



Call for Proposals 2022

"Development of new analytic tools and pathways to accelerate diagnosis and facilitate diagnostic monitoring of rare diseases"

Guidelines for Applicants

Submission deadline for pre-proposals: February 16th, 2022 at 2 PM (CET)

For further information, please visit us on the web:

<http://www.ejprarediseases.org/>

Or contact:

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1. Application Process

1.1 Registration

Research consortia who intend to submit a transnational project proposal should **register as soon as possible** via the electronic proposal system: <https://ptoutline.eu/app/ejprd22> . Please fill in the data sheet in the system. The same data sheet can be used for the submission of pre-proposals and full proposals (if invited).

1.2 Pre- and Full Proposals

There will be a **two-stage submission procedure for joint applications**: a pre- and full proposal stage. In both cases, one joint proposal document (in English) shall be prepared by the partners of a joint transnational proposal and must be submitted by the coordinator only to the JCS via the electronic submission system: <https://ptoutline.eu/app/ejprd22> . Proposals must be prepared using the templates provided on the EJP RD web page (www.ejprarediseases.org). Proposals not conforming to template instructions (including length and format) will be rejected.

You will not need to submit a paper version of your proposal; however, both the **electronic pre-proposals and full proposals need to be signed** (electronic signature or a scanned copy of the signature page will be accepted).

Joint pre-proposals (in English) must be received by the JCS in an electronic version no later than February 16th, 2022 at 2:00 p.m. Central European Time (CET).

Full proposals (in English) must be received by the JCS in an electronic version no later than June 15th, 2022 at 2:00 p.m. Central European Summer Time (CEST).

1.3 Rebuttal stage

Please note that project coordinators will be provided with the opportunity to study the assessments of external reviewers and comment on their evaluations of full proposals (for details see section 7.3 in the “Call text” document).

2. Advice for preparing your proposal

Carefully read the “Call Text” and this “Guidelines for Applicants” document, including the call aim, evaluation criteria and national eligibility criteria and requirements.

Proposals not conforming to the following may be rejected without review:

- Make sure that your proposal falls into the scope of the call (Section 4 of the call text)
- Make sure that your proposal fulfils the eligibility criteria of the call (Section 5 of the call text)
- Make sure that all consortium members have understood the national eligibility criteria and requirements (Annex 1) and that they fulfil these criteria

- Make sure that all consortium members contacted their national representative and confirmed eligibility with their respective funding organizations in advance of submitting an application (see Annex 1)
- Prepare your proposal in advance and enter the requested information on the submission site as soon as possible to avoid possible overloading on the submission deadlines
- Use the proposal templates provided on the EJP RD website (www.ejprarediseases.org)
- Respect the length limitations of each section in the proposals

3. Project description

Applicants will describe and justify the following elements:

The elements marked with a “” will have to be developed only for full proposals*

Background, present state of the art in the research field

- Need for research rationale: description of the unmet need that is addressed by the proposed work, rationale of the rare diseases chosen
- Present state of the art, recent insight from literature
- Preliminary results obtained by the consortium members

Objectives and hypothesis

- Main and secondary hypothesis

Soundness and pertinence

- Innovative aspects, originality, novelty
- Social care and public health interest
- Applicants should include information about other ongoing development work and explain why their approach should be supported*.

Workplan & methodology (*highlighting feasibility*)

- Research strategy
- Methodologies justification and presentation
- Enrollment: study location(s), inclusion/exclusion criteria, total number of corresponding patients followed by partners and collaborators of the project.
- Number of participants calculation (if applicable): description, justification, expected response rate.
- Statistical power (if applicable): appropriate statistical methods description, name and affiliation of the responsible biostatistics' expert.
- *Quality monitoring: risk management, contingency plans (identification of possible bottlenecks and go/no go steps).

Impact

- Results: description of expected results and their implementation
- Impact: description of the potential impact of the expected results on the addressed unmet need
- Benefits: description of individual and collective benefits that could be expected

***Valorization, translation in practice**

- Effective measures to exploit and disseminate the project results, to communicate the project, and to manage research data
 - o Present / future position with regard to intellectual property rights, both within and outside the consortium
 - o Scientific communication (articles, presentations...): description of plan, tools and responsibilities for communication towards clinical community
 - o PAO/Public communication: description of plan, tools and responsibilities for communication towards PAOs, patients, any concerned people
- Innovative potential: relevant application for rare diseases care
- Translatability: opportunities to exploit the methodology and/or expected results for other rare and non-rare diseases
- Sustainability: description of plan for sustainability of infrastructures or resources initiated by the project, follow-on funding and/or draft study plans past the grant end, articulation with other existing research infrastructures**.

***Ethical and legal issues, data management**

- Ethical and legal issues management plan description, including:
 - o the recruitment of participants (e.g. direct/indirect incentives for participation, the risks and benefits for the participants etc.)
 - o the material collection (e.g. sensitive or personal data etc.)
 - o ensuring the wellbeing of the children involved
 - o ensuring consentSee H2020 Guidance "How to complete your ethics self-assessment" that can be found here:
http://ec.europa.eu/research/participants/data/ref/h2020/grants_manual/hi/ethics/h2020_hi_ethics-self-assess_en.pdf
- GDPR management: plan description, name and affiliation of the Data Protection Officer (DPO).
- Data management strategy: plan description to make research data findable, accessible, interoperable and re-usable (FAIR).

Work packages, timeline and budget

- Description of the aims/work packages: synopsis and timeframe, including project coordination and management as well as *innovation management activities
- * Justification of requested budget: rational distribution of resources in relation to project's activities, partners responsibilities and time frame; please also specify co-funding from other sources necessary for the project if applicable
- Diagram which compiles the work plan, timeline, sequencing of work packages, contribution of the partners to each work package and their interactions (Gantt chart, Pert or similar)

Responsibilities and workloads

- For each research partner: competence and experience in the field(s) of the proposal (previous work in the field, specific expertise); responsibilities in each work package; *ongoing or submitted research grants.
- For PAO/patient representative: role and contribution, access to and engagement of patients, responsibilities in each work package.
- Added values: complementarity of the participants within the consortium, benefit of transnational collaboration
- *Management plan: operating and coordination methods

PAOs engagement/involvement

**Those elements will have to be developed only for full proposals*

***The use of **existing European health research infrastructures** and/or **IRDiRC recognized resources** is strongly encouraged when appropriate: e.g. research infrastructures established as a European Research Infrastructure Consortium (ERIC) or identified on the roadmap of the European Strategy Forum on Research Infrastructures (ESFRI). Projects are invited to identify the existing European research data infrastructures that may be used and how these may be mobilized, in particular for long-term data curation and preservation, when needed (in accordance with EU and [IRDiRC recommendations](#)). The following ESFRI European Research Infrastructures and European/international projects or their results may be of use to consortia:*

- [BBMRI](#) Biobanking and Biomolecular Resources Research Infrastructure
- [ELIXIR](#) The European Life Sciences Infrastructure for Biological Information and [ELIXIR Research Data Management Kit \(RDMkit\)](#)
- [INFRAFRONTIER](#) European Infrastructure for Phenotyping, Archiving and Distribution of Mouse Models
- [INSTRUCT](#) Integrated Structural Biology Infrastructure for Europe
- [EU-OPENSREEN](#) European high-capacity screening network
- [EATRIS](#) European Infrastructure for Translational Medicine
- [Rd-Connect](#) An integrated platform connecting databases, registries, biobanks and clinical bioinformatics for rare disease research
- [IRDiRC recognized resources](#)
- [Matchmaker Exchange](#) Federated platform to facilitate the matching of cases with similar phenotypic and genotypic profiles
- [Horizon 2020 FAIR Data Management Plan](#) Annex 1
- [Orphanet Rare Disease Ontology](#)
- [Human Phenotype Ontology](#)
- [Recommendations for Improving the Quality of Rare Disease Registries](#)

**** The [EJP RD's Resource Finder](#) provides scientific partners with a vast number of existing research data and services grouped into categories and represented as 11 `nodes` in the mindmap.*

4. Early Career Researchers (ECRs)

4.1 Definition

ECRs are defined as per the regulations of the European Research Council criteria for starting grants. In short, the researcher must have been awarded their first doctoral degree (**PhD**) **two to seven years prior to the pre-proposal submission deadline**. Extensions to this period are allowed (with documentation) in the case of reasonably justified career breaks: absence for maternal, paternal or long-term sick leave, and compulsory military service.

For medical doctors (or applicants holding a degree in medicine), an MD is not by itself considered equivalent to a PhD award. To be considered an ECR, these applicants must provide the certificates of both a medical doctor degree and a PhD, or proof of an appointment that requires doctoral equivalency (e.g. post-doctoral

fellowship or professorship appointment). MD applicants that do not hold a PhD must have been awarded their **MD four to nine years prior to the pre-proposal submission** deadline. For medical doctors who have been awarded both an MD and a PhD, the date of the earliest degree that makes the applicant eligible will be used to calculate eligibility. Extensions to this period will be given (with documentation) for clinical training periods up to a maximum of four years.

4.2 Eligibility of ECRs

The following dates must be provided by Early Career Researchers so that their eligibility can be evaluated according to their respective regional/national regulations. **This information must be present in the CV in the pre- and full proposal forms.**

Medical doctors with PhD

Medical Studies: *indicate dates (start and end) of your studies (year and month)*

End of studies: *indicate date of your medical certificate*

PhD Time: *indicate dates (start and end) of your PhD time (year and month)*

PhD: *indicate date of your PhD certificate*

Appointment: *indicate dates (start and end) of the appointment that requires doctoral equivalency (e.g. post-doctoral fellowship or professorship appointment), only if applicable*

Medical doctors without PhD

Medical Studies: *indicate dates (start and end) of your studies (year and month)*

End of studies: *indicate date of your certificate*

Appointment: *indicate dates of the appointment that requires doctoral equivalency (e.g. post-doctoral fellowship or professorship appointment)*

Other Early Career Scientists with PhD

Studies: *indicate dates (start and end) of your studies (year and month)*

End of studies: *indicate date of your certificate*

PhD Time: *indicate dates (start and end) of your PhD time (year and month)*

PhD: *indicate date of your PhD certificate*

Other Early Career Scientists without PhD

Studies: *indicate dates (start and end) of your studies (year and month)*

End of studies: *indicate date of your certificate*

Appointment: *indicate dates of the appointment that requires doctoral equivalency (e.g. post-doctoral fellowship or professorship appointment)*

Reasons for Extensions, if applicable

Clinical Training: *indicate dates (start and end) of clinical training (year and month);*

Parental leave: Women: *number of children (1.5 years are given per child; in case of longer maternal leave, please indicate the exact dates); Men: indicate exact dates of paternal leave (per child)*

Career Break: *indicate dates (year and month) of other career breaks: long-term sick leave, compulsory military service, carer's leave*

5. Financial and Legal Issues

5.1 Funding model and Call governance

The EJP RD JTC 2022 Funding Partners have agreed to launch a joint call using the **“virtual common pot” funding mode**. This means that national/regional funding will be made available through national/regional funding organizations according to national/regional funding regulations.

ISCIII (Spain) is acting as Joint Call Secretariat (JCS) to assist the Call Steering Committee (CSC), and the national/regional funding bodies during the implementation of the call.

The JCS will be responsible for the administrative management of the call. It will be the primary point of contact referring to the call procedures between the research consortia, the funding organizations (CSC), and the peer reviewers. The project **coordinator will be the point of contact for the JCS** during the application procedure and is responsible for forwarding this information to other partners.

CSO-MOH (Israel) and FNRS (Belgium) will be responsible for the follow-up phase until the funded research projects have ended.

5.2 Widening for the inclusion of under-represented or under-subscribed countries

5.2.1 Definition of widening

For proposals invited to the full proposal stage, there will be a widening step to provide the **opportunity to add partners** to the consortium (up to a maximum total of 8, see section 5.4 *Consortium Makeup* of the Call Text). This step will allow for the addition of partners from participating countries that are usually underrepresented in the call, as well as those undersubscribed (countries without any selected applicants for the 2nd stage). This inclusion will not be considered as a fundamental change between pre- and full proposal. **Inclusion of new research teams is not mandatory. The new teams included should bring an added value and expertise to the projects.**

5.2.2 Process

A list of countries eligible for this widening procedure will be published on the EJP RD website after completion of the 1st stage of evaluation and sent to the coordinators that are invited to write a full proposal.

The relevant national funding agencies may produce a list of research teams that could provide additional expertise to projects. **For this, the title, pre-proposal abstract, and composition of the consortium will be shared with potentially interested research teams.** The JCS will then provide this list to the coordinators of projects invited to the full proposal stage and give them the option of adding them to the existing consortium.

The coordinator/partners of projects invited to the 2nd stage of evaluation can also inquire themselves about suitable partners from among listed countries. The new prospective partner must then contact their national funding agency to confirm their eligibility.

In all cases, the final decision on whether to take a new research team on board will be taken by the project consortium. The rules concerning the maximum number of partners in a consortium and the maximum of two research teams per country must

still be respected. Furthermore, the new research team must be eligible for the national funding agency. For this purpose, national funding agencies from underrepresented or undersubscribed countries may indicate that only national research teams that were already involved in pre-proposals (and thus are eligible) are allowed to make use of this widening step.

5.3 EJP RD Mentoring programme

Applicants that are invited to submit a second stage proposal are strongly encouraged to make use of the EJP Rare Diseases Mentoring Programme. This completely free service offered by EJP RD matches your project with mentors that have expertise in applied research and translational development. During full proposal development the service will provide independent feedback and advice to applicants in critical methodological areas. This will help to ensure robust and reliable study results so that the project – if successful – has the best chances of advancing towards implementing the results for patient care. These areas include:

- Appropriate use of analytical technologies and read-outs;
- Appropriate sampling methods and sample handling;
- Statistical methodology;
- Good practice in validation and follow-on study planning;
- Risk management strategies for the development process.

Applicants should indicate in the pre-proposal form if they wish to make use of the Mentoring service. If they are invited to full proposal submission, the Mentoring team will soon after contact the applicants to schedule a planning call, after which mentors will be assigned to the project based on the individual needs.

Please note that in order to provide a reliable and useful service, applicants must request the mentoring service no less than 4 weeks before the full proposal submission deadline.

5.4 Funding contracts

Each project includes several partners (including a project coordinator) as beneficiaries. **Each partner will have a separate funding contract/letter of grant with their respective national/regional funding organizations, and according to their regulations.**

Changes to the composition of research consortia or budget cannot occur within the contract/letter of grant without thorough justification. Minor changes will be handled by the relevant national/regional funding agency. In the case of major changes, an independent expert may be consulted to help with the final decision of the funding organizations. **Research partners must inform the JCS and the respective funding bodies of any event that might affect the implementation of the project.**

5.5 Project start and consortium agreement

Consortium members of projects selected for funding **must fix a common project start date**, which will be the reference date for yearly and final reports and extensions. This common project start date must appear in the **Consortium Agreement (CA)**.

The project consortium partners must sign a CA for cooperation. For reference see the [DESCA 2020 Model Consortium Agreement](#). It is recommended that the CA be signed by all relevant parties before the official project start date. Please note that national/regional regulations may apply concerning the requirement for a CA (please contact your national/regional contact point or check Annex 1). This consortium agreement must be made available on request to the relevant EJP RD JTC 2022 funding organizations.

The purpose of the CA shall be:

- to underpin the collaboration and provide research partners with mutual assurance on project management structures and procedures, and their rights and obligations towards one another
- to assure the CSC that the research consortium has a satisfactory decision-making capability and is able to work together in a synergistic manner

The following subjects should be addressed by the CA (at minimum):

- purpose of and definitions used in the CA
- names of organisations involved
- common start date of the research project
- organisation and management of the project
- role and responsibilities of the research consortium coordinator and the research partners: person in charge, their obligations and key tasks, conditions for their change
- deliverables (transnational reports and, if relevant, requirements for national reports where coordination is required)
- resources and funding
- confidentiality and publishing
- intellectual property rights (how this issue will be handled between research partners)
- 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)

5.6 Ownership of intellectual property rights

Results and Intellectual Property Rights (IPR) resulting from projects funded through the EJP RD JTC 2022 will be owned by the beneficiaries' organizations according to national/regional rules on IPR. In the case of joint development of intellectual property, consortium partners will resolve this issue internally using their consortium agreement and relevant legal guidelines and taking into account their relative contributions.

The results of the research project and IPR created should be actively exploited and made available for use, whether for commercial gain or not, in order for public benefit to be obtained from the knowledge created.

The funding organizations shall have the right to use documents, information and results submitted by the research partners and/or to use the information and results for their own purposes, provided that the owner's rights are kept and taking care to specify their origin.

5.7 IRDiRC policies and guidelines

The project partners are expected to follow [IRDiRC policies and guidelines](#).

5.8 European and International standards

The submitted proposals must respect relevant European and international standards including:

- [H2020 ethics manual](#) for research projects,
- [The Declaration of Helsinki](#) Ethical Principles for Medical Research Involving Human Subjects,
- *The General Data Protection Regulation (GDPR): the European Regulation (EU) 2016/679 on the protection of natural persons with regard to the processing of personal data and on the free movement of such data:*
- <https://publications.europa.eu/en/publication-detail/-/publication/3e485e15-11bd-11e6-ba9a-01aa75ed71a1/language-en>; [The EC Directive 2010/63/EU](#) on the protection of animals used for scientific purposes,
- [European Research Council Guidelines on Implementation of Open Access to Scientific Publications and Research Data](#),
- To make research data findable, accessible, interoperable and re-usable (FAIR), a [data management](#) strategy is mandatory in the full proposal. [Example questions for a data management strategy](#).
- General ethical and legal requirements: Ethics is an integral part of research. Ethics should be embedded in the research and considered from the outset, and although legal and regulatory considerations may vary across different countries, EJPRD will only fund proposals which comply with national and international ethical standards, rules and legislations
- International Ethical Guidelines for Biomedical Research Involving Human Subjects CIOMS-WHO (2016);
- Oviedo Convention and its Additional Protocol on human rights and biomedicine, concerning biomedical research (2005);
- COUNCIL OF EUROPE COMMITTEE OF MINISTERS. Recommendation CM/Rec (2016)6 of the Committee of Ministers to member States on research on biological material of human origin (Adopted by the Committee of Ministers on 11 May 2016).

5.9 Publication of Results

Each beneficiary must ensure open access (free of charge, online access for any user) to all peer-reviewed scientific publications relating to their results, if this is compliant with national/regional funding regulations.

Funding recipients must ensure that all outcomes (publications, etc.) of transnational EJP RD projects include a proper acknowledgement of EJP RD and the respective national/regional funding partner organizations. This includes the display of the EJP RD logo when possible.

Unless the EC requests or agrees otherwise or unless it is impossible, any dissemination of results (in any form, including electronic) must:

1. display the EU emblem and

2. include the following text: "This project has received funding from (Name of funding agency) partner of the EJP RD. The EJP RD initiative has received funding from the European Union's Horizon 2020 research and innovation programme under grant agreement N°825575"

When displayed together with another logo, the EU emblem must have appropriate prominence. For the purposes of the obligations under this Article, the beneficiary may use the EU emblem without first obtaining approval from the Agency. This does not however give it the right to exclusive use. Moreover, the beneficiary may not appropriate the EU emblem or any similar trademark or logo, either by registration or by any other means.

6. General Data Protection Regulation

The following Data Privacy Notice applies

By submitting an application to the call JTC2022, applicants consent to the use, processing and retention of their data for the purposes of:

- processing and evaluating the application 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 Funding Party's relationship with them.
- analysing and evaluating the call;
- reporting to the European Commission/ Research Executive Agency (REA) on the Co-funded call.
- providing aggregate data to national and European surveys and analyses.
- complying with audits that may be initiated by the Funding Parties and the European Commission (or its agencies).

The members of the EJP RD consortium may share an applicant's data with third parties (some of which may be based outside the European Economic Area) in relation to the above activities including evaluators, auditors and the European Commission (or its agencies).

The members of the EJP RD consortia 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. The members of the EJP RD consortia may also link the data that applicants provide in their application with future data that applicants provide as part of the ongoing management and reporting on a call award which may be awarded to them.

Data on Funding Parties including contact details of FC members and National Contact Points/Regional Contact Points are kept for the purpose of the call communication. The information will be published with prior consent of the respective management bodies.

ANNEX 1: Country and Region-Specific Guidelines

MEDICAL RESEARCH FUTURE FUND (MRFF), AUSTRALIA

It is strongly advised that all applicants contact their EJP RD National/Regional Contact Point in good time before the submission of a proposal

Country	Australia
Funding organization	Medical Research Future Fund (MRFF) https://www.health.gov.au/initiatives-and-programs/medical-research-future-fund
National contact person	Michael Nutt (National Health and Medical Research Council (NHMRC) – Managing Organisation) Michael.Nutt@nhmrc.gov.au
Funding commitment	\$1 Million AUD
Overheads	Overheads are not eligible costs for MRFF
Anticipated number of fundable research partners	3
Maximum funding per grant awarded to a partner	\$300,000 AUD
Eligibility of a partner as a beneficiary institution	MRFF Eligible Organisations may apply for and receive funding
Eligibility of costs, types and their caps	All eligibility rules including budget requirements are outlined in the MRFF 2022 Joint Transnational Call Grant Opportunity Guidelines available on GrantConnect
Submission of the proposal at the national level	Yes
Submission of financial and scientific reports at the national level	Yes
Further guidance	Refer to the MRFF 2022 Joint Transnational Call Grant Opportunity Guidelines available on GrantConnect

AUSTRIAN SCIENCE FUND (FWF), AUSTRIA

It is strongly advised that all applicants contact their EJP RD National/Regional Contact Point in good time before the submission of a proposal

Country	Austria
Funding organisation	Fonds zur Förderung der Wissenschaftlichen Forschung (FWF) / Austrian Science Fund http://www.fwf.ac.at
National contact person	Stephanie Resch Phone: +43 (1) 505 67 40-8201, E-mail: stephanie.resch@fwf.ac.at Anita Stürtz Phone: +43 (1) 505 67 40-8206, E-mail: anita.stuertz@fwf.ac.at
Funding commitment	0,6M€
Overheads	Overheads are not eligible costs for FWF.
Anticipated number of fundable research partners	2
Maximum funding per grant awarded to a partner	For scientists funded by the FWF, the funding is limited to “project-specific costs, i.e. personnel and non-personnel costs that are essential to carry out the project and that go beyond the resources made available from the research institution’s infrastructure, according to the general FWF Funding Guidelines published at https://www.fwf.ac.at/fileadmin/files/Dokumente/Antragstellung/Einzelprojekte/p_application-guidelines.pdf The FWF does not finance infrastructure or basic equipment at research institutions. Overheads may not be requested. Subcontracts must be well justified, i.e. must represent the only or the most economical way to have the work performed, please contact the FWF directly for clarification of individual cases. The current FWF salary scale (http://www.fwf.ac.at/en/research-funding/personnel-costs/) indicates the salaries that may be requested.
Eligibility of a partner as a beneficiary institution	Individual researcher, working in any kind of non-profit organisation: e.g. University, University hospital, Non-university research institute Please refer also to the general FWF Funding Guidelines: http://www.fwf.ac.at/fileadmin/files/Dokumente/Antragstellung/Einzelprojekte/p_application-guidelines.pdf available on: http://www.fwf.ac.at/en/research-funding/application/international-programmes/joint-projects-era-nets/
Additional specific rules	Please note that the number of ongoing/approved/submitted projects in which one researcher can serve as principal investigator is limited to three in the Stand-Alone Projects Programme, International Programmes (including ERA-Net projects!), Clinical Research and Arts-Based Research Programmes. Principal investigators who already have three ongoing/approved/submitted projects will not be permitted to submit another application within those programmes until 12 months before the end of one of their ongoing projects. You are strongly advised to contact the national representative in case you may be affected by this regulation. https://www.fwf.ac.at/fileadmin/files/Dokumente/Antragstellung/project_number_limit.pdf

Conditions for PAO funding	Not applicable
Submission of the proposal at the national level	<p>FWF Submission:</p> <p>In addition to the application at the call secretariat administrative data (in accordance with the FWF guidelines for stand-alone projects) must be submitted online to the FWF at https://elane.fwf.ac.at/. This is required already at the pre-registration stage via the programme category "IK – International Projects (preproposal, deadline 16. February 2022 / 02:00 p.m. CET)".</p> <p>For the full proposal stage applicants must choose the programme category "I – International Projects" (full proposal, deadline 15. June 2022 / 02:00 p.m. CET).</p> <p>Both steps are mandatory.</p> <p>For submissions to be valid, the cover sheet generated at the end of the online submission process must be printed out and signed. It can then either be sent to the FWF by conventional mail (FWF, Sensengasse 1, 1090 Vienna) or scanned in, given a digital signature and sent to the FWF (office@fwf.ac.at) as an e-mail attachment. Detailed information may be found under the Internet http://www.fwf.ac.at/fileadmin/files/Dokumente/Antragstellung/Internationale_Programme/i_infosheet-era-net.pdf</p>
Further guidance	<p>http://www.fwf.ac.at/en/research-funding/application/international-programmes/joint-projects-era-nets/</p>

RESEARCH FOUNDATION FLANDERS (FWO), BELGIUM, FLANDERS

It is strongly advised that all applicants contact their EJP RD National/Regional Contact Point in good time before the submission of a proposal

Country	Belgium (Flanders)
Funding Organization	The Research Foundation - Flanders (FWO)
National Contact Person	<p>Toon Monbaliu (FO) eranet@fwo.be +32 (0)2 550 15 70</p> <p>Kristien Peeters (SBO) eranet@fwo.be +32 (0)2 550 15 95</p>
Funding Commitment	0,35 m. EUR
Overhead	Overhead has to be included – see category 'Eligibility of costs, types and their caps'.
Anticipated number of fundable research partners	1
Maximum funding per grant awarded to a partner	350.000 EUR (overhead included)
Eligibility of a partner as a beneficiary institution	<p>Both the FWO Strategic Basic Research (SBO) and junior/senior research project (FO) funding channels are integrated in this call, each with specific regulations. It is, in the light of the projects eligibility, of utmost importance to respect their particular regulations. For example when it comes to the mandatory valorisation aspect for the SBO projects (see 'additional conditions for FWO funding' below).</p> <p><u>Who can be eligible for FWO funding?</u> The eligibility of institutions and its researchers can be verified in the relevant regulations: - For junior/senior research projects, see articles 10-12 - For Strategic Basic Research, see articles 4-8</p> <p><u>Additional conditions for FWO funding:</u></p> <ol style="list-style-type: none"> 1. When the strategic basic research channel (SBO) would be the appropriate source of funding, we ask researchers to provide us with a 'valorisation plan' before the pre-proposal submission deadline. There is no fixed format and one A4 page should suffice. What the FWO wants to know is i) how the valorisation within Flanders - and potentially internationally – will take place and ii) which Flemish actors are involved in this. This information can be submitted to the general eranet@fwo.be email address.

	<ol style="list-style-type: none"> 2. SBO projects aiming at the development of a spin-off company are not eligible here. 3. Researchers have to inform the central research coordination units, at their host institutions, about their participation. 4. One and the same researcher can only participate in 2 different research projects/consortia when applying for FWO funding, within the same call. Double funding is not allowed. 5. Projects may last up to 36 months, which implies the funding has to be budgeted and spent accordingly. 6. The PI, for each of the participating institutions from Flanders, must hold an appointment that fully covers the duration of the research project. <p>ERA-NET participation does not interfere with the 'regular' project submission framework, and is consequently not taken into account for calculating the max. available number of new applications and running projects combined.</p>
<p>Eligibility of costs, types and their caps</p>	<p>The regular FWO cost categories from the (junior/senior) 'research project' or SBO project funding channels are eligible:</p> <p>The maximum requested budget per partner amounts to 350.000 EUR (incl. overhead). Beware, the funding rules differ per FWO funding channel (FO and SBO):</p> <ul style="list-style-type: none"> - FO: a 6% structural overhead should be calculated on the direct costs. E.g., a practical example: when the sum of all costs (personnel, consumables, travel, etc.) amounts to 300.000 EUR, then the overhead will be 18.000 EUR (6% of 300.000 EUR) and the total requested cost 318.000 EUR. This total requested cost may never exceed 350.000 EUR (for further detailed financial information, see chapters 6, 7 and 8 in the project regulations). - SBO: The SBO cost model applies. Generally, a 17% overhead rate is applicable.
<p>Conditions for PAO funding</p>	<p>PAO funding only possible as subcontractor.</p>

Submission of the proposal at the national level	<p>No submission at the national/regional level is required. However, if SBO, a valorisation plan has to be submitted.</p>
Further guidance	<p>It is always strongly advised to contact the FWO before submission, in order to verify the eligibility of the researchers and avoid ineligible projects/research consortia (see 'National Contact Persons' above).</p> <p>Information available at:</p> <ul style="list-style-type: none"> - Call page for European programmes - Junior/senior research projects (FO) - SBO research projects (SBO)

FUND FOR SCIENTIFIC RESEARCH-FNRS (F.R.S.-FNRS), BELGIUM, FRENCH SPEAKING COMMUNITY

It is strongly advised that all applicants contact their EJP RD National/Regional Contact Point in good time before the submission of a proposal

Country	Belgium
Funding organisation	Fund for Scientific Research – FNRS (F.R.S.-FNRS)
Management organisation	Fund for Scientific Research – FNRS (F.R.S.-FNRS)
National contact person	<p>Dr. Florence Quist Phone: +32 (0)2 504 9351 E-mail: florence.quist@frs-fnrs.be</p> <p>Joël Groeneveld Phone: +32 (0)2 504 9270 E-mail: joel.groeneveld@frs-fnrs.be</p>
Funding commitment	0,2 Mio€
Overheads	“Overhead” is not an eligible cost. If the project is selected for funding, these costs will be subject to a separate agreement between the institution of the beneficiary and the F.R.S.-FNRS.
Anticipated number of fundable research partners	1
Maximum funding per grant awarded to a partner	200.000 €
Eligibility of project duration	Maximum 3 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 4 years but should remain within the 200.000 € budget maximum (cf. PINT-Multi regulations , art. III.3, second paragraph)
Eligibility of a partner as a beneficiary institution	All eligibility rules and criteria can be found in the PINT-Multi regulations . It is strongly advised to contact the F.R.S.-FNRS prior to submission regarding the eligibility criteria.
Eligibility of costs, types and their caps	All eligibility rules and criteria can be found in the PINT-Multi regulations . It is strongly advised to contact the F.R.S.-FNRS prior to submission regarding the eligibility criteria.
Conditions for PAO funding	Participating Belgian patients organizations could be financed via subcontracting, provided that the criterion for subcontracting detailed in the PINT-MULTI regulations are fulfilled (see art. III.3).

Submission of the proposal 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 EJP RD call 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.
Submission of other information at the national level	N/A
Submission of financial and scientific reports at the national level	Financial reporting must be submitted to the F.R.S.-FNRS
Further guidance	PINT-MULTI regulations , e-space

CANADIAN INSTITUTES OF HEALTH RESEARCH- INSTITUTE OF GENETICS (CIHR-IG), CANADA

It is strongly advised that all applicants contact their EJP RD National/Regional Contact Point in good time before the submission of a proposal

Country	Canada
Funding organisation	Canadian Institutes of Health Research, Institute of Genetics (CIHR-IG)
National contact person	<p>Jennifer Vineham Phone: +1 343 552-2760 Email: jennifer.vineham@cihr-irsc.gc.ca</p> <p>Etienne Richer Email: Etienne.Richer@cihr-irsc.gc.ca</p>
Funding commitment	<p>CIHR: CAD \$1,350,000, Ataxia Canada: CAD \$450,000 Total funding available: CAD \$1,800,000 CAD \$150,000 per year per project maximum.</p>
Overheads	Not an allowable cost.
Anticipated number of fundable research partners	4 projects
Eligibility of project duration	3 years
Eligibility of a partner as a beneficiary institution	
Eligibility of costs, types and their caps	<p>Eligibility of principal investigator or other research team member Academia, Clinical, Public Health https://cihr-irsc.gc.ca/e/50805.html#g-3</p> <p>Investigator (early career) For this competition only: a researcher who, at the time of application, has held a full time, independent research appointment, for a period of 0 to 7 years (84 months) *.</p> <p>All time spent in research appointments/positions will be taken into consideration when determining eligibility irrespective of time spent in a clinical component or other duties (i.e., administrative, academic, etc.). Should an applicant hold or have held a part-time appointment/position, CIHR will count that time as 50% (e.g., a one-year</p>

	<p>part-time appointment/position will count for 6 months towards the maximum). Leaves of absence will be considered in the calculation of eligibility (i.e., are credited as twice the amount of time taken) and should be included in the Employment section under Leaves of Absence in your Common CV.</p> <p>*Please note that due to the impact of COVID-19 on early career researchers, CIHR temporarily adjusted the period of eligibility for an ECR. All those who held ECR status as of March 1, 2020 – or who secured their first academic appointment after this date – will have their status extended to 0-84 months from 0-60 months.</p> <p>CIHR will closely monitor the pandemic and its impact on ECRs overall and on specific groups with the intent that further interventions may be warranted.</p> <p>Eligibility of costs, types and their caps https://www.nserc-crsng.gc.ca/InterAgency-Interorganismes/TAFA-AFTO/guide-guide_eng.asp</p>
Conditions for PAO funding	<p>Canadian patient advocacy organizations (PAOs) are not eligible to participate in the role of Nominated Principal Applicant (NPA) nor Principal Applicant (PA). However, it is possible for a PAO to be represented by an individual in the role of co-applicant or collaborator. In this case, the NPA may request funds in their budget to support the activities of the PAO representative on the project.</p>
Submission of the proposal at the national level	<p>Short application as per CIHR Funding Opportunity (link to follow). Summary of the applications potentially relevant for funding by Ataxia Canada will be made available to that organization to confirm relevancy of the proposals to their mandate.</p>
Submission of other information at the national level	<p>NA</p>
Submission of financial and scientific reports at the national level	<p>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.</p>
Further guidance	<p>NA</p>

FONDS DE RECHERCHE DU QUÉBEC - SANTÉ (FRQS), QUÉBEC (CANADA)

It is strongly advised that all applicants contact their EJP RD National/Regional Contact Point in good time before the submission of a proposal

Country	QUÉBEC (CANADA)
Funding organisation	FONDS DE RECHERCHE DU QUÉBEC - SANTÉ (FRQS) – https://frq.gouv.qc.ca/
National contact person	Maxime Beaudoin, Programs Manager Tel: (+1) 514-873-2114 ext.4369 Maxime.beaudoin@frq.gouv.qc.ca
Funding commitment	<p>\$450, 000 CAD (~0.32 Mio. €) available for Quebec researchers.</p> <p>The total maximum amount that can be requested in support of a Quebec/Canada component is \$150,000 CAD per year for up to 3-years from all Canadian funding sources CIHR-IG, FRQS and their funding partners.</p> <p>Funds are subject to availability of funds voted annually to FRQS by the National Assembly of Québec and FRQS Board of Directors' approval.</p>
Overheads	Overheads means "frais indirects de recherche" and will be managed separately by the FRQS. <u>They should not</u> be included in the requested budget.
Anticipated number of fundable research partners	<p>Anticipated number of funded research groups: 1-2*</p> <p>FRQS is providing funding to support up to one project as outlined in the call text.</p> <p>*Canadian funders agencies (FRQS and CIHR-IG) will be working together to maximize participation from the Canadian research community.</p>
Eligibility of project duration	Up to 3-years
Eligibility of a partner as a beneficiary institution	<p>Quebec applicants must meet the eligibility criteria of FRQ research grants.</p> <p>Eligible institutions are Quebec Universities or Institutions within Quebec's health and social services network. Please consult the full list of Quebec Eligible managing institutions.</p>

<p>Eligibility of costs, types and their caps</p>	<ul style="list-style-type: none"> • Operational costs (research personnel, consumables, animals) • Costs related to scientific and ethical evaluation (clinical research projects) • Coordination-related cost (project administration and travel expenses for attending joint meetings) • Costs related to knowledge translation and translation • Conference attendance (up to 5% per year of the grant amount starting the first year with proper justification) <p>There is NO support for salaries of investigators or equipment. There is NO supplement for the Coordinator of an international consortium.</p> <p>Further information about eligible and ineligible expenses is available in section 8 of the FRQ Common General Rules.</p> <p><i>Overheads means "frais indirects de recherche (FIR)" are not an eligible expense. It will be managed separately.</i></p>
<p>Conditions for PAO funding</p>	<p>FRQS cannot fund Quebec/Canada PAOs directly nor be an eligible partner (Investigator(s)) on the application.</p> <p><i>However, it is possible for a PAO to be represented by an individual in the role of <u>Individual contributor or collaborator</u>. In this case, the Quebec Principal Investigator (PI) may request funds in their budget to support the activities of the PAO representative on the project or they could be supported as subcontractors.</i></p>
<p>Submission of the proposal at the national level</p>	<p><u>Non-extensive requirements</u></p> <p>Quebec Principal investigator (PI) <u>must submit a short application form</u> through FRQS e-portfolio. Quebec Co-investigator(s) has to consent to be part of this application before the institutional approval. CCV of all the investigators must be updated with the most recent information for the eligibility check. Managing institutions approval must be done lastly which automatically activates the final submission.</p> <p>\$CAD Budget form will be requested.</p> <p>Please consult the FRQ webpage of JTC2022 EJP RD program for complementary information about the requirements.</p> <p>Documents sent via mail or e-mail will NOT be accepted.</p>

	<p>FRQS short application form will follow the exact Call deadlines.</p>
<p>Submission of other information at the national level</p>	<p>*Quebec Investigators applying for funding from FRQS and CIHR must also complete an abbreviated CIHR form <u>at the Proposal stage only (second stage)</u> in order to be eligible to receive funding from both agencies. Please contact CIHR representative for more information.</p>
<p>Submission of financial and scientific reports at the national level</p>	<p><u>Yes, according to regional/national regulations.</u></p> <p>Scientific reports according to EJP RD template and requirements.</p> <p>Minimum requirements such as electronic Annual financial reporting according to FRQ Common General Rules and Grant Regulations.</p> <p>A Final scientific report under FRQ template will be requested at the very end of the project.</p> <p><i>The FRQS reserves the right to request any additional or complementary information related to the research project granted.</i></p> <p>Note: Further requirements will be stated in the official FRQS Letter of Grant addressed to the selected Quebec Principal Investigator(s).</p>
<p>Further guidance</p>	<p>Basic research ethics training is mandatory for all recipients of an FRQS grant when their part of research project that involve human beings. PI and Co-PI(s) on the project must therefore successfully complete levels 1 and 3 of MSSS Ethics online training by the Ministère de la Santé et des Services sociaux. Post-doctorate on the project are also encouraged to complete this training.</p> <p>5.5 Research Ethics and Conformity (FRQ Common General Rules) Funding holders and managing institutions must demonstrate the highest standards of research ethics and scientific integrity. All research projects involving human beings, including biological material (body parts, products, tissues, cells or genetic material from a human body, of a living or dead person) or administrative, scientific or descriptive data from human beings, usually require the approval of the research ethics board of the institution under whose authority or auspices the project is conducted (or a research ethics board recognized by</p>

	<p>that institution). In the case of grants, the managing institution is responsible for obtaining all necessary ethics approvals. For complementary information about FRQ standards adopted please consult FRQ Ethics and integrity section.</p>
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MINISTRY OF EDUCATION, YOUTH AND SPORTS (MEYS), CZECH REPUBLIC

It is strongly advised that all applicants contact their EJP RD National/Regional Contact Point in good time before the submission of a proposal

Country	Czech Republic
Funding organization	Ministry of Education, Youth and Sports (MEYS) www.msmt.cz
National contact person	Judita Klosaková MSMT, Department of Research and Development Phone : +420 234 811 504 E-mail : judita.klosakova@msmt.cz
Funding commitment	0.6 M€
Overheads	Eligible costs for a Czech participant involved in a project consortium are defined by § 2 of the Act No. 130/2002 Coll. on Support of Research, Experimental Development and Innovation from Public Funds and on Amendment to Some Related Acts. The maximum indirect costs set for the present call are 25 % (flat rate) of direct costs without the sub-contracting.
Anticipated number of fundable research partners	(2-3)
Maximum funding per grant awarded to a partner	No restriction
Eligibility of a partner as a beneficiary institution	The participants from the Czech Republic in the projects □ consortia must meet the criteria of the research and knowledge-dissemination organisation (hereinafter referred to as "research organisation") in accordance with the Framework for State Aid for Research and Development and Innovation (2014/C 198/03). These might be public universities, public research institutes and/or another entities classified as research organisations. It is obligatory that the Czech participants involved in the projects □ consortia prove compliance with the eligibility criteria and fulfilment of the conditions set by § 18 of the Act No. 130/2002 Coll. on Support of Research, Experimental Development and Innovation from Public Funds and on Amendment to Some Related Acts by means of a Statutory Declaration .
Eligibility of costs, types and their caps	Eligible costs for a Czech participant involved in a project consortium are defined by § 2 of the Act No. 130/2002 Coll. on Support of Research, Experimental Development and Innovation from Public Funds and on Amendment to Some Related Acts. The maximum indirect costs set for the present call are 25 % (flat rate) of direct costs without the sub-contracting. The aid intensity for activities carried out by a research organisation might be at the level of 100 % provided that the research organisation complies entirely with requirements stipulated by the Article 2.1.1 "Public funding of non-economic activities" of the Framework for State Aid for Research and Development and Innovation (2014/C 198/03) and proves it by means of the above-mentioned Statutory Declaration . Should the above-stated criteria not be fulfilled by the Czech participant, funding rates will be adjusted appropriately by the Ministry of Education, Youth and Sports and will reach the level of 100 % for fundamental/basic research activities, 50 % for applied research activities and 25

	<p>% for experimental development activities. For further information on the eligibility cost please see https://www.msmt.cz/vyzkum-a-vyvoj-2/ejp</p>
Conditions for PAO funding	<p>The MEYS does not fund PAOs/patient representatives.</p>
Submission of the proposal at the national level	<p>It is obligatory:</p> <ul style="list-style-type: none"> -that the Czech participants involved in the projects □ consortia prove compliance with the eligibility criteria and fulfilment of the conditions set by §18 of the Act No. 130/2002 Coll. on Support of Research, Experimental Development and Innovation from Public Funds and on Amendment to Some Related Acts by means of a Statutory Declaration. -that each Czech participant in a project consortium is requested to specify the costs related to the envisaged R&D activities in detail by using the Eligible Costs Specification. <p>The template and detail information will be available on websites of the Ministry of Education, Youth and Sports dedicated to the EJP RC JTC 2022.</p> <p>All of the requested documentation (i.e. Statutory Declaration and Eligible Costs Specification) shall be sent by each Czech participant in a project consortium to the Ministry of Education, Youth and Sports no later than the date of the official submission deadline for pre-proposals, both by electronic correspondence and data box/post.</p> <p>The electronic version of requested documentation shall be sent to the address of electronic correspondence: Judita.Klosakova@msmt.cz. One signed and stamped hard copy (by the statutory representative of research organisation) of requested documentation shall be submitted as well following the instructions stipulated on websites of the Ministry of Education, Youth and Sports.</p>
Further guidance	<ul style="list-style-type: none"> • https://www.msmt.cz/vyzkum-a-vyvoj-2/ejp

ACADEMY OF FINLAND (AKA), FINLAND

It is strongly advised that all applicants contact their EJP RD National/Regional Contact Point in good time before the submission of a proposal

Country	Finland
Funding organisation	Academy of Finland (AKA) http://aka.fi
National contact person	<p>Heikki Vilen Heikki.vilen@aka.fi +358 29 533 5135</p> <p>Rita Rinnankoski-Tuikka rita.rinnankoski-tuikka@aka.fi +358 29 533 5096</p>
Funding commitment	0,6M€
Overheads	According to full cost model SOCIAL
Anticipated number of fundable research partners	2
Maximum funding per grant awarded to a partner	Maximum funding is 300 000 € / consortium (full cost model, Academy's 70 % share of expenses so maximum total budget is ~428 570 €) – if there are several Finnish partners in the same consortium, they must share this among themselves.
Eligibility of a partner as a beneficiary institution	Eligibility rules for the applicants are the same as with Academy project funding: https://www.aka.fi/en/research-funding/apply-for-funding/calls-for-applications/apply-now2/academy-project-funding-2021-all-research-fields/ . AKA funds only Finnish research organizations (universities, public research institutes, university hospitals). Applicant needs to have an affiliation in such organization. Private not-for-profit research organizations such as VTT and CSC are also eligible – please check with the contact person.
Additional specific rules	NA
Conditions for PAO funding	AKA cannot fund PAOs directly – minor expenses (such as travel costs to a few meetings) can be included in the budget of a Finnish academic partner
Submission of the proposal at the national level	At the plan of intent stage of the call all Finnish applicants are advised to contact the national contact person. Submission of a national proposal will only be required in autumn 2022 from projects to be funded.

Submission of financial and scientific reports at the national level	Yes, according to national regulations.
Further guidance	NA

FRENCH NATIONAL RESEARCH AGENCY (ANR), FRANCE

It is strongly advised that all applicants contact their EJP RD National/Regional Contact Point in good time before the submission of a proposal

Country	France
Funding organisation	French National Research Agency (Agence Nationale de la Recherche –ANR-) http://www.agence-nationale-recherche.fr
National contact person	Health & Biology Department Agence Nationale de la Recherche –ANR 50 avenue Daumesnil - 75012 Paris, France Florence Guillot Email: EJPRDcall@anr.fr
Funding commitment	3.11 M€ Funding limits apply per partner for this call: Each partner may be granted up to 300 000 € as a coordinating partner or 250 000 € as a non-coordinating partner . The maximum amount that can be requested by French partners per project is 500 000 € The minimum funding amount per partner is 15 000 €.
Overheads	The ANR heading for “overheads” in the ANR funding breakdown is «frais d’environnement». 13% 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 ”)
Anticipated number of fundable research partners	10-12
Eligibility of project duration	2-3 years
Eligibility of a partner as a beneficiary institution	<p>Eligible institutions:</p> <ul style="list-style-type: none"> - Public research organisation or related-one^l such as EPST, EPIC, universities, university hospitals, non-university research institutes, foundations (max. rate of support: 100% of marginal costs, for organisations funded at “marginal cost”) - Enterprises: large & SMEs (max. rate of support: 45% of total costs for SMEs & 30% for larger companies) <p>Additional eligibility criteria:</p> <ul style="list-style-type: none"> - The coordinator (if from a French organisation) must belong to a public research organisation. - ANR will not provide double funding to finance projects or part of projects that have been funded through other national and international calls. ANR will cross-check the proposals submitted to ensure they have not been submitted to the ANR through other calls.

Eligibility of costs, types and their caps	Eligible costs include (but are not limited to) the following: personnel costs for temporary contracts; small equipment; consumables and animal costs; travel; and sub-contracting, if necessary, to carry out the proposed activities (sub-contracting costs max 50% of requested budget per partner). Eligible costs depend on the type of partner and consortium makeup. Please refer to the ANR Funding regulations for more details.
Conditions for PAO funding	French PAO can be funded as a partner if they perform research activities. Before applying, contact ANR for the rate of support your organisation is eligible to. Be aware that PAOs are often private entities and thus can generally not request a 100% support rate. Otherwise, French PAO can be funded as sub-contractor of a French partner. The maximum amount to sub-contract to a PAO is 20 k€ per project.
Submission of the proposal at the national level	No. Please note that some funding agencies that request submission at national level may be made available upon request applications (pre-proposals and full proposals) after the publication of the funding decision (i.e. SRC).
Submission of other information at the national level	No. However, please contact the national contact point for the ANR to enquire about their eligibility before submitting a proposal. In the proposal, justification of all costs must be provided especially sub-contracting, other direct costs...
Submission of financial and scientific reports at the national level	Financial reporting: must be completed according to ANR regulations, and the funding contract that beneficiaries will have to sign. Scientific reports: individual scientific reports are not required. However, ANR funded partners should contribute to the project report to be submitted by the coordinator of the project to EJP RD. These reports will be the basis for validation of yearly advancements of the project by ANR. Consortium agreement signed by all consortium's members and a data management plan must be provided.
Further guidance	<u>Règlement financier</u> Please read the Modalities for French applicants for this call on the ANR website.

FRENCH FOUNDATION FOR RARE DISEASES (FFRD), FRANCE

It is strongly advised that all applicants contact their EJP RD National/Regional Contact Point in good time before the submission of a proposal

Country	France
Funding organisation	Foundation for Rare Diseases (Fondation Maladies Rares) https://fondation-maladiesrares.org/eng/
National contact person	Fondation Maladies Rares Plateforme Maladies rares 96 rue Didot - 75014 Paris, France aap-bio@fondation-maladiesrares.com Diana Désir-Parseille - Phone : (33) (0) 1 58 14 22 81
Funding commitment	100 000€
Overheads	Overheads are not eligible costs.
Anticipated number of fundable research partners	TBD
Maximum funding per grant awarded to a partner	20 000€
Eligibility of project duration	3 years max
Eligibility of a partner as a beneficiary institution	Eligible institutions: - Public research institutes such as EPST, EPIC, universities, university hospitals, non-university research institutes (max. rate of support: 100% of marginal costs)
Eligibility of principal investigator or another research team member	The coordinator (if from a French institution) must belong to a public research organisation.
Eligibility of costs, types and their caps	Personnel costs for temporary contracts; small equipment; consumables and animal costs; travel; and sub-contracting, if necessary to carry out the proposed activities. Overheads are not eligible costs.

Conditions for PAO funding	No funding of PAOs
Submission of the proposal at the national level	No
Submission of other information at the national level	No
Submission of financial and scientific reports at the national level	Yes. Financial reporting is submitted to FFRD financial modalities and must be followed according to the contract that will be signed with the future beneficiaries. Scientific reports: individual scientific reports are not required. However, French partners should contribute to the central report to be submitted by the coordinator of the project.
Further guidance	

BMBF/PT-DLR, GERMANY

It is strongly advised that all applicants contact their EJP RD National/Regional Contact Point in good time before the submission of a proposal

Country	Germany
Funding organisation	German Federal Ministry for Education and Research (BMBF) www.gesundheitsforschung-bmbf.de
Management organisation	German Aerospace Center, DLR Project Management Agency (DLR-PT) www.pt-dlr.de
National contact person	<p>German Aerospace Center DLR Project Management Agency Health Division Clinical Research, University Medicine, Digital Health Heinrich-Konen-Straße 1 53227 Bonn Germany</p> <p>Dr. Katarzyna Saedler Dr. Michaela Fersch Dr. Ralph Schuster</p> <p>Tel: +49228-38212453</p> <p>E-Mail: SelteneErkrankungen@dlr.de</p>
Funding commitment	3 Mio€
Overheads	Overheads refer to "Gemeinkosten" (applicable for Helmholtz-centres and Fraunhofer-Society) as well as "Projektpauschale" (applicable for universities and university hospitals). The "Projektpauschale" generally will amount to 20% of the applied total project expenditure. For further information on the "Projektpauschale" please refer to https://foerderportal.bund.de/easy/module/profi_formularschrank/download.php?datei1=1750 (Pos. 0865) or contact the German national contact point for this EJP RD call.
Anticipated number of fundable research partners	Partners in about 10 projects

Maximum funding per grant awarded to a partner	Max. 300.000 EUR per consortium including overheads (i.e. if two German partners participate in a consortium, the sum of funding requested by both groups must not exceed 300.000 EUR)
Eligibility of project duration	Maximum 3 years

NATIONAL RESEARCH, DEVELOPMENT AND INNOVATION OFFICE (NKFIH), HUNGARY

It is strongly advised that all applicants contact their EJP RD National/Regional Contact Point in good time before the submission of a proposal

Country	Hungary
Funding organization	Ministry of Innovation and Technology
Management organization	National Research, Development and Innovation Office (NKFIH) http://nkfi.gov.hu/ ; http://nkfi.gov.hu/for-the-applicants
National contact person	National Research, Development and Innovation Office, Kéthly Anna tér 1, Budapest, H-1077, Hungary Dr. Előd Nemerkenyi Assistant of International Affairs, Department of Research and Development, NKFIH Phone: +36 1 8963987 E-mail: elod.nemerkenyi@nkfi.gov.hu Dr. Gábor Tóth head of unit, Unit for Medical and Biological Sciences, Department of Research and Development, NKFIH Phone: +36 1 8961727 E-mail: gabor.toth@nkfi.gov.hu
Funding commitment	300.000 €
Overheads	10% of the total costs of the project. Applicants should consult NKFIH '2019-2.1.7-ERA-NET' call regulations for details.
Anticipated number of fundable research partners	2
Maximum funding per grant awarded to a partner	Up to 150.000 €. If more than one partner applies from Hungary, their total requested funding should not exceed 150.000 euros.
Eligibility of project duration	Up to 3 years
Eligibility of a partner as a beneficiary institution	Universities, academic and public research institutions, public health institutions (university or non-university hospitals and clinics). An SME or a non-profit organization is eligible if its main activity is research according to its deed of foundation [category: 'research and knowledge-dissemination organisation' – see <i>Commission Regulation (EU) No. 651/2014 Article 2 (83)</i>]. All eligibility rules and criteria can be found in the '2019-2.1.7-ERA-NET' call regulations. It is strongly advised to contact NKFIH prior to submission regarding the eligibility criteria.
Eligibility of costs, types and their caps	100% of eligible research-related costs for basic (exploratory) research. The maximum indirect costs (overhead) are 10% of total costs. The maximum funding of 150.000 € per project includes the overhead. Detailed list of eligible costs (basically personnel, consumables, animals, equipment, travel, subcontracts, overhead) and guidelines to prepare the budget plan can be found in the call text and guideline of NKFIH '2019-2.1.7-ERA-NET'

	call (https://nkfi.gov.hu/palyazoknak/nkfi-alap/era-net-ejp-cofund-2019-217-era-net/palyazati-felhivas-2019-217-era-net).
Eligibility of principal investigator	The principal investigator must hold a Ph.D., D.Sc., or equivalent degree and be employed by an eligible institution. Researchers cannot participate in more than one proposal submitted to the same joint transnational call.
Conditions for PAO funding	No funding of PAOs.
Submission of the proposal at the national level	Hungarian applicants are strongly requested to contact NKFIH to confirm eligibility before submitting a proposal. Basic information should be provided to NKFIH, including applicant name and institution, as well as an estimation of the requested budget. Upon the EJP RD funding decision a proposal should be formally submitted to NKFIH in its electronic proposal system (EPTK). This is necessary for funding and managing the project by NKFIH.
Submission of financial and scientific reports at the national level	Yes, according to national regulations.

HEALTH RESEARCH BOARD (HRB), IRELAND

It is strongly advised that all applicants contact their EJP RD National/Regional Contact Point in good time before the submission of a proposal

Country	Ireland
Funding organization	Health Research Board, HRB, www.hrb.ie
National contact person	Dr. Amanda Daly Dr. Louise Drudy Email: eujointprogrammes@hrb.ie
Funding commitment	€370.000
Overheads	In accordance with the HRB Policy on Usage of research overheads , the HRB will contribute to the indirect costs of the research through an overhead payment of 30% of Total Direct Modified Costs (TDMC excludes student fees, equipment and capital building costs) for laboratory or clinically-based research and 25% of Total Direct Modified Costs if desk-based research.
Anticipated number of fundable research partners	1-2
Maximum funding per grant awarded to a partner	Up to €370,000 in total
Eligibility of a partner as a beneficiary institution	Lead applicants (Principal Investigators) from Ireland working in a recognized HRB Host Institution (Policy on Approval of HRB Host Institutions) are eligible for HRB funding in this call. Please see HRB Funding Schemes for HRB Guidance and FAQ for eligibility details.
Eligibility of costs, types and their caps	Funding available is inclusive of overheads and pension contributions <ul style="list-style-type: none"> • Salary related costs • Small equipment costs (€10,000) • Travel • Direct running costs (including costs for PAO participation) • FAIR data management costs • Dissemination and knowledge exchange costs • Overheads

Conditions for PAO funding	Patient advocacy organisations (PAOs) are not eligible to participate as a partner (Lead applicant). However, it is possible for the Lead applicant to request funds in their budget to support the activities of a PAO representative on the project. Eligible costs include consultation workshops, costs of participation in advisory groups, travel expenses, payments for time reviewing material (in line with the Host institutions policies) etc.
Submission of the proposal at the national level	For full proposal stage: Irish partners are asked to provide a copy of the submitted full proposal to the HRB National contact person following submission to the European Portal. Applicants from Ireland who are part of consortia invited to submit Full Applications will have to submit further budget and deliverables templates for the HRB. Templates will be provided by the HRB to invited applicants
Further guidance	<p>Applicants from Ireland must consult the HRB Guidance and FAQs for this call, for important eligibility information: HRB Funding Schemes</p> <p>Irish Partner(s) are not eligible for HRB funding for:</p> <ul style="list-style-type: none"> • Proposals involving basic biomedical research • Proposals seeking to evaluate a pilot or feasibility study <p>Proposals from Irish partners that include Human Embryonic Stem Cell Research will be deemed ineligible.</p>

CHIEF SCIENTIST OFFICE, MINISTRY OF HEALTH (CSO-MOH), ISRAEL

It is strongly advised that all applicants contact their EJP RD National/Regional Contact Point in good time before the submission of a proposal

Country	Israel
Funding organization	Chief Scientist office, Ministry of Health (CSO-MOH) http://www.health.gov.il/
National contact person	Dr. Irit Allon Phone: +972-2-5082167 Email: Irit.allon@moh.health.gov.il
Funding commitment	Up to 300.000 Euros
Overheads	10% of the entire project
Anticipated number of fundable research partners	Up to 2
Maximum funding per grant awarded to a partner	Up to 140,000 Euros (Additional EUR 20,000 for coordination of a consortium)
Eligibility of a partner as a beneficiary institution	Position in a university, research center or hospital. Research authority must approve position prior to submission.
Eligibility of costs, types and their caps	Materials and consumables; Travel (up to 10%); No salaries for applicants; No heavy equipment, Institutional overhead -10%
Conditions for PAO funding	No funding of PAOs
Submission of the proposal at the national level	Prior to submission of the pre-proposal to EJP RD, Israeli researchers need to submit to CSO-MOH an IL abstract approved by their research authority including budget distribution. The IL abstract will contain the project title, acronym and partners and will elaborate the part of the Israeli group in the project . IL abstract is <u>not</u> the abstract of the entire project. No submission of IL abstract can result in declaration of the consortium as ineligible. Researchers cannot apply for more than one grant from any ERA-Net funded by CSO-MOH or submit more than one proposal to any programme.

Further guidance

If the application involves human or animal experiments, bioethics approvals must be submitted with the application or up to 4 months later.

Please see detailed instructions at www.health.gov.il/research-fund

ITALIAN MINISTRY OF HEALTH (MoH-IT), ITALY

It is strongly advised that all applicants contact their EJP RD National/Regional Contact Point in good time before the submission of a proposal

Country	Italy
Funding organisation	Italian Ministry of Health www.salute.gov.it
Management organisation	Italian Ministry of Health General Directorate for Innovation & Research in Health
National contact person	Italian Ministry of Health Gaetano Guglielmi g.guglielmi@sanita.it Chiara Ciccarelli +39 06 5994 3919 c.ciccarelli@sanita.it Research.eu.dgric@sanita.it Anna Ceccarelli +39 06 5994 3835 a.ceccarelli-esterno@sanita.it
Funding commitment	1.5 M€
Overheads	Overhead (maximum 10% of the requested fund).
Additional documents required	The Italian Ministry of Health will check for the pre-eligibility of the applicants before the submission of the pre-proposals to speed up the eligibility check process. To this end, it is mandatory that the applicants fill out and return a pre-eligibility check form (sent to all IRCCSs) through the IRCCS Scientific Directorate or ISS Directorate of Human and Economic Resources using the WFR System (Code ER) before the submission of their pre-proposals to the Joint Call Secretariat. The form, completed and duly signed, has to be returned at least 10 working days before the pre-proposal submission deadline. Applicants will receive a written notification of their eligibility status. The pre-eligibility form can be downloaded here: http://www.salute.gov.it/imgs/C_17_pagineAree_4441_listaFile_itemName_0_file.pdf

Maximum funding per grant awarded to a partner	Max. 250.000 EUR per consortium (i.e., if two IRCCSs participate in a consortium, the sum of funding requested by both groups must not exceed 250.000 EUR)
Eligibility of project duration	Maximum 3 years
Eligibility of a partner as a beneficiary institution	Only Scientific Institutes for Research, Hospitalisation and Healthcare (IRCCS) are eligible. No academic and industrial partners are eligible.
Eligibility of costs, types and their caps	<p>Only the costs generated throughout the duration of the project can be eligible.</p> <ul style="list-style-type: none"> • Personnel (only ad hoc contracts/consultants/fellowships, max 50% of the requested fund). • Travel costs and subsistence allowances (max 10% of the requested fund) only if associated with training <ul style="list-style-type: none"> ◦ activities linked to the project. • Equipment (rent/leasing only, no limits), consumables (no limits), dissemination of results (publications, <ul style="list-style-type: none"> ◦ meetings/workshops etc.- max 1% of the requested fund). • Data handling and analysis (no limits). • Transfer of eligible funds abroad for leasing, sub-contracts, etc. is not allowed <p>Sub contracts are not allowed except in case of absolute necessity and to fund the Italian PAOs (see below); the costs for sub contracts need to be authorized by the It MoH in advance, following a detailed request. In this case, the pre eligibility must be requested 20 working days before the deadline of the call.</p>
Conditions for PAO funding	<p>Italian PAOs can be funded for a PAO a sub contractor through and IRCCS's budget if they fulfil the eligibility criteria of the EC. The maximum cost eligible for a sub contract is 25.000 Euros</p> <p>Italian PAOs can still participate in Consortia as "Collaborators" with their own funds</p>
Submission of the proposal at the national level	No
Submission of other information at the national level	After the joint EJP RD 2022 peer review has been completed and the final (scientific) ranking list has been performed and endorsed by the Call Steering Committee, the Ministry of Health will invite the principal investigators of the projects approved for funding to enter the formal national negotiations (according to national regulations). The funding of this projects are under the Ricerca Corrente IRCCS rules.
Submission of financial and scientific reports at the national level	Submission of an annual scientific and financial reports at the national level could be required according to the rules of the Ministry of Health Ricerca Corrente - IRCCS

Further information

The pre-eligibility form can be downloaded here:

http://www.salute.gov.it/imgs/C_17_pagineAree_4441_listaFile_itemName_0_file.pdf

REGIONAL FOUNDATION FOR BIOMEDICAL RESEARCH (FRRB), ITALY, (LOMBARDY)

It is strongly advised that all applicants contact their EJP RD National/Regional Contact Point in good time before the submission of a proposal

Country	Italy
Funding organization	Fondazione Regionale per la Ricerca Biomedica - Regional Foundation for Biomedical Research (FRRB)
National contact person	Paola Bello, Marcello De Amico Via Taramelli 12, 20124 – Milano Tel: +39 02 67650174 bandi@frb.it
Funding commitment	€ 1.500.000,00
Overheads	Up to 20% flat rate calculated on direct costs – Subcontracting costs excluded.
Anticipated number of fundable research partners	3-4
Maximum funding per grant awarded to a partner	Maximum € 500,000 per project. MAXIMUM TWO PARTNERS FROM LOMBARDY PER PROJECT. (If there are two Lombardy partners in the same consortium, the amount of 500,000 will be shared)
Eligibility of a partner as a beneficiary institution	<p>Eligible applicants:</p> <ul style="list-style-type: none"> •Public or Private Italian IRCCS (Scientific Institutes for Health Research, Hospitalization and Health Care) •Public Health Care Providers (ASST) •Universities (only in in partnership with one IRCCS, public or private, or ASST located in Lombardy and requesting funding to FRRB) •Research Institutes (only in in partnership with one IRCCS, public or private, or ASST located in Lombardy and requesting funding to FRRB) <p>All applicants must be located in Lombardy and their activities should take place in Lombardy. Enterprises and for-profit Organisation are NOT eligible.</p> <p>A Principal Investigator (PI) cannot simultaneously hold more than one FRRB active grant. PIs who are currently FRRB grant holders cannot apply to the EJP RD JTC 2022 unless their project is closed before the deadline for EJP RD JTC 2022 pre-proposals. A project is considered closed when the final financial and scientific reports have been sent to FRRB. This rule applies only to PIs (grant holders), not to their team members.</p>
Eligibility of costs, types and their caps	<p>Direct costs:</p> <ul style="list-style-type: none"> •Personnel (for public IRCCS and ASST, ONLY staff recruited specifically on the project) •Consumables, animals purchase, maintenance and breeding. •Equipment (on hire or eligible amortization rate).

	<ul style="list-style-type: none"> •Travel: max 10% of the total direct costs (<i>overheads and subcontracting costs excluded</i>) •Publications (only open access): max 5% of the total direct costs (<i>overheads and subcontracting costs excluded</i>). •Overheads: 20% flat rate calculated on direct costs (<i>Subcontracting costs excluded from this calculation</i>). •Subcontracting: max 20% of the total direct costs (<i>overheads costs excluded</i>) •Other direct costs: please include here other costs, including those related to patient involvement (insurance, reimbursement, etc.). <p>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.</p> <p>Only costs generated over the lifetime of the project will be considered eligible.</p>
Conditions for PAO funding	PAO are not eligible for FRRB funding
Submission of the proposal at the regional level	<p>It is not necessary to send the proposal to FRRB. However, FRRB requires a Pre-eligibility form.</p> <p>According to internal procedures, Regional Foundation for Biomedical Research (FRRB) will carry out an eligibility check to potential applicants prior to the submission of the pre-proposals.</p> <p>The eligibility check will be based on the verification of a dedicated form ("Pre-eligibility form"), also available on the FRRB institutional website, to be returned, by email, to FRRB (bandi@frb.it), duly completed and signed by the Principal Investigator at least 10 working days before the pre-proposal submission deadline.</p> <p>FRRB will provide feedback on the "<i>Pre-eligibility form</i>", ONLY in case of major non-eligibility issues.</p> <p>Principal Investigators (PIs) who submit a proposal without sending the "<i>Pre-eligibility form</i>" to FRRB beforehand will be automatically excluded.</p> <p>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.</p> <p>Information and instructions on how to fill the Pre-Eligibility check form will be published on the dedicated webpage (https://www.frrb.it/it/ejp-jtc-2022)</p> <p>Following the award, Lombardy beneficiaries will be requested to submit annual scientific and financial reports.</p>
Further guidance	Administrative and financial guidelines will be provided by FRRB in due time to the contact persons of the funded organisations.

TUSCANY REGION (RT/TuscReg), TUSCANY (ITALY)

It is strongly advised that all applicants contact their EJP RD National/Regional Contact Point in good time before the submission of a proposal

Country / Region	Italy
Funding organisation	Tuscany Region http://www.regione.toscana.it/
Regional contact person	Donatella Tanini Phone : +39 055 4383256 Teresa Vieri Phone : +39 055 4383289 Email: ejprare@regione.toscana.it Office for Health Research and investments, Directorate for Health, welfare and social cohesion Tuscany Region
Funding commitment	Up to 300.000 euros
Overheads	Up to 10% of the direct cost of the project, intended to cover the general cost of the institution that hosts the research team.
Anticipated number of fundable research partners	2-3
Maximum funding per grant awarded to a partner	Up to 300.000 euros
Eligibility of project duration	Up to 3 years
Eligibility of a partner as a beneficiary institution	A. Authorities of the Tuscany Health Service-SST (Local Health Authorities, University Hospitals) and the SST bodies that carry out institutional research activities (Fondazione Toscana Gabriele Monasterio and ISPRO Institute for Study, Prevention and Networking Oncology) located in the territory of Tuscany. B. Universities and other research institutes located in the territory of Tuscany. NB: Institutions referring to point B. are eligible only in partnership with institutions referring to point A.

<p>Eligibility of principal investigator or other research team member</p>	<p>The Principal Investigator must be affiliated to one of the eligible bodies</p>
<p>Eligibility of costs, types and their caps</p>	<p>Only costs generated over the lifetime of the project will be considered eligible.</p> <ul style="list-style-type: none"> - Personnel (ad hoc temporary contracts ONLY); - Consumables (no limit); - Equipment (on hire/leasing or eligible amortisation rate ONLY); - Travel (Up to 10% of the requested fund) Travel expenses and subsistence allowances associated with activities only linked to the project; - Other direct costs : <ul style="list-style-type: none"> o dissemination of results (publications, organization of meetings/workshops etc.- up to 5% of the requested fund); o data handling and analysis (no limit) o patients costs - subcontracting (up to 20%of the direct costs of the projects) - <i>Overheads</i> (Up to 10% of the direct cost of the project excepted subcontracting).
<p>Conditions for PAO funding</p>	<p>PAO cannot be directly funded by Tuscany Region in the framework of this call.</p>
<p>Submission of the proposal at the regional level</p>	<p>Yes Tuscany Region will grant an eligibility clearance to the potential applicants prior to the submission of their pre-proposals. The eligibility check will be performed by Tuscany Region offices after receiving a dedicated form (available on Tuscany Region institutional web-site or on request to ejprare@regione.toscana.it) duly filled and signed by the Tuscan Principal Investigator and by the legal representative of the beneficiaryThe form should be sent to Tuscany Region (ejprare@regione.toscana.it), at least, 10 working days before the pre-proposal submission deadline.</p>
<p>Submission of other information at the regional level</p>	<p>No</p>
<p>Submission of financial and scientific reports at the regional level</p>	<p>Yes/Submission of intermediate/final scientific and financial reports at the regional level could be required according to regional agreement</p>
<p>Further guidance</p>	<p>Financial guidelines will be published in due time on Tuscany Region's website.</p>

RESEARCH COUNCIL OF LITHUANIA (LMT), LITHUANIA

It is strongly advised that all applicants contact their EJP RD National/Regional Contact Point in good time before the submission of a proposal

Country	Lithuania
Funding organization	Lietuvos mokslo taryba (LMT) / Research Council of Lithuania http://www.lmt.lt
National contact person	Dr. Živilė Ruželė Phone: (+370) 676 14383, E-mail: zivile.ruzele@lmt.lt
Funding commitment	0.15M€
Overheads	Up to 30 % from those direct costs - personnel, subcontracting, contractual research, consultancy.
Anticipated number of fundable research partners	1
Maximum funding per grant awarded to a partner	100-150K€ (up to 100K€ for consortium partner or up to 150K€ for coordinator)
Eligibility of a partner as a beneficiary institution	Eligible for funding institutions are Lithuanian research and higher education institutions included in the Register of Education and Research institutions and public healthcare institutions. Beneficiary institution 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).
Eligibility of costs, types and their caps	Only costs generated during the lifetime of the project, related to project are eligible: staff, travel, consumables, subcontracts, contractual research, consultancy, equipment and instruments, dissemination of results, data handling and analysis, overheads (up to 30 % from the listed direct costs - staff, subcontracts, contractual research, consultancy).
Conditions for PAO funding	PAO can be a subcontractor or a 'project partner' of the eligible beneficiary institution (see section Eligibility of a partner as a beneficiary institution)
Submission of the proposal at the national level	No
Further guidance	Further information: https://www.lmt.lt/lt/mokslo-finansavimas/era-net-ir-kitos-koordinavimo-veiklos/europos-jungtine-programa-retos-ligos/3033 The proposals are submitted by the researcher(s) together with the eligible beneficiary institution. The beneficiary institution employs the principal investigator to work in the project and his workload must be at least 20 hours multiplied by the number

of months to execute the project. Hourly rates approved by the Chairman of the Lithuanian Research Council must be applied for the personnel costs. All other general rules for competitive funding of Research Council of Lithuania apply: <https://www.e-tar.lt/portal/lt/legalAct/0a8bead0577611e9975f9c35aedfe438/asr>

NATIONAL RESEARCH FUND (FNR), LUXEMBOURG

It is strongly advised that all applicants contact their EJP RD National/Regional Contact Point in good time before the submission of a proposal

Country / Region	Luxembourg
Funding organisation	Luxembourg National Research Fund - FNR www.fnr.lu
National contact person	Dr. Sean Sapcariu 2, avenue de l'Université L-4365 Esch-sur-Alzette Telephone: +352 691 362 831 Email: sean.sapcariu@fnr.lu
Funding commitment	0,30 M€
Overheads	
Anticipated number of fundable research partners	2 research partners
Maximum funding per grant awarded to a partner	The maximum funding cannot be larger than the funding commitment of the country
Eligibility of project duration	3 years
Eligibility of a partner as a beneficiary institution	Beneficiary institutions must be accredited by the Ministry in charge of public sector research. See website for details (https://www.fnr.lu/fnr-beneficiaries/) .
Eligibility of principal investigator or other research team member	Principle Investigators must follow the following guidelines: http://storage.fnr.lu/index.php/s/g4OPmRwEYhYwRkZ/download <ol style="list-style-type: none"> 1. He/she must have a proper employment contract with the eligible beneficiary institution at the starting date of the project. 2. The employment contract must last for the full duration of the research project. 3. He/she must be an experienced researcher who holds a doctoral degree at the date of the submission of the proposal.

Additional eligibility criteria	Luxembourgish principal investigators cannot be involved in more than 2 proposals submitted to this call.
Eligibility of costs, types and their caps	<p>Personnel costs; Consumables; Equipment (only depreciation costs); Travel (according to travel plan); Subcontracting (up to 25% of direct costs - needs detailed justification, includes all external services, project core activities cannot be subcontracted); Indirect costs</p> <p>Please see INTER application guidelines for more information (https://www.fnr.lu/funding-instruments/inter/)</p>
Conditions for PAO funding	FNR can fund PAOs which are eligible beneficiaries of FNR funding. For further information, please contact the FNR.
Submission of the proposal at the national level	All joint applications must also be submitted to the FNR by the Luxembourg-based scientist, along with the FNR INTER documents. This must be done no later than 5 days after the lead agency deadline, and must be done via the FNR Online Grant Management System.
Submission of other information at the national level	<p>The FNR requires the following other documents to be submitted to the FNR's grant management system :</p> <ul style="list-style-type: none"> - INTER Budget form, INTER Project plan, Gantt Chart
Submission of financial and scientific reports at the national level	The FNR expects annual reports and a final report for all projects funded through this call.
Further guidance	https://www.fnr.lu/fnr-international-cooperation/

NATIONAL CENTRE FOR RESEARCH AND DEVELOPMENT (NCBR), POLAND

It is strongly advised that all applicants contact their EJP RD National/Regional Contact Point in good time before the submission of a proposal

Country	Poland
Funding organization	National Centre for Research and Development (NCBR)
National contact person	Dr Marcin Chmielewski Department for International Cooperation, ul. Nowogrodzka 47a, 00-695 Warszawa, Poland Tel: (+48) 22 39 07 109, (+48) 571 226 666 marcin.chmielewski@ncbr.gov.pl
Funding commitment	0.6 Mio. €
Overheads	Maximum 25% of eligible project costs (excluding subcontracting)
Anticipated number of fundable research partners	1 - 3
Maximum funding per grant awarded to a partner	Maximum 200 000 € per project, regardless of the number of Polish partners in the project consortium.
Eligibility of a partner as a beneficiary institution	<p>Following entities are eligible to apply:</p> <ul style="list-style-type: none"> • Micro, Small, Medium and Large enterprise. • Research organization. • Group of entities (within the meaning of art. 37 section 1, point 1a of The Act of 30 April 2010 on the National Centre for Research and Development, published in Journal of Laws item 1861, 2020); <ul style="list-style-type: none"> • Organization must be registered in Poland. • For enterprises it is strongly advised to state in the Pre-proposal application form the KRS number of the enterprise and the size of the enterprise (micro/small, medium, large). • A condition for the participation of a group of entities as the Applicant in the competition is its formal existence on the date of submission of the pre-proposal, confirmed by its members concluding, at least conditionally, agreement on the creation of a group of entities. • Please note that group of entities counts as two project partners from Poland (it meets the limit on the number of participants from the same country, please refer to call text for details).
Eligibility of costs, types and their caps	<p>The eligible costs shall be the following:</p> <ol style="list-style-type: none"> 1. personnel costs (researchers, technicians and other supporting staff to the extent employed on the research project); 2. operating costs including costs of instruments, equipment, technical knowledge, patents, costs for buildings and land, costs of materials, supplies and similar products incurred directly as a result of the research activity.

	<p>3. cost of contractual research, costs of consultancy and equivalent services used exclusively for the research activity; this cost type cannot account for more than 70% of all eligible costs of a project; the subcontracting can be obtained from consortium partner only in justified case, this need will be verified by a national expert panel.</p> <p>4. additional overheads incurred indirectly as a result of the research project; those costs cannot account for more than 25% of eligible project costs and are counted as a multiplication by percentage given above and the rest of direct costs, excluding subcontracting (3); It means $4 = (1+2) * 25\%$.</p>																				
	<p>Funding quota of Polish participants can be up to 100% for universities or research organizations. In the case of enterprises, funding quota will be decided on a case-by-case basis depending on the size of the company, type of research/development, risk associated with the research activities and commercial perspective of exploitation, under the Regulation of the Minister of Science and Higher Education of 19 August 2020 on criteria and rules on granting state aid by the National Centre for Research and Development, published in Journal of Laws item 1456, 2020.</p>																				
	<table border="1"> <thead> <tr> <th></th> <th>Large Enterprises</th> <th>Medium Enterprises</th> <th>Small Enterprises</th> <th>Universities and research organizations</th> </tr> </thead> <tbody> <tr> <td>Fundamental/Basic Research</td> <td>Not eligible</td> <td>Not eligible</td> <td>Not eligible</td> <td>Not eligible</td> </tr> <tr> <td>Industrial/Applied Research</td> <td>Up to 50+15 (max 65 %)</td> <td>Up to 50+10+15 (max 75 %)</td> <td>Up to 50+20+15 (max 80 %)</td> <td>Up to 100 %</td> </tr> <tr> <td>Experimental development</td> <td>Up to 25+15 (max 40 %)</td> <td>Up to 25+10+15 (max 50 %)</td> <td>Up to 25+20+15 (max 60 %)</td> <td>Up to 100 %</td> </tr> </tbody> </table>		Large Enterprises	Medium Enterprises	Small Enterprises	Universities and research organizations	Fundamental/Basic Research	Not eligible	Not eligible	Not eligible	Not eligible	Industrial/Applied Research	Up to 50+15 (max 65 %)	Up to 50+10+15 (max 75 %)	Up to 50+20+15 (max 80 %)	Up to 100 %	Experimental development	Up to 25+15 (max 40 %)	Up to 25+10+15 (max 50 %)	Up to 25+20+15 (max 60 %)	Up to 100 %
		Large Enterprises	Medium Enterprises	Small Enterprises	Universities and research organizations																
Fundamental/Basic Research	Not eligible	Not eligible	Not eligible	Not eligible																	
Industrial/Applied Research	Up to 50+15 (max 65 %)	Up to 50+10+15 (max 75 %)	Up to 50+20+15 (max 80 %)	Up to 100 %																	
Experimental development	Up to 25+15 (max 40 %)	Up to 25+10+15 (max 50 %)	Up to 25+20+15 (max 60 %)	Up to 100 %																	
<p>Only Industrial/Applied Research and Experimental Development will be funded. Other type of activities (e.g. coordination, dissemination, management) is not eligible for funding as separate research tasks in the project schedule.</p>																					
Conditions for PAO funding	Funding is only available for project partners, meeting eligibility criteria given above.																				
Submission of the proposal at the national level	Polish Participants will be informed and invited to submit Polish proposal once the international evaluation and the ranking list will be established.																				
Further guidance	Please refer to full call documentation.																				

SLOVAK ACADEMY OF SCIENCES (SAS), SLOVAKIA

It is strongly advised that all applicants contact their EJP RD National/Regional Contact Point in good time before the submission of a proposal

Country	Slovakia
Funding organization	Slovak Academy of Sciences (SAS)
National contact person	Zuzana Cernakova, PhD. International Cooperation Dpt., SAS Phone: +421257510118 Email: cernakova@up.upsav.sk
Funding commitment	120,000 €
Overheads	Up to 20% of the direct costs
Anticipated number of fundable research partners	1
Maximum funding per grant awarded to a partner	120,000 €
Eligibility of a partner as a beneficiary institution	<p>Only research institutes and/or centres of the Slovak Academy of Sciences are eligible organisations for funding by the SAS (up to 100%). The main applicant must have an employment contract with the SAS institute/centre on behalf of which the application is being submitted. If his/her contract is on a part-time basis, it must be for more than 50% of standard working time. All members of the applicant's team except doctoral students must, too, have employment contracts with the same or another SAS institute/centre.</p> <p>Applicants from other Slovak R&D centres (universities and/or other organisations from Slovakia) can join project consortia only as collaborators who must secure their own funding.</p>
Eligibility of costs, types and their caps	Funding available for eligible Slovak researchers is up to 120,000 EUR per project (i.e. 40,000 EUR per year) in accordance with the SAS Presidium's resolution no. 136 (of 14 October 2021), of which 45,000 EUR is an in-kind contribution (spoluúčasť) of the respective SAS institute or centre in the form of permanent salaries. This must be declared in a Letter of Commitment sent to the national contact point by the application deadline. A template will be published alongside the Call announcement at www.sav.sk in the 'International Cooperation' section (Medzinárodná spolupráca).

	<p>Costs other than the in-kind contribution (Personnel costs, Consumables, Travel costs, Equipment, Other direct costs, Overheads) up to 75,000 EUR must comply with specific rules and limits outlined in the financial rules for awarding SAS grants for international research projects available at:</p> <p>https://www.sav.sk/?lang=sk&doc=services-news&source_no=25&news_no=7569</p> <p>Applicants are strongly encouraged to read the said document carefully and to contact the national contact point before submission in order to ensure compliance.</p>
Conditions for PAO funding	<p>The SAS does not fund PAOs/patient representatives.</p>
Submission of the proposal at the national level	<p>Submission of an application at the national level will be required once the international evaluation and the ranking list have been performed and endorsed by the Call Steering Committee. Only the Slovak partners of the projects recommended for funding will be invited to submit the national-level application. The final decision on funding of the Slovak partners must be approved by the SAS Presidium.</p>
Further guidance	<ul style="list-style-type: none"> • www.sav.sk • 133 Act of February 19, 2002, on the Slovak Academy of Sciences • Financial rules for awarding SAS grants for international research projects

INSTITUTE OF HEALTH CARLOS III (ISCIII), SPAIN

It is strongly advised that all applicants contact their EJP RD National/Regional Contact Point in good time before the submission of a proposal

Funding Organisation	National Institute of Health Carlos III Instituto de Salud Carlos III (ISCIII) www.isciii.es
National Funding Programme	Acción Estratégica en Salud (AES 2021) http://www.isciii.es/ISCIII/es/contenidos/fd-investigacion/fd-financiacion/convocatorias-ayudas-accion-estrategica-salud.html
National Contact Point for the 10th call of E-RARE	Ignacio Baanante Email : ibaanante@isciii.es Tel : (+34) 9182 22576 Maria Druet Email: mdruet@isciii.es Tel: (+34) 91822 2530
Initial funding pre-commitment	2 M€ <i>8-12 groups tentatively envisaged to be funded.</i>
Maximum funding per awarded Spanish project partner	Maximum funding from ISCIII per awarded Spanish project partner r <ul style="list-style-type: none"> • Up to 175,000 € per partner (overheads included) • Up to 250,000 € per coordinator ((overheads included)
Eligible institutions	<ul style="list-style-type: none"> • Hospitals, primary health care or public health administration of the Spanish National Health System (SNS) These institutions may manage research via a foundation regulated in accordance to the Spanish Act 50/2002, of December 26th (a copy of the foundation's statutes may be submitted). • Accredited Health Research Institutes (Institutos de Investigación Sanitaria acreditados, IIS) Accredited according to the RD 339/2004, of February 27th or RD 279/2016 (These institutions may manage research via a foundation regulated according to the Spanish Act 50/ 2002, of December 26th)

<p>Additional eligibility criteria</p>	<p>https://eng.isciii.es/eng.isciii.es/QuienesSomos/IIS/Paginas/Acreditacion.html</p> <ul style="list-style-type: none"> • CIBER. Team members applying to the call must be from at least two groups belonging to CIBER in two different home institutions and one of these two should be a Hospital, primary health care or public health administration of the Spanish National Health System (SNS) or Accredited Health Research Institutes (Institutos de Investigación Sanitaria acreditados, IIS). Please contact Cristina Rodríguez (cristina.rodriguez@ciberisciii.es) for more information related to CIBER's eligibility. • Academia or Other Research Centers. These entities can only participate if they apply together with Hospitals, primary health care or public health settings of the Spanish National Health System (SNS), or Accredited Health Research Institutes (Institutos de Investigación Sanitaria acreditados, IIS) in the same proposal. It is not allowed to apply independently, thus there must be two beneficiary Spanish institutions requesting funding to ISCIII in the same proposal. • Institutions belonging to the INB/ELIXIR-ES (The Spanish Institute of Bioinformatics) as long as the participants are those from the Institute of Bioinformatics. <p>PLEASE NOTE:</p> <ol style="list-style-type: none"> Applicants from ISCIII are eligible. Eligibility criteria from AESI 2021 apply. Durations of national grants are up to 3 years. Same institution cannot participate with more than one partner in the same project proposal. Only one PI per beneficiary institution may be funded within the same proposal. IPs with ongoing EJP RD projects in 2022 can not apply to the current call unless the alive project or the new application is as Coordinator There is no other incompatibility with AES 2021. A given PI can apply only once to this call.
<p>Eligibility of PI and team members</p>	<ul style="list-style-type: none"> • Principal Investigators (PI) can only participate in one project proposal per call. • Principal Investigators (PIs) belonging to any IIS could apply from the IIS or from any of the institutions belonging to the IIS • The Principal Investigator (PI) and all members of the research group must belong to the eligible institution or be affiliated to CIBER or an IIS.

	<p>Excluded personnel as Principal Investigator (PI):</p> <ul style="list-style-type: none"> • Those undergoing a postgraduate training in Health Specialization (MIR, EIR, FIR, QIR, BIR, PIR, RFIR) • Those undergoing research training (e.g. PhD students, or “Río Hortega” contracts). • Researchers contracted by a RETIC/RICOR. • Those undergoing postdoctoral training (e.g. “Sara Borrell” or “Juan de la Cierva” contracts).
<p>Eligible costs</p>	<ul style="list-style-type: none"> • Personnel costs for temporary employment contracts (scholarships are not eligible), with a maximum of 36 PM in total for the personnel contracts altogether. • Current costs, small scientific equipment, disposable materials, travelling expenses and other costs as included in AES 2022 that can be justified as necessary to carry out the proposed activities. • Overheads, according to AES 2022. • National applications will be required by ISCIII. Spanish Applicants should periodically check in the web page of ISCIII if they are qualified. ISCIII may not send invitations to the mandatory national phase. • Double funding of the same concept is not allowed.
<p>Conditions for PAO funding</p>	<p>The ISCIII does not fund PAOs/patient representatives.</p>
<p>National phase</p>	<ul style="list-style-type: none"> • National applications will be required by ISCIII. Spanish Applicants should periodically check in the web page of ISCIII if they are qualified. ISCIII may not send invitations to the mandatory national phase. • Double funding of the same concept is not allowed. <p>Due to administrative and legal regulations, the National Institute of Health Carlos III declares the end of September 2022 as national deadline for the decision on fundable project consortia which include Spanish partners to be funded by ISCIII. Any concerned applicant in a proposal for which no final decision has been made by the deadline, could be declared not fundable by ISCIII.</p>
<p>Requirements on data and repositories</p>	<ul style="list-style-type: none"> • Researchers funded by ISCIII must make public the human genomic data, as well as relevant data (phenotype and exposition data) generated inside the funded project and will use open access repositories. Researchers must also make public all the necessary information for the interpretation of these genomic data, including lab protocols, data instruments survey tools. Regarding genomic data it is understood association of complete genomes (GWAS), matrixes of de polymorphism of a single nucleotide (SNP) and sequence of genome, and transcriptomic, metagenomic, epigenomic and

	<p>gene expression data. The researchers whose projects are funded by ISCIII are recommended to store their scientific data at the "ELIXIR Core Data Resources" or if non-European repositories or data bases they must be certified by ELIXIR or the US National Center for Biotechnology Information (NCBI).</p> <ul style="list-style-type: none"> ISCIII may not fund project that requires the construction of new repositories without decommissioning plans or ensured sustainability after the project's end.
<p>Requirements for clinical studies</p>	<p>Spanish groups participating in a proposal performing a clinical study are encouraged to contact and include as members of the team personnel from the Clinical Research Unit (Unidades de Investigación Clínica y Ensayos Clínicos - UICEC) of their institutions. These Units belong to ISCIII's platform that supports Clinical Research and participate in ECRIN-ERIC. Find here the list of UICECs. For additional information please contact: sectec.scren.hcsc@salud.madrid.org or Tel.: (+34) 91 330 38 58</p>
<p>Acknowledgements</p>	<p>Any publication resulting from the granted projects must acknowledge "Award no. XX by ISCIII through AES 2021 and within the European Joint Programme Rare Diseases framework" even after the end of the project.</p>

SWEDISH RESEARCH COUNCIL (SRC), SWEDEN

It is strongly advised that all applicants contact their EJP RD National/Regional Contact Point in good time before the submission of a proposal

Country	Sweden
Funding organization	Swedish research council, SRC www.vr.se
National contact person	Louise Rügheimer E-mail: Louise.rugheimer@vr.se Phone: +46(0)8 122 13 618
Funding commitment	20 MSEK, approximately 1 820 000€
Overheads	The grant amount includes indirect costs.
Anticipated number of fundable research partners	2-4
Maximum funding per grant awarded to a partner	For Swedish participation in a consortium, the maximum amount that may be applied for is 4 500 000 SEK (approximately 450 000 EUR) for a consortium with 1 Swedish partner, or 6 000 000 SEK (approx. 600 000 EUR) if the consortia contain two Swedish partners. Use the exchange rate of 1 EURO=10,13 SEK to calculate actual grant amounts for the application
Eligibility of a partner as a beneficiary institution	Not applicable
Eligibility of costs, types and their caps	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.
Conditions for PAO funding	

<p>Submission of the proposal at the national level</p>	<p>All Swedish project leaders participating in the call for support from the Swedish Research Council shall also submit a parallel application using the Swedish Research Council's application system Prisma. The application form in Prisma can be reached from the call text at the SRC website: Swedish and English</p> <p>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 will result in the Swedish partner being declared ineligible.</p> <p>All Swedish applicants must communicate with the EJP RD national contact person regarding their intention to participate in the call, before submission of the consortium application.</p>
<p>Further guidance</p>	<p>See national call texts for all national requirements: Swedish and English</p>

VINNOVA, SWEDEN

It is strongly advised that all applicants contact their EJP RD National/Regional Contact Point in good time before the submission of a proposal

Country	Sweden
Funding organization	Vinnova www.vinnova.se
National contact person,	Gunnar Sandberg, gunnar.sandberg@vinnova.se +46 8 4546445
Funding commitment	9MSEK, 900 000 Euro.
Overheads	
Anticipated number of fundable research partners	3-5
Maximum funding per grant awarded to a partner	The maximum amount of funding for Swedish participation is 3 million SEK for 1 Swedish partner and 4.5 million SEK for 2 Swedish partners. The consortia need to include at least one Swedish partner from industry when applying for funding from Vinnova.
Eligibility of a partner as a beneficiary institution	The Swedish participation applying for funding from Vinnova should have at least one partner from industry. Universities, public research institutes, healthcare providers and industry. For more information see Vinnova.se
Eligibility of costs, types and their caps	Universities, public research institutes and public healthcare providers may receive funding of up to 100 % of their eligible costs, provided that the project is part of their non-economic activities. Large companies can apply for 20 % of their eligible costs. Small and medium sized companies can apply for 70 % of their eligible costs The eligible cost are defined in: Terms and conditions for Vinnova funding Vinnova
Conditions for PAO funding	Approved administrating organisations see Vinnova.se

<p>Submission of the proposal at the national level</p>	<p>General condition for grant application (Find the right funding Vinnova) Swedish applicants must also apply via the Vinnova web portal (Find the right funding Vinnova), where the Swedish project partner/s should add the requested information and upload the EJP-RD JTC 2022 project plan. Please follow the instructions on the national call page. If you have any questions don't hesitate to email or call one of the Vinnova contact persons.</p>
<p>Further guidance</p>	<p>General condition for grant application (Find the right funding Vinnova)</p>

SWISS NATIONAL SCIENCE FOUNDATION (SNSF), SWITZERLAND

It is strongly advised that all applicants contact their EJP RD National/Regional Contact Point in good time before the submission of a proposal

Country	Switzerland
Funding organisation	Swiss National Science Foundation (SNSF)
National contact person	Dr Tobias Braun Division Biology and Medicine Wildhainweg 3, P.O. Box, CH-3001 Bern Phone: +41 31 308 21 67 tobias.braun@snf.ch
Funding commitment	1 Mio Swiss Francs (equivalent to approx. 0.9 Mio €)
Overheads	Overhead costs may not be included in the Swiss project budget. Overhead contributions, calculated on the basis of the total research funding given to a particular institution through all SNSF funding instruments, are paid directly to the applicant's institution on a yearly basis.
Anticipated number of fundable research partners	3-4, each Swiss applicant may be partner in only one EJP RD JTC 2022 proposal (Art.7.3, SNSF Regulations on Project Funding).
Eligibility of a partner as a beneficiary institution	n.a.
Eligibility of principal investigator or other research team member	<p>Where not otherwise specified, the SNSF Funding Regulations, in particular, the SNSF Regulations on Project Funding apply:</p> <ul style="list-style-type: none"> • SNSF Funding Regulations • General Implementation Regulations for the Funding Regulations • SNSF Regulations on Project Funding <p>All Swiss partners in EJP RD projects must meet the eligible criteria for applicants in SNSF Project Funding. Swiss partners who have not previously obtained a project grant from division Biology and Medicine must contact the national contact point to confirm their eligibility as an applicant prior to submitting a proposal to the EJP RD JTC 2022.</p> <p>Foreign members of the international consortia applying for funding through the EJP RD JTC 2022 cannot be declared as "project partners" in the sense of Art. 11.2 of the SNSF Funding Regulations and may not receive any funding through the Swiss partner.</p> <p>Article 17 of the SNSF Funding Regulations applies, i.e. EJP RD proposals with overlapping funding periods with ongoing SNSF grants are only allowed if the two research projects are thematically distinct and pursue different goals.</p> <p>Grants given to Swiss partners will be managed according to SNSF Funding Regulations.</p>
	Please note: The SNSF exclusively funds research conducted for non-commercial purposes. Pursuant to the Swiss 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 full academic freedom.
Eligibility of costs, types and their caps	<p>According to the SNSF Regulations on Project Funding (article 8), the following costs may be covered:</p> <ul style="list-style-type: none"> - the salaries of scientific and technical staff in research projects within the scope of the salary ranges and rates prescribed by the SNSF; - material costs that are directly related to the research work, namely material of enduring value, expendable items, field expenses, travel expenses, third-party charges, cost of computing time and data as well as of providing open access to research data; - direct costs incurred through the use of research infrastructure linked to the research work; - costs for the organisation of conferences and workshops in connection with the funded research; - costs for national and international cooperation and networking activities carried out in connection with the funded research.
Conditions for PAO funding	According to our eligibility criteria, PAO are <u>not</u> eligible as partners.
Submission of the proposal at the national level	Swiss partners are required to submit the pre-proposal and the full proposal to www.mySNF.ch together with the submission of the respective proposals to the EJP RD Joint Call Secretariat. For this, Swiss partners need a personal account on www.mySNF.ch . The SNSF office may ask Swiss partners to submit supplemental information as needed.
Submission of financial and scientific reports at the national level	Yearly financial reports for the use of SNSF funds and a scientific report at the end of the project.
Further guidance	Consortia including Swiss partners must submit a data management plan (DMP) which complies with the SNSF policy on open research data .

NETHERLANDS ORGANIZATION FOR HEALTH RESEARCH AND DEVELOPMENT (ZonMw), THE NETHERLANDS

It is strongly advised that all applicants contact their EJP RD National/Regional Contact Point in good time before the submission of a proposal

Country	The Netherlands
Funding organization	ZonMw, The Netherlands organisation for health research and development, PO Box 93245, 2509 AE The Netherlands, https://www.zonmw.nl/nl/
National contact person	Sonja van Weely, PhD Kirsten Wilkens, MSc Email: ERareJTC2018@zonmw.nl (preferred) Tel. +31 70 349 5467/5220 Tel. +31 70 515 0390
Funding commitment	1.6 million euro
Overheads	Overheads are not eligible costs for ZonMw
Anticipated number of fundable research projects	6-7 projects
Maximum funding per grant awarded to research partner or research project	Up to 250,000.- euro for a Dutch research project partner or coordinator for a 3-year project proposal. In case a project consists of two Dutch research partners (only possible if one partner classifies as Early Career Researcher), the total amount of the ZonMw funding for the project is still maximized to 250,000.- euro.
Eligibility of a partner as a beneficiary	Category A. Dutch research organisations as defined in EU state aid legislation. ¹ Category B. Other Dutch rare diseases centers of expertise that are recognised by the Dutch Ministry of Health. • Please read the eligibility of costs for categories A and B very carefully below.

¹ Framework for state aid for research and development and innovation (2014/C 198/01), Article 15(ee).

<p>research institution</p>	<ul style="list-style-type: none"> • Max. 1 application as coordinator is allowed. • A specific Dutch researcher is allowed to take part in max. 2 applications. • 1 Dutch researcher per application is allowed; a second Dutch researcher in an application is only allowed in case it concerns an Early Career Researcher (see 4.1 in the Guidelines for Applicants). • The track record of the principle investigator (PI) is part of the assessment. • Cofinancing (in cash or in kind) is encouraged.
<p>Eligibility of principal investigator or other research team member</p>	<p>The principle investigator (PI) should have (or get upon granting of the project) an employment contract at the eligible institution for at least the duration of the project; the PI does not need to have a permanent position at the institute. A signed letter from the department head or other responsible official of the institute has to be submitted to ZonMw at the deadline of application of the full proposal in which information on the employment contract of the PI is indicated. Furthermore, in this letter the department head or other responsible official should also guarantee that the applicant will have the time and facilities to perform the research properly and according to plan. The PI should show strong commitment to (the results of) the project.</p>
<p>Eligibility of costs, types and their caps</p>	<ul style="list-style-type: none"> • Aid for the concerning activities to the organisations in category A does not result in state aid according to the Framework for State Aid for Research and Development and Innovation. Organisations in category A must meet the criteria of the research and knowledge-dissemination organisation (hereinafter referred to as "research organisation") in accordance with the Framework for State Aid for Research and Development and Innovation (2014/C 198/03). • Aid to organisations in category B is state aid and will be granted under the General Block Exemption Regulation: EC REGULATION No 651/2014, section 25 ('GBER'). All relevant conditions of the GBER apply, including but not limited to: <ul style="list-style-type: none"> • section 1.4 (no outstanding recovery order following a previous Commission decision, no aid to undertakings in difficulty) • section 1.5 (non violation of Union law by means of conditions or financing method), • section 8 (cumulation) • consideration 18 (the work on the aided project or activity starts only after the beneficiary has submitted a written application for the aid) • The aid intensity depends on the Technology Readiness Level (TRL) of the activities of the Dutch partner, which should be clearly specified in a separate document to be sent to ZonMw 1 week before the deadline of application of the full proposal as well as the other mandatory declarations: <ul style="list-style-type: none"> ▪ Declaration order for recovery of state aid (in Dutch). ▪ Declaration company not in difficulty.

- Declaration of accumulation of state aid.

Eligible costs of research projects executed by organisations in **category A** are (see the [ZonMw grant terms and conditions from 1st July 2013](#)):

- personnel. Scientific personnel has to be appointed at a scientific institution in The Netherlands.
- consumables, animals,
- equipment,
- travels,
- costs for dissemination of results (implementation)
- In most cases (e.g., in case of university/university medical centers) overhead is not allowed and the salary scales of [VSNU](#) (universities) or [NFU](#) (University Medical Centres) have to be used.

Please use the ZonMw budget formats as basis for the [budget calculations](#).

Eligible costs of research and development projects executed by organisations in **category B** shall be allocated to a specific category of research and development and may be the following:

- (a) personnel costs: researchers, technicians and other supporting staff to the extent employed on the project;
- (b) costs of instruments and equipment to the extent and for the period used for the project. Where such instruments and equipment are not used for their full life for the project, only the depreciation costs corresponding to the life of the project, as calculated on the basis of generally accepted accounting principles are considered as eligible.
- (c) other operating expenses, including costs of materials, supplies and similar products, incurred directly as a result of the project.

National Funding rates

Funding quota of Dutch participants can be up to 100% for organisations in **category A**.

The funding quota for organisations in **category B** will be decided on a case-by-case basis depending on the size of the company, type of research/development in accordance with GBER, section 25.5 and 25.6.

Fundamental/ Industrial/Experimental development will be funded. Other type of activities (e.g. coordination, management) is not eligible for funding as separate research tasks in the project schedule.

Conditions for PAO funding

Patient advocacy organisations (PAO) that are located in the Netherlands can be eligible and may apply for maximum of 20,000.- euro subsidy for joining a project of three years under the de minimis aid ([EC Regulation 1407/2013](#)).

All relevant conditions of this Regulation apply, including but not limited to:

Art. 3.2: The total amount of de minimis aid granted in the Netherlands to a single PAO shall not exceed 200,000.- euro over any period of three fiscal years. The PAO part of the grant application will be rejected if:

1. one of the situations referred to in Article 1 of the de minimis regulation occurs which would preclude application of the de minimis regulation; or
2. the de minimis ceiling would be exceeded by allocation of the requested grant funding; or
3. your application does not meet the conditions of the de minimis regulation in some other way.

- If several patient organisations located in the Netherlands are involved in the project, the maximum of 20,000,- euro has to be divided. The de minimis regulation has to be used for every PAO in a project.
- A specific PAO is allowed to take part in max. 2 applications.
- A PAO cannot be coordinator of a project.

Eligibility PAOs

Eligible PAOs are defined as not-for-profit organisations located in the Netherlands, which are patient focused, and where patients and/or carers and/or family members of patients represent a majority of members in governing bodies. These are:

- Umbrella organisations (e.g. representing either European organisations and/or national umbrella organisations for rare diseases);
- European rare disease specific organisations (i.e. representing national organisations or individual patients on rare diseases) and
- National rare disease specific organisations.

PAOs has to fulfil the following criteria and this will be checked at the eligibility phase of the preproposal. Please send the following information to ERareJTC2018@zonmw.nl before **February 16, 2022**:

(1) to prove that you follow these criteria,

(2) information on the proposed activities of the PAO mentioned in the preproposal,

(3) include a

https://www.zonmw.nl/fileadmin/zonmw/documenten/Subsidieoproepen/Staatssteun/Verklaring_de-

[minimissteun.pdf](#) (in Dutch) in which you list the de minimis aid that you have received in the two previous tax years and in the current tax year.

Legitimacy:

- Represent rare diseases according to EU prevalence criteria (5/10 000) as defined in the: EU Regulation on Orphan Medicinal Products (1999), Commission Communication on Rare Diseases 2008), Council Recommendation on an Action on Rare Diseases (2009), and Directive on Patients' Rights in Cross-Border HealthCare (2011)
- the organisation should be formally established and registered as a not-for-profit organisation in The Netherlands
- **Mission/objectives:** the organisation shall have its mission/objectives clearly defined and should agree to have it/them published on the EJP RD website.
- **Activities:** the organisation shall have, as part of its activities, a specific interest in rare diseases which should be documented (e.g. through a report published on the organisation website).
- **Representation:** the organisation shall be representative of rare disease patients within the Netherlands or throughout the EU/EEA.
- **Structure:**
 - The organisation should have governing bodies which includes a majority of rare disease patients or family members of rare disease patients.
 - Includes in its governing structure a designated representative legally authorised to sign a contract with ZonMw
- **Accountability:**
 - With proven activities such as rare disease patient support and/or advocacy activities and/or rare disease research
 - Statements and opinions of the organisation should reflect the views and opinions of its members and adequate consultation procedures with those members should be in place. In particular, the organisation should ensure that the appropriate flow of information is in place to allow dialogue both ways: from and towards its members.
- **Transparency:**
 - The organisation shall be financially independent, particularly from the pharmaceutical industry (max. 50% of funding from several companies) and disclose to ZonMw its sources of funding both public and private by providing the name of the bodies and their individual financial contribution, both in absolute terms and in terms of overall percentage of the organisation budget. Any relationship with corporate

	<p>sponsorship should be clear and transparent. This information shall be communicated to ZonMw on an annual basis.</p> <ul style="list-style-type: none"> The organisation shall publish on its website the registered statutes, sources of funding, and information on their activities. <ul style="list-style-type: none"> Communication To facilitate communication, one contact person shall be identified for each PAO. <p>Eligible costs are</p> <p>(a) personnel for the specific research project;</p> <p>(b) travels;</p> <p>(c) dissemination of results (implementation).</p> <p>Non-eligible costs are office and IT equipment (workstation, mobile phone, tablets, etc.) and overhead.</p>
<p>Submission of the proposal at the national level</p>	<ul style="list-style-type: none"> Submission of the full proposal to ZonMw will be carried out once the international evaluation and the ranking list have been performed and endorsed by the Call Steering Committee and European Commission. ZonMw will send a letter to invite the selected researcher or PAO to submit the granted full proposal. The Dutch consortium partners in honoured consortia have to comply with ZonMw procedures for granted projects (e.g. uploading via ProjectNet or AIMS - including the ZonMw budget format, and reporting annually). Scientific personnel has to be appointed at a scientific institution in The Netherlands. Granted consortia with a Dutch partner have to draw up and sign a Consortium Agreement in which also the intellectual property rights are incorporated. A final draft version of the Consortium agreement (approved by all parties but not yet signed) will be required in order to assess conformity with applicable European state aid law, IP conditions and the General Terms and Conditions Governing Grants of ZonMw as of July 2013. If the Consortium agreement is rejected, the funding by ZonMw cannot be granted. For more details and conditions: https://www.zonmw.nl/en/research-and-results/co-financing/grants-and-collaborationscontributions-from-third-parties/ Before the start of the granted project the Dutch researcher needs to compose a data management plan and complete key items to explain how to make the data collection from the Dutch part of the research project FAIR.

	<p>If a co-financer is not included in the consortium agreement, a signed Letter of Commitment needs to be submitted to ZonMw with the application. For more details (in Dutch): ZonMw financiële aspecten bij samenwerking.</p>
Further guidance	<ul style="list-style-type: none">• Collaboration with patient organisations is recommended; see also 5.5 in the Call text.• The ZonMw grant terms and conditions from 1st July 2013) apply for Dutch consortium partners.

THE SCIENTIFIC AND TECHNOLOGICAL RESEARCH COUNCIL OF TURKEY (TUBITAK), TURKEY

It is strongly advised that all applicants contact their EJP RD National/Regional Contact Point in good time before the submission of a proposal

Country	Turkey
Funding organization	The Scientific and Technological Research Council of Turkey, https://tubitak.gov.tr/
National contact person	Dr. Jale Şahin Phone: +90 312 298 1796 E-mail: jale.sahin@tubitak.gov.tr , uidb@tubitak.gov.tr
Funding commitment	0,5 M Euro
Overheads	Overheads are eligible costs only for academy and public institutions and subjected to the terms and conditions stated in TUBITAK 1071 Programme.
Anticipated number of fundable research partners	3-4 projects
Maximum funding per grant awarded to a partner	<p>If all the team/teams from Turkey are involved in the international project proposal submitted to EJP RD JTC2022 as partners, the budget requested from TUBITAK can be max 1,000,000 TL per project excluding Institution Share and Project Incentive Bonus (PI), and a partner can ask for:</p> <ul style="list-style-type: none"> • max 720.000 TL for public institutions and organizations, higher education institutions, training and research hospitals, • max 1.000.000 TL for private institutions <p>If one of the team(s) from Turkey is included as the coordinator* in the international project proposal submitted to EJP RD JTC2022, the budget requested from TÜBİTAK can be max 1,500,000 TL per project excluding Institution Share and Project Incentive Bonus (PI), and a coordinator can ask for:</p> <ul style="list-style-type: none"> • max 1.000.000 TL for public institutions and organizations, higher education institutions, training and research hospitals,

	<ul style="list-style-type: none"> max 1.500.000 TL for private institutions. <p>* In the relevant project, only one organization from Turkey can take part as a coordinator, or one organization can take part as a coordinator and other organizations/organizations can take part as partners. The specified budget caps will be applied in these cases.</p>
Eligibility of project duration	Maximum 3 years (36 month)
Eligibility of a partner as a beneficiary institution	Legal body: university, Public institutions and organizations and training and research hospitals, Large enterprises and SMEs.
Eligibility of costs, types and their caps	Personnel, consumables, equipment, travel, consultancy & service procurement.
Conditions for PAO funding	PAOs are not eligible for funding. However, the project coordinator can make a payment to a PAO only if the PAO is able to bill the provided service.
Submission of the proposal at the national level	<p>Yes</p> <p>Applicants from Turkey must make a national application through TUBITAK application system: http://uidb-pbs.tubitak.gov.tr/.</p> <p>For further information please contact to national contact person.</p>
Submission of other information at the national level	<p>Yes, for proposals selected for funding.</p> <p>Turkish partners in the projects selected for funding are obliged to provide Ethics Approval Certificate and/or Legal Permission Licences and other related documents (if necessary).</p>
Submission of financial and scientific reports at the national level	Yes, according to national regulations.

Further guidance

Please check the [guide](#) for national application rules.