EP PerMed Strategic Research and Innovation Agenda (SRIA) and Draft Proposal Presentation

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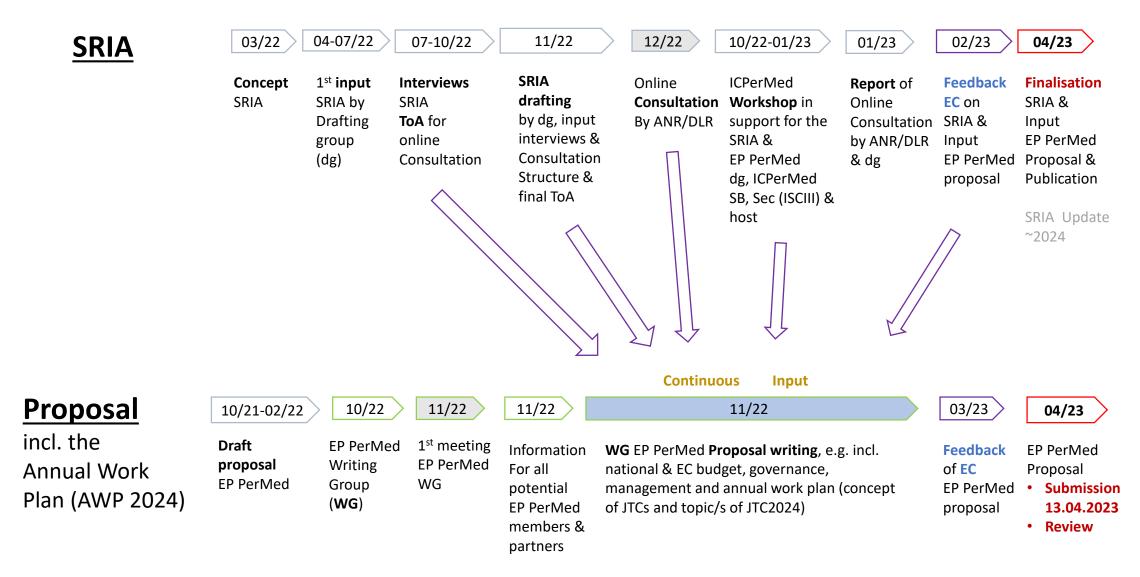
EP PerMed Drafting Group

Strategic Research and Innovation Agenda (SRIA)

General context

- All European Partnerships have to develop a SRIA
- The EP PerMed proposal will be directly connected/refer to the SRIA
- The EP PerMed activities (e.g. Joint Transnational Calls, and other activities as trainings, events, etc.) will be based on the SRIA and EP PerMed proposal

EP PerMed SRIA and Proposal development - timeline



dg: EP PerMed drafting group and EC/MS delegates; ToA: Triplets of Action; GA: Grant Agreement, CA: consortia agreement;
WG: EP PerMed proposal writing group; JTC: Joint Transnational Call; [SB: Steering Board and Sec: Secretariat ICPerMed]

START end of 2023

Strategic Research and Innovation Agenda (SRIA) – concept developed

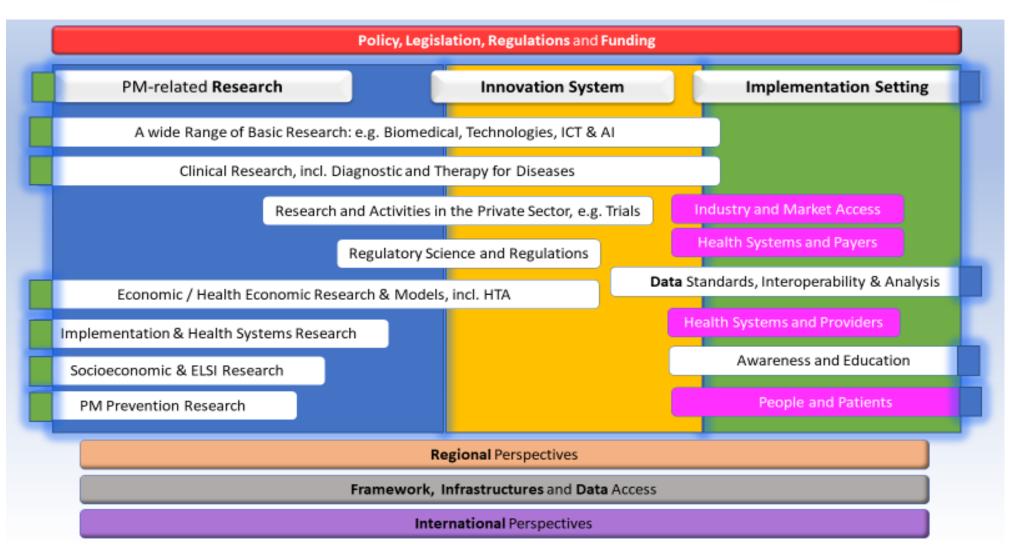
The SRIA will address for example the following topics:

- Personalised Medicine related **research** in all related areas
- Innovation system and Personalised Medicine
- The **implementation** setting and Personalised Medicine
- <u>Regional</u> and <u>international</u> perspectives of PM
- Framework, <u>infrastructures</u> and data access for PM approaches

Strategic Research and Innovation Agenda (SRIA) – concept developed







Strategic Research and Innovation Agenda (SRIA) – content development – 1st steps

The **SRIA** will identify gaps, needs and challenges as well as opportunities and visions which need to be addressed, overcome or reached to foster PM for the benefit of patients and citizens within sustainable health systems.

Process:

- Expertise of all involved initiatives & partners and existing strategic documents
- Organisation of activities to gather information and ideas for EP PerMed and its SRIA:
 - Around **70 interviews** with international PM experts and stakeholders were completed by the Drafting Group
 - So called: "Triplets of Action, short ToA" were identified
 - An Online Consultation with the 1st set of 47 ToA was performed (12/22)

General structure for the « Triplets of Action, ToA »

Challenge	Should describe the status quo, indicate/indication what is still lacking, why this is the case and what negative consequences this has.
Objective	Should indicate what exactly should be done to address the challenge in terms of specific actions
Outcome	Should describe how the situation will have improved once the actions described in the objectives have been successfully implemented.

Online Consultation with the 1st set of 47 ToA was performed (12/22)

Structure of the Consultation

The Triplets of Action (ToA) were presented in 4 categories:

- Personalised Medicine related research (ToA numbers 1.1 1.17)
- Innovation System and Personalised Medicine (ToA numbers 2.1 2.8)
- Personalised Medicine Implementation Setting (ToA numbers 3.1 3.16)
- Overarching Activities (ToA numbers 4.1 4.6)

Contribution to the consultation:

- Validation and urgency of the presented ToAs
- Selection of 5 most important ToAs per section
- Selection of 5 most important ToAs considering all 47 ToAs
 - How 15M€ would be distributed amongst the selected 5 ToAs
- Possibility to propose new or refine presented ToAs

Online Consultation with the 1st set of 47 ToA was performed (12/22)

Outcome of the Consultation

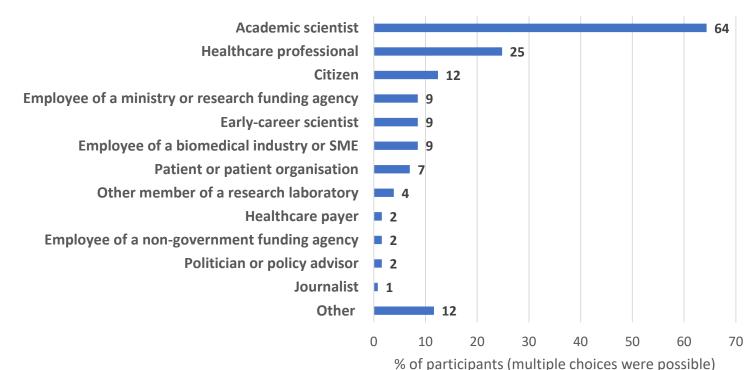
First outcomes presented today

Consultation report will be published soon (in preparation)

General statistics

Open consultation was open from 21 November – 31 December 2022 Prepared by EP PerMed Drafting Group, disseminated via ICPerMed and ERA PerMed webpages and social media, special ICPerMed newsletter, emails to ICPerMed bodies (ExCom, AB, SHF) and to ICPerMed related CSA coordinators, through national channels and by EC

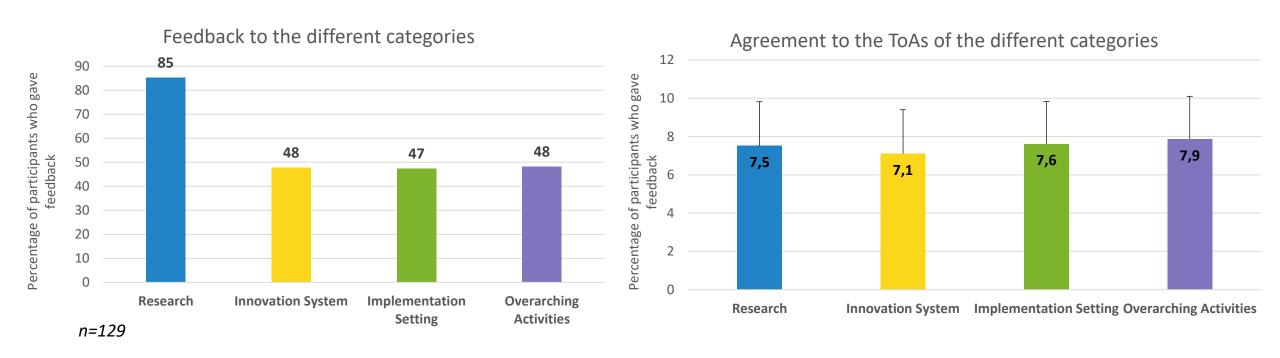
- Number of interested viewers: 714
- Number of submitted and analysed surveys: 129
- Participants from 21 countries







Summary of analyses



Ranking of ToAs per category

		Triplet of Action 1.7 "New targets for personalised therapies making use of an improved	72%	Research
Research	Research			Innovation System
	(n=110)		55%	Implementation Setting
			55%	Overarching Activities
Innovation Syst (n=61)	In a continue Constant	Triplet of Action 2.5 "Early cooperation between public research and the private sector"	75%	
	(11-01)	Triplet of Action 2.1 "Expanded knowledge on value for PM"	<mark>61%</mark>	
Implementat Setting (n=6 Overarchin Activities (n=		Triplet of Action 3.3 "Training and Education of healthcare professionals"	51%	
	· · · · · ·			
	Setting (n=61)	Triplet of Action 3.2 "Test Beds in hospitals"	41%	
	Quararahing	Triplet of Action 4.5 "Network of national and regional innovation hubs"	84%	
	•		•	
	Activities (II=02)	Triplet of Action 4.3 "Connected large-scale health databases"	82%	

Top-ranked ToAs over all categories

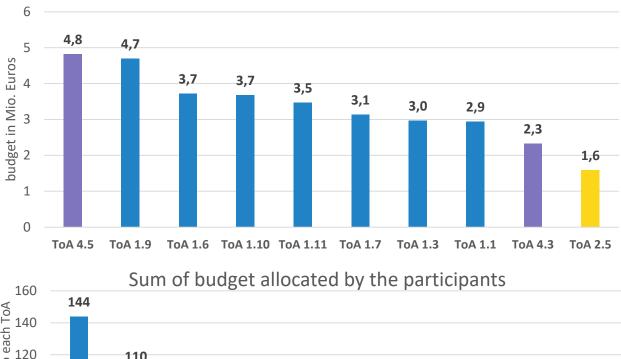
Group 1	ToA 1.1	A collaborative approach between pre-clinical and clinical research
	ToA 1.7	New targets for Personalised therapies making use of an improved understanding of disease mechanisms
	ToA 1.6	More biomarker evidence for PM

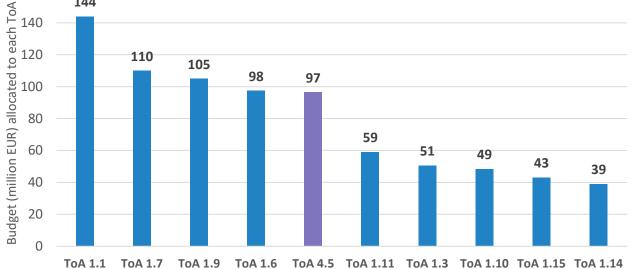
Group 2	ToA 1.9	Single-cell technologies in combination with AI and ML for PM
	ToA 4.5	Network of national and regional innovation hubs
	ToA 1.11	Medical cohorts for collecting high-quality health and molecular data
	ToA 1.3	New treatment modalities for PM
	ToA 2.5	Early cooperation between public research and the private sector

Group 3	ToA 4.3	Connected large-scale health databases
	ToA 1.10	Broader biomarker approaches to enable more informed health decisions
	T. A 4 F	

ToA 1.5 Active involvement of patients in PM research

Mean budget allocated to the 10 favoured ToAs





Note: first ToA of Category 3 Implementation followed on place 13/11: ToA 3.2 with 32 Mio. € in total and a mean budget allocated of 3.27Mio. €

Suggestions for new ToAs

Early detection of non-communicable disease

Improvement of patients' adherence to therapy

Predictive patient-derived disease models for PM*

Technology Platforms for accelerating and de-risking of Advanced Therapy Medicinal Product (ATMP) innovations**

Need to promote equity in PM clinical studies

Improve equity in health and wellbeing across individuals and groups

High-throughput high-content profiling workflows for early stage evaluation of safety / toxicity (and off-target reactivity) of drug candidates

*10x nominated, **2x nominated

- In addition 10 ideas/topics for ToAs were provided but not elaborated
- Suggestions for adaptations of 13 ToAs

Research Innovation System Implementation Setting Overarching Activities

Strategic Research and Innovation Agenda (SRIA) – content development – this workshop

- Additional support for the SRIA development and EP PerMed activities by this workshop
- The workshop is the last input for the SRIA development before its finalisation by the Drafting Group
- We are looking forward to fruitful discussions in the panels and active participation of all workshop participants

Thank you for your attention

You are also welcome to contact us or other representatives of the SRIA <u>Drafting Group</u> or the proposal <u>Writing Group</u> in the case of questions!

Contact: <u>EPPerMed@dlr.de</u>

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