

Guidance for the development and conduct of ERS Clinical Research Collaborations (CRC)



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1 Introduction

The European Respiratory Society (ERS) contributes to the coordination of research activities in respiratory medicine across Europe, by encouraging new initiatives, producing guidelines, supporting and disseminating information, developing joint documents with other major associations and/or international scientific societies, and other research activities including partnerships with Industry Partners.

ERS may support the work of a Clinical Research Collaboration (CRC) in areas of respiratory medicine where a pan-European multi-centre network of principal investigators aims to advance clinical and translational research. It may also integrate non-European countries as well as non-ERS members, representing different scientific disciplines which complement the network with multispecialty, multidisciplinary and multifunctional know-how and expertise.

If the application is successful, the CRC is expected to follow the ERS rules as described in chapter 7.

2 Definitions

2.1 ERS CRC

The aims of an ERS CRC are to promote the exchange of research ideas among clinicians and affiliated scientists in Europe and/or globally; to plan, conduct, evaluate and publish clinical and translational studies; to gain eligibility for network funding; to build an infrastructure for prospective clinical research and to agree on standardised approaches to address specific research needs. Outputs of a CRC would typically be scientific products, such as original articles for publication in scientific journals or abstract submission for presentation at scientific congresses. Clinical research resulting in results presentations and scientific session proposals for the ERS International Congress are encouraged.

Approved ERS CRCs are not legal entities. ERS aims to endorse projects under the current CRC rules and facilitate CRCs' developments. ERS CRCs are embedded in ERS activities and projects.

Clinical studies are health-related research studies in humans that follow a pre-defined (research) protocol. Multicentre research studies proposed by the CRC can be observational or interventional, prospective or retrospective. The types proposed can assess any combination of treatment, prevention, diagnostic, screening, health economics, implementation or quality of life studies. These studies can provide a better understanding of the disease, including treatments and clinical practice patterns by generating innovative, new or alternative interventional and practice approaches.

2.2 Patient and public input

The ERS Science Council recognises that patient and public input into CRCs is important, strongly encouraged, and likely to help to:

- Underpin the activities and outputs of the collaboration with patient experience.
- Ensure that the work of the CRC addresses key issues of concern to patients or that may have been overlooked by healthcare professionals.
- Build on the real-world experience of patients to ensure that proposed clinical studies and trials are feasible and acceptable to patients, and are likely to influence clinical practice.
- Provide input from individuals across Europe to ensure factors such as access, equity and cost are taken into account.
- Optimise patient access to, and support for, the resulting outputs of the CRC.

The European Lung Foundation (ELF; www.europeanlung.org) would welcome contact from any CRC chairs keen to investigate ways in which patient input could enhance their work. ELF has a long experience in coordinating patient engagement and an established network of patient organisations and Patient Advisory Groups with members from across Europe, who are keen to support CRC activities. Options include patient input into research priority-setting, patient representatives joining the CRC steering committee (e.g Patient Chair or steering committee member), patient input into clinical study design and outcome measures, patient review of participant information sheets, consent forms



and ethics documentation, patient consultation (including surveys and focus groups), patient input into governance and ethics discussions, and the development of a patient version of outcome documents.

To discuss possible options for patient input, please contact ELF - Patient involvement and engagement (info@europeanlung.org). [Find out more about ELF's patient input process on their website \(https://europeanlung.org/en/about-us/our-patient-input-process/\)](https://europeanlung.org/en/about-us/our-patient-input-process/).

2.3 Development and dissemination of novel research protocols

The CRC should decide on priority research questions related to where there is an unmet medical need, and where there is a need to gather real world evidence/data in a multi-centre research study. Based on the identified questions, the CRC will develop and share new research protocols on clinically orientated topics of the relevant disease area and lead the development of priority research projects in the field.

Each year, the ERS Science Council identifies key areas where there are potential gaps in the CRC portfolio by providing a 'highlight notice' and encourages proposals in the identified topic(s).

3 Application and approval process

All CRC proposals are evaluated on scientific excellence and potential impact. The CRC proposals aim to describe the purpose and structure of the CRC and show how the objectives will be achieved. The CRC proposals can be submitted by several ERS members belonging to one or more scientific assembly/group.

3.1 Online Platform

Applications and all supporting documentation should be submitted in English and online *via* the ERS CRC application platform (<https://www.ersnet.org/research/clinical-research-collaborations>). The web browser Google Chrome is advised to be used for a smooth fill in of information and upload of all documents.

The application is considered as complete when it contains:

1. The CRC online application form duly filled (see Appendix 1 for the description of requested fields)
2. One page CV with the 5 main relevant publications of both Chairs, justifying their expertise in the field and their role/responsibility in the CRC proposal.
3. Lay Summary (500 words max) to be shared with patients; it should be written in such a way that a non-scientist can easily understand the aims, outcomes and potential relevance to future practice.
4. The annual Declaration of Interest (DoI) signed by both Chairs. This document will be requested by the ERS office through the myERS platform after the submission deadline.

The deadline for submitting applications and documents is 15th of October each year without extension.

3.2 Evaluation and approval process

Each application will go through the following evaluation and approval process:

1. An administrative check for the completeness and eligibility will be conducted for all received CRC applications by the ERS Office.

A CRC proposal is considered as eligible to be sent for peer-review if the below selected criteria are met:

- The project membership is multicentric. Significant participation of European countries is expected. Participation of institutions based outside of Europe can be included.
- It focuses on disease areas of respiratory medicine or related issues.
- The objectives and ambitions of the proposal is to develop a work programme and does not restrain itself to a single clinical study or trial.
- At least one of the Chairs is an ERS member.



The project membership diversity i.e. the inclusion of patient(s) or carer representative(s), ERS early career members (<40 years old), gender balance (see ERS Diversity and Inclusion policy: <https://www.ersnet.org/ers-diversity-and-inclusion>) is considered an asset for the proposal but will not prevent the application from being considered eligible if only this last point is not met. Please refer to section 7.2 *CRC Composition* for additional details.

The ERS CRC Director is entitled not to send a proposal to peer-review, should any of these criteria not be fulfilled.

2. The ERS CRC Director will select a minimum of three independent reviewers, who are members of the CRC Working Group, to comment on the content of the application. Note that the reviewers might be excluded if there is a conflict of interest (CoI) through contributing directly or indirectly to the CRC project application.
3. The reviewers will evaluate the scientific excellence of the eligible CRC proposals and the specific contribution that the CRC will offer to the respiratory community. Plans to obtain sustained external funding will also be evaluated during the reviewing process.
 - CRC proposals for renewal WITH ERS funding support will follow the same application and reviewing process as new applications. However, other reviewing criteria may be used measuring the success of the CRC in the previous funding period and the relevance of project continuation. Therefore, applicants should underline in the application form the achievements so far, the expected future developments as well as the plans to make the CRC sustainable (*i.e.* external source of funding of the CRC if deemed necessary, future CRC chair leadership).
 - Note that for CRCs which have secured substantial external source of funding AND wish to apply for renewal, ERS CRC funding may not be guaranteed (see section 7.5 *CRC Tier Model - Tier 3*).
 - CRC proposals addressing a 'highlight notice' topic(s) will follow the same reviewing process as the other applications. These proposals will not be favoured during the review process but for applications evaluated of equal excellence, topics including highlight notices would be prioritized.
4. ELF will also review the applications for the level and appropriateness of patient involvement. Patient reviewers may also be consulted during the reviewing process. When patient involvement has not been included, ELF may suggest at the review stage how patient involvement could benefit the CRC and propose ways of involving patients.
5. Based on the reviewers' assessment, the ERS CRC Director with the consultation of the CRC Working Group will make acceptance/rejection recommendations of the CRC proposals.
6. The ERS Science Council decides based on the recommendation of the ERS CRC Director on approval or rejection of the CRC applications. It may request scientific and logistical changes as appropriate. The ERS Science Council may also make recommendations on membership of the CRC. The ERS Science Council reserves the right 1/ to reduce the funding request or 2/ to propose approval of the CRC without ERS funding and grant the "ERS CRC" badge only.
7. The ERS Science Council decision has to be finally endorsed by the ERS Executive Committee.

The time taken for the reviewing and approval procedure may vary and can take around four (4) months (until February of the following year) for a decision. The applicants of successful and unsuccessful CRC proposals will receive a notification letter by email.

4 Project Development

For all approved CRC proposals (both new proposals and renewals), a kick-off teleconference will be organised with the CRC chairs, the CRC director, a nominated CRC Link person and a member of ERS office in order to describe the terms and conditions of their project recognition, discuss logistic aspects, and, eventually, funding and supplementary background materials prior to the start (or continuation) of their projects. ELF will approach CRC chairs to discuss plans for patient involvement following the kick-off teleconference, if relevant.

CRC chairs will be asked to sign a CRC agreement to settle the terms of collaborations between ERS and the CRC chairs.



Funding for CRCs is not to be considered as an ERS grant; it is funding that ERS has earmarked for up to a three (3)-year period for routine expenses during project development. Funding for ERS-supported projects will be managed by the ERS office in Lausanne. All expenditure charged to the ERS project accounts must be approved by the ERS office in advance.

Annual reporting is expected to the ERS CRC Director and CRC Working Group to assess project advances according to the CRC agreement and to review the *per annum* budget spent (see section 7.7 *Reports*). Any substantive changes in the goals, strategy or the budget of a CRC must be reported to the ERS office at any time of the ERS CRC funding period and should be approved by the ERS Science Council.

All official ERS policies including those on travel, publications, conflicts of interest, CME, and others must be followed during all phases of the project. Failure to comply with these will result in immediate cessation of the CRC status. The ERS CRC Director and the Chair of the ERS Science Council will inform the ERS Secretary General who will act accordingly.

5 Publications

Results of CRCs should lead to manuscripts for publication in relevant peer-reviewed scientific journals. Submission to an ERS journal, such as the European Respiratory Journal (ERJ), are encouraged and will be subject to the normal procedures for peer review. Guidance for manuscript preparation and submission to the ERJ can be found at: <https://erj.ersjournals.com/authors/instructions>

If during or after the research project the CRC steering committee considers the development of Guidelines, Statements, or Technical Standards relevant to the CRC topic, then, an official application through the ERS Task Forces funding scheme, under the supervision of the ERS Guidelines Director, should be submitted.

Preliminary results from these clinical research studies may lead to submission of abstracts for presentation at scientific congresses and submission for presentation at the ERS International Congress is encouraged.

Note that any statement or other official communication in the name of ERS must be approved by the ERS Executive Committee before publication.

CRCs are encouraged to consider how to communicate their outputs to patients and the public, for example, through the production of a lay summary or factsheet to accompany a scientific publication. The European Lung Foundation can support the production of patient-facing materials. Please also view the section 7.3 *Funding*.

6 Education

The allocated ERS CRC funds cannot be used to cover educational activities. If during the conduct of the research project, educational activities (including webinars), research seminars, or other ERS dissemination methods are thought to be relevant, a separate application through the appropriate scheme is advised with cross-CRC initiatives particularly encouraged.

7 Rules for CRCs

7.1 CRC Governance

The CRC should be governed *via* its own steering committee, which should be composed of the CRC chairs, academic ERS members and any other non-member academic stakeholders. Patient representatives could be members of the CRC steering committee. The CRC steering committee refers to their nominated CRC Link person and the CRC Director who, supported by the CRC Working Group, reports to the ERS Science Council.

7.2 CRC Composition

A CRC is a multidisciplinary research network and expected to be primarily composed of principal investigators who are ERS members. The initiative should represent a significant proportion of pan-European countries to ensure a wide perspective and representativeness. The presence of ERS early career members (< 40 years old), the gender balance (see [ERS Diversity and Inclusion policy](#)) as well as patient representatives is strongly encouraged.



It may also integrate non-European countries as well as non-ERS members, representing other scientific disciplines which would complement the network with multispecialty, multidisciplinary and multifunctional know-how and expertise. CRC members can join the project later whenever deemed necessary or suitable.

After six (6) years, the ERS Science Council would encourage to revise the CRC chairs/ Leadership to promote emerging leaders and include new perspective to the CRC. A succession strategy plan and progress status will be requested in the annual progress reports (see *section 7.7 Reports*).

7.3 Funding

The approved CRC will be funded up to €15,000 per year by ERS for a maximum of three (3) years, allocating a total budget of €45,000.

Allocated funds by ERS will be administered by the ERS office according to the terms of the CRC agreement. For the use of the ERS CRC funding regarding meeting expenses, refer to *section 7.9 Meetings*.

The release of the *per annum* ERS CRC funds is dependent on successful review of the annual progress report (*section 7.7 Reports*).

ERS does not have the financial means to fund the research or clinical studies and/or trial that the network intends to conduct. If appropriate and relevant, grant applications for research and/or network funding should be submitted to the EU and/or other funding agencies with the aim of funding the activities of the CRC.

ERS provides independent funding to ELF to facilitate patient involvement at a basic level. However, patient participation to CRC meetings has to be funded by the ERS CRC funds and sufficient budget should be considered.

ELF may also facilitate patient involvement at a higher level but there are charges for this. See '[ELF Levels of Involvement](#)' to see what patient input activities can be covered and likely costs (e.g. creation of patient surveys). These higher levels of support from ELF may be a more realistic option when external funding is secured (e.g. industry, academic grants).

Please note that ELF has limited capacity to support CRCs at higher level and decisions on support available will be discussed with each CRC.

7.4 CRC Link person

Once the CRC is endorsed, a member of the CRC Working Group is nominated by the CRC Director as a CRC Link person and will have the responsibilities outlined below towards the CRC project:

1. Provide close support to the CRC research project during the first year at the initial/set-up stages of the project.
2. Oversee the CRC project advance and delivery according to the CRC proposal.
3. Provide insights, problem solving support, and facilitate the CRC progress.
4. Link for any potential collaboration with other CRCs.
5. Provide feedback to ELF on patient engagement.
6. Review the CRC annual progress reports and evaluate progresses compared to the initial CRC application and/or previous CRC annual progress reports.
7. Provide any recommendations or highlight any corrective actions to be taken if deemed necessary.
8. Report to CRC Director and/or the rest of the CRC Working Group any potential CRC project issues or risks to be raised at any project stage.

CRC chairs should liaise with the CRC Link person throughout the duration of the CRC for any type of support and invite him/her to join teleconferences for project progress status that the CRC chairs may have with the rest of the CRC project.

CRC chairs can meet face to face with their CRC Link person during the ERS International Congress.

Note that the CRC Link person should not have a conflict of interest (CoI) with the CRC project.



7.5 CRC Tier Model: Funding partners engagement and ERS Research Agency

When the CRC project is endorsed, three (3) different tiers/layers of ERS engagement can be applied by the CRC chairs. The level of engagement is mainly related to additional fund-raising activities and involvement of the ERS Research Agency.

Industry partnerships with CRCs are encouraged where considered appropriate and beneficial for the delivery of the aims of the CRC. In case of Tier 2/3 and to ensure coordination between CRCs and other ERS initiatives and industry, the partnerships must be discussed with the CRC Director.

In case of engagement with Industry, ERS would act as Neutral broker between the Industry partners and the CRC consortium which includes the below characteristics:

- Transparency
- Inclusion
- Agreed benefits
- Agreed milestones
- Reporting
- Control of funding

Transition to one tier to the next level is possible depending on the development of the CRC project.

Any exceptions to the conditions laid out in each tier (for example where an existing set up makes it difficult to fully agree to all conditions) should be agreed by the ERS Science Council Chair and CRC Director and made clear to the concerned parties.

| TIER 1 | |
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| Goal | Provide financial support to the set up and/or maintenance of a CRC research network where the researchers wish to receive seed funding from ERS but wish to run the project independently of the ERS Research Agency. |
| CRC Funds | The CRC funds would be held and managed by the ERS office to cover the expenditure mentioned in the CRC proposal. |
| Conditions | <ol style="list-style-type: none"> 1. CRC chairs are responsible for ensuring the approved budget is not over-spent. 2. Salary expenditure could be anticipated for administrative staff, NOT for expert stakeholders involved in the project. 3. ERS office is not involved in the conduct of the project. ERS cannot be held responsible for any actions taken by the CRC research network. In case of litigation, ERS is not liable. 4. For any current or future collaboration with any third-party funders independently of the ERS Research Agency (i.e. pharmaceutical and medical device companies, other medical or research societies, foundations...), the CRC chairs cannot claim to be an ERS Research Agency supported CRC but should state they have received a financial support from ERS to support their project. 5. ELF ensures basic services linked to patient involvement in the project. See ELF Levels of Involvement. |
| TIER 2 | |
| Goal | Bring a CRC research network to a status where they are able to obtain through the support of ERS-Research Agency substantial funding needed to deliver and achieve major objectives of the CRC. |
| CRC Funds | The CRC funds would be held and managed by the ERS office to cover the expenditure mentioned in the CRC proposal. |



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| Conditions | <ol style="list-style-type: none">1. In case of collaboration with funding partners, negotiations and contracting activity will be coordinated by ERS office.<ol style="list-style-type: none">a. As per ERS policy, contribution from a minimum of two different industry companies is expected.b. All external funding for the CRC will be collected and managed through the ERS office with a retention of 10-percent (10%) yearly overheads from the industry funds spent to cover the indirect costs of the ERS office.c. The CRC chairs will lead the discussions on the scientific content and the ERS office will lead the contractual and financial matters.d. The CRC chairs will have to provide a milestones-based project summary including benefits offered to funding partners, a description of the governance structure, a data access policy, a publication policy, as well as a detailed budget and a plan how the budget will be used. This will have to be approved by the ERS office, the ERS CRC Director and ERS Science Council chair.e. The CRC chairs should always involve the ERS office in their contact with potential funders and particularly the Industry to ensure there is one single communication channel through ERS office regarding funding and all other terms of the contractual agreement.f. In case contractual relationships are already in place this condition could be submitted to modifications.g. Any new contractual agreement with funding partners will be between ERS and the funding Partner only and the application for funding will follow the agreed process set out by the ERS office.h. The same conditions apply for any funding partner. However, the requirements of the different funders need to be taken into consideration during the negotiation period.2. The CRC should not start incurring costs outside of the €15,000 Euros per year (€45,000 Euros in total) original ERS CRC funding agreement until the contracts with the funding partners for the wider projects outlined by the project summary have been finalized.3. ERS would provide through the ERS Research Agency basic operational and project management support i.e. organise regular TCs and meetings, support in setting up documentation (project plans and budgets) and monitoring project delivery, suggest service providers, ensure links with other CRCs or ERS research activities (e.g. congress, research seminars, fellowships, task forces...). This will be adapted depending on the needs of the projects.4. Additional responsibilities of the ERS Research Agency would be negotiated on a per project basis, before starting the negotiation process with funding partners and have to be covered by the additional CRC funds.5. Funds can be used to develop a website for the CRC and a project specific logo to reflect its own identity. Project-specific website could be managed through ELF if sufficient funding is available and ELF has capacity.6. An ERS CRC should use ERS logo made specially for ERS CRCs that would act as a kite mark to demonstrate that ERS ensures all the benefits promised to funding partners.7. CRC should recognise ERS for its financial support on all material, presentations and publications. Use of ERS logo needs to be approved in advance by the ERS office.8. Meetings can be organized through the ERS office on request (<i>see section 7.9 Meetings</i>). |
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| | <p>9. ELF ensures basic services linked to patient involvement in the project - plus further activities for an additional cost. Any activities / tasks you would like ELF to facilitate should be discussed and agreed upfront with ELF, the CRC chairs and the ERS office following the CRC kick-off teleconference and that direct and indirect costs shall be covered by the CRC funds if deemed necessary. ELF would submit a detailed plan and budget that would need approval of the CRC Chairs and the ERS office. See ELF Levels of involvement for further details.</p> |
| TIER 3* | |
| Goal | <p>To ensure that the timelines and deliverables agreed with the funding partners are met as well as the adequate use of the funds.</p> <p><i>*This model is only applicable once relationship with funding partners is established. Therefore, this model is not applicable at the time of the endorsement of the CRC.</i></p> |
| CRC Funds | <p>The CRC Funds AND the additional external Funds received from funding partners would be held and managed by the ERS office.</p> <p><i>Note: for CRCs which have secured substantial external source of funding AND wish to apply for renewal, ERS CRC funding may not be guaranteed.</i></p> |
| Conditions | <p>The following additional conditions from Tier 2 enter into forces:</p> <ol style="list-style-type: none"> 1. The ERS office will collect and manage funding for the CRC and will retain 10-percent (10%) yearly overheads from the funds received from funding partners to cover the direct and indirect costs of the ERS office for his neutral broker role as described in section 7.5. 2. ERS could have a broader role in the CRC through the ERS Research Agency mainly in project management to ensure that sub-projects activities are met - outside of overall contract requirements with fund providers. This should be covered by the additional external funds received from funding partners. 3. The CRC chairs will have to comply with the conditions of the collaboration agreement with the Funding Partners, in terms of the milestones, objectives of the project, access to data and reporting terms. 4. The allocation of the additional external funds is decided by the CRC chairs. Involvement of external consultants should be approved by ERS and be limited to tasks that could not be performed by the CRC chairs, CRC members, ERS early career members, ERS office or academic institutions. Involvement of external consultants should be kept to the minimum and supported by good arguments. 5. The allocated funds from funding partners can NOT be used for the payment of the salary of experts stakeholders (i.e. CRC chairs, principal investigators) and patients representatives for their involvement in the project. However, the funds could be used to cover the operational staff costs (e.g. Ph. D students, biostatistician, data scientist,...) involved for the completion of the CRC tasks defined in the budget plan. 6. The allocated funds from industrial funding partners can NOT be used for the development of Guidelines, Statements, or Technical Standards. 7. The ERS Research Agency will set up the appropriate separated contracts with the participating academic institutions/partners for the contribution to the expenses in relation to the CRC tasks defined in the budget plan. ERS is a non-for-profit organisation and recommends to each participating institution to not apply overheads or limit overheads to a maximum 20% rate. |



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| | <p>8. Any change from the original plans especially to the milestones and deliverables, the budget and the plan on how to spend it as well as the funding partners' benefits have to be discussed with ERS office, the CRC Director and ERS Science Council chair prior to implementation.</p> <p>9. ELF ensures basic services linked to patient involvement in the project - plus further activities for an additional cost. Any activities / tasks you would like ELF to facilitate should be discussed and agreed upfront with ELF, the CRC chairs and the ERS office and that direct and indirect costs shall be covered by the additional external funding. ELF would submit a detailed plan and budget that would need approval of the CRC chairs and the ERS office. See ELF Levels of involvement for further details.</p> |
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7.6 Duration of support

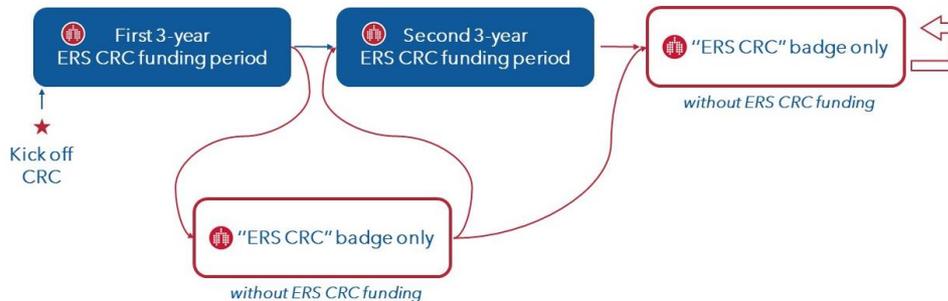
The approved CRC will follow different phases of funding with the applicable conditions:

1. **Initial ERS CRC funding support:** The proposed CRC would be financially supported by ERS for an initial duration of three (3) years, based upon the deliverables defined in the CRC agreement.
2. **Renewal WITH ERS CRC funding support:** CRC renewal over the initial duration of three (3) years might be considered upon reapplication and will be evaluated according to section 3.2, clause 7. If the CRC chairs apply for renewal with direct ERS funding, the option for a CRC to receive ERS CRC funds is applicable for **only one additional period of three (3) years**, giving a maximum of six (6) years of funding.
 - Note that if the ERS CRC funding support has been discontinued during/after the initial period of three (3) years and renewed with the "ERS CRC" badge only (see below), the CRC is still eligible to apply for renewal WITH ERS funding support at a later stage.
3. **Renewal of the "ERS CRC" badge WITHOUT ERS CRC funding support:** When the CRC has reached the end of a funding period, the CRC chairs could apply for renewal without direct funding through ERS to keep the "ERS CRC badge" allowing access to the benefits linked to CRCs (e.g. section 7.9 *Meeting facilities* and 7.11 *Promotion*) for an additional three (3) year period. In such a case, the CRC rules continue to apply especially the clause 7.7 *Reports* and 7.12 *Liability*.
 - The "ERS CRC" badge renewal application is managed through the annual report process at the end of a funding period i.e. **end of March** (see clause 7.7 *Reports*).
 - Reviewing criteria measuring the success of the CRC in the previous period and the relevance of project continuation may be used in order to approve the renewal of the CRC with the "ERS CRC" badge only. Therefore, the CRC chairs should underline the expected future developments, the plans to make the CRC sustainable and a suitable succession planning for the next three (3) year period.
 - If interested, the CRC chairs could apply for the renewal of the "ERS CRC" badge at the end of each three (3) year period without a limited number of renewals applications.
4. **End of ERS CRC funding period:**
 - At the end of an ERS CRC funding period, any left-over of the CRC funding can be retrieved back to ERS even if the CRC is renewed with the "ERS CRC" badge only for the next additional period of three (3) years.
 - If the CRC is not renewed (with ERS funding support or with the "ERS CRC" badge only), then ERS will terminate the CRC and will remove all the benefits linked to CRCs. If applicable, any left-over budget will be cancelled and retrieved back to ERS.

The below figure describes the different life-courses of a CRC with ERS CRC funding support or "ERS CRC" badge.



CRCs different lifecourses with
ERS CRC funding or ERS CRC badge



7.7 Reports

CRC chairs will provide an annual progress report, explaining 1/ the achievements and completion of expected deliverables or any deviation from the initial plans, 2/ their sustainability and succession planning strategies, 3/ the patient engagement strategy and 4/ a detailed description of the expenses, to be sent to the ERS office (scientific@ersnet.org) usually **by the end of March each year**.

Any "ERS CRC" badge renewal applications will be reviewed at the same time of the annual progress report i.e. at the end of the three (3) year period. The CRC Director, the CRC Working Group and ELF will review the status of all ongoing CRCs and will provide feedback, including recommendations if deemed necessary.

For CRCs that have run for at least six (6) years, additional reviewing criteria may be used to measure the success of the CRC and the relevance to continue the "ERS CRC" badge each year. These CRCs are expected to produce at least 2 publications over the last 2 years (no editorial/review) OR 1 publication in a journal with an Impact Factor > 10 in order to continue the "ERS CRC" badge the following year. This criterium shall not constitute an exclusive criterium, however the reasons why if it has not been met should be indicated to the CRC Working Group for evaluation.

The progress status of all running CRCs (i.e. CRCs with ERS funding support AND CRCs with "ERS CRC" badge only) will be reported to the ERS Science Council.

The release of the *per annum* ERS CRC funds or the continuation of the "ERS CRC" badge are dependent on successful review of the annual progress report. ERS reserves the right to reduce the amount of funding if the amounts allocated in the previous year has not been fully spent or not used within the ERS CRC guidelines or to withdraw the "ERS CRC" badge in case of non-compliance to the terms of the CRC agreement.

7.8 Regulatory and ethical issues

The institution of investigators from the CRC would need to endorse the legal sponsor responsibility of any clinical studies and trials conducted in the frame of the CRC. ERS will not endorse the legal sponsor responsibility for any clinical studies or trials.

The initiative should seek Ethics Committee approval on clinical studies/trials in at least one Ethical committee per country involved.

All projects involving clinical trials should comply with the relevant local and international specific regulations and as a requisite to be considered, provide proof of having the necessary insurances in place, as well as an EudraCT number in the case of drug-based research.

7.9 Meetings

The ERS CRC funding cannot be used to cover meeting expenses during or in relation to the ERS International Congress (travel, registration or accommodation), except for patient representatives as defined in the relevant ERS travel policy. If the CRC is approved, the ERS office can offer meeting facilities (room with audiovisual equipment) at the ERS International Congress. Meeting room request form will be sent by the ERS office to CRC chairs in May each year.



If funds are used for meetings organised outside of the ERS International Congress through the ERS office, then, the [ERS CRC and Task Force travel policy](#) for reimbursement will apply. Claims for reimbursement of expenses must be accompanied by the relevant receipts.

Meeting facilities also exist at the ERS office in Lausanne, Switzerland and Brussels, Belgium with a limited number of ~15 participants. In such case, the meeting facilities are free of charge for the CRC but Travel/Accommodation/Catering expenses will be charged on the ERS CRC funds.

For meetings organised at ERS office or to an external place, logistical support (travel/accommodation) can be provided upon request to ERS office up to 3 months before the meeting.

7.10 Joint CRC with other organisations

Applicants are encouraged to seek for collaboration with other organisations. In such a case, it should be highlighted in the CRC proposal. If collaboration with another organisation is approved, a written agreement will need to be established by all parties outlining the major terms of collaboration in accordance with ERS policies. This will include details of how the expenses and outcomes of the project will be handled.

7.11 Promotion

The CRC can be promoted through ERS promotional channels and a dedicated webpage can be set up on the ERS website. The initiative can use the ERS logo on its website and for its correspondence during the period for which it has been endorsed.

7.12 Liability

Each CRC, through its chairs, has total responsibility related to all actions and activities undertaken using the ERS name and logo. These actions and activities should be in compliance with the CRC agreement. ERS should be informed in advance of any action or activity performed by the CRC (e.g. involving ERS name and logo) which is not part of the approved CRC agreement. Failure to inform ERS will result in immediate cessation of the CRC recognition and funding.

7.13 Intellectual property

Each CRC proposal should clearly define the sharing between all partners involved, the ownership or sharing of intellectual property or any other product or outcome from the work performed under the CRC. This should also include any subsequent financial revenue.

7.14 CRC assets sharing

The ERS CRC umbrella intends to establish a collaborative spirit and the ERS CRC programme encourages sharing of project assets between CRCs. Therefore, ERS expects CRC chairs to assess thoroughly requests made through the ERS Office to share any results of the CRC (existing patient data, registries, biobanks or samples collected) within the frame of the CRC as well as protocols, SOPs or templates with other researchers to answer additional research questions or to support the setup of other projects.



8 Appendix 1 - CRC online application: Checklist of the requested fields/documents

This appendix represents a checklist of the different requested fields of the CRC online application form and /documents which would support you to prepare your CRC application and submit it to the ERS CRC application platform.

Only CRC applications received through the ERS CRC application platform will be considered for review. Any exceptional case must be discussed upfront with the ERS Office by contacting scientific@ersnet.org before the submission deadline of October 15th.

| Description | Completed |
|--|--------------------------|
| CRC online application form (sections 1 to 3 describe the requested fields to be completed online) | |
| 1. CRC Description | |
| 1.1 Title | <input type="checkbox"/> |
| 1.2 Lead Assembly and Groups relevant for your proposal: And Tracks: <input type="checkbox"/> Airway diseases <input type="checkbox"/> Interstitial lung diseases <input type="checkbox"/> Paediatric respiratory diseases <input type="checkbox"/> Pulmonary vascular diseases <input type="checkbox"/> Respiratory critical care <input type="checkbox"/> Respiratory infections Sleep and breathing disorders <input type="checkbox"/> Thoracic oncology | <input type="checkbox"/> |
| 1.3 Summary (250 words max) | <input type="checkbox"/> |
| 1.4 <input type="checkbox"/> New application or <input type="checkbox"/> Renewal WITH funding support | <input type="checkbox"/> |
| 1.5 Background and relevance <i>In case of application renewal, please underline the achievements so far.</i> | <input type="checkbox"/> |
| 1.6 Objectives <i>In case of application renewal, please highlight adaption of objectives in line with project advances.</i> | <input type="checkbox"/> |
| 1.7 Deliverables and Publications <i>Note that any statement or other official communication in the name of the Society must be approved by ERS before publication.</i> | <input type="checkbox"/> |
| 1.8 Action plan and deliverables timelines <i>Indicate the yearly deliverables expected <u>for the three years</u> duration of the CRC funds, listed chronologically.</i> | <input type="checkbox"/> |
| 1.9 Risks <i>Indicate the top 3 risks foreseen along with the mitigation actions to monitor each risk.</i> | <input type="checkbox"/> |
| 1.10 Patient input factors <i>If public and patient input is being considered, please explain the role you envisage for patient input, how you think this could add value to your CRC, and what form(s) of input would be appropriate/desirable. Please outline your plans for patient input, including key activities and the ways in which patient perspectives will be incorporated into the other aspects of the CRC. We recommend contacting ELF in advance to discuss plans for involving patients and the support available: info@europeanlung.org</i> | <input type="checkbox"/> |
| 2. Stakeholders | |
| 2.1 Proposed Chairs | <input type="checkbox"/> |



| | |
|---|--------------------------|
| <ul style="list-style-type: none"> - Chair 1 AND Chair 2: Name (first, last), Title, Institution, Department, mailing address, Post code and City, country, email address and telephone. | |
| <p>2.2 Members of the CRC</p> <ul style="list-style-type: none"> - Name, Specialty, institution, City, country <i>[enter as many you have member]</i> <p><i>We recommend to consider an adequate diversity of the CRC membership. The presence of ERS early career members (< 40 years old) and the gender balance is strongly encouraged.</i></p> | <input type="checkbox"/> |
| <p>2.3 ELF involvement <input type="checkbox"/> Yes <input type="checkbox"/> No</p> <p><i>If public and patient input is being considered, please explain the role you envisage for patient input, how you think this could add value to your CRC, and what form(s) of input would be appropriate/desirable. Please include details of key activities and the ways in which patient perspectives will be incorporated into the CRC. In case you do not see a role for patient involvement, at the review stage ELF may suggest how patient involvement could benefit the CRC and propose ways of involving patients.</i></p> | <input type="checkbox"/> |
| <p>2.4 Other Parties <input type="checkbox"/> Yes <input type="checkbox"/> No</p> | <input type="checkbox"/> |
| <p>If yes, if the proposal is for a joint CRC with (an)other society-ies, please indicate:</p> <ul style="list-style-type: none"> - Name(s) of the Society-ies and: Nature of the contributions (ie. Financial, support) <i>[enter as many you have identified Society]</i> | <input type="checkbox"/> |
| <p>If yes, if industry partnership is foreseen, please indicate:</p> <ul style="list-style-type: none"> - Name(s) of the industrial company-ies and: Nature of partnership (ie. Financial, support) <i>[enter as many you have identified industrial company]</i> | <input type="checkbox"/> |
| 3. Budget Details | |
| <p>3.1 Expenditure details</p> <p>Source (i.e. Meetings, salaries, events, etc) / Amount (€) / comments <i>[enter as many you have identified source to spend budget]</i></p> <p><i>After approval, the ERS will support the CRC for the three years duration period allocating a total budget of 45,000.00 Euros. Please provide a detail of the sources of expenditure which will be covered by the CRC funds.</i></p> <p><i>Salary expenditure is anticipated for administrative staff, NOT for the salary of expert stakeholders involved in the project</i></p> | <input type="checkbox"/> |
| Lay Summary (500 words max) | <input type="checkbox"/> |
| Chair 1_ One page CV with his/her 5 main relevant publications | <input type="checkbox"/> |
| Chair 2_ One page CV with his/her 5 main relevant publications | <input type="checkbox"/> |
| Chair 1_ Declaration of Interest (DoI) | (*) |
| Chair 2_ Declaration of Interest (DoI) | (*) |

**The Declaration of Interest (DoI) of both Chairs will be requested by the ERS office through the myERS platform after the submission deadline. Both Chairs should ensure to reply back to the ERS office in order to have the application fully completed.*