligibility decisions will not be handled by the JCS and need to be addressed to the respective national/regional funding organisation.

10.2- Evaluation criteria

1. **Excellence**

- Clarity and pertinence of the objectives (pre-proposal and full proposal)

- Credibility of the proposed approach and methodology, in relation to the research objectives (pre-proposal and full proposal)

- Soundness and research base of the concept (pre-proposal and full proposal)

- Novelty, potential to advance the field, timeliness, and innovation (pre-proposal and full proposal)

- Scientific excellence of the consortium (pre-proposal and full proposal)

2. Impact

- Impact of the proposal to achieve the objectives of the call topic (pre-proposal and full proposal)

- Potential of the expected results for public health, and animal health, agriculture, or environmental benefit (including economic viability/sustainability where appropriate) (pre-proposal and full proposal)

- Relevance and consideration of the One Health concept (pre-proposal and full proposal)

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- Potential for fostering a longer-term international network of researchers. For example, bringing together specific know-how and/or innovative technologies, gathering a critical mass of patients or biological material, sharing of resources (models, databases, biobanks, etc.), and international comparisons (pre-proposal and full proposal)

- Potential reach of the project results, including dissemination and communication measures. Accessibility of the proposed innovative strategy (different geographical areas, different populations including low-resource or underserved populations) (full proposal only)

- Appropriateness of end-user and stakeholder participation/engagement, for example, policy makers, industry, patient organisation, health and veterinary care, farmers, etc. (full proposal only)

3. Quality and efficiency of the implementation

- Coherence and effectiveness of the work plan, including appropriateness of the allocation of tasks within the given timeframe (pre-proposal and full proposal)

- Adequate distribution of the tasks between the project partners considering the required expertise (pre-proposal and full proposal)

- Strength of the transnational collaboration (balanced geographical distribution of the tasks) (pre-proposal and full proposal)

- Integration of social, economic, equity and cultural dimensions into the proposed research (pre-proposal and full proposal)

- Quality of the proposed Open Science practices, data management, Intellectual Property management, and Freedom to Operate where appropriate (full proposal only)

- Appropriateness of the management and governance structures and procedures, including risk and innovation management (full proposal only)

- Potential exploitation (including strategy to identify and address potential barriers) and relevance of the outcomes of the findings beyond the current project (long term strategy) (full proposal only)

- Contingency plan, including risk assessment and mitigation (including of unforeseen circumstances) (full proposal only)

- Justification of the requested budget and cost-effectiveness of the project (appropriate distribution of resources in relation to project's activities, partner responsibilities and time frame) (full proposal only)

Please note:

Proposals not relevant to the call topics and objectives (out of the scope) will not be funded, independently of their scientific quality. The decision if a project is in/out of scope will be taken by the reviewers and the evaluation panel in the pre-proposal stage.

10.3- Scoring system

Evaluation scores will be awarded for the three main criteria (Excellence, Impact and Implementation), and not singularly for the different aspects listed therein, although these different aspects will be taken into consideration in scoring the main criteria.

The weight of each of the three main criteria (Excellence, Impact and Implementation) is equal.

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The scoring system:

0: Failure. The proposal fails to address the criterion in question or cannot be judged because of missing or incomplete information.

1: Poor. The proposal shows serious weaknesses in relation to the criterion in question.

2: Fair. The proposal generally addresses the criterion, but there are significant weaknesses that need corrections.

3: Good. The proposal addresses the criterion in question well, but few improvements are possible.

4: Very good. The proposal addresses the criterion very well, but minor improvements are possible.

5: Excellent. The proposal successfully addresses all aspects of the criterion in question; there are no suggestions for improvement.

In order for an application to be considered fundable, the threshold score for individual criteria is set at three (3) (of a maximum of five (5)). The overall threshold for the score for all three criteria together is set at ten (10). The maximum score that can be reached from all three criteria together is fifteen (15) points.

11- Responsibilities of the Grantees

Post-award responsibilities of the grantees are described in the section "Responsibilities of the grantees JTC 2027". Applicants should keep in mind these obligations while drafting their budget and planning their workload.

12- Other information

12.1- Financial modalities

Funding is initially granted for a maximum of three years in accordance with national/regional regulations and applicable legal provisions.

Project partners will be funded by their relevant national/regional funding organisation. Therefore, eligible costs, funding rules and the type of studies allowed will vary between the respective funding organisations. Thus, it is recommended that each project partner defines their own budget in accordance with the funding rules of their own country/region. Please note that some funding organisations may limit the number of national/regional applicants per proposal or may limit the maximum budget national/regional applicants can request within a consortium.

For information on the specific funding rules and eligibility criteria of the national/regional funding organisation:

- read the National Rules and Requirements carefully

- in addition, applicants are strongly advised to reach out to their relevant funding organisation before applying; please note that for some countries/regions it might be mandatory.

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Any budget change between the pre-proposal and the full proposal **must be approved** by the national/regional funding organisations before submission (provided that it complies with the national/regional rules, that the oversubscription of the funding organisation does not surpasses the threshold, that the newly requested budget still complies with the eligibility rules regarding the composition of the consortium and that the changes are duly justified and related to the scope and ambition of the proposal).

Please note If a partner is found to be non-eligible at any step of the process by one of the funding organisations, the entire proposal could be rejected without further review**.**

12.2- Social and gender equity, cultural sensitivity and economic viability

It is important that consortia and research proposals are founded upon principles of social and gender equity, cultural sensitivity and economic viability. Consortia are highly encouraged to apply these principles to the composition, leadership and management of research projects. Especially where Low- and Middle-Income countries are involved in the proposal, the impact to improving health and wellbeing should be considered.

Where relevant, research projects are expected to apply an intersectional and multi-dimensional approach by integrating sex, gender and other individual and population-level determinants of health (such as age, socio-economic status, ethnicity) into the project's design, implementation, monitoring, evaluation and knowledge translation activities.

Research projects are expected to consider individual- and population-level determinants of health when collecting and analysing data to design and/or implement interventions in ways that are accessible and affordable to target beneficiaries, to systematically capture and report on sex, gender, and other relevant factors in the project research outputs, and to meaningfully engage the participation of targeted marginalised groups in the research activities.

12.3- General Data Protection Regulation (GDPR)

By submitting an application, the applicants acknowledge that the personal data contained in the proposal will be processed in accordance with Regulation (UE) 2016/679 General Data Protection Regulation (GDPR) based on article 6.1 (e) and (c) () and for the purposes described below.

Personal data is processed exclusively for:

- processing and evaluating the pre- and full proposal where processing shall be lawful only if and to the extent that processing is necessary for the performance of a task carried out in the public interest or in the exercise of official authority vested in the controller;
- administering any subsequent funding award;
- managing the relationship between the applicant and the Funding Partner Organisations;
- analysing and evaluating the call;
- providing aggregated data to national and European surveys and analyses on the funded projects;
- complying with audits that may be initiated by the Funding Partner Organisations and the European Commission (or its agencies).

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Personal data may be shared with Funding Partner Organisations and authorised third parties, including external scientific evaluators, strictly for the purposes described above. Some of these recipients may be located outside the European Economic Area (EEA).

Where personal data is transferred outside the EEA, such transfers shall be carried out in accordance with Chapter V of the GDPR, including adequacy decisions of the European Commission where applicable, or appropriate safeguards such as Standard Contractual Clauses.

The following Funding organisations outside the European Economic Area will use their national data protection rules:

- Australia (NHMRC)
- Canada (CIHR)
- Israel (CSO-MOH)
- Moldova (ANCD)
- Switzerland (SNSF)
- Turkey (TUBITAK)
- the United Kingdom (UKRI)

By the time of the call launch, the European Commission issued adequacy decisions for personal data protection laws in Israel, Switzerland and the United Kingdom.

Funding Partner Organizations and third parties may link the data that applicants provide in the application with national, bibliographic or external research funding data which is available through public subscription-based databases (e.g. Scopus, Web of Science, etc.) or other national/open datasets.

12.4- Confidentiality

The content of the pre-proposals and full proposals received under this call is deemed to be confidential, except for consortia including project partners applying for funding from the Swedish funding organisation (SRC). For those consortia, the applications (pre- and full proposals) may be made available upon request after the respective call deadlines.

Responding to a EUP OHAMR call for proposals, both as project coordinator or project partner, gives the EUP OHAMR, the European Commission and the Funding Partner Organisations the right to use and store the information submitted for analysis of the call success rate, national response rate, etc.

The proposals will be handled confidentially by the JCS and by the national/regional funding organisations. In selecting the international experts for the PRP, the JCS shall endeavour to avoid any possible conflicts of interest. Each expert will sign a declaration of confidentiality and confirm the absence of conflicts of interest. In case of a conflict of interest the reviewer will be withdrawn from evaluating the respective proposal. Conflicts of interest are managed and recorded throughout the evaluation process.

Accepting a EUP OHAMR grant award and associated grant contract from a national funding organisation gives the EUP OHAMR, the European Commission and Funding Partner Organisations the right to store, share, and analyse information on beneficiaries and consortia (rules may differ between different countries). Composition of the awarded consortia (Principal investigators, Institution) as well as the title, acronym and abstract of funded projects will be published and openly accessible. No data will be shared with third parties or commercial entities without the formal consent of the project

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coordinators, except for consortia including project partners applying for funding from the Swedish funder (SRC). For those consortia, the applications (pre- and full proposals) may be made available upon request after the respective call deadlines.

13- Contact Point

Joint Call Secretariat (JCS):

DLR Project Management Agency

PD Dr. Barbara Junker / Dr. Yen Ly-Sauerbrey / Dr. Patricia Ruiz Noppinger

Tel. +49 228 3821-1326

EUP-OHAMR@DLR.de

14- History of Change

Date	Description

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National Rules and Requirements

Important note to applicants: Applications to EUP OHAMR joint transnational calls can require the submission of additional information on national funding platforms. All applicants must have fulfilled **both joint and national requirements for an application to be eligible**.

This is only a summary. Refer to national websites and contact the respective contact person for full details.

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Australia - NHMRC

Full name of the funding organisation:	**National Health and Medical Research Council**
National/ Regional Financial Commitment:	**4 000 000 AUD**
Expected number of funded projects:	**8**
Contact details: (person (s), e-mail, telephone):	**NHMRC Research Help Centre + 61 2 6217 9000**
	international@nhmrc.gov.au

Maximum/ Minimum funding per grant Awarded to a project partner: Up to $600,000 CAD per grant Awarded to a project partner.

Exchange Rate: 1 Euro = 0.6103 AUD (Reserve Bank of Australia, 7 August 2026)

Eligible Call Topics: NHMRC will fund projects under all topics – 1, 2 and 3.

Eligible One Health Settings: Projects addressing human health will be eligible for funding. NHMRC will not fund research on animal health, plants, food, and/or environment that do not include human health research activities.

Eligible institutions: Institutions must be a NHMRC Administering Institution to be eligible to receive and administer NHMRC funding.

Eligible experimental approaches and disciplines: All types of studies or experimental approaches are admissable.

Eligibility criteria for the Principal Investigator:

Applications for NHMRC funding are subject to the general eligibility requirements as set out in the *2027 NHMRC-European Partnership on OneHealth Antimicrobial Resistance Grant Opportunity Guidelines* when available and within the NHMRC Funding Agreement. Applicants should read all relevant reference material.

Applicants from Australia intending to apply for funding via NHMRC should notify the national contact point by email to signal an intention to join a consortium, via international@nhmrc.gov.au.

Eligible costs: Applicants should Refer to NHMRC Direct Research Costs guidelines:www.nhmrc.gov.au/funding/manage-your-funding/funding-agreement

There is no minimum value for grants funded through this grant opportunity. The amount of funding for each EUP OHAMR grant will be based on assessment of the requested budget. Applications must clearly justify the requested duration and budget and how they will support the proposed outcomes of the research. Internal NHMRC budget review will consider this information and NHMRC may alter the duration and/or budget to ensure the research aims and objectives can be achieved while ensuring value with money. An altered budget does not alter the scope of the proposed research activity

Submission of the proposal (or other information) at the national level: Full proposals must be submitted through the NHMRC grant management system.

Further guidance: The total provisional funding allocation for this funding call is $4 million AUD. The NHMRC funding allocation will be distributed to successful grants until the $4 million funding allocation

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is exhausted. NHMRC's Research Committee reviews and advises indicative budget amounts to be awarded across individual funding schemes annually.

Australian applicants should refer to the NHMRC website and GrantConnect for specific information on this Grant Opportunity when it becomes available.

NHMRC funding is subject to governmental approval processes.

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Austria-FWF

Full name of the funding organisation:	**Fonds zur Förderung der Wissenschaftlichen Forschung**
National/ Regional Financial Commitment:	**1 400 000 €**
Expected number of funded projects:	**3-4**
Contact details: (person (s), e-mail, telephone):	**Inge Unfried** **inge.unfried@fwf.ac.at** **+43 676 83487 8210** **Lukas Koller** **Lukas.koller@fwf.ac.at** **+43 676 83487 8222**

Addendum: For the submission of applications, the FWF application guidelines effective as of November 16, 2026—including the cost guidelines—shall apply, and any conflicting provisions in the National Annex shall not apply.

The provisions set out in this national annex are binding for all FWF applicants and apply in addition to the requirements of the respective European Partnership.

FWF applications must meet all eligibility criteria at both the pre-proposal and full proposal stages.

Principal Investigators are requested to contact the relevant FWF contact person(s) well in time before the submission deadline if they have questions regarding the national project or its submission.

Maximum/ Minimum funding per grant awarded to a project partner:

The FWF generally does not have any minimum limit. The maximum limit per project submitted to the FWF is 450.000 €. Projects that exceed the maximum limit will not be eligible. This limit includes the annual valorization of personnel costs as well as 5% general project costs (calculated automatically in the cost breakdown in elane).

Project Duration:

Max 36 months - as specified by JTC2

Exchange Rate: NA

Eligible Call Topics: Topic 1 and 2

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Eligible One Health Settings: Human Health, Animal Health, Environment and Plant Health

Eligible institutions:

Austrian research institutions must be registered in the FWF's Research Institution Portal before the submission of the pre-proposal. The proposed research must be carried out in Austria under the auspices of the Austrian lead research institution. Applications are submitted by the research institution where the project will be carried out. Please refer to section 1.3.1-1.3.2 of the FWF Application Guidelines for more details.

Eligible experimental approaches and disciplines:

Funding may be requested for projects in basic research that are clearly defined, innovative, with plausibly described objectives and methods, and are limited in duration (see FWF Application Guidelines section 1.2).

Eligibility criteria for the Principal Investigator:

The principal investigator must be employed at the Austrian research institution applying for funding with an employment of at least 5 hours/week at the time the project is scheduled to begin and throughout the entire duration of the project (subject to an employment contract/Dienstvertrag).

The principal investigator must have appropriate scientific qualifications and eligibility as described in section 1.3 and 1.4 of the FWF Application Guidelines and sufficient time resources to carry out the proposed research. Applicants must comply with the fwf_project_number_limit.pdf.

Eligible costs:

The budget specified in the national submission must correspond to this consortium partner's budget submitted to the EP and be calculated on the basis of the applicable FWF personnel cost rates at the time of submission in both the pre-proposal and full proposal stages. Minor deviations resulting from updated FWF personnel cost rates between stages are permissible.

To ensure consistency, project costs should be calculated at elane (FWF) prior to inclusion in the consortium proposal. The FWF does not fund institutional or indirect overhead costs.

The "General Project Costs" automatically calculated in the elane cost breakdown on the basis of the requested FWF cost categories are an FWF-specific flat rate and may be reported under the "Overheads" heading in the EP budget forms for technical reasons, even though they do not constitute overheads in the FWF sense.

Project-specific costs are eligible for funding. These include personnel and non-personnel costs that are needed to carry out the project and that are not included in the infrastructure provided by the research institution (see section 2.3 of the FWF Application guidelines). The FWF does not finance the infrastructure or basic equipment of research institutions.

Submission of the proposal (or other information) at the national level:

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In addition to the application at the Call Secretariat, administrative data must be submitted online to the FWF at elane (user manual) - for both the pre-proposal and the full proposal stage. FWF submissions of the pre- and the full proposal before each specific deadline are both mandatory.

For the FWF submission of pre-proposals applicants must choose the program category "PIK – International Projects preproposal" deadline Tuesday, 2nd February 2027,14:00 CET (local time) and upload the following documents as individual annexes:

- CV of the FWF principal investigator(according to the FWF Application guidelines section 2.2.3 – Annex 3).
- Proof of publication output of the FWF principal investigator (according to FWF Application Guidelines section 2.2.4).
- A detailed justification of costs must be submitted as a mandatory attachment to the application at the pre- and full proposal stage. For further details, please refer to the FWF Application guidelines - section 6.1 Appendix A.
- The pre-proposal submission at the FWF must include a short statement (max. 1 page) that describes the basic research aspects of the project part to be financed by the FWF.

IMPORTANT: Please note that applications are only valid once they have been approved for submission by the research institution (PROFI) in the submission portal in due time before the specified deadline.

For the full proposal stage applicants must choose the program category "KIN– International Projects" (Deadline Thursday, 24th June 2027, 14:00 CET, local time). Further information will be communicated by the FWF to the applicants invited for full proposal submissions.

Further guidance:

It is recommended to contact the FWF contact person in case of questions regarding the national project.

If there is any uncertainty about general application requirements or about accounting for career interruptions, the FWF recommends contacting the FWF Office or the FWF Equal Opportunities and Diversity in Research Funding unit in good time before submitting the application to confirm that all requirements are met and that any career interruptions can be accounted for. In cases of doubt, the appropriate decision-making bodies of the FWF shall decide on applicants' eligibility.

Projects that do not comply with the FWF submission guidelines are considered as invalid.

FWF submissions during the pre- and the full proposal stages are both mandatory.

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Belgium-FNRS

Full name of the funding organisation:	**Fonds de la Recherche Scientifique**
National/ Regional Financial Commitment:	**300 000 €**
Expected number of funded projects:	**1**
Contact details: (person (s), e-mail, telephone):	**Dr. Maxime Bonsir +32 2 504 92 36** **Dr. Florence Quist +32 2 504 93 51** **international@frs-fnrs.be**

Maximum/ Minimum funding per grant Awarded to a project partner:

Maximum 300 000 € for a total period of three years. If the project involves the recruitment of a PhD student, the project duration of the F.R.S.-FNRS sub-project could be up to four years (see PINT-MULTI Regulations for details).

Exchange Rate: NA

Eligible Call Topics: Topic 1, 2, and 3

Eligible One Health Settings: Human Health, Animal Health, Environment and Plant Health

Eligible institutions:

All eligibility rules and criteria can be found in the PINT-MULTI Regulations.

Eligible experimental approaches and disciplines:

Please note that the F.R.S.-FNRS only funds Basic research (low Technology Readiness Level) carried out in a research institution from the "Fédération Wallonie-Bruxelles". The F.R.S.-FNRS will not fund industrial partners or any activity related to the private sector. Nevertheless, partners funded by the F.R.S.-FNRS can be in a consortium where there are also partners from the private sector.

Clinical research is eligible for funding by the F.R.S.-FNRS under the condition that the funded research is oriented towards the understanding of fundamental mechanisms, excluding clinical trials with purely therapeutic or industrial goals

Eligibility criteria for the Principal Investigator:

All eligibility rules and criteria can be found in the PINT-MULTI Regulations.

Eligible costs:

All eligibility rules and criteria can be found in the PINT-MULTI Regulations.

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Please do not include overhead costs in the budget, as these are covered through a separate agreement between the FNRS and the host research institution in the event that the project is funded.

Please check the Practical guide on costs for any other questions.

Submission of the proposal (or other information) at the national level:

Applicants to F.R.S.-FNRS funding must provide basic administrative data by submitting an administrative application on e-space **within 5 working days after the general deadline of OH-AMR to be eligible**. Please select the "PINT-MULTI" funding instrument when creating the administrative application. Proposals invited to the second stage will be able to complete the pre-proposal form and provide information for the full proposal upon validation by the F.R.S.-FNRS

Further guidance: PINT-MULTI regulations, e-space

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OH AMR

Belgium-FWO

Full name of the funding organisation:	**Fonds Wetenschappelijk Onderzoek-Vlaanderen**

National/ Regional Financial Commitment:	**700 000€**
Expected number of funded projects:	**2-3**

Contact details: (person (s), e-mail, telephone):	**Toon Monbaliu (FO)**	europe@fwo.be
	+32 (0)2 550 15 70	
	Kristien Peeters (SBO)	europe@fwo.be
	+32 (0)2 550 15 95	

Maximum/ Minimum funding per grant Awarded to a project partner:

Maximum 350.000 EUR per project (overhead included). If several FWO-funded partners are involved in one project, the funding must be shared between them.

Exchange Rate: NA

Eligible Call Topics: Topic 1, 2, and 3

Eligible One Health Settings: Human Health, Animal Health, Environment and Plant Health

Eligible institutions:

Academic. In this call, FWO deploys two of its regular funding channels: Fundamental Research Projects (FO) and Strategic Basic Research Projects (SBO). Researchers should choose the appropriate channel based on their project type, with eligibility details available in the regulations on the FWO website.

Eligible experimental approaches and disciplines:

Fundamental and strategic basic research in all scientific disciplines. Projects aiming at the development of a spin-off company are not eligible.

Clinical research and innovative intervention studies (including medical devices, technologies and tools for prevention, diagnosis and treatment of human and animal diseases) are eligible only if they address a fundamental scientific question.

Eligibility criteria for the Principal Investigator:

In this call, the PI can be a coordinator on one proposal or a partner on up to two proposals. Participating in this call does not affect FWO's national project submission limits. The PI must have an

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appointment covering the full project duration. PIs who become emeritus during the application year or project period are ineligible, i.e. art. 10 §7 of the regulations FO does not apply.

Eligible costs:

Different cost models and overhead calculations apply to each channel (FO vs. SBO). For overhead calculation, apply a structural rate to total costs: FO projects use 6% and SBO projects use 17%. For example, an SBO project costing 250,000 EUR amounts to 292,500 EUR with a 17% overhead, staying within the budget cap of 350,000 EUR. On FWO's e-portal, enter only the actual cost; FWO will add the overhead. The project has a duration of 36 months and all allocated funds must be spent within this timeframe. There are no automatic extensions granted, nor may any unused funds be carried forward after the project's end date, i.e. article 28 of the regulations FO and article 14 of the regulations SBO do not apply.

Submission of the proposal (or other information) at the national level:

Applicants for FWO funding must submit a **mandatory administrative application** through FWO's e-portal. Select "Research projects – European programme fundamental research" for FO projects or "Research projects – European programme strategic basic research" for SBO projects. If multiple Flemish partners request FWO funding, include all relevant partner details in a single e-portal submission. The national submission deadline matches the joint transnational call's preproposal stage. However, to confirm eligibility, it is recommended to consult the FWO administration at least one week prior. Failure to comply with these requirements may result in ineligibility.

Further guidance:

To avoid any potential issues, we encourage you to contact the FWO administration in advance. We are pleased to assist you in ensuring the eligibility of your project proposal and consortium.

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Belgium-SPW

Full name of the funding organisation:	**Service Public de Wallonie**
National/ Regional Financial Commitment:	**1 000 000 €**
Expected number of funded projects:	**1-2**
Contact details: (person (s), e-mail, telephone):	**Fleur Roland fleur.roland@spw.wallonie.be** **Cédric Morana cedric.morana@spw.wallonie.be**

Maximum/ Minimum funding per grant Awarded to a project partner: NA

Exchange Rate: NA

Eligible Call Topics: Topic 2 and 3

Eligible One Health Settings: Human Health, Animal Health, Environment and Plant Health

Eligible institutions: Universities, Higher education institution, Research Centres, Large companies, SME's

Eligible experimental approaches and disciplines: Applied Research.

SPW can cover pre-clinical trials but not clinical trials.

Eligibility criteria for the Principal Investigator: The Walloon partners should include at least one company established in Wallonia. The research budget of the Walloon companies must account for at least 40% of the total research budget of all Walloon partners.

Eligible costs: The Walloon "guide des dépenses éligibles" is applicable.

Submission of the proposal (or other information) at the national level: Walloon applicants must contact SPW at least 2 weeks before submission deadline. They must also submit a proposal on the regional submission platform: ONTIME.

Further guidance:

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- Applicants to SPW funding must submit their pre-proposal on the regional application platform ONTIME. Proposals invited to the second stage must also be submitted on the same platform. The submission deadlines are the same as the general deadline of the OH-AMR call.
- Applicants who want to submit a proposal are requested to contact SPW at least 2 weeks before the submission deadline.
- Participants must be companies, universities/higher education institutions, accredited or certified research centers, established in the Walloon Region and conduct R&D activities within the project.
- Walloon partners in the consortium must include at least one company, and the research budget of the Walloon partner companies must account for at least 40% of the total research budget of all Walloon partners.
- Participants must be based in Wallonia, and the Walloon companies must have an operational unit in Wallonia.
- Participants cannot receive any other public funding for the same activities.
- Participants must have fulfilled their obligations under any previous support granted by the Region.
- Companies must not be in difficulty according to the European Union guidelines.
- Projects must focus solely on civilian technologies, products, processes, and services.
- Participants must present an innovative R&D project that has a positive impact on the Walloon economy and aligns with the priority of the regional Smart Specialization Strategy (S3 Wallonia).
- Participants must demonstrate their ability to carry out the tasks assigned to them in the project, to exploit its results, and to have a positive socio-economic and sustainable development impact on Wallonia.
- The project cannot start at a TRL (Technology Readiness Level) lower than 3.

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Full name of the funding organisation:	**Canadian Institutes of Health Research**
National/ Regional Financial Commitment:	**$600 000 CAD**
Expected number of funded projects:	**1**
Contact details: (person (s), e-mail, telephone):	**CIHR Contact Centre support-soutien@cihr-irsc.gc.ca +1 613-954-1968**

Maximum/ Minimum funding per grant Awarded to a project partner:

Up to $600,000 CAD per grant Awarded to a project partner.

Exchange Rate: 1 Euro = 1.6058 (Bank of Canada rate as September 10, 2026)

Eligible Call Topics:

Consistent with the One Health approach embedded within the Pan-Canadian Action Plan on AMR and prioritized within CIHR-III's Strategic Plan for 2021-2026, and CIHR's commitment to advancing the objectives of the Management Action Response Plan on Antimicrobial Resistance projects addressing Call projects must be relevant to Call Topics 1, 2, or 3 are eligible for CIHR funding, provided they include a clear and substantive human health research component. For Topic 3, eligibility is limited to projects in which environmental research activities are explicitly linked to, and intended to generate knowledge or outcomes relevant to, human health. Projects focused solely on animal health, plant health, food systems, or environmental settings without a demonstrated human health connection are not eligible for CIHR funding. Projects may include only one (1) Canadian component

Eligible One Health Settings:

CIHR can only fund research groups working on Human Health through this call.

Eligible institutions:

Details regarding eligible applicants for a given competition will be specified in the funding opportunity on ResearchNet. Individuals in the Nominated Principal Applicant role must be affiliated with a CIHR eligible Canadian postsecondary institution and/or their affiliated institutions; non-governmental organisations with a research or knowledge translation mandate; individuals working with municipal, provincial, and/or territorial governments are also eligible where the research proposed is not already funded by that Government of Canada sector.

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Eligible experimental approaches and disciplines:

Projects involving pre-clinical studies or clinical trial pilots are eligible under this Funding Opportunity.

Eligibility criteria for the Principal Investigator:

Individuals in the Nominated Principal Applicant (NPA) role must have their substantive role in Canada for the duration of the requested grant term. Note that CIHR is not prescriptive regarding the duration of time that a NPA must physically reside in Canada, as this falls under the purview of applicable policies of the administering institution, employment terms and conditions, and or collective agreements Appointments and/or positions that can be renewed prior to the end of the requested grant term are eligible at the discretion of the administering institution.

Projects receiving a CIHR grant must comply fully with the CIHR Funding Policies. Policies and guidelines cover areas such as Applicant Responsibilities, Official Languages policy, Access to Information and Privacy Acts. For more information, please refer to Tri-Council Policy Statement: Ethical Conduct for Research Involving Humans (TCPS 2) and Tri-Agency Framework: Responsible Conduct of Research.

Eligible costs:

Applicants should review the Use of Grant Funds Section of the Tri-Agency (CIHR, NSERC and SSHRC) Guide on Financial Administration for a complete listing and description of allowable costs and activities.

Submission of the proposal (or other information) at the national level:

Canadian applicants will need to submit both the pre and full proposals through ResearchNet in addition to submitting to the Joint Call Secretariat.

Further guidance:

Applications submitted to CIHR require applicant consent and institutional approval (if applicable) to the use and disclosure of full application and nominative information for relevance review and funding decisions at the time of application.

- The applicant will be required to submit all reporting requirements to CIHR in addition to the JCS.
- The Nominated Principal Applicant will be required to submit an Electronic Final Report to CIHR. This online report will be made available to the Nominated Principal Applicant on ResearchNet at the beginning of the grant funding period and can be filled in as the research progresses.
- All reports may be shared with partners supporting the grant.
- The Nominated Principal Applicant must have successfully completed one of the sex- and gender-based analysis training modules available online through the CIHR Institute of Gender and Health and have submitted a Certificate of Completion (see How to Apply section). Select

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and complete the training module most applicable to your research project. Applicants are encouraged to review the "How to integrate sex and gender in research" section on the CIHR website.

CIHR is committed to promoting the inclusion and advancement of groups underrepresented in science as one of the means to enhance excellence in research, training and knowledge translation. CIHR's position on equity, diversity and inclusion (EDI) is available in the Tri Agency Statement on Equity, Diversity and Inclusion. Additional guidance can be found on the Best practices in Equity, Diversity and Inclusion in Research webpage.

It is noted that CIHR does not retain or claim any rights to IP in relation to research that if funds. Accordingly, the Canadian researchers retain full freedom in negotiating the Partner Consortium Agreement (PCA) required, including whether or not to accept the IP conditions. The Nominated Principal Applicants do NOT need to send a signed copy of the PCA to CIHR. The proposals will be funded based on the ranking list recommended by the Peer Review Panel and decided by the Call Steering Group. The final funding decision will be made by the national/regional funding organizations and will be subject to budgetary considerations with the goal of optimal usage of the available budget. Approved joint transnational teams may receive an across-the-board cut to the budget, if necessary, to maximize the number of funded opportunities. For full details of CIHR's requirements, please refer to the Funding Opportunity on ResearchNet.

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Czech Republic- MZCR & AZVCR

Full name of the funding organisation:	**Ministerstvo Zdravotnictvi Ceske Republiky (MZCR)**
National/ Regional Financial Commitment:	**500 000 €**
Expected number of funded projects:	**2**
Contact details: (person (s), e-mail, telephone):	
Monika Kocmanova; National coordinator on Health-related European Partnerships (AZVCR); Email: monika.kocmanova@azvcr.cz; Phone: + 420 778 973 186	
Olga Laaksonen; Head of the Science, Research, and Subsidies for Education Unit (MZCR); Email: olga.laaksonen@mzd.gov.cz; Phone: +420 604 786 141	

Maximum/ Minimum funding per grant Awarded to a project partner:

The maximum funding must not exceed EUR 250,000.

Exchange Rate: Convert amounts into Czech crowns (CZK) using the Czech National Bank (CNB) exchange rate valid on the last day for submitting full proposals.

Eligible Call Topics: Topic 1, 2, and 3

Eligible One Health Settings: Human Health

Eligible institutions: Research Organisations, Enterprises, Patient Organisations. All eligibility rules and criteria can be found on the Czech Health Research website (Výzva 2027 – AZV ČR). It is **recommended** to contact the responsible person at the Czech Health Research Council (prior to submission regarding the eligibility criteria).

Conditions for PO funding – Patients organisations can receive direct funding if they take an active role in the project´s research activities. This means they must contribute to specific research objectives (for example, by being involved in one or more work packages) and these types of research activities must be clearly described in their **statues.**

Definition of ECR - A natural person engaged in research who, in the year of submission of the Project proposal to the call for proposals, has received their Ph.D. academic title or its equivalent in the past 8 years, or has obtained it no later than the date of conclusion of the Contract/issuance of the Project Decision. If the Proposer has been on maternity or parental leave, has suffered a long-term illness, or has interrupted his/her scientific career for similar objective reasons, the time limit of 8 years from the award of the academic degree of Ph.D. or its equivalent is increased by this period. These facts (award of the degree, parental leave, etc.) shall be documented by the Applicant by means of a Sworn Statement.

Eligible experimental approaches and disciplines:

The Ministry of Health of the Czech Republic (MZCR) is not permitted to finance projects focused exclusively on basic research. It may, however, support projects that combine basic and applied research, provided that applied research plays the principal role.

Eligibility criteria for the Principal Investigator:

The PI must hold a Ph.D. degree (for more information, please refer to the national document *"Podmínky pro zapojení českého uchazeče do výzvy",* available on the AZVCR website). MZCR/AZVCR sets a criterion for the **maximum workload**, **which must not exceed 1.00 FTE within a single submitted project**. There is **no minimum workload requirement for the PI** in relation to European Partnerships (EP) in health-care area imposed by the national funding authority (MZCR). However, the allocated FTE must correspond to the actual workload required for the project. Within the EP, the assigned FTE is subject to verification by the institution or organisation employing the PI. Therefore, the provisions of Act No. 262/2006 Coll., the Labour Code, must be respected. AZV CR does not impose any additional FTE-related restrictions within the framework of the EP.

Eligible costs:

All eligibility of costs, types and their caps can be found on the Czech Health Research Council (Výzva 2027 – AZV ČR). It is **strongly recommended** to contact the responsible person at the Czech Health Research Council (prior to submission regarding the eligibility criteria).

Submission of the proposal (or other information) at the national level:

Prior to submission of the pre-proposal to EUP OH AMR, Czech researchers need to submit to the Czech Health Research Council the following documents:

1. **Sworn Statement of a Legal Entity / Natural Person (mandatory)**
2. **Sworn Statement for a Research Organisation (if relevant)**
3. **Sworn Statement of composition consortium (only if SMEs or industry are involved in the project proposal from the Czech side)**
4. **Application Form**

Czech partners are required to complete a **national Application Form**, providing basic information about the applicant and any co-applicant(s), if applicable, including their respective budgets. It is necessary to list all national institutions involved in the project due to funding requirements.

All these documents are available on the website at the Czech Health Research Council AZV ČR –Výzva 2027 – AZV ČR.

Prior to submission of the full proposal to EUP OH AMR, Czech researchers need to submit to the Czech Health Research Council the following documents:

1. **Documents** related to **professional competence**, depending on the nature of the project, must be provided in the form of a **Sworn Statement**, which will be available on the website at the Czech Health Research Council AZV ČR –Výzva 2027 – AZV ČR.
2. **Updated Application Form**

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Czech partners are required to complete the **updated national Application Form**, providing basic information about the applicant and any co-applicant(s), if applicable, including their respective budgets. It is necessary to list all national institutions involved in the project due to funding requirements.

According to Czech regulations, the main Czech applicant will sign a grant agreement with the national funding authority (MZCR) and, if there are any other Czech co-applicant(s), will subsequently enter into a cooperation agreement with them.

At the international level (pre- or full proposal), it is preferable to list only one Czech partner – the main applicant. If needed, it is possible to list more than one partner (in accordance with the call rules); however, at the national level, there will be one main Czech applicant while the remaining national institutions will act as co-applicants. Together, they must share the allocated project budget among themselves.

The total project budget must not exceed EUR 250,000.

In case the projects of Czech participants are recommended for funding based on the results of the international evaluation and after the approval of the representatives of the funding authorities of the countries participating in the EUP OH AMR calls, the Ministry of Health of the Czech Republic / the Czech Health Research Council may ask the successful Czech participants to submit additional documents in order to issue a decision on the provision of purpose-special support according to the rules established by the Ministry of Health of the Czech Republic / Czech Health Research Council.

Further guidance: Výzva 2027 – AZV ČR

IMPORTANT: All projects will be funded under the Programme for the Support of Applied Health Research for the Years 2024–2030 and must therefore start on 1 January 2028 and be completed no later than 31 December 2030, as the programme itself ends at the end of that year.

In the Consortium Agreement, it must therefore be stated that due to the national funding conditions applicable to the Czech partner, the project duration for the Czech partner differs from that of the other consortium partners. The Czech partner must start its project activities on 1 January 2028 and complete them no later than 31 December 2030, without any extension possible.

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Denmark-IFD

Full name of the funding organisation:	**Innovationsfonden, Innovation Fund Denmark**
National/ Regional Financial Commitment:	**1 600 000 €**
Expected number of funded projects:	**2-5**
Contact details: (person (s), e-mail, telephone):	**Marie Friis Madsen**
	marie.friis.madsen@innofond.dk, +4561905059

Maximum/ Minimum funding per grant Awarded to a project partner:

Min: 75 000 EUR

Max: 400 000 EUR /650 000 for all Danish partners in a project

Exchange Rate: 7,5 Danish Crown for 1 Euro

Eligible Call Topics: Topic 1, 2, and 3

Eligible One Health Settings: Human Health, Animal Health, Environment and Plant Health

Eligible institutions: All

Eligible experimental approaches and disciplines:

Eligibility criteria for the Principal Investigator:

Eligible costs: See Guidelines for International Collaborations

Submission of the proposal (or other information) at the national level:

Danish applicants will be invited to e-grant, where they must upload declarations.

Further guidance:

Guidelines for International Collaborations

Full name of the funding organisation:	**Regionaal- ja Põllumajandusministeerium**
National/ Regional Financial Commitment:	**100 000 €**
Expected number of funded projects:	**1**
Contact details: (person (s), e-mail, telephone):	**Maarja Malm, maarja.malm@agri.ee +372 5670 2945**

Maximum/ Minimum funding per grant Awarded to a project partner:

Maximum funding per partner is 100 000 EUR.

Exchange Rate: NA

Eligible Call Topics: Topic 1

Eligible One Health Settings: Human Health, Animal Health, Environment, Plant Health

Eligible institutions:

The Host Institution may be any legal entity that is registered and located in Estonia and has an Estonian bank account. Restrictions to eligible institutions and applicants may however be stated in the general call conditions which also apply to Estonian participants.

The Host Institution (the final recipient) is the institution to which the grant will be allocated.

After the submission deadline (in case of two-stage application, after the preproposal deadline) and upon the notice from the Funding Organisation, the Host Institution must confirm to the Funding Organisation in writing that the Estonian sub-project can be carried out on their premises in Estonia and that they will employ the Principal Investigator during the sub-project, should the project receive funding.

If the Host Institution is an undertaking, the State aid and de minimis aid regulations must be taken into account.

Eligible experimental approaches and disciplines: REM cannot fund innovation.

Eligibility criteria for the Principal Investigator:

The Principal Investigator (PI) is a researcher who acts as the Estonian team leader in the project proposal. He/she will be responsible for how the grant is used and how the Estonian part in the project is executed. The PI must be clearly identified in the proposal.

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The Principal Investigator:

1. must have an updated public profile in the Estonian Research Information System (ETIS) by the submission deadline (in case of two-stage application, by the preproposal deadline);

2. must hold a doctoral degree or an equivalent qualification. The degree must be awarded by the submission deadline (in case of two-stage application, by the preproposal deadline) at the latest;

3. must have published at least three articles that comply with the requirements of Clause 1.1 of the ETIS classification of publications, or at least five articles that comply with the requirements of Clauses 1.1, 1.2, 2.1 or 3.1, within the last five calendar years prior to the proposal submission deadline (in case of two-stage application, prior to the preproposal deadline). Patents are equalled with publications specified under Clause 1.1. A monograph (ETIS Clause 2.1) is equalled with three publications specified in Clause 1.1 if the number of authors is three or fewer. If the applicant has been on maternity, paternity, or parental leave, in compulsory military service, or if there have been other exceptional circumstances (e.g., serious illness), they can request the publication period requirement to be extended by the relevant period of time.

If the Principal Investigator has received the PhD degree outside Estonia, its correspondence to an Estonian doctoral degree must be recognised by either the Estonian ENIC-NARIC Centre or the Host Institution in accordance with the Regulation of the Government of the Republic of April 6, 2006, No. 89 "Evaluation and academic recognition of documents proving foreign education and the name of the qualification awarded in the foreign education system terms and conditions of use". The Funding Organisation may ask for a relevant Evaluation Report.

If several Estonian institutions participate in a proposal, all institutions must have a Principal Investigator who meets the national eligibility requirements.

In the same call, a Principal Investigator may participate in no more than two proposals, in one of which she/he may act as coordinator (lead PI).

Eligible costs:

The budget contains costs directly related to the implementation of the project (direct costs) and indirect (overhead) costs. The direct costs must be used to carry out the project and be separately identifiable.

Direct costs:

1. Personnel costs are monthly salaries (along with all state taxes, contributions, and compensations arising from law) of the project participants, calculated according to their commitment and in proportion to their total workload at their Host Institution.

2. Other direct costs are:

- travel costs that may cover expenses for transport, accommodation, daily allowances and travel Insurance. If the project is funded from the European Regional Development Fund (Mobilitas 3.0) resources, travel costs are eligible only for travels abroad;

- consumables and minor equipment directly and fully related to the project;

- publication and dissemination of project results;

- organising meetings, seminars or conferences (e.g. room rent, catering, equipment rental and related costs);

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- fees for participating in scientific forums, conferences and other events directly and fully related to the project;

- patent costs;

- all other costs that are identifiable as clearly required for carrying out the project (e.g. translation, copy editing, webpage hosting, postage costs etc.) and are directly and fully related to the project.

Indirect costs (overhead) are costs that cannot be identified as specific costs directly linked to the performance of the action and/or should cover the general expenses of the Host Institution related to the management of the grant. Office consumables and costs for equipment and services intended for general use (e.g., phone bills, copy service, printer) should be covered from the indirect costs. Indirect costs are 15% of the personnel costs of the Estonian sub-project.

Subcontracting costs are direct costs. Subcontracting costs should cover only additional or complementary research related tasks (e.g. analyses, conducting surveys, building a prototype, etc.) performed by third parties. Subcontracting costs should not be included in the overhead calculation. The activities and budget should be described in the proposal. Core project tasks should not be subcontracted. Subcontracting costs may not exceed 15% of the total costs of the Estonian sub-project.

Double funding of activities is not acceptable (i.e. the same cost item may not be reimbursed from different funding sources).

If several Estonian institutions participate in one proposal, the sum of their requested budgets may not exceed the maximum contribution per proposal of the respective national Funding Organisation indicated in the call documents.

Submission of the proposal (or other information) at the national level: NA

Further guidance:

Only research projects (basic or applied) within TRL range 1–6 are funded.

National Eligibility Criteria for grant applications in partnership calls for transnational research projects (https://etag.ee/wp-content/uploads/2026/11/Vastavusnouded-RV-uhiskonkurssidel_-EN_nov2026.pdf)

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Full name of the funding organisation:	**Agence Nationale de la Recherche**
National/ Regional Financial Commitment:	**1 800 000 €**
Expected number of funded projects:	**8**
Contact details: (person (s), e-mail, telephone):	**Dr Sophie GAY/ Dr Yue XIAO** EUPOHAMR_calls@agencerecherche.fr +33 1 78 09 80 39 / +33 1 73 54 82 41

Maximum/ Minimum funding per grant Awarded to a project partner:

Maximum amount per project if two French partners are involved: 450 000 €

Maximum funding per partner: 200 000 € (Increased to 350 000 € for coordinators)

Minimum amount per partner: 15 000 €

Eligible Call Topics: Topic 1, 2 and 3

Eligible One Health Settings: Human Health, Animal Health, Environment and Plant Health

Eligible institutions: Research organisations and undertakings

ANR may fund research organisations and undertakings, as defined by the EC regulation on State aid for research, development and innovation (see the ANR Funding regulations at actual costs for further reference). Only research organisations that have their primary establishment in France may be funded. As for undertakings, ANR may fund those that have their real head office in an EU member State and an establishment (primary or secondary) in France.

Companies in difficulties are excluded from ANR subventions (More details here: ANR-RF-Fiche-EED-2021.pdf).

Eligible experimental approaches and disciplines: ANR may finance fundamental research, industrial research and experimental developments in all research disciplines. ANR can cover pre-clinical trials but not clinical trials.

Eligibility criteria for the Principal Investigator:

To be eligible for this call as Principal Investigator, researchers must hold a research position in an eligible institution covering the duration of the project. The salary of the Principal Investigator will not be covered by the present grant. Researchers applying to this call as Principal Investigator while not having a permanent position must imperatively send the documents (working contract covering the

duration of the project, chair announcement covering the duration of the project, commitment of the employer) proving their eligibility by mail to the ANR contact point (EUPOHAMR_calls@agencerecherche.fr) before the closing of the call. Researchers do not need to head their own lab, or group while applying. If a researcher does not head its own research group/lab, the signature of the head of the department should be included in the letter of intent, even at the pre-proposal stage.

Eligible costs:

ANR may finance fundamental research, industrial research and experimental developments. Funding rates vary based on the type of research and the type of partners requesting support, please refer to the funding regulations of ANR at https://anr.fr/fr/rf/ to construct the project budget and fill the form related to economical activities to identify your funding rate.

Standard ANR funding rules apply for eligible costs, unless stated otherwise in the Annex « Modalités pour les partenaires sollicitant une aide de l'ANR ». These rules are specified in ANR's "ANR Funding regulations" details ("at actual costs" version https://anr.fr/fileadmin/documents/2025/ANR-RF-072025-cout-reel.pdf). An explanatory note regarding the eligible costs is also available at: https://anr.fr/fr/rf/fiche-couts/

Eligible costs include (but are not limited to) the following: personnel costs (for temporary contracts only for organisations funded at "coût marginal"); small equipment; consumables and animal costs; travel; and sub-contracting costs (max 50% of requested budget per partner). Detailed information, factsheets, FAQs and other resources are available at https://anr.fr/fr/rf/

The ANR heading for "overheads" in the ANR funding breakdown is «frais d'environnement». 13,5% of the total eligible costs must be applied for if the partner belongs to a public research organisation (or other organisation funded at "marginal" costs), or up to 68% of the total personnel costs and 7% of other costs for partners funded at full economic cost (such as enterprises) (cf " règlement financier ")

Submission of the proposal (or other information) at the national level:

At the pre-proposal stage, Principals Investigators not having a permanent position and requesting ANR funding must submit by e-mail the document proving that their salary will be covered by a funding source other than the OHAMR Grant (see above for more details).

At the full-proposal stage, applicants requesting ANR funding must submit their administrative and financial data on ANR's submission tool.

Applicants are strongly advised to first prepare and complete their budget in the ANR submission platform, and then report/forward to the coordinator the same figures in the EUP OHAMR submission platform (PT-outline). This approach will ensure consistency between both submissions and reduce the risk of discrepancies or errors in the financial data.

For more details, please refer to the dedicated EUP OHAMR 2027 call page on the ANR website: https://anr.fr

Further guidance:

At the granting stage, partners selected for funding by ANR may have to adjust and complete administrative and financial data provided on the ANR platform.

Administrative reporting documents (financial reports or certificate of closure, depending on the type of beneficiary, must be communicated to ANR according to the provisions of ANR Funding regulations ("at actual costs" version).

Access to genetic resources and benefit sharings: Funded partners participating in projects falling within the scope of the regulations on access to genetic resources and benefit-sharing (Nagoya protocol) will be required to provide evidence to demonstrate compliance with these obligations, and must ensure that all data relating to such genetic resources or associated traditional knowledge are kept in order to demonstrate that the necessary due diligence has been exercised.

In case the project is selected for funding, one of the partners funded by ANR must be designated as « référent ANR ». This partner will be the lead contact point with ANR for the scientific and administrative aspects. This partner must communicate to ANR the required Scientific reports, the Consortium Agreement, Data management plans according to the funding contract and the EUP OHAMR obligations and fill the requested data on the OASIS platform during the course of the project.

In keeping with the national "PPST" policy (scientific and technological potential protection), applicants to ANR should consult their local "FSD" (security and defence officer, where available) on their project before applying. Applications to ANR may be forwarded to the HFSD of the French Ministry of research and higher education for screening. A negative appraisal by the HFSD may cause ANR to reject the proposal.

The above-mentioned terms and conditions are only summarized translations of those entailed in the ANR Funding regulations and in the Appendix "Modalités pour les Partenaires sollicitant une aide de l'ANR". In case of a conflict of interpretation between the terms and conditions stated in this annex and the "Modalités de participation" and "Règlement financier", the latter shall prevail.

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Germany- BMFTR & DLR

Full name of the funding organisation:	**Bundesministerium für Forschung, Technologie und Raumfahrt (BMFTR)** **Deutsches Zentrum für Luft- und Raumfahrt Projektträger (DLR)**
National/ Regional Financial Commitment:	**3 700 000 €**
Expected number of funded projects:	**10**
Contact details: (person (s), e-mail, telephone):	**Barbara Junker, Patricia Ruiz, Yen Ly-Sauerbrey** **EUP-OHAMR@DLR.de +49 228 3821-1326**

Maximum/ Minimum funding per grant Awarded to a project partner: 350 000 €

Exchange Rate: NA

Eligible Call Topics: Topic 1, 2 and 3

Eligible One Health Settings: Human Health, Animal Health, Environment and Plant Health

Eligible institutions:

- Universities
- University hospitals
- Non-university research institutes
- Industry

Note: industry is funded with a maximum of 50-60% of their costs.

Eligible experimental approaches and disciplines: The funding organization will not fund clinical trials.

Eligibility criteria for the Principal Investigator: None

Eligible costs: Personnel, Consumables, Animals, Subcontracts, Equipment, Travel, Overheads.

Overheads refer to “Gemeinkosten” (applicable e.g. for Helmholtz centres and Fraunhofer-Society) as well as “Projektpauschale” (applicable for universities and university hospitals).

Individual project coordinators/partners may request up to 350 000 Euro including overheads. A project consisting of two German partners may request a maximum of 550 000 Euro including overheads.

Submission of the proposal (or other information) at the national level: Not necessary

Further guidance: For further details please refer to the national guidelines.

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Hungary-NKFIH

Full name of the funding organisation:	**Nemzeti Kutatási, Fejlesztési Innovációs Hivatal**
National/ Regional Financial Commitment:	**300 000 €**
Expected number of funded projects:	**1-2 (The number of projects funded depends on the requested amount in the submitted applications.)**
Contact details: (person (s), e-mail, telephone):	**Zsuzsanna Kürti ncp@nkfih.gov.hu +36 1 795 9500**

Maximum/ Minimum funding per grant Awarded to a project partner:

300 000 EUR / project

Exchange Rate: NA

Eligible Call Topics: Topic 1, 2, and 3

Eligible One Health Settings: Human Health, Animal Health, Environment and Plant Health

Eligible institutions:

Institution of higher education, Other budgetary research institution, Other budgetary institution (outside education or research), Enterprise-based research organisation, Enterprise (non-research type), Non-profit research organisation, Other non-profit organisation (outside research), HUN-REN Hungarian Research Network.

All eligibility rules and criteria can be found in the national call. **Please consult the national call for participation eligibility and funding intensity: Call for proposals to support Hungarian organisations successfully participating in the joint international calls for proposals of European Partnerships in Horizon**:
https://nkfih.gov.hu/palyazoknak/nkfi-alap/horizont-europa-europai-partnersegek-magyar-szervezetek-tamogatasa-2024-121-he-partnerseg/palyazati-felhivas

Applicants are advised to verify the applicable funding intensity and eligibility conditions in the relevant National Call.

Eligible experimental approaches and disciplines: All.

Eligibility criteria for the Principal Investigator:

For details, please refer to the Guide for Applicants for the **2024-1.2.1-HE_PARTNERSÉG** national call:
https://nkfih.gov.hu/palyazoknak/nkfi-alap/horizont-europa-europai-partnersegek-magyar-szervezetek-tamogatasa-2024-121-he-partnerseg/palyazati-felhivas

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Eligible costs:

Personnel (temporary, permanent), subcontracting, services, travel, consumables, equipment, coordination, communication and dissemination, overhead (20%), overhead for travel related activities (5%). Please consult the national call for cost eligibility.

https://nkfih.gov.hu/palyazoknak/nkfi-alap/horizont-europa-europai-partnersegek-magyar-szervezetek-tamogatasa-2024-121-he-partnerseg/palyazati-felhivas

Submission of the proposal (or other information) at the national level:

The Hungarian partners of proposals recommended for funding have to submit an application for the national call after having been informed of the international funding decision.

For details, please refer to the Guide for Applicants for the **2024-1.2.1-HE_PARTNERSÉG** national call: https://nkfih.gov.hu/palyazoknak/nkfi-alap/horizont-europa-europai-partnersegek-magyar-szervezetek-tamogatasa-2024-121-he-partnerseg/palyazati-felhivas

Further guidance:

The Guide for Applicants for **2024-1.2.1-HE_PARTNERSÉG** national call is applicable.

https://nkfih.gov.hu/palyazoknak/nkfi-alap/horizont-europa-europai-partnersegek-magyar-szervezetek-tamogatasa-2024-121-he-partnerseg/palyazati-felhivas

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Ireland-DAFM

Full name of the funding organisation:	**Department of Agriculture, Food and the Marine**
National/ Regional Financial Commitment:	**200 000€**
Expected number of funded projects:	**1**
Contact details: (person (s), e-mail, telephone):	**Dr. Willie Ryan willie.ryan@agriculture.gov.ie**

Maximum/ Minimum funding per, grant awarded to a project partner:

Maximum of €200,000 per project, subject to funding budget.

Exchange Rate: NA

Eligible Call Topics:

Topic 2: Understand and characterise the role of host commensal micro-organisms (including fungi) in AMR dynamics and develop solutions aiming at selective decolonising of resistant organisms in the host (human and animals) to prevent transmission.

Eligible One Health Settings: Animal Health, Environment and Plant Health

Eligible institutions:

Research Performing Organisations (RPO) as described in National Annex Call_Guidelines_for_Irish_Applicants_GEH_011025.pdf

Eligible experimental approaches and disciplines:

Call_Guidelines_for_Irish_Applicants_GEH_011025.pdf

Eligibility criteria for the Principal Investigator:

Permanent staff member of RPO as described in National Annex Call_Guidelines_for_Irish_Applicants_GEH_011025.pdf

Eligible costs:

as described in National Annex Call_Guidelines_for_Irish_Applicants_GEH_011025.pdf

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Submission of the proposal (or other information) at the national level:

further information from **willie.ryan@agriculture.gov.ie**

Further guidance:

Additional Guidance can be found in National Annex
Call_Guidelines_for_Irish_Applicants_GEH_011025.pdf

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Full name of the funding organisation:	**Taighde Eireann – Research Ireland**
National/ Regional Financial Commitment:	**600 000€**
Expected number of funded projects:	**1-2**
Contact details: (person (s), e-mail, telephone):	**Suzan Gunbay**
	suzan.gunbay@researchireland.ie
	General mailbox eu-cofund@researchireland.ie

Maximum/ Minimum funding per grant Awarded to a project partner:

Up to €330,000 direct costs* for Irish-based researchers applying as a project partner

Up to €405,000 direct costs* for Irish-based researchers applying as a project coordinator

*The maximum total award, including 30% overhead contribution, will be €430,000, for a partner and €530,000 for applicants who take on the role of coordinator.

Exchange Rate: NA

Eligible Call Topics: Topic 1, 2, and 3

Eligible One Health Settings: Human Health, Animal Health, Environment and Plant Health

Eligible institutions:

Irish Host Research bodies eligible for Research Ireland funding.

Please refer to Research Ireland's Policies and Guidance for the list of eligible Research Performing Organisations: **Eligibility Information**

Eligible experimental approaches and disciplines:

All disciplines are eligible for Research Ireland funding.

Clinical trials and investigations are not eligible for funding by Research Ireland for the Irish-based applicant. Where clinical studies are proposed within the workplan of a proposal, study sites and the delineation of activities among proposal partners should be made clear.

Eligibility criteria for the Principal Investigator:

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Only an academic partner or coordinator based in an eligible Irish Host Research body may apply for Research Ireland funding.

The Irish-based applicant must:

- hold a PhD or equivalent qualification for at least 3 years by the pre-proposal deadline. The official date is defined as the day, month and year that the degree was conferred i.e., the month and year printed on the official PhD certificate.

AND

- be a member of the academic staff of an eligible Research Body (permanent or with an active contract that covers the period of the grant)

OR

- be a contract researcher with a contract that covers the period of the grant, who is recognised by the eligible Research Body as an independent investigator and will have an independent office and research space for which he/she will be fully responsible for at least the duration of the Research Ireland grant

OR

- be an individual who will be recognised by the eligible Research Body upon receipt of the grant as an academic staff or as a contract researcher as defined above. The applicant does not necessarily need to be employed by the Research Body at the time of the application submission.

AND

- be an author on at least three international peer-reviewed articles. Only original research publications, and not review articles or other secondary research literature, are acceptable.

Please refer to the Research Ireland call webpage for more information on eligibility criteria. Please note that Research Ireland may contact applicants directly to confirm eligibility post submission.

Eligible costs:

Funding is provided for up to 100% of eligible costs. The following indicates the maximum levels of funding that may be requested:

Up to €330,000 direct costs* for Irish-based researchers applying as a project partner

Up to €405,000 direct costs* for Irish-based researchers applying as a project coordinator

*The maximum total award, including 30% overhead contribution, will be €430,000, for a partner and €530,000 for applicants who take on the role of coordinator.

Eligible costs

1. **Salary-related costs** for research personnel. Please use current **Research Ireland Team Member Salary Scales**. The Irish partner cannot request their own salary or buy-out.
2. **Equipment costs** up to a maximum value of €10K
3. **Travel costs** with consideration for Research Ireland's **Guidance for Sustainable Travel Policy**
4. **Direct running costs** (materials and consumables)
5. **Dissemination and knowledge exchange costs**

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6. **Subcontracting costs** are considered an eligible budget category however strong justification for subcontracting must be provided and pre-approved directly with Research Ireland in advance of proposal submission.

Unless otherwise stated, all rules regarding listed eligible costs apply as defined within Research Ireland's grant budget policy. Please refer to this policy closely in the development of the budget request.

Submission of the proposal (or other information) at the national level:

Please give a brief notification of your intent to submit a pre-proposal through email to **eu-cofund@researchireland.ie** before the submission deadline. Within the notification, please include the following information:

- Call topic
- List of project partners
- Irish Host institution*
- Total budget request to Research Ireland
- Whether you intend to apply as a coordinating or non-coordinating partner

*Please note it is also the responsibility of the applicant to notify the Research Office of their Host Institution of their intention to submit a pre-proposal.

Further guidance:

State Aid: Applicants are advised that funding awarded by Research Ireland under the One Health Antimicrobial Resistance (OHAMR) Partnership Programme will be subject to, and must comply with, State aid rules and the conditions of the EU Commission General Block Exemption Regulation (GBER). Funding will be awarded to successful applicants under Article 25, in respect of aid for research and development projects. For further details please consult: Taighde Éireann-Research Ireland Research and Innovation Scheme 2021-2026

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Ireland-HRB

Full name of the funding organisation:	**Health Research Board (HRB)**
National/ Regional Financial Commitment:	**530 000 €**
Expected number of funded projects:	**1-2**
Contact details: (person (s), e-mail, telephone):	**Orla Morton, HRB-JTCs@hrb.ie**

Maximum/ Minimum funding per grant Awarded to a project partner:

Project Partners: €330,000 direct costs.

Project Coordinators: €405,000 direct costs (with the additional €75,000 for coordination-specific activities).

For more details, please visit https://www.hrb.ie/funding-scheme/eup-ohamr-2027/

Exchange Rate: NA

Eligible Call Topics: Topic 1 and Topic 3

Eligible One Health Settings: All One Health settings (Human, Environment, Animal, Plant) are eligible, provided the research focus clearly demonstrates its relevance and connection to Human Health.

Eligible institutions:

Lead applicants must be affiliated with approved HRB Host Institutions. See also the HRB Policy on Approval of Host Institutions (https://www.hrb.ie/wp-content/uploads/2024/10/List-of-Approved-HRB-Host-Institutions.pdf).

Eligible experimental approaches and disciplines:

All Irish partners who are undertaking feasibility and/or interventional studies must adhere to the HRB Clinical Trial and Interventions Research Governance Policy (https://www.hrb.ie/wp-content/uploads/2024/05/HRB-Clinical-Trials-and-Interventions-Governance-Policy-1.pdf).

Eligibility criteria for the Principal Investigator:

For more details, please visit https://www.hrb.ie/funding-scheme/eup-ohamr-2027/

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Eligible costs:

For more details, please visit https://www.hrb.ie/funding-scheme/eup-ohamr-2027/

Submission of the proposal (or other information) at the national level:

The Host Institution must send the application to HRB-JTCs@hrb.ie within three working days following the submission deadline, along with the completed Host Institution sign off form. Eligibility form must be completed for new applicants. For more details, please visit https://www.hrb.ie/funding-scheme/eup-ohamr-2027/

Further guidance:

Please see HRB's dedicated funding scheme page on the HRB website for more detailed guidance and FAQs specific to applicants based in Ireland.

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Israel-CSO-MOH

Full name of the funding organisation:	**Ministry Of Health - Chief Scientist Office**
National/ Regional Financial Commitment:	**360 000 €**
Expected number of funded projects:	**2**
Contact details: (person (s), e-mail, telephone):	**Dr. Ronit Meyuhas**
	ronit.meyuhas@moh.gov.il +97225082159

Maximum/ Minimum funding per grant Awarded to a project partner:

0.36M (up to 2 projects. 140K per project + additional 40K per project coordinators)

Exchange Rate: NA

Eligible Call Topics: All with relation to human health.

Eligible One Health Settings:

Human health, Animal health and Environmental health (Only with relation to human health).

Eligible institutions:

Hospitals, clinics, laboratories, academic and public research institutions.

Eligible experimental approaches and disciplines:

Standard National Grant Conditions apply.

Eligibility criteria for the Principal Investigator:

PI should hold a Ph.D., M.D., D.M.D., D. Sc or equivalent degree and employed by an eligible institution.

Eligible costs:

Materials and consumables; Travel (up to 10%); No salaries for PIs; No heavy equipment, Institutional overhead 10%.

Submission of the proposal (or other information) at the national level:

Prior to submission, researchers will submit to CSO-MOH an abstract approved by their research authority including budget distribution. No submission of abstract can result in declaration of the consortium as ineligible. If the application involves human or animal experiments, bioethics approvals must be submitted with the application or up to 4 months later. Research will not be funded simultaneously by CSO-MOH on more than one grant (Era-NET or national). Researchers can not apply for more than one grant from any ERA-NET funded by CSO-MOH or submit more than one proposal for any programme.

Further guidance:

Please see detailed instructions of application at the national level and reporting at https://www.gov.il/he/service/era-net-instructions-for-israeli-researchers

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Full name of the funding organisation:	Fondazione regionale per la Ricerca Biomedica
National/ Regional Financial Commitment:	**€ 1,500,000.00**
Expected number of funded projects:	**3/4**
Contact details: (person (s), e-mail, telephone):	
Fabio Rondini fabio.rondini@frrb.it	
+39 02 6765 0163	
Giulia Maria Rossignolo: giuliamaria.rossignolo@frrb.it	
+39 02 6765 0159	

Maximum/ Minimum funding per grant Awarded to a project partner:

€ 500,000.00

Exchange Rate: NA

Eligible Call Topics: Topic 1, Topic 2, Topic 3

Eligible One Health Settings:

Human health

Eligible institutions:

Eligible applicants must be located in Lombardy:

1. Public or Private Italian IRCCS (Scientific Institutes for Health Research, Hospitalization and Health Care). They can apply as Partners or Coordinators of a Consortium.

2. Public Health Care Providers (ASST). They can apply as Partners or Coordinators of a Consortium.

3. Agenzie di Tutela della Salute (ATS). They can apply as Partners or Coordinators of a Consortium.

4. Azienda Regionale Emergenza Urgenza (AREU). They can apply as Partners or Coordinators of a Consortium.

5. Research Institutes - only in partnership with one of the organisations above (1,2,3,4). They can apply as Partners of a Consortium.

6. Universities - only in in partnership with one of the organisations above (1,2,3,4). They can apply only as Partners of a Consortium.

Enterprises and for-profit organisations are NOT eligible.

Eligible experimental approaches and disciplines:

Only translational and clinical research projects will be funded; fundamental research is not eligible. For definitions of basic, translational, and clinical research, please refer to the document published on the FRRB webpage dedicated to the OH AMR JTC 2027 call for proposals.

Eligibility criteria for the Principal Investigator:

A Principal Investigator (PI) cannot simultaneously hold more than one FRRB grant. PIs who are currently FRRB grant holders cannot apply to a new JTC unless their project is closed before the start of a new project. A project is considered closed when the final financial and scientific reports have been sent to FRRB.

This rule applies only to PIs, not to team members.

Personnel costs of PIs who have a permanent contract with their own organisation are NOT eligible.

Eligible costs:

Direct costs:

• Personnel (for public IRCCS and ASST, ATS and AREU, ONLY staff recruited specifically on the project).

Personnel costs of PIs who have a permanent contract (contratto indeterminato) with their own organisation are NOT eligible.

• Consumables, animals purchase, maintenance and breeding.

• Equipment (on hire or eligible amortization rate).

• Travel: max 10% of the total direct costs (overheads and subcontracting costs excluded)

• Publications (only open access): max 5% of the total direct costs (overheads and subcontracting costs

excluded).

• Other direct costs: please include here other costs, including those related to patient involvement (insurance, reimbursement, etc.).

• Subcontracting: max 20% of the total direct costs (overheads costs excluded)

Indirect costs:

• Overheads: 20% flat rate calculated on direct costs (Subcontracting costs excluded from this calculation).

FRRB will require the submission of a financial audit certificate together with the final financial report. This cost, to be included under the “Subcontracting” category will be eligible up to a maximum of € 8,000.00.

Only costs generated over the lifetime of the project will be considered eligible.

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Submission of the proposal (or other information) at the national level:

According to internal procedures, FRRB will run a check on the potential applicants prior to the submission of the pre-proposals.

The pre-eligibility check will be based on the verification of a dedicated form ("Pre-eligibility check form"), available on the FRRB dedicated webpage, to be completed and uploaded on the FRRB platform (https://bandi.frrb.it/login) by the Principal Investigator at least 10 working days before the pre-proposal submission deadline.

FRRB will provide feedback on the "Eligibility check form" ONLY in case of major non-eligibility issues.

In addition, FRRB provides an excel sheet to help applicants abide by FRRB funding rules. This form is meant to support the PIs in the elaboration of the proposal budget, but it does not need to be sent to FRRB.

Once the funded projects have been announced, FRRB will invite the PIs' host institutions of the projects approved for funding to sign a funding agreement.

Further guidance:

Administrative and Financial guidelines will be provided by FRRB on the website in due time

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Full name of the funding organisation:	**Ministero della Salute**
National/ Regional Financial Commitment:	**1 000 000 €**
Expected number of funded projects:	**2**
Contact details: (person (s), e-mail, telephone): Francesca Martorina, f.martorina@sanita.it, int-dgric@sanita.it , +39 06 5994 2240	

Maximum/ Minimum funding per grant Awarded to a project partner: 400.000 EUR. In case two eligible partners (IRCCS) are involved in the Consortium, the total amount must be shared between them. No more than two partners may receive funding under the same project.

Exchange Rate: NA

Eligible Call Topics: Topic 1, 2, and 3

Eligible One Health Settings:

Funding eligibility is limited to researchers working in clinical research from the IRCCS. However, projects may encompass activities across the different One Health domains — including Human Health, Animal Health, Plant Health, and the Environment — as long as they are relevant to the clinical research objectives.

Eligible institutions:

Only IRCCS (Istituti di Ricovero e Cura a Carattere Scientifico) researchers are eligible to apply.

Not eligible for funding: Universities, other research Institutes, companies.

Eligible experimental approaches and disciplines:

The MOH-IT can cover research teams working on clinical studies. These studies should be completed by the end of the project. Any requests for a no-cost extension cannot exceed a cumulative duration of 12 months.

Eligibility criteria for the Principal Investigator:

- A Principal Investigator (PI) cannot participate in more than one 2027 Joint Transnational Call (JTC) funded by the Italian Ministry of Health. If the PI is already the beneficiary of other active projects funded by the Italian Ministry of Health within European JTC schemes, the researcher is not allowed to submit a new application.

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- A PI holding an active IT-MoH grant may submit a proposal to a new JTC only if the ongoing project has been formally closed prior to the pre-proposal submission deadline. A project is deemed closed only upon submission and formal acceptance by the Italian Ministry of Health of both the final financial report and the final scientific report through the Workflow Research platform. If the ongoing project is expected to conclude within six months from the JTC pre-proposal deadline, the PI must submit a formal declaration stating that no request for project extension will be made.
- These eligibility restrictions apply exclusively to individuals serving as Principal Investigators. Co-investigators, collaborators, and other project team members who are not designated as PIs are not subject to these limitations.
- No more than two Italian PIs are eligible to apply for the same project.

Eligible costs:

Direct Costs:

- Personnel (only temporary contracts or permanent contracts for the number of hours dedicated to the project) Max. 60%.
- Consumables/Supplies.
- Equipment (only on leasing or rent).
- Travel (Max. 30%).
- Dissemination activities (Max. 1%).
- Publication costs: Max. 2%; Open Access Max 5%.

Other direct costs such as:

- Patients' recruitment costs.
- Animals/Model costs.
- Coordination costs only for Coordinator

Indirect Costs:

Overhead (Max.10%, of direct costs). Other indirect costs are not eligible. Transfer of eligible funds abroad is not allowed.

Subcontracts: (max 10%) they are allowed ONLY upon approval by Italian Ministry of Health presenting via Workflow research platform (ER communication code) a request together with the pre-eligibility check form, 20 working days before the pre-proposal submission deadline. Subcontracting cannot involve the core activities of the project.

Submission of the proposal (or other information) at the national level:

- Applicants are required to complete and submit a pre-eligibility check form to the Italian Ministry of Health using the Workflow Research platform (ER communication code), before submitting their pre-proposal to the Joint Call Secretariat (JCS). The completed form has to be submitted no later than 20 working days before the pre-proposal submission deadline.

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- If subcontracting is envisaged, it must be explicitly requested and pre-approved by the Italian Ministry of Health as part of the eligibility clearance process. Such requests must be submitted alongside the pre-eligibility check form.
- Changes to the project acronym or budget declared in the pre-submission eligibility check are not permitted.
- The mandatory pre-eligibility check form and the GDPR Article 13 Information Notice can be downloaded at the following link: https://www.salute.gov.it/new/it/tema/sistema-ricerca-del-ssn-enti-e-finanziamenti/internazionalizzazione-della-ricerca-biomedica/
- Applicants will receive written notifications of their eligibility status.
- Changes to the PI, the project acronym or budget declared in the pre-submission eligibility check are not permitted.

National phase:

- After completion of the JTC peer review process and endorsement of the final (scientific) ranking list by the Call Steering Committee, the Italian Ministry of Health will contact the PIs of the awarded projects for the necessary follow-up actions.
- Submission of final scientific and financial reports at Italian Ministry of Health will be required.

Further guidance:

Further information on the rules and guidelines of the Italian Ministry of Health can be found at https://www.salute.gov.it/new/it/tema/sistema-ricerca-del-ssn-enti-e-finanziamenti/internazionalizzazione-della-ricerca-biomedica/

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Latvia-LZP

Full name of the funding organisation:	**Latvijas Zinātnes padome**
National/ Regional Financial Commitment:	**600 000 €**
Expected number of funded projects:	**1-2**
Contact details: (person (s), e-mail, telephone):	**Laura Kunga-Jēgere, laura.kunga-jegere@lzp.gov.lv, +371 26102855**

Maximum/ Minimum funding per grant Awarded to a project partner: up to **100 000 EUR** per year (up to 300 000 EUR for one partner in one project). If there are two Latvian participants in one project, each can receive up to 100 000 EUR per year. No more than two Latvian partners may be in one project.

Exchange Rate: NA

Eligible Call Topics: Topic 1, 2, and 3

Eligible One Health Settings: Human Health, Animal Health, Environment and Plant Health

Eligible institutions: scientific organisations (universities, research institutions) and small, medium or large enterprises.

Universities and research organisations must be a legal entity based in Latvia and registered in the Register of Scientific Institutions.

Private legal entities (enterprises) must be a legal entity with no financial irregularities, based in Latvia and registered in the Register of Enterprises, and they must submit financial reports from the last two years.

Eligible experimental approaches and disciplines:

All approaches and disciplines are eligible, except LZP cannot cover clinical trials.

Eligibility criteria for the Principal Investigator:

No specific eligibility criteria.

Eligible costs:

1. Direct costs: personnel costs, travel costs, subcontracts (up to 25% from total direct costs), equipment (only depreciation costs), materials and other direct costs such as consumables, publication costs etc.
2. Indirect costs: can reach a maximum of 25% from total direct costs, excluding subcontracting costs.

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Double funding of the same project, as a whole or in part, is prohibited, whether the funding comes from LZP or from another source.

Scientific organisations can receive up to 100% national funding, enterprises – up to 80% state aid.

Submission of the proposal (or other information) at the national level:

Not necessary.

Further guidance:

Further information on the conditions for receiving funding can be found on the LZP website: www.lzp.gov.lv

The funding of RTD activities is provided pursuant in accordance with: - the Regulation of the Council of Ministers of the Republic of Latvia No 259 on the procedure for providing support for participation in international cooperation programs for research and technology (adopted on 26 June 2015), - the COMMISSION REGULATION (EU) No 651/2014 of 17 June 2014 declaring certain categories of aid compatible with the internal market in application of Articles 107 and 108 of the Treaty.

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Full name of the funding organisation:	**Lietuvos Mokslo Taryba**
National/ Regional Financial Commitment:	**300 000 €**
Expected number of funded projects:	**1-2**
Contact details: (person (s), e-mail, telephone):	**Živilė Ruželė, zivile.ruzele@lmt.lt, +370 676 14 383**

Maximum funding per grant Awarded to a project partner: 150 000 €

Maximum funding per grant Awarded to a project coordinator: 200 000 €

Maximum funding per project Awarded to two partners/coordinator and partner: 200 000/250 000 €

Exchange Rate: NA

Eligible Call Topics: Topic 1, 2, and 3

Eligible One Health Settings: Human Health, Animal Health, Environment and Plant Health

Eligible institutions: academic (research and study institutions), public health and care institutions (e.g. university hospitals)

Beneficiary academic institution must have a status of Lithuanian research and study institution and must be included in the Register of Education and Science Institutions.

Beneficiary institution (grant holder) manage the state budget funds allocated to the project following the rules stated in the legal acts, as well as representing the project partners (if applicable 'project partner' means public or private legal entity that, together with the eligible institution, created the conditions for project implementation).

A private legal entity registered in the Republic of Lithuania may be a project partner for a beneficiary institution. Research Council of Lithuania may allocate funding to this partner in accordance with Commission Regulation (EU) 2023/2831 of 13 December 2023 on the application of Articles 107 and 108 of the Treaty on the Functioning of the European Union to de minimis aid.

Eligible experimental approaches and disciplines:

All approaches and disciplines are eligible.

Eligibility criteria for the Principal Investigator:

Principal Investigator must be a PhD holder.

Principal investigators from Lithuania cannot be involved in more than 1 proposal submitted to this call.

Principal investigator may be involved in the EUP OHAMR grants while being already funded for another project at the national level.

Eligible costs:

Only costs generated during the lifetime of the project, related to the project activities, are eligible. Eligible cost types: personnel, consumables, subcontracting, equipment and instruments, other direct costs, costs for dissemination of results, data handling and analysis, overheads (up to 20 % from direct costs).

More details about eligibility of costs:

https://www.e-tar.lt/portal/lt/legalAct/0a8bead0577611e9975f9c35aedfe438/asr

Submission of the proposal (or other information) at the national level:

The submission of the proposal at the national level is not required.

Following funding decision, grant signing institution and the PI must complete and submit the national document (the template can be found following this link) containing this information: more detailed planed budget, foreseen dissemination and communication activities and expected outputs from project results with the granted research team contribution (scientific papers, patents, etc.)

Further guidance:

For any information, please refer to contact person. All information about the call is published on LMT website under Calls webpage.

General information for applicants submitting proposals to European Partnerships calls can be found here.

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Full name of the funding organisation:	**Xjenza Malta**
National/ Regional Financial Commitment:	**500 000 €**
Expected number of funded projects:	**1-2**
Contact details: (person (s), e-mail, telephone):	**Ms Ana Cuca ana.cuca@gov.mt/+356 23602322**
	Ms Luca Halasz luca.halasz@gov.mt/ +356 23602330
	Cc: eusubmissions.xjenzamalta@gov.mt

Maximum/ Minimum funding per grant Awarded to a project partner:

The maximum amount that national partner/s can jointly request per project is €500,000.

Exchange Rate: NA

Eligible Call Topics: Topic 1, 2, and 3

Eligible One Health Settings: Human Health, Animal Health, Plant Health and Environment

Eligible institutions:

1. Malta-based applicants that are Eligible Undertakings, with an Operating Base in Malta, planning to carry out Fundamental Research, Industrial Research and/or Experimental Development projects and must either be:
   a) A partnership constituted under the Companies Act, being a partnership en nom collectif, en commandite or a limited liability company; or
   b) Be duly registered as a co-operative society under the Co-Operative Societies Act, or
   c) Professional body; or
   d) NGOs; or
   e) Non-profit making entities (including Foundations).

will be eligible for funding subject to the terms and conditions laid out in the latest version of the National Rules for Participation (State Aid).

2. Any Public Entity or Public Research or Knowledge-Dissemination Organisation registered in Malta, that do not carry out an economic activity within the meaning of Article 107 TFEU, will be eligible for funding subject to the terms and conditions laid out in the latest version of the National Rules for Participation (Non-State Aid).

Eligible experimental approaches and disciplines: All approaches and disciplines.

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Eligibility criteria for the Principal Investigator:

Researchers, technicians, and supporting staff, to the extent that they are employed on the relevant project or activity by the eligible Malta-based entity. Further information can be found in the detailed National Rules accessible from the Xjenza Malta website: https://xjenzamalta.mt/media/funding-schemes/

Eligible costs:

Eligible costs and rates of funding depend on the type of the Malta-based entities and the funding route chosen (caps apply).

Eligible costs include the following: personnel; instruments, specialised equipment, and research consumables; IP and knowledge transfer activities; travel and subsistence; subcontracted activities; overheads and other operating expenses.

Further information can be found in the detailed National Rules accessible from the Xjenza Malta website: https://xjenzamalta.mt/media/funding-schemes/

Submission of the proposal (or other information) at the national level:

The national application form together with the required annexes can be downloaded from the Xjenza Malta website (https://xjenzamalta.mt/media/funding-schemes/) and must be sent to eusubmissions.xjenzamalta@gov.mt by the national deadline specified in the detailed National Rules.

Further guidance:

Further information can be found in the detailed National Rules accessible from the Xjenza Malta website: https://xjenzamalta.mt/media/funding-schemes/

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Full name of the funding organisation:	National Agency for Research and Development
National/ Regional Financial Commitment:	100 000 €
Expected number of funded projects:	1-2
Contact details: (person (s), e-mail, telephone):	
Mrs Natalia Bolocan, (373) 22 29 28 26, natalia.bolocan@ancd.gov.md	
Mrs Aurelia Hanganu, (373) 22 23 49 20, aurelia.hanganu@ancd.gov.md	
Ms Iana Tihon, (373) 22 27 27 91, iana.tihon@ancd.gov.md	

Maximum funding per grant Awarded to a project partner: 50 000 €

Exchange Rate: 1 EUR = 20,02 MDL (08/10/2026)

Eligible Call Topics: Topic 1,2 and 3

Eligible One Health Settings: All One Health settings: Human Health, Animal Health, Plants, and Environment

Eligible institutions:

- Beneficiary institutions eligible for funding must be public organizations that operate within the fields of research and innovation. This eligibility is in line with Articles 95 and 96 of the Science and Innovation Code of the Republic of Moldova No. 259/2004 and with points 3, 6, and 19 of Government Decision No. 382/2019, which outlines the Methodology for financing projects in these domains.
- Eligible institutions must not be declared insolvent or have their bank accounts blocked by an irrevocable court decision.
- Eligible institutions must have the legal right to carry out research and innovation activities as stipulated in its incorporation documents/statutes.
- Eligible institutions must not have violated any terms of previously signed contracts for research and innovation funding.

Eligible experimental approaches and disciplines: NARD can cover all eligible experimental approaches and disciplines, provided that all activities comply with applicable national legislation and regulatory requirements, e.g., the Law No. 15 of 19 June 2025 on Medicinal Products regarding clinical trials.

Eligibility criteria for the Principal Investigator:

- The Principal Investigator must hold a doctoral or habilitated doctoral degree in a relevant scientific field and be employed by a research institution within the Republic of Moldova.
- A person may not serve as Principal Investigator for more than one proposal within the same call, in order to prevent conflicts of interest and ensure equitable distribution of leadership responsibilities.
- The Principal Investigator's scientific experience must be relevant to the proposed project, as demonstrated by published works and other scientific activities carried out over the past 5-10 years.

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Eligible costs:

Applicants can apply for the following types of costs under their project proposals:

- **Personnel Costs:** This includes the remuneration of research team members, which must comply with public sector salary regulations and legislative frameworks. The remuneration of research team members must not exceed 70% of the total eligible costs. Salaries for administrative, financial, and technical staff may account for up to 5%.
- Travel costs must not exceed 10% of the total eligible costs.
- **Other Costs:** This includes various additional expenses, for example: participation fees for events, intangible assets (including patent acquisition expenses), procurement of raw materials, consumables (including reagents), animals, and laboratory inventory necessary for project experiments.

Non-eligible Costs:

- Costs related to the maintenance and development of public research infrastructure used in the project.

Expenses are incurred only after the contract has been signed and registered by the Contracting Authority. The budget must clearly detail the allocation of state funds and co-financing, with calculations supported by estimates. Proposals that do not meet these criteria, or that include costs for activities already funded or in the process of being funded from other sources, will be declared ineligible. Additionally, no modifications or improvements to proposals will be accepted after submission.

Submission of the proposal (or other information) at the national level:

The submission of the proposal at the national level is not requred.

Proposals are submitted in two stages:

- Pre-proposal by Monday 1st February 2027, and
- Full proposal by Wednesday 23rd June 2027

Further guidance:

For further information, please refer to the contact person and the website: ancd.gov.md.

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Full name of the funding organisation: Zorgonderzoek Nederland (ZonMw)
National/ Regional Financial Commitment: €2.500.000. The budget is made available by NWO (Dutch Research Council) and ZonMw.
Expected number of funded projects: 6
Contact details: (person (s), e-mail, telephone): Karen 't Hooft, Anouk Warmerdam E: AMR@zonmw.nl, T: +31 70 349 5001

Maximum/ Minimum funding per grant Awarded to a project partner:
Call topic 1: A maximum of € 350.000 per consortium with Dutch partner(s).
Call topic 3: A maximum of € 400.000 per consortium with Dutch partner(s).

Two Dutch partners in the consortium: max. € 350.000 (topic 1)/400.000 (topic 3) in total for the 2 partners together.

Exchange Rate: NA

Eligible Call Topics:
Call topics 1 and 3

Eligible One Health Settings: Indicate here if your funding organisation can fund all the different One Health settings (Human Health, Animal Health, Plants, Environment), or only a subset of them (i.e. the funding organisation X can only fun research groups working on Human Health through this call).
All One Health settings (Human Health, Animal Health, Plants, Environment)

Eligible institutions:
The following organisations based in the Netherlands are eligible for funding:

- Research organisations, that meet the definition of a research organisation as referred to in EU state aid legislation may receive the research grant (Framework for state aid for research and development and innovation (2014/C 198/01), i.e. the R&D&I framework. See: EUR-Lex - 52014XC0627(01) - EN - EUR-Lex

Note: To be eligible for ZonMw funding as a *research organisation* under the European **R&D&I Framework**, the legal status of the organisation must be established in advance to ensure that no unlawful state aid is granted. This is done through two routes*:
* *See fast-track procedure described below for organisations already recognised by ZonMw as a research organisation within the past 12 months (since January 2026).*

- **Route 1: Direct admission for Universities and University Medical Centres (UMCs)**
  The following organisations are exempt from the eligibility assessment and may submit a grant application directly:

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- Universities are recognised as research organisations by virtue of their status, in accordance with the R&D&I Framework.
- University Medical Centres (UMCs) confirm their status as research organisations by submitting a declaration. The declaration must be issued by, or on behalf of, an authorised signatory registered with the Chamber of Commerce (KvK). The declaration form is only required in the full proposal stage and can be submitted to kwalificatieOO@zonmw.nl with AMR@zonmw.nl in cc before the submission deadline of the full proposals.

- **Route 2: Eligibility Assessment for Other Organisations**
  All other organisations that wish to apply for funding as a research organisation, must submit the following documents to ZonMw prior to the pre-proposal stage:
  1. A fully completed ZonMw Research Organisation Eligibility Assessment Questionnaire (in Dutch).
  2. A digitally certified extract from the Dutch Chamber of Commerce (KvK) Trade Register, issued no more than three months prior to submission.
  3. The organisation's deed of incorporation and/or current articles of association.
  4. The most recent annual financial statements (preferably from 2025), including an auditor's report, where the organisation is legally required to prepare such statements.

  Applicants may also submit additional documents that are relevant to the assessment of their status as a research organisation. ZonMw may request further information if the documents provided do not offer sufficient clarity for the assessment. In that case, applicants will be given one additional two-week period to submit these documents.

  Deadline for route 2: as soon as possible but no later than **9 December 2026 at 14:00 CET** by email to **kwalificatieOO@zonmw.nl**, stating the title of this grant call (EUP OHAMR Joint transnational call 2027) in the subject line and with **AMR@zonmw.nl** in cc. For more information about the assessment, please visit the website or contact ZonMw via kwalificatieOO@zonmw.nl and AMR@zonmw.nl.

  Organisations will be notified of the outcome of the assessment within two weeks of receipt of all the documents.

*** Fast-track procedure**

Organisations recognised by ZonMw as research organisations within the past 12 months (since January 2026), where articles of association and activities remained unchanged, may follow the fast-track procedure. The organisation must confirm this by submitting the designated declaration to kwalificatieOO@zonmw.nl with AMR@zonmw.nl in cc before the submission deadline of the pre-proposals. A new eligibility assessment is not required. The outcome of the completed assessment must still be valid at the time the pre-proposal is submitted. Organisations unsure of their status can contact ZonMw via AMR@zonmw.nl before the submission deadline of the pre-proposals.

Consequences of inability to confirm status:

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If the documents are not submitted on time, or if ZonMw cannot positively confirm the organisation's status as a research organisation, the organisation cannot submit an application in the capacity of a research organisation.

Eligible experimental approaches and disciplines:
No particular conditions. ZonMw follows the requirements presented in the call text.

Eligibility criteria for the Principal Investigator:
No particular criteria, ZonMw follows the criteria presented in the call text.

Eligible costs:
Relevant project expenses:

- Salary-related costs according to most recent NFU or VSNU table. Note that PhD positions cannot be applied for in this call, due to the maximum project duration of 3 years.
- Travel costs
- Material/equipment and consumer goods
- Dissemination and knowledge exchange costs
- Data management/data steward
- Open access costs with a maximum of € 5,000 per project

For project-specific material costs up to 25% of the grant amount can be allocated. Subsequently, up to 50% of the material budget can be used for outsourcing. Otherwise, parties other than mentioned above can only participate as a non-funded partner, and need to secure their own funding in order to be involved in a project.

ZonMw does **not** cover overhead costs.

The budget of Dutch partners must comply with the ZonMw Grant Terms and Conditions: https://www.zonmw.nl/en/news-and-funding/funding/grant-conditions-and-finances/

State Aid
No funding will be awarded by ZonMw if this would or could constitute unlawful state aid (section 107 of the Treaty on the Functioning of the EU (TFEU). Funding awarded to research organisations[1] for the execution of non-economic research activities does not constitute state aid. Under this call, proposals may be submitted for projects involving fundamental research[2], industrial research[3], and/or experimental development[4].

Submission of the proposal (or other information) at the national level:
National submission
In case the joint proposal is recommended for funding, national applicants will have to submit a formal application through MijnZonMw. Selected applicants will be invited by email by ZonMw to submit their application. Awards will be subject to standard ZonMw Grant Terms and Conditions, including all applicable rights, conditions, and obligations. For more information, please visit the website or see the infographic.

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Obligations for ZonMw funded projects:

- Compliance with ZonMw procedures, including submission via MijnZonMw, use of the ZonMw budget format, and periodic reporting. More information and documents that must be submitted via 'MijnZonMw' can be found here.
- A Consortium Agreement, including intellectual property provisions, and a Data Management Plan are required. A signed copy of the Consortium Agreement must be submitted to ZonMw within 12 months after project starts.

Additional eligibility criteria

- An applicant may request funding for one project only (as part of an international consortium) in this call for proposals, either as main applicant or co-applicant within an application for ZonMw funding.
- Each application (i.e. the Dutch component of an international consortium) must have only one main applicant (i.e. the Dutch partner or coordinator), who is responsible for both the scientific and financial management to ZonMw.

Further guidance:

- ZonMw will avoid double funding and will not finance projects or part of projects that have been funded through other calls. ZonMw will cross-check the proposals submitted to ZonMw through the national and international calls for possible demands of double funding.
- All publications resulting from scientific research wholly or partly funded by ZonMw must immediately be made available in Open Access (without embargo), in accordance with the ZonMw Open Access policy. ZonMw accepts various Open Access routes. In addition to articles, ZonMw encourages recipients to make other types of scientific publications available in Open Access. Make sure to consult the ZonMw Open Access publication and Data management policies.
- For more information, please consult the national contact person by sending an email to AMR@zonmw.nl.

[1] As defined in EU state aid legislation: Framework for state aid for research and development and innovation (R&D&I Framework) (2014/C 198/01)
[2] Article 25, paragraph 2 (a) of the GBER
[3] Article 25, paragraph 2 (b) of the GBER
[4] Article 25, paragraph 2 (c) of the GBER

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Norway-RCN

Full name of the funding organisation:	**Research Council of Norway**
National/ Regional Financial Commitment:	**1 340 000 €**
Expected number of funded projects:	**2 -3**
Contact details: (person (s), e-mail, telephone):	**Hilde G. Nielsen, hgn@rcn.no, +47 40 92 22 60**

Maximum/ Minimum funding per grant Awarded to a project partner:

- Research Council of Norway do not operate with minimum funding per grant.
- Within a single project, the maximum funding is 350 000 €. If the Norwegian participant has a coordinator role, the maximum funding is about 450 000 €.
- If there are more than one Norwegian partners in a consortium they have to share the maximum RCN's funding per project between them.
- Depending on the volume of submitted and eligible projects, up to 25 % additional funding may be allocated to the call to fund additional projects on the ranking list.

Exchange Rate:

- For funded projects, the contractual budget will be in Norwegian kroner (NOK) using the exchange rate from the pre-proposal deadline. The official exchange rate can be found on the websites of the European Central Bank (https://www.ecb.europa.eu/stats/policy_and_exchange_rates/euro_reference_exchange_rates/html/eurofxref-graph-nok.en.html).

Eligible Call Topics: Topic 1, 2, and 3

Eligible One Health Settings:

- The entire value chain for research and innovation should be safeguarded.
- All One Health settings (human, animal, plan, environment, social sciences) related to human health are qualified.

Eligible institutions:

- Norwegian universities, university colleges, hospitals, research institutes, public sector, user organisations, NGOs, SME and other private industry.
- Organisations excluded from funding:
    - The Research Council cannot award support to an enterprise that is defined as an "undertaking in difficulty" under the state aid rules (see the "Definition of 'undertaking in difficulty'" on our website).
    - "Enkeltpersonforetak", that is Norwegian companies with sole proprietorship, cannot participate as coordinator, but can participate as subcontractors.

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- Additional eligibility criteria:
    - All applicants and partners must comply with the State Aid rules. All projects are to be carried out as effective collaboration between the partners. Undertakings (companies) that participate in the consortium must also not receive indirect state aid in the form of advantageous conditions for cooperation with the research institutions taking part in the consortium. SME or industry/large enterprise partners are funded with up to 50% of their eligible project costs. See *Conditions for awarding state aid* (https://www.forskningsradet.no/en/state-aid) for more details.
    - The Research Council of Norway does not award personal doctoral scholarships. Costs for doctoral (PhD) and postdoctoral fellowship positions must be included in the project application.
    - For postdoctoral fellowships, duration of the support is limited to a minimum of three years and a maximum of four years. The overhead cost is included in the rates for personnel.

Eligible experimental approaches and disciplines:

- The entire value chain for research and innovation should be safeguarded.
- All one health settings (human, animal, plan, environment, social sciences) related to human health are qualified.
- Clinical studies are eligible, please read the Requirements and guidelines for registration and disclosure of medical and health-related studies involving human participants.

Eligibility criteria for the Principal Investigator:

- Principal investigators must come from an approved Norwegian research organization: https://www.forskningsradet.no/en/financing/research-organisations/approved-research-organisations/ .

Eligible costs:

- Payroll expenses, consumables, operating expenses, network measures.
- The participation must follow RCN's General Terms and Conditions for R&D Projects.
- See *What to enter* in the project budget (https://www.forskningsradet.no/en/financing/how/budget). Note that the cost category "Procurement of R&D services" will not be used in this call.
- Funding to Norwegian SMEs and Industry will be provided according to the State aid rules. See Conditions for awarding state aid (https://www.forskningsradet.no/en/state-aid/) for more details.
- For funded projects, the contractual budget will be in Norwegian kroner (NOK) using the exchange rate from the pre-proposal deadline.

Submission of the proposal (or other information) at the national level:

- Data Management Plan must be in place to sign the contract with RCN for funded projects.

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- The Research Council is a driving force in making research as open as possible. Participation must follow RCN´s policy for Open Science.
- Submission of the proposal (or other information) at the national level: If the proposal is granted, information about national registration will be given.

Further guidance:

- Writing grant applications

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Poland-NCN

Full name of the funding organisation:	**Narodowe Centrum Nauki**
National/ Regional Financial Commitment:	**1 000 000 €**
Expected number of funded projects:	**3 - 4**
Contact details: (person (s), e-mail, telephone):	Alicja Dyląg, alicja.dylag@ncn.gov.pl, +48 532 086 494 Dr Anna Fiust, anna.fiust@ncn.gov.pl

Maximum/ Minimum funding per grant Awarded to a project partner: NA

Exchange Rate: 1 EUR = 4,2401 PLN

Eligible Call Topics: Topic 1, 2, and 3

Eligible One Health Settings: Human Health, Animal Health, Environment and Plant Health

Eligible institutions:

Proposals may be submitted by the entities specified in Article 27 (1) of the Act on the National Science Centre (NCN Act).

Eligible experimental approaches and disciplines:

All approaches and scientific disciplines are eligible.

Eligibility criteria for the Principal Investigator:

The principal investigator of the Polish research team shall hold at least a PhD degree when submitting the national proposal to NCN. Other eligibility criteria are described in Chapter IV of the UNISONO Resolution no. 76/2026 of the Council of National Science Centre, dated September 10, 2026.

Eligible costs:

Applicants may apply for funding of costs relevant, necessary, and directly connected to the proposed research project, including:

- **Salaries and scholarships** (including post-doc positions and scholarships/salaries for students and PhD students).

Please note: Limits apply depending on the size of the research team - please refer to the UNISONO Resolution. If the principal investigator is also the coordinator of an international consortium submitting a joint proposal the budget of salaries and scholarships may be increased.

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- **Research equipment, devices and software:** applicants may seek funding for research equipment up to PLN 500 000 per unit.
- **Other direct costs:** materials and consumables, outsourcing and subcontracting, business trips, visits and consultations, collective investigators, other costs;
- **Overheads:** Indirect costs of open access of up to 2% of direct costs that may be spent on open access to publications and research data; other indirect costs of up to 20% of direct costs that may be spent on costs that are related indirectly to the research project, including the cost of open access to publications and research data.

Submission of the proposal (or other information) at the national level:

At the pre-proposal stage Polish applicants are not required to submit any documents to NCN. At the full proposal stage, Polish applicants must submit their national proposals in the OSF submission system.

Further guidance:

For full details of NCN requirements, please refer to the OHAMR 2027 Call Announcement published on the NCN website.

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Portugal - FCT

Full name of the funding organisation: Fundação para a Ciência e a Tecnologia, I.P.

National/ Regional Financial Commitment: 400.000 €

Expected number of funded projects:2-4

Contact details: (person (s), e-mail, telephone):

Marta Abrantes (marta.abrantes@ai2portugal.pt**) +351 213911596**

Joana Pinheiro (joana.pinheiro@ai2portugal.pt**) +351 213911567**

Maximum funding per grant Awarded to a project partner:
The maximum amount of funding to be requested to FCT by a consortium with a **Portuguese Main Applicant** is **150 000,00 €**.
The maximum amount of funding to be requested to FCT by a consortium with a **Portuguese Project Applicant** is **100 000,00 €.**
If more than one Portuguese applicant participating in the **same international consortium** applies for funding by FCT, the **combined funding** demanded by all the Portuguese applicants **may not exceed the maximum financial threshold for proposals with a Portuguese Coordinator** (150 000,00€) or with a Portuguese **Partner** (100 000,00€). Portuguese Coordinator and/or Partners in the same international consortium will therefore have to share the funding that will be granted by FCT.
For information on funding rates, see no. 2 of article 7 of FCT Regulation.

Eligible Call Topics: All call topics are eligible.

Eligible One Health Settings: FCT may fund projects in any of the different One Health settings (Human Health, Animal Health, Plants, Environment).

Eligible institutions: For information on the type of beneficiaries eligible for FCT funding under this call, see article 3 of FCT Regulation. For information on the criteria of beneficiaries' eligibility, see article 5 of FCT Regulation.

Eligible experimental approaches and disciplines: Not applicable.

Eligibility criteria for the Principal Investigator: The transnational rules indicated in this call text apply.

Eligible costs: For the purposes of defining the budget, the terms defined in article 8 of FCT Regulation apply to eligible expenses and in article 9 to non-eligible expenses.
Excluded from the range of eligible expenses are the **salaries and other remuneration supplements** of teachers, researchers and other staff with a previously established indefinite contract with the Public Administration. Expenditure on **adapting buildings and facilities** is limited to a maximum of 10% of the project's total eligible expenses. The **project's indirect costs** are based on the application of a flat rate of 25% of the direct eligible costs.

Submission of the proposal (or other information) at the national level: **Statement of Commitment –** available on FCT's webpage dedicated to this call:

- Within 10 working days after the deadline for submitting the pre-proposal, a **Statement of Commitment** duly signed by the Researcher in Charge (partner and/or coordinators) and by the legal representant of the Portuguese Proposing Institution must be sent to the Contact Point.

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- The stamp or white seal of the Portuguese Proposing Institution will not be required on a digitally signed Statement of Commitment.

Portuguese applicants of transnational consortia that do not apply for funding from FCT do not need to submit the Statement of Commitment to FCT.

Further guidance:

Applications requesting funding from FCT under this call will be subject to Regulation on projects funded solely by national funds, published in Regulation No. 999/2016, in its current wording, that is, as amended and republished by Regulation No. 5/2024 of 3 January, and corrected by Rectification Statement No. 366/2024/2, published in the *Diário da República*, 2nd series, No. 100, of 23 May 2024, and by all other applicable national and European Union legislation.

For information on the criteria of projects' eligibility, see article 6 of FCT Regulation.

The percentage of time dedicated to transnational projects will **not** be added to the percentage of time dedicated to existing national projects.

Please consult FCT's webpage dedicated to this call.

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Full name of the funding organisation:	**Centrum Vedecko Technickych Informacii Slovenskej Republiky**
National/ Regional Financial Commitment:	**300 000 €**
Expected number of funded projects:	**1-2**
Contact details: (person (s), e-mail, telephone):	**Libuša Ďaďová, European Partnerships Coordinator** libusa.dadova@cvtisr.sk **+421 915 808 814** **Marek Babušiak, European Partnerships Coordinator** marek.babusiak@cvtisr.sk

Maximum/ Minimum funding per grant Awarded to a project partner:

The maximum funding amount is currently 300 000 EUR per Slovak partner. These amounts may be further increased by up to 15 000 EUR if a Slovak partner submits a patent, trademark, or design application, and/or publishes project results in an internationally indexed (peer-reviewed) scientific journal (e.g., WoS/Scopus).

Exchange Rate: NA

Eligible Call Topics: Topic 1, 2, and 3

Eligible One Health Settings: Human Health, Animal Health, Environment and Plant Health

Eligible institutions:

Legal entities established in the Slovak Republic, such as public or private research and academic institutions, higher education institutions, SMEs, public sector entities, and other relevant organizations actively involved in research, development, and innovation.

- Research institutions (e.g. the Slovak Academy of Sciences and its institutes)
- Academic sector (e.g. universities and higher education institutions)
- Public administration bodies and organizations established by them, including local and regional government authorities
- Non-governmental non-profit organizations
- Cluster organizations
- Private sector entities (entrepreneurial/business sector)

Eligible experimental approaches and disciplines:

Eligibility criteria for the Principal Investigator:

Eligible costs:

- Personnel costs (salaries of researchers, technicians and other support staff employed by the beneficiary, to the extent that they are directly involved in the project, salaries of project management personnel and other essential positions necessary for the implementation and coordination of the project;
- Costs of instruments and equipment;
- Costs for contract research, technical knowledge and patents purchased or licensed from external sources under market conditions, as well as costs for consultancy and equivalent services used exclusively for the project.
- All costs must be fully utilized and accounted for by the end of 2029.

General eligibility rule:

All expenditures incurred by Slovak project participants must comply with:

- Programme Slovakia, specifically Priority 1P1 Science, Research and Innovation, Specific objective RSO1.1: Development and enhancement of research and innovation capacities and the uptake of advanced technologies, Measure 1.1.3: Support for international cooperation in the field of research, development and innovation
- The provisions of the State Aid Scheme to Support Partnerships in the Field of Research, Development and Innovation under the Programme Slovakia;
- Strategy for Financing the ERDF, ESF+, CF, FST, and ENRAF 2021–2027.
-

Slovak applicants are advised to thoroughly familiarize themselves with the Guide for Applicants in European Co-funded Partnerships and Guide to Eligibility of Expenses under the Programme Slovakia, as the planned co-financing will be provided through this programme. It is further recommended that at least 30% of the Slovak applicant's total budget be dedicated to the actual performance of research activities.

Submission of the proposal (or other information) at the national level:

Submission of pre-proposal: Submit pre-proposals to the OHAMR Call Secretariat only.

Submission of full proposal: After having been informed about the international funding decision, CVTI SR will require also submission of separate application for national funding into the national submission platform.

Further guidance:

All Slovak applicants are strongly advised to contact the CVTI SR's contact points before submitting their proposals.

The proposed project activities must be in line with the priorities defined in the Research and Innovation Strategy for Smart Specialisation of the Slovak Republic 2021-2027 (SK RIS3 2021+), which serves as the strategic framework for research, development and innovation investments in Slovakia.

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All Slovak entities must have their contractual financial matters settled with CVTI SR by the end of 2029. Therefore, the project duration is limited to maximum 24 months.

Relevant national documents:

Programme Slovakia, Research and Innovation Strategy for Smart Specialisation of the Slovak Republic 2021-2027 (SK RIS3 2021+), State Aid Scheme to Support Partnerships in the Field of Research, Development and Innovation under the Programme Slovakia.

Useful links:
Programme Slovakia
SK RIS3 2021+
Strategy for Financing the ERDF, ESF+, CF, FST, and ENRAF 2021–2027

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Spain-IACS

Full name of the funding organisation:	Instituto Aragonés de ciencias de la Salud (IACS)
National/ Regional Financial Commitment:	150 000 €
Expected number of funded projects:	1-2
Contact details: (person (s), e-mail, telephone):	Elena Portero - eportero.iacs@aragon.es

Maximum/ Minimum funding per grant Awarded to a project partner:

No minimum funding per project.

Maximum 150.000€ participating as coordinator and 100.000€ participating as partner.

Exchange Rate: NA

Eligible Call Topics: All

Eligible One Health Settings: Human health, animal health and environmental health

Eligible institutions: Public Research institutions that carry out their activity in Aragon.

Eligible experimental approaches and disciplines: IACS will not fund clinical trials.

Eligibility criteria for the Principal Investigator: PI must be linked, through a civil servant, statutory or labour relationship, with the applicant institution for the whole project duration.

Coordinating PIs may only participate in this role in a single project in this call.

Eligible costs: Personnel costs related to personnel specifically hired for the project by the applicant institution, including salaries, employer Social Security contributions, legally established compensations and other duly justified expenses derived.

Travel, accommodation and subsistence costs directly incurred as a result of the research project, exclusively for people who are part of the project research team or hired under the funded project. Applicants must follow their respective internal travel policies.

Consumables, services costs and expenses related to the publication in open access and dissemination of project results, duly justified and necessary for the successful completion of the project.

Registration fees for congresses or conferences for the presentation and dissemination of the results.

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It is compulsory to include the cost of an external independent financial audit certificate up to a maximum of € 1.500

Indirect costs 15% of total direct costs

Submission of the proposal (or other information) at the national level: N.A.

Further guidance:

All members of the research team must be linked through a civil servant, statutory or labour relationship with the applicant institution.

More than one partner from Aragon may participate in the same project.

Double funding of the same concept is not allowed.

The projects must respect the fundamental principles established in national and international declarations, protocols and conventions on research ethics, as well as respect the requirements established in national and regional legislation in the field of biomedical research, development and innovation, personal data protection and bioethics.

Project funding will be subject to compliance with the relevant subsidy regulations and other applicable legislation at both the national and Aragon regional levels.

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Spain-AEI

Full name of the funding organisation:	**Agencia Estatal de Investigación**
National/ Regional Financial Commitment:	**800 000 €**
Expected number of funded projects:	**4-5**
Contact details: (person (s), e-mail, telephone):	**Ana Revilla; amr@aei.gob.es**

Maximum/ Minimum funding per grant Awarded to a project partner:

The following funding limits for a three-year project are considered eligibility criteria. Proposals not respecting these limits could be declared ineligible.

Two partners per project asking to the AEI are allowed only if one of them is the coordinator of the international proposal. The budget distribution in this case should be balanced.

This condition applies also if more partners are involved funded by other Spanish funders (ISCIII, IACS).

Maximum funding per project:

One AEI applicant (partner):

CD (€)*: 140.000

CI (25%) (€)*: 35.000

TOTAL (€)*: 175.000

One AEI applicant – (coordinator)

CD (€)*: 220.000

CI (25%) (€)*: 55.000

TOTAL (€)*: 275.000

One coordinator and one applicant- (balanced budgets)

CD (€)*: 260.000

CI (25%) (€)*: 65.000

TOTAL (€)*: 325.000

Additional funding for substantial experimental tasks (per project)

CD (€)*: 30.000

CI (25%) (€)*: 7.500

TOTAL (€)*: 37.500

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* The direct costs must be rounded to the thousands in the application. Indirect costs are 25% of direct costs requested.

Exchange Rate: NA

Eligible Call Topics: Topic 1, 2, and 3

Eligible One Health Settings: Human Health, Animal Health, Environment and Plant Health

Eligible institutions:

The AEI will fund non-profit research organizations (such as universities, research centres, technological centres and other private non-profit institutions performing RDI activities in Spain). They must have been previously beneficiaries of any of the AEI calls, and they must ensure contractual relationship with the Principal Investigator (PI) during the whole duration of the project.

Be aware that applicants from the Accredited Health Research Institutes (IIS), hospitals, primary health care or public health administration of the Spanish National Health System (SNS) and CIBER must apply for funding to the Instituto de Salud Carlos III (ISCIII), also participating in this call.

After the evaluation process and based on their budgetary availability and requested funding of selected projects, AEI and ISCIII reserve the right to exchange applicants to each other to optimize the available funds, provided the respective eligibility rules are met.

IMPORTANT: Spanish legal entities which are part of mixed centres will be considered as a unique beneficiary, and thus the maximum funding should not exceed the limits per proposal established above.

Although not fundable by the AEI, the private sector is encouraged to participate in consortia with Spanish academic groups, using their own funds or applying to national or regional calls by CDTI or regional innovation agencies. This is especially encouraged when there are for profit companies from other countries in the consortium or for projects coordinated by Spanish PIs.

Eligible experimental approaches and disciplines:

Only excellent proposals with exclusively RDI activities will be funded by the AEI. Entire communication work packages, without research associated, are not eligible costs for AEI.

The final funding will take into account the transnational evaluation of the collaborative proposal, the scientific quality of the Spanish group, the added value of the international collaboration and the financial resources available.

Clinical trials are eligible up to phase 1 (please contact the national contact point for further details).

Eligibility criteria for the Principal Investigator:

Eligibility criteria for PIs:

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PIs must be eligible according to the PCI general requirements 4Sep26 of the corresponding PCI call (read PCI2026_1 call as an example) and read carefully the "General Conditions Trans Calls 2026" in Oportunidades de Colaboración

Eligible costs:

- Please, **consult "*Artículo 8. Conceptos financiables*"**, especially paragraph ***a) "Costes de Personal"*** in PCI2026_1 call resolution as a reference, since eligible costs will be similar.
- Contracts for supporting personnel (gross remuneration and social security contributions) related to the implementation of the funded project are eligible costs. The contract must specify the percentage of the hired person's time dedicated to the project. The costs of permanent staff linked to the beneficiary entity or members of the research team will not be considered eligible costs.
- Direct costs such as current costs, disposable materials, travelling expenses, coordination costs, and other costs that can be justified as necessary to carry out the proposed activities are eligible.
- Indirect costs (overheads) are eligible costs (25% of total direct costs, including subcontracting).
- Subcontracting should not exceed 25% of total requested budget.
- Clinical trials are eligible up to phase 1, with a maximum of 50% of the total budget.

Submission of the proposal (or other information) at the national level:

Funding Programme:

The framework for this funding action is the "Plan Estatal de Investigación Científica, Técnica e Innovación 2024-2027". On a national level, the call will be managed by the "Subdirección de Programas Horizontales e Internacionalización" of the AEI.

Instrument for funding

The instrument for funding the Spanish groups is the call for "Proyectos de Colaboración Internacional (PCI)". Please, read carefully PCI2026_1 call resolution as an example.

Submission of the pre and full proposal at the national level:

PIs and beneficiaries are strongly encouraged to check eligibility before submitting a pre-proposal, since no changes will be accepted afterwords. No PI or beneficiary changes will be accepted between pre and full proposal as well as during the national call phase.

2) **Within one week after the pre-proposal submission** in the partnership website, the Spanish PI must ALSO submit a copy of the international joint pre-proposal at the AEI application website. The Spanish **PI** and the "**Representante Legal**" of her/his institution **must duly sign the application electronically. Failure to provide both required signatures will result in the pre-proposal being declared ineligible and excluded from the evaluation process.**
2) The "**Declaración Responsable**"***** document can be downloaded from Oportunidades de Colaboración website and must be uploaded through the AEI application website. This document is **mandatory** and constitutes an essential requirement for the proper processing of the application.

Applicants should include both the PI's full name (with both surnames, if applicable) and the **full legal name of their institution** as it is stated in SISEN – Sistema de registro de Entidades.

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Further guidance:

Submission of financial and scientific reports at the national level:

Financial and scientific reports are required as indicated in the national PCI2026_1 call resolution since will be similar.

Acknowledgement:

Any publication or dissemination activity resulting from the granted projects must acknowledge funding by the "Agencia Estatal de Investigación" according to AEI's web guidelines.

Beneficiaries are obliged by these requirements and those of the international call.

Data Protection:

By submitting a grant application, the applicants consent to communication of the data contained in the application to other public administrations, with the aim of further processing of the data for historical, statistical or scientific purposes, within the framework of the Organic Law 3/2018, of December 5, on Personal Data.

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Full name of the funding organisation:	**Instituto de Salud Carlos III**
National/ Regional Financial Commitment:	**500 000 €**
Expected number of funded projects:	**2-3**
Contact details: (person (s), e-mail, telephone): Astrid Valencia Quiñónez; ma.valencia@isciii.es	
+34 918222227	

Maximum/ Minimum funding per grant Awarded to a project partner:

Only one eligible beneficiary per consortium may apply for funding from ISCIII.

If this Spanish partner is not the Coordinator of the transnational project: 200.000 € (overheads included).

If this Spanish partner is the Coordinator of the transnational project: 275.000 € (overheads included).

Exchange Rate: NA

Eligible Call Topics: Topic 1, 2, and 3

Eligible One Health Settings: Human Health (eligible for funding). Animal Health (ISCIII can only fund research groups working on AH related to zoonotic diseases). Environment (eligible only if the research has a direct connection to human health)

Eligible institutions: ·Hospitals, primary health care centers, or public health administration bodies within the Spanish National Health System (SNS). These institutions may manage research activities through a foundation regulated under Spanish Act 50/2002, of December 26th. A copy of the foundation's statutes may be submitted, if applicable.

·Accredited Health Research Institutes (Institutos de Investigación Sanitaria acreditados, IIS). Institutes aaccredited under RD 279/2016. These institutions may manage research activities through a foundation regulated by Spanish Act 50/ 2002, of December 26th. A list of accredited IIS can be found at ISCIII website.

·CIBER. Only one Principal Investigator (PI) per consortium may be eligible by ISCIII. To meet the eligibility criteria, the research team must include members from at least two different CIBER groups based in separate home institutions. At least one of these two institutions must be either a hospital, a primary health care or a public health administration body within of the SNS or an IIS. For further information regarding CIBER´s eligibility, contact directly pai@ciberisciii.es

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Applicants not related with the National Health System from non-profit research organizations such as universities, OPIs, technological centres and other private non-profit institutions performing RDI activities in Spain need to apply for funding to the Agencia Estatal de Investigación (AEI), following their eligibility criteria. It is highly recommended to this type of members to integrate in their consortium other Spanish clinical partner (IIS/SNS/CIBER) eligible for funding by ISCIII.

Those declared by AES 2027 as ineligible to receive funds by ISCIII. Specifically for this call, the following entities are not eligible: National Technological Centres and National Centres for Supporting Technological Innovation. These are defined as those registered under Royal Decree 2093/2008, of 19 December

Please be aware that in 2027, some Institutions may be declared ineligible to receive funds from ISCIII under this call. Spanish applicants are advised to consult the ISCIII website and the National Contact Point of this call to confirm institutional eligibility.

The same beneficiary institution cannot participate as more than one Principal Investigator (PI) within the same project proposal.

Eligible experimental approaches and disciplines: Spanish research groups involved in the implementation of a clinical trial within the proposal are strongly recommended to include in their team a representative from their scientific node of the EU Clinical Trials Network (SCReN or ECRIN-ERIC). If such a node does not exist at their institution, it is recommended to include a staff member from their institution´s Clinical Research Supporting Platform (Unidad de Investigación Clinica- UIC).

Eligibility criteria for the Principal Investigator:

·PIs must hold a PhD degree. This is a mandatory requirement.

·Each PI may only participate in one project proposal per call.

·PIs affiliated with an IIS, may apply ONLY through the IIS.

·The PI and all research team must be affiliated with institutions that are eligible under the terms of this call.

·Only one PI per beneficiary institution may be funded within the same proposal.

·PIs currently coordinating or participation in an ongoing International Collaboration (PCIN) project funded by the JPI AMR, with an end date after the 31st of December 2027, are no eligible to apply for this call as PI. This incompatibility applies only to the PI, not to other team members. This restriction does not apply in the case that the PI participate as coordinator in the new application or in the ongoing project.

·Additional incompatibilities may apply. Please consult the AES 2027 for full details.

The following categories of researchers are not eligible to apply as PI:

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·Individuals undergoing postgraduate training in health specialization programs (e.g. MIR, EIR, FIR, QIR, BIR, PIR, RFIR)

·Individuals undergoing research training, such as PhD students or those with a “Río Hortega” contract.

·Individuals in postdoctoral training, such as those holding “Sara Borrell” or “Juan de la Cierva” contracts.

·Researchers contracted under RICORs or platforms funded by ISCIII.

Eligible costs:

-Personnel must be contracted under the appropriate professional category required for the development of the project (e.g. superior technician, BSc (Grado), MSc (Máster), or PhD (Doctor)). All contracts must be align with the official salary tables published on the ISCIII website/ AES 2027. No deviation from these salary tables will be accepted. Personnel cost must strictly to the defined amounts, neither higher and lower salaries will be eligible.

-Contracts for PhD students must be framed with the National Subprogramme for Training (scholarships are not eligible).

- Personnel costs will not be eligible when corresponding to contracts for PhD students and PhD (doctor) under the frame of Art. 21 and Art. 22 of Law 14/2011, of 1st June, following the specifications established in AES2027

-Personnel costs will be eligible for a maximum of 36 person-months in total, across all personnel contracts in the project.

-Contracts may cover the full duration or a specific part of the project period, depending on project needs.

-Personnel costs will not be eligible if they correspond to civil servants or equivalent personnel, as defined in Article 3.4 of the AES 2027 whether they are employed by the beneficiary institution or are members of the research team.

-Personnel costs will be eligible when corresponding to contracts under the frame of Art. 23bis of Law 14/2011, 1st June, following the specifications established in AES 2027.

Other eligible costs may include: current costs expenses, small scientific equipment, disposable materials, travel and accommodation expenses, complementary expenses (use of central and general research support services of the beneficiary institution) cost related to publication and dissemination of results and additional costs included in the AES 2027, provided they are clearly justify as necessary for carrying out the proposed activities.

• Overheads (indirect cost) are eligible up to a maximum of the 25% of the total direct costs, in accordance with the AES 2027.

• Double funding of the same concept is not allowed. Cost already covered by other grants or funding sources may not be charged to this project.

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Submission of the proposal (or other information) at the national level:

•Full proposal applicants selected must submit a national application to ISCIIII, in accordance with the timeline and procedures outlined in the AES 2027, except for those whose proposals have been declared ineligible for funding by Joint Call Secretariat prior to the national application deadline.

•Due to administrative and legal regulations, ISCIII has set 31 of October 2027 as the national deadline for the Call Steering Committee to decide on the fundable project consortia that include Spanish partners requesting funding from ISCIII. Any national applicant whose proposal is declared fundable after this date will be considered ineligible for funding from ISCIII.

Further guidance:

• Submission of a pre-eligibility form needed at national level

In order to expedite the eligibility check process, it is mandatory for all the applicants to submit the PI´s CVA-ISCIII. This document must be sent by the PI by via email before the pre-proposal submission deadline to the following address: ma.valencia@isciii.es. The email subject line must include the PI´s full name and the proposal acronym.

• Additional clause regarding available grant

After the evaluation process and based on their budgetary availability and requested funding of selected projects, AEI and ISCIII may exchange applicants to each other to optimize the available funds, provided the respective eligibility rules are met.

• Requirements on data and repositories

Researchers funded by ISCIII are required to make public the human genomic data, as well as relevant data (including phenotype and exposition data) generated within the funded project, by depositing them in open access repositories. In addition, researchers must provide all necessary information to enable the interpretation of these genomic data, including laboratory protocols, data instruments, and survey tools. For the purpose of this requirement, genomic data refers to the association of complete genomes (GWAS), single nucleotide polymorphism (SPN) matrices, genome sequences, as well as transcriptomic, metagenomic, epigenomic, and gene expression data. Researchers funded by ISCIII are recommended to store their scientific data at the “ELIXIR Core Data Resources". If non-European repositories or databases are used, they must be certified by ELIXIR.

ISCIII may no fund projects that require the creation of new repositories and/or databases if they do not include a decommissioning plan or clear strategy to ensure sustainability beyond the end of the funding period.

• Acknowledgements

Any publication, database, product, or event – whether protected by intellectual property rights (IPR)or not- resulting from the funded projects must be include the following acknowledge: “Award no. XX by ISCIII through the AES 2027 and within the EUP OHAMR framework”. This acknowledgment must be included even after the project has ended, and must also incorporate additional

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acknowledgments that may be specifically requested by ISCIII. For further information, please consult ISCIII's ROR

Links:

List of IIS: https://www.isciii.es/en/institutos-investigacion-sanitaria-presentacion

ELIXIR Core Data Resources: https://elixir-europe.org/platforms/data/core-data-resources

CVA-ISCIII: https://www.isciii.es/documents/20119/33967/Curriculum+Vitae+Normalizado+Abreviado+manual+AES_2023.pdf/1374cc8e-7657-4c95-8ab2-46f95a67aaca

ISCIII's ror: https://ror.org/00ca2c886

OH AMR

Sweden-SRC

Full name of the funding organisation:	**Vetenskapsrådet, Swedish Research Council**
National/ Regional Financial Commitment:	**15 000 000 SEK (approx. 1 385 000 Euro)**
Expected number of funded projects:	**3-5**
Contact details: (person (s), e-mail, telephone):	**Emmanouil Tsakoumis,** emmanouil.tsakoumis@vr.se **+46 (0) 73 302 13 54**

Maximum/ Minimum funding per grant awarded to a project partner:

Minimum 1 200 000 SEK (approx. 110 000 Euro) in total per Swedish partner in a project.

Maximum 3 500 000 SEK (approx. 321 000 Euro) in total per consortium for Swedish participation with maximum 2 Swedish partners.

Consortia coordinated by a Swedish organisation may request an additional maximum of 500 000 SEK (approx. 45 000 Euro) for the whole duration of the project.

Exchange Rate:

Use the exchange rate of 1 Euro=10,81 SEK to calculate actual grant amounts for the application.

Eligible Call Topics: All topics (Topic 1, Topic 2, Topic 3)

Eligible One Health Settings: All settings (Human Health, Animal Health, Plant Health and Environment)

Eligible institutions:

An administrating organisation approved by the Swedish Research Council (such as: higher education institutions (academic), regions, public agencies, research institutes and other organisations performing research).

Eligible experimental approaches and disciplines:

The Swedish Research Council can cover all disciplines.

Eligibility criteria for the Principal Investigator:

- The applicant must be an individual researcher holding a PhD. Only researchers at an administrating organisation approved by the Swedish Research Council may apply. Please refer to general applicant eligibility requirements found here.

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- The applicant should not have an ongoing JPIAMR or EUP OHAMR project grant, or any other project grant concerning the same project funded by the Swedish Research Council at the start of the grant period.
- All Swedish applicants are encouraged to communicate with the EUP OHAMR national contact person regarding their intention to participate in the Call, before submission of the consortium application.
- You can only take part in one consortium within this call, either as coordinator or partner.
- The number of Swedish partners requesting funding from the Swedish Research Council in one consortium could not be more than two (2).
- No funding of industrial partners.
- All Swedish project leaders participating in the Call for support from the Swedish Research Council must also submit a parallel application using the Swedish Research Council's application system, Prisma. The application form in Prisma can be reached from the national call text at the Swedish Research Council's website.

Parallel application is a mandatory eligibility criterion. Failure to submit the parallel application to the Swedish Research Council before the deadline of the Prisma call may result in the Swedish partner being declared ineligible.

Eligible costs:

The project grant may be used to fund all types of project-related costs, such as salaries (including your own salary, however no more than corresponding to the person's activity level in the project), running costs (such as consumables, travel including stays at research facilities, publication costs and minor equipment), premises and depreciation costs.

Grants may not be used for scholarships. If a doctoral student participates, project funds may not be paid out as salary during teaching or other departmental duties.

Submission of the proposal (or other information) at the national level:

A parallel application must be submitted in the Swedish Research Council's application system, Prisma. See above for more information.

Further guidance:

See national call texts in Swedish and English for all national requirements.

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Switzerland-SNSF

Full name of the funding organisation:	**Schweizerischer Nationalfonds zür Forderung der Wissenschaftlichen Forschung**
National/ Regional Financial Commitment:	**1'364'285 CHF**
Expected number of funded projects:	**4-5**
Contact details: (person (s), e-mail, telephone):	**Clémence Le Cornec**
	amr@snf.ch - +41 31 308 22 60

Maximum/ Minimum funding per grant Awarded to a project partner:

Minimum funding per award: CHF 100'000 / project

Maximum funding per award: CHF 300'000 / project

Exchange Rate:

Please use the monthly average exchange rate from the European Central Bank (Swiss franc (CHF)). Please note that the budget submitted on the SNSF Portal must be in CHF. You only need to convert to EUR for the budget submitted to the OH AMR call platform.

Eligible Call Topics: Topic 1, 2 and 3

Eligible One Health Settings: Human Health, Animal Health, Plant Health and Environment

Eligible institutions:

The SNSF exclusively funds research conducted for purposes that are not directly commercial. Pursuant to the Research and Innovation Promotion Act RIPA and the legal framework of the SNSF, no research grants are awarded if the relevant research is conducted for directly commercial purposes or if the persons involved in the research work do not enjoy scientific independence. The SNSF only funds research conducted within eligible Swiss-based non-commercial research institutions. Commercial institutions are not eligible.

Eligible experimental approaches and disciplines: All

Eligibility criteria for the Principal Investigator:

Applicants must comply with the SNSF Funding Regulations and the SNSF Regulations on project funding.

Participation of Swiss-based partners requesting financial support from the SNSF is restricted to one project (Art.6.3, SNSF Regulations on project funding). They may, however, participate in other consortia projects as self-financed partners.

Proposals with overlapping funding periods with ongoing SNSF projects are only approved if the research projects pursue different goals (Article 17 of the SNSF Funding Regulations).

Swiss-based investigators are not permitted to join research consortia as full partners at the stage of full-proposal submission. However, replacement of an existing Swiss-based, full partner may be permitted if duly justified. Swiss-based investigators joining as self-funded partner at full-proposal submission stage is permitted.

Eligible costs:

Eligible costs are outlined in the SNSF Funding Regulations (Art. 28) and the SNSF General Implementation Regulations (Section 2).

Project overhead costs cannot be applied for. They are calculated on the basis of the research funding acquired by eligible institutions under eligible funding schemes. Overhead contributions are paid in retrospect at a flat rate to the institutions of the SNSF awardees.

Submission of the proposal (or other information) at the national level:

Mandatory, parallel submission of pre- and full-proposal via the SNSF Portal

Swiss-based full consortium partners who apply for SNSF funding to participate in this Call must submit pre-proposals and full proposals **via the SNSF Portal** at the same submission deadline of the consortium application. These submissions are mandatory and do not replace the submission of the consortium application to the Call Secretariat.

Pre-proposal forms are created by selecting "European Partnership: OH AMR:". In case the pre-proposal is invited to a full proposal, a notification will be received by email. The original pre-proposal application will be accessible again for adjustments. Once all the necessary updates are made, the **Full-proposal** can be submitted.

In case of multiple, Swiss-based partners participating in the same consortium, only one application is to be submitted on the SNSF Portal, whereby one Swiss-based partner must act as "corresponding applicant" and the other Swiss-based partners are to be listed as "other applicants".

Self-funded partners and international partners of the consortium applying for funding at different funding agencies from the SNSF and cannot be declared as "project partners" in the sense of article 11.2 of the SNSF Funding Regulations. For the submission via the SNSF Portal, they are to be declared as "consortium members" under the tab research plan instead and must apply for their funding at their respective research funding organisations.

Further guidance:

Data management plan

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Applicants will have to complete the DMP on *my*SNF once the project is approved, regardless of whether a DMP is requested by the consortium. The DMP has to cover the research data, which are collected, observed, generated or reused in the Swiss part of the project and has to comply with the SNSF Open Research Data Policy.

Grant management

Grants will be managed according to standard SNSF rules described in SNSF Funding Regulations. Yearly financial reports for the use of SNSF funds must be submitted via *mySNF.* As a final scientific report, the SNSF requests the submission of the final scientific report submitted to the OH AMR Call Secretariat. No other scientific report is requested.

Information available at:

- SNSF Funding regulations
- General Implementation Regulations
- SNSF Regulations on Project Funding

Please contact the SNSF administrative offices should anything be unclear.

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Türkiye-TUBITAK

Full name of the funding organisation:	**Turkiye Bilimsel ve Teknolojik Arastirma Kurumu**
National/ Regional Financial Commitment:	**500 000 €**
Expected number of funded projects:	**3-4**
Contact details: (person (s), e-mail, telephone):	
	Damla AVCI **damla.avci@tubitak.gov.tr; +90 312 298 12 92**
	Gurkan BOZKURT
	gurkan.bozkurt@tubitak.gov.tr; +90 312 298 11 60

Maximum/ Minimum funding per grant Awarded to a project partner:

- **Max. funding per partner (Universities, Training and research hospitals, Public organizations and institutions):** 125.000 €
- •**Max. funding per private organization partner:** 200.000 €

Exchange Rate: When calculating the equivalent in foreign currency of expenses denominated in Turkish Lira (TRY), the Central Bank of the Republic of Türkiye (CBRT) effective foreign exchange selling rate applicable on the relevant application date shall be used as the basis. The amounts stated in budget forms completed in different currencies during international project applications, as well as changes in exchange rates, shall not affect the budget item-based limits specified in this document in Turkish Lira. Projects approved for funding shall be implemented taking into account the limits specified in this document.

Eligible Call Topics: Topic 1, 2, and 3

Eligible One Health Settings: Human Health, Animal Health, Environment and Plant Health

Eligible institutions:

- Universities
- Training and research hospitals
- Public organizations and institutions
- Private Organizations

*Applications submitted by foundations, associations and their economic enterprises, cooperatives, unions, sole proprietorships, and ordinary partnerships are not eligible.

Eligible experimental approaches and disciplines:

All research areas are eligible for support, except for clinical trials.

Eligibility criteria for the Principal Investigator:

- If employed by a higher education institution or a training and research hospital, they must hold a doctoral degree (PhD) or an equivalent qualification.

- If employed by research infrastructures operating under Law No. 6550, other public institutions and organizations, or private organizations, a minimum of a four-year undergraduate university degree is sufficient.

- They must be permanent/full-time employees of the institution where the project will be carried out (including higher education institutions, training and research hospitals, research infrastructures operating under Law No. 6550, other public institutions and organizations, or private organizations). Retired individuals and postdoctoral researchers who are not employed by any institution or organization are exempt from this requirement.

Eligible costs:

TÜBİTAK 1071 Program** rules are eligible.

Submission of the proposal (or other information) at the national level: Yes

For the international pre-proposal stage, National application procedure is applied as well.

Further guidance:

https://tubitak.gov.tr/en

https://tubitak.gov.tr/sites/default/files/2026-03/1071_Usul_Ve_Esaslari_79_sayili_YK_Islenmis_Hali.pdf

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OHAMR

United-Kingdom-MRC-UKRI

Full name of the funding organisation:	**Medical Research Council - United Kingdom Research and Innovation (MRC - UKRI)**
National/ Regional Financial Commitment:	**Subject to conditions of eligibility and peer review being fully met £1 million will be available to UK researchers.**
Expected number of funded projects:	**3-4**
Contact details: (person (s), e-mail, telephone):	**Kathryn Davidson Infectionsandimmunity@mrc.ukri.org**

Maximum/ Minimum funding per grant Awarded to a project partner:

Individual consortia may request a maximum of 345 000 € (£300,000) of funding from UK government (MRC-UKRI and DHSC) per application - funding requests can be split as appropriate across the remits of MRC-UKRI and DHSC but must not exceed 345 000 € (£300,000) in total; each application should clearly articulate which aspects of the funding request are and how much funding is requested targeted to each funder. This maximum amount refers to the 80% fEC value and not the 100% fEC value.

Exchange Rate: £1: €1.15

Eligible Call Topics: Topic 1

Eligible One Health Settings:

MRC, part of UK Research and Innovation (UKRI), are able to support all research areas that fall within the remit of MRC. For Topic 1, MRC-UKRI funding should contribute to novel understanding of AMR emergence or transmission relevant to human health. **Potential applicants are strongly advised to contact the MRC National contact (Infectionsandimmunity@mrc.ukri.org), in advance of making an application, to resolve any eligibility queries.**

Eligible institutions:

Applicants must be eligible to receive UKRI research funding and this opportunity is open to organisations following standard eligibility rules with one exception as below; details of eligibility for MRC funding can be found on the UKRI website Organisation eligibility – UKRI. Please note that industrial partners may **not** request costs from MRC-UKRI. Please note that any organisations that are using the UKRI Horizon Europe Participant Identification Code (PIC, which is 906446474) are **not eligible for funding** in this OHAMR call.

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Eligible experimental approaches and disciplines:

All work must fall within the specific remit of MRC – work primarily within the remit of the other UKRI councils will **not be supported** and **applicants are strongly encouraged to contact the MRC prior to submission to confirm remit and suitability**.

All experimental approaches, including *in vitro*, *in silico* and *in vivo* pre-clinical are eligible for funding.

Eligibility criteria for the Principal Investigator:

Potential applicants are strongly advised to contact the Call Secretariat, or the MRC-UKRI contact, in advance of making an application, to resolve any eligibility queries.

For the purposes of this call, a 'partner' requesting funding from MRC-UKRI is a legal entity. Multiple researchers from the same legal entity form a single partner, but only one of these researchers should be named on the OHAMR application form.

We welcome collaboration with industrial or other non-academic partners where appropriate; however, costs incurred directly or otherwise by these partners cannot be met by MRC and these partners cannot claim funding except through specific subcontracting arrangements (see https://www.ukri.org/publications/mrc-guidance-for-applicants/application-and-policy-guide-for-the-ukri-funding-service/). Similarly, as part of this transnational call, MRC-UKRI cannot meet costs incurred by international partners – such work must be costed directly to the relevant national funder.

Eligible costs:

- The MRC-UKRI component of applications should use full economic costings (fEC). The total amount requested must be the 80% fEC value. The submitting organisations must agree to find the balance of fEC for the project from other resources. In the 'financial plan' section of the OHAMR application form, the 'sum requested' is the 80% fEC amount. The 'total' is the 100% fEC amount plus any further resources or in kind contributions.
-
- Eligible costs include project-related costs incurred after the award start date, including:
  - Project leads and Project co-Leads time
  - Research and technical staff
  - Estates and Indirect costs
  - Animal costs
  - Travel and Subsistence
  - Equipment
  - Consumables
  - Recruitment and advertising costs for staff directly employed on the project
  - Costs related to research data management
  - NHS Research costs.
-
- Ineligible costs include:
  - International costs
  - NHS support and NHS treatment costs
  - PhD students
  - Publication costs, including open access publication costs.
  - Industrial partners
  -

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- For more information regarding eligible costs, please see the MRC Guidance for Applicants.

Submission of the proposal (or other information) at the national level:

- As well as the OHAMR application form, MRC-UKRI applicants must also complete a UK Budget Proforma at both the pre-proposal and full-proposal stage (please email Infectionsandimmunity@mrc.ukri.org to receive a form).
    - Costs should be included in pounds sterling (GBP) on the UK budget proforma (which can be downloaded from the call entry in the Funding Finder on UKRI's website) and included on the OHAMR application form in Euros using an exchange rate of £1:€1.15.
    - Applicants should include a statement on the UK budget proforma to confirm the exchange rate used, and that costs are entered at 80% fEC according to standard Research Council funding policy.
    - Applicants should include separate proformas for costs requested from MRC-UKRI and from DHSC.
- Applications that incur excess treatment costs for studies involving patients will be required to complete a SoECAT form if invited for a full application. Please see the MRC Guidance for Applicants for further information.

Please submit these additional forms separately to MRC-UKRI by emailing Infectionsandimmunity@mrc.ukri.org (including the consortium acronym and number); do not add them to the OH AMR consortium application submissions.

Please note successful UK partners in transnational consortia will be required to upload a single application to UKRI within one month of the notice of award. This is a purely administrative step and will not be peer-reviewed again; this must exactly replicate the UKRI component of the OHAMR application. A single UKRI award will be issued to all UK partners within a consortium, who will be responsible for disseminating the funding as per the application. The UK partners must therefore identify which researcher is the lead UK applicant, who will be responsible for submitting the UKRI application and whose institution will be responsible for disbursing UKRI funds to any other UKRI-funded partners.

Awards are subject to UKRI Terms and Conditions for funding. Award letters will include any additional terms and conditions specific to the call.

Further guidance:

For further details please refer to: MRC guidance for applicants – UKRI

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OHAMR

United-Kingdom- DHSC

Full name of the funding organisation:	**Department of Health and Social Care (DHSC)**
National/ Regional Financial Commitment:	**Subject to conditions of eligibility and peer review being fully met £0.5 million will be available to UK researchers.**
Expected number of funded projects:	**1-2**
Contact details: (person (s), e-mail, telephone):	**Kathryn Davidson** Infectionsandimmunity@mrc.ukri.org
Please note that DHSC funds will be administered by MRC-UKRI	

Maximum/ Minimum funding per grant Awarded to a project partner:

Individual consortia may request a maximum of €345,000 (£300,000) of funding from UK government (MRC-UKRI and DHSC) per application – funding requests can be split as appropriate across the remits of MRC-UKRI and DHSC and each application should clearly articulate which aspects of the funding request are and how much funding is requested targeted to each funder. This maximum amount refers to the 80% fEC value and not the 100% fEC value.

Exchange Rate: £1: €1.15

Eligible Call Topics: Topic 1

Eligible One Health Settings:

DHSC are able to support all research areas that fall within the remit of the National Institute for Health and Care Research (NIHR). Work funded through DHSC should generate evidence to support public health, healthcare or policy decisions aimed at reducing the burden of AMR, including evaluation of interventions, stewardship approaches, infection prevention and control measures, or other actionable measures to reduce AMR transmission.

Potential applicants are strongly advised to contact the National contact (Infectionsandimmunity@mrc.ukri.org), in advance of making an application, to resolve any eligibility queries.

Eligible institutions:

MRC-UKRI are administering funds on behalf of DHSC. Applicants must be eligible to receive UKRI research funding and this opportunity is open to organisations following standard eligibility rules with one exception as below; details of eligibility for MRC funding can be found on the UKRI website Organisation eligibility – UKRI. Please note that industrial partners may **not** request costs from MRC-UKRI. Please note that any organisations that are using the UKRI Horizon Europe Participant Identification Code (PIC, which is 906446474) are **not eligible for funding** in this OHAMR call.

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Eligible experimental approaches and disciplines:

All work must fall within the specific remit of the NIHR. Applicants are strongly encouraged to contact the national contact prior to submission to confirm remit and suitability. Please note that DHSC funding cannot be requested for research involving animals or animal tissues; all DHSC funding must focus on human tissues, participants or data with a clear trajectory towards patient benefit.

Eligibility criteria for the Principal Investigator:

Awards will be made through MRC-UKRI on behalf of DHSC. Potential applicants are strongly advised to contact the Call Secretariat, or the national contact, in advance of making an application, to resolve any eligibility queries.

For the purposes of this call, a 'partner' requesting funding from DHSC is a legal entity. Multiple researchers from the same legal entity form a single partner, but only one of these researchers should be named on the OHAMR application form.

Applicants who intend to collaborate with industrial or other non-academic partners should note that any costs incurred, direct or otherwise, by these partners cannot be met by DHSC and that these partners cannot claim funding.

Eligible costs:

- The DHSC component of applications should use full economic costings (fEC). The total amount requested must be the 80% fEC value. The submitting organisations must agree to find the balance of fEC for the project from other resources. In the 'financial plan' section of the OHAMR application form, the 'sum requested' is the 80% fEC amount. The 'total' is the 100% fEC amount plus any further resources or in kind contributions.
-
- Eligible costs include project-related costs incurred after the award start date, including:
  - Project leads and Project co-Leads time
  - Research and technical staff
  - Estates and Indirect costs
  - Travel and Subsistence
  - Equipment
  - Consumables
  - Recruitment and advertising costs for staff directly employed on the project
  - Costs related to research data management
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-
- Ineligible costs include:
  - International costs
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- For more information regarding eligible costs, please see the MRC Guidance for Applicants.

Submission of the proposal (or other information) at the national level:

- As well as the OHAMR application form, DHSC applicants must also complete a UK Budget Proforma at both the pre-application and full application stage (please email Infectionsandimmunity@mrc.ukri.org to receive a form).
    - Costs should be included in pounds sterling (GBP) on the UK budget proforma (which can be downloaded from the call entry in the Funding Finder on UKRI's website) and included on the OHAMR application form in Euros using an exchange rate of £1:€1.15.
    - Applicants should include a statement on the UK budget proforma to confirm the exchange rate used, and that costs are entered at 80% fEC according to standard Research Council funding policy.
    - Applicants should include separate proformas for costs requested from DHSC and from MRC-UKRI.
- Applications that incur excess treatment costs for studies involving patients will be required to complete a SoECAT form if invited for a full application. Please see the MRC Guidance for Applicants for further information.

Please submit these additional forms separately to MRC-UKRI by emailing Infectionsandimmunity@mrc.ukri.org (including the consortium acronym and number); do not add them to the OH AMR consortium application submissions.

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Awards are subject to UKRI Terms and Conditions for funding. Award letters will include any additional terms and conditions specific to the call.

Further guidance:

For further details please refer to: MRC guidance for applicants – UKRI

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Responsibilities of the Grantees JTC 2027

1- Granting Arrangements

Partners from the projects approved for funding will subsequently enter into granting arrangements with the relevant funding organisations, according to their applicable grant awarding process and will be funded directly by the respective Funding organisations. Partners from the projects approved for funding must fix **a common scientific project start date**, **which will be the reference date for the progress reports**. Projects are expected to start **between December 2027 and April 2028**.

2- Intellectual Property

The ultimate goal of the EUP OHAMR is to bring together national research efforts in order to make better use of public Research and Development (R&D) resources and to tackle common global challenges more effectively in selected key areas.

For EUP OHAMR activities to contribute effectively to socioeconomic progress, the results of the research activities must be exploited. This requires appropriate identification and protection of the intellectual property being generated and effective knowledge transfer. Any particular protection and exploitation strategy should be agreed before the research activities start. The ten principles of Socially Responsible Licensing should be part of this strategy (https://research-and-innovation.ec.europa.eu/research-area/industrial-research-and-innovation/eu-valorisation-policy/knowledge-valorisation-platform/thematic-focus/socially-responsible-licensing-tool-increase-societal-value-creation-ri_en).

Depending on the nature of the research and on the interests of the different parties, if there are opportunities for exploitation, it is recommended that parties decide in advance on either adopting a common exploitation strategy or leaving exploitation of results to the party best placed to commercialise it, with appropriate compensation mechanisms for the contributing parties. Please see section below for a link to a simplified consortium agreement template, available on the DESCA website. National rules and regulations may apply, please consult the national rules.

3- Consortium agreement

The partners of each funded project **are required to set up and sign a Project Consortium Agreement** (PCA) in order to deal with issues related to the role, tasks and responsibilities within the consortium, the protection of intellectual property, and where applicable how the consortium will address the ten principles of Socially Responsible Licensing. The coordinator is responsible for providing the PCA signed by all partners to the EUP OHAMR secretariat (reporting@ohamr.eu) when requested during the mid-term progress reporting. Upon request, this consortium agreement must be made available to the concerned funding organisations. The project consortium is strongly encouraged to sign the PCA before the official project start date and, in any case, the PCA should be signed **no later than six months after the scientific project start date**. The PCA needs to be in accordance with the national funding rules of the respective funding partner organisations - see national rules. Please note that certain funding organisations may need the signed PCA to release the funds.

The PCA must address (as a minimum), the following points:

- common start date and duration of the research project and the duration of the PCA;

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- organisation and management of the project;
- role, tasks, and responsibilities of each partner;
- the resources and funding;
- confidentiality and publishing;
- Intellectual Property Rights (if applicable);
- how the ten principles of Socially Responsible Licensing will be addressed (if applicable);
- decision making within the consortium;
- handling of internal disputes;
- the liabilities of the research partners towards one another (including the handling of default of contract).

Any issues regarding funding are a bilateral matter between each project partner and the relevant funding organisation and should be excluded from the PCA.

Please see the DESCA website for further information and templates of a simplified consortium agreement under the Horizon Europe Framework.

4- Open Science

EUP OHAMR requires grant holders to make data and research findings resulting from their project available following the principle "as open as possible, as closed as necessary". To this end, EUP OHAMR supports research consortia in implementing Open Science practices, focusing on data management according to FAIR principles (Findable, Accessible, Interoperable, Re-usable), and sharing research findings via open access publications. The open science requirements for EUP OHAMR funded projects are in line with the guidelines for research data management of Horizon Europe.

Data Management

Data management is mandatory for EUP OHAMR funded projects. It involves first that applicants are strongly encouraged to look for options for re-using existing data, standards, tools, and infrastructures, particularly those familiar to the AMR research community, to enhance interoperability.

Secondly, applicants must ensure that they plan the appropriate activities for their project, enabling them to make data available as soon as possible and re-usable for verifying results, and future research and innovation. Their plan must include the necessary expertise on data stewardship[1] and sufficient budget to be able to manage research data, in line with the FAIR principles. Grant holders will receive more details on the requirements later in the process.

Note that all requirements and guidelines referring to 'data', also apply to other resources that are used to perform the research, and are needed to re-use or validate the data and published research findings. These resources include for instance physical research outputs (such as biological samples, molecular derivatives), research software, (meta)data standards and tools.

Capacity and community building

The EUP OHAMR plans to organize online-awareness and -training workshops for granted projects. These activities will be organized throughout the course of the partnership. Attendance (on-line) to the workshops will be mandatory for the coordinators and partners of the funded projects (FAIR awareness) and for the project's data stewards and/or other members of the project group who are involved in data-related tasks (data FAIRification).

[1] Data stewardship is the responsible planning and executing of all actions on digital data before, during and after a research project, with the aim of optimising the usability, reusability and reproducibility of the resulting data.

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The aim is to improve knowledge and skills in creating re-usable and machine-actionable (i.e. FAIR) data, and to build an OHAMR-community of data-experts. Funded projects benefit from the workshops as they lead to a common understanding on data FAIRification and its benefits in research, an equal playing field for researchers, and preventing the reinvention of the wheel by individual research groups.

The EUP OHAMR as a whole, benefits as the workshops facilitate the research community to profit optimally from reusable data and emerging AI-approaches to advance knowledge generation on OHAMR.

What needs to be done in the application phase?

Applicants must check the requirements for data management and data sharing of the relevant national funder. EUP-OHAMR specific requirements will be provided later in the process, when grants are awarded.

Applicants do not need to deliver a data management plan (DMP) yet. However, already in this phase, they must plan carefully what is needed for re-using existing data (including the permission required therefore), and for managing data (and other resources) to become findable and re-usable for future users. The application must particularly include:

- A plan to involve experts on data stewardship, who are facilitated to dedicate their expertise to the project to realise data re-usability. The data-experts must also be able to participate in workshops aimed at building FAIR data-expertise and community engagement.
- Sufficient budget to enable the data-experts to be involved in the project.
- Sufficient budget to make use of tools, services and infrastructures to realise planned and required data re-usability. Be aware of the requirement to provide access to research data in trusted repositories.

What needs to be done once the project is funded?

Once the project is funded, a DMP must be developed. The Project Coordinator must prepare a DMP **no later than six months** after the scientific project start date. The project coordinator is responsible for sending an updated DMP with the midterm and the final term progress reports of the project to the EUP OHAMR secretariat (reporting@ohamr.eu). The Horizon Europe template for DMP should be used. Grant holders will receive further information, when grants are awarded.

Finalising data management: At the end of the project, research data and other outputs, software and other tools or instruments necessary to validate the publications' conclusions must be made available by depositing in a trusted repository.
The final DMP must be completed and delivered, including information on how resources can be found, where they are stored, and access conditions.

Open Access Publications

Beneficiaries must disseminate results, including scientific publications, to the public as soon as possible. Peer-reviewed scientific publications must be made available in open access. Guidelines and options to comply to the requirements are available on How to comply with Horizon Europe mandate for publications.

Peer-reviewed publications issued from EUP OHAMR funded projects must be open access by depositing the final version or peer-reviewed manuscripts in a trusted repository. For journal articles,

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a Creative Commons Attribution (CC BY) or equivalent open license should be requested to the editor. For publishing long-texts, Creative Commons Attribution Non-Commercial/Non-Derivatives licenses are also allowed. In addition, the research outputs, tools, or instruments necessary to validate the publications' conclusions should be deposited in a trusted repository.

Each participant may also be required to comply with the Open Access policy of its funding organisation (See country-specific information in national rules). To find out if a scientific journal complies with open access, check on the Directory of Open Access Journals or the Journal Checker tool. All research projects funded by EUP OHAMR are eligible to publish on Open Research Europe (ORE), the Platform of the EC at no cost.

5- Project Monitoring and reporting

Overall project monitoring will be the responsibility of the EUP OHAMR Secretariat (reporting@ohamr.eu). On behalf of the project consortium, the coordinator is required to submit progress reports (mid-term and final-term) as to be outlined in the monitoring policy of the transnational projects supported under the EUP OHAMR.

The monitoring outputs and outcomes from the reports will be collected and made accessible to all funding organisations. In addition, the monitoring of each funded project will also be done **through follow-up meetings with expert panel to assess the outcomes and impact of projects**. The EUP OHAMR Secretariat will contact the coordinator requesting the reports as per required timeline and will provide to provide detailed information on the reporting needs.

Other than the progress reports, grantees have an obligation to provide the EUP OHAMR with updated information of the consortium and its results, if requested, even beyond the tenure of the projects.

6- Ethics

Each funded consortium must have all necessary ethics approvals for research on animals, and/or research involving human subjects or data/samples obtained from human subjects according to national/regional law and regulation and in compliance with EU Horizon Europe rules before initiation of such research. Applications for ethics approval and ethics approvals should be made available immediately to the EUP OHAMR Secretariat upon request. The EUP OHAMR may perform an ethics review of the research at any time (evaluation and/or follow-up of the funded projects).

Project coordinators must inform the EUP OHAMR Secretariat as well as the funders supporting the project if ethics approvals are denied. The notification should be communicated no more than 14 calendar days after the rejection and the proposed rescue plan (new request for ethics approval, modification of the workplan/ project scope) must be approved by the funders supporting the project.

Any partner of a consortium in breach of research ethics regulations will subject the whole project for re-evaluation by all funding organisations of the project resulting in potential inhibition of all activities, withdrawal of funds, cancelling of contracts, and /or legal action or other sanctions according to national law.

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7- Changes to on-going grants

The funded projects may need to request changes in the project consortium (for e.g. change of organisation, change of coordinator/partner) or in the project workplan due to circumstances beyond their control that might prevent them from conducting research and thereby hinder implementation of their project during its tenure. These changes should be **exceptional and supported by reasonings due to scientific or administrative constraints**. Substantial changes must be approved by the different funding organisations involved in the project. The project coordinator should inform the EUP OHAMR Secretariat (postaward@ohamr.eu) as soon as possible using the project change request form. The EUP OHAMR Secretariat will inform the relevant funding organisations, who will decide upon the proper action to be taken.

8- Communication

Coordinators of the funded projects are required to deliver, upon request, an abstract or a short video presenting their project or a summary of their project results suitable for communication and dissemination purposes.

The project coordinators should be available to participate in meetings/workshops/podcast with the aim of:

• disseminating project results;

• developing a joint strategy to coordinate and facilitate integration of the planned activities of the EUP OHAMR;

• supporting the uptake of the research results by stakeholders;
• communicating results across EUP OHAMR.

Importantly, all funding recipients must ensure that all research outcomes (i.e. publications, tools, software, databases) of transnational EUP OHAMR funded projects include proper acknowledgement of the EUP OHAMR and the respective funding partner organisations using the text below:

"This project received funding from [name of funding organisations, or an acknowledgment as requested by your national/regional funding organisations] under the umbrella of the European Partnership on One Health Antimicrobial Resistance (EUP OHAMR) (GA N° 101217154 of the EU Horizon Europe Research and Innovation Programme)."

For any oral presentation, the EUP OHAMR logo, the logo of the national/regional funders as well as the EU emblem "co-funded by the European Union" should be displayed (with the same size for each logo).

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