



norecopa

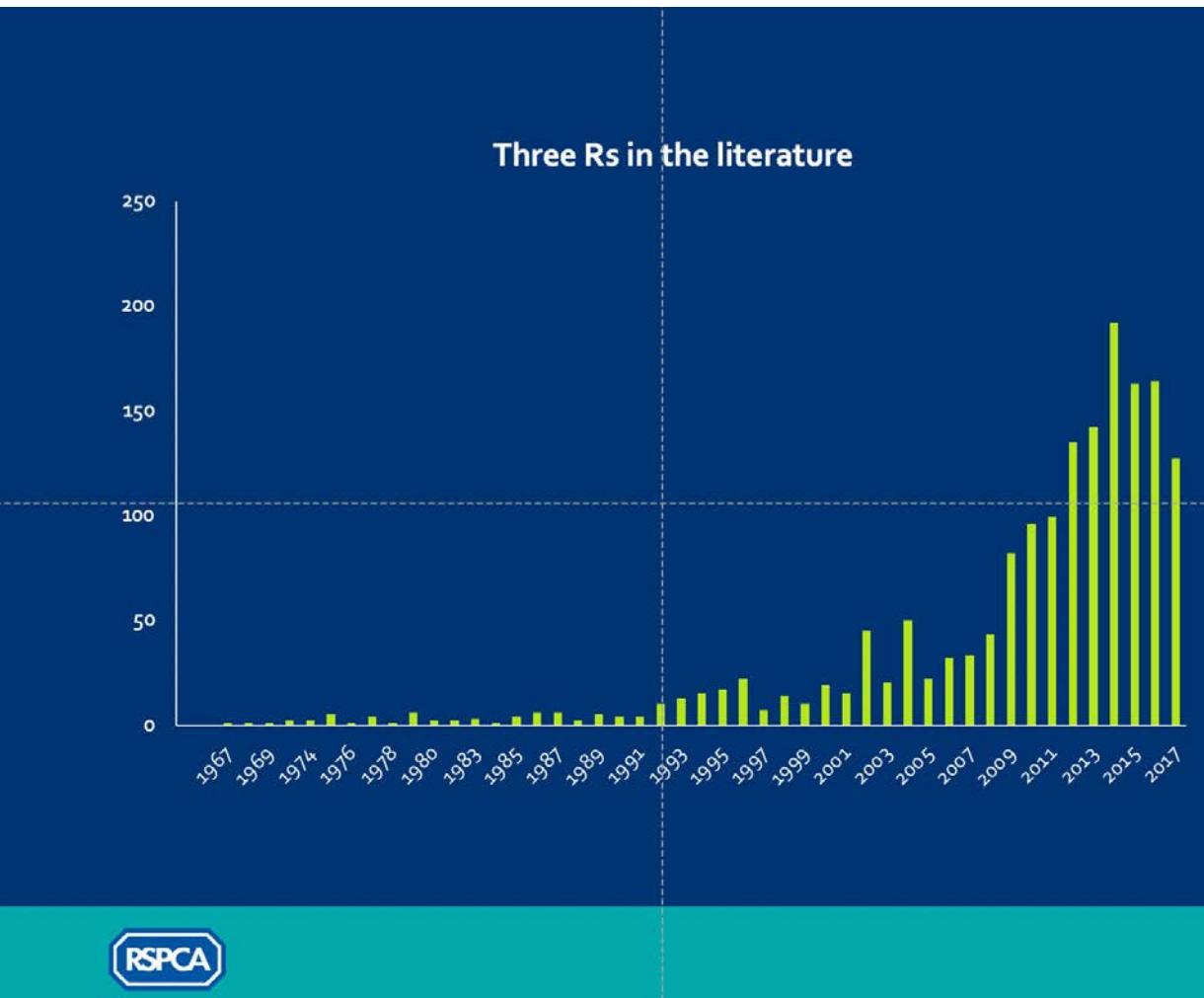
norecopa.no

Congratulations to the

FIN3R Centre!

norecopa.no/FIN3R





norecopa.no/media/7985/elliot-lilley-101017.pdf

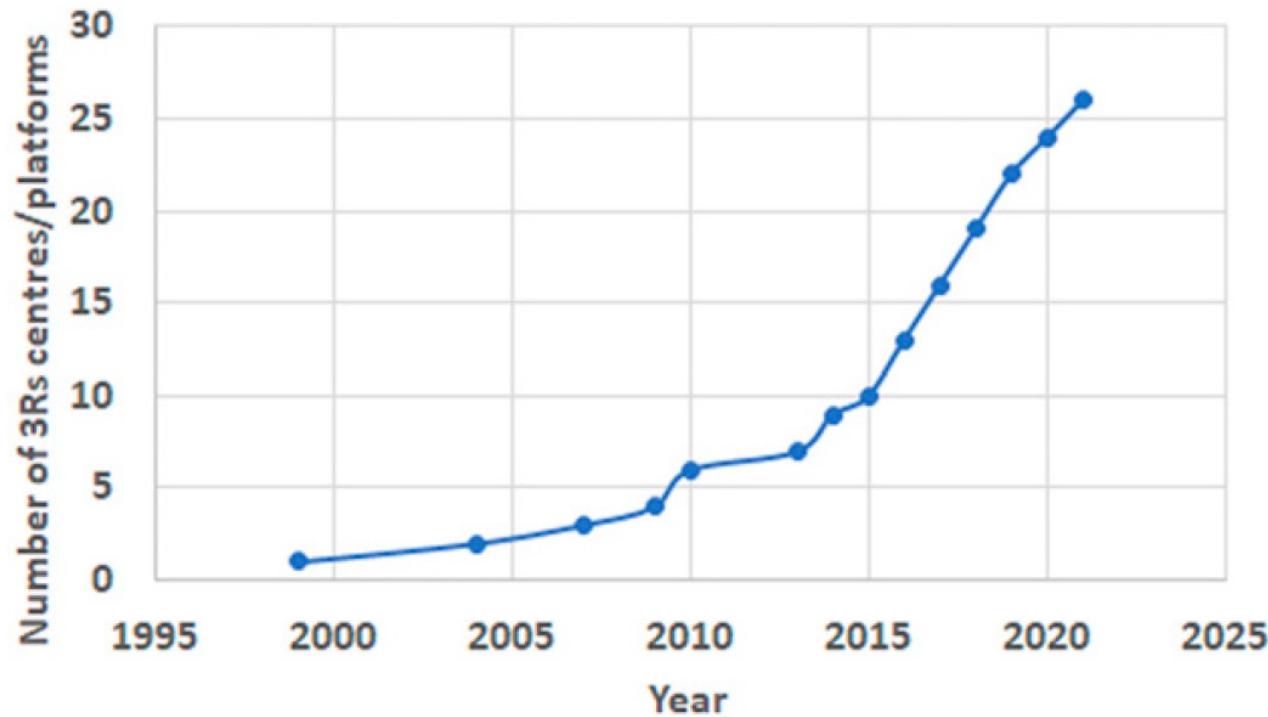


Figure I. The cumulative increase in the number of Three Rs centres and platforms in Europe over recent years.

Neuhaus *et al.*, 2022

50th Anniversary Article

The Rise of Three Rs Centres and Platforms in Europe*

Alternatives to Laboratory Animals
2022, Vol. 0(0) 1–31
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- Centres
- [Replacement](#) i
 - [Reduction](#) i
 - [Refinement](#) i
 - [ecopa](#) i
- Associations
- [ACURET](#) i
 - [AFLAS \(includes South Korea\)](#) i
 - [Culture of Care Network](#) i
 - [ecopa](#) i
 - [EU-NETVAL](#) i
 - [EU3Rnet](#) i
 - [FELASA](#) i
 - [FESSACAL](#) i
 - [Scand-LAS](#) i
 - [Concordat on Openness](#) i



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Is Fin3R:

- An animal welfare organisation?
- A society for the interests of scientists?
- Part of the authorities' monitoring and regulatory system?
- A funder of 3R research & development (budget)?

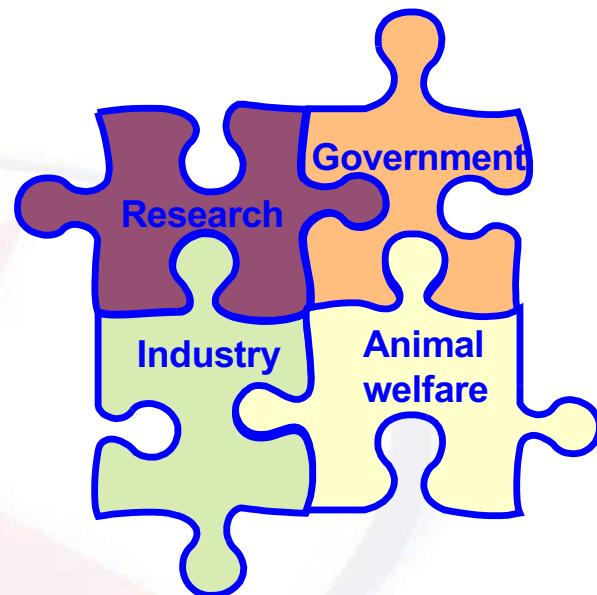
Norecopa =

*“Norway’s **Consensus Platform** for the
Replacement, Reduction and Refinement of Animal Experiments”*

European Consensus-Platform for Alternatives
ecopa.eu



Recognises National Consensus Platforms (NCPs) with 4 stakeholders
equally represented:



Other stakeholders? e.g. human patients



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Relationship to Fincopa, FICAM, FinLAS, FALAV, Finnish Reproducibility Network, National Committee, TOKES, and other 3R centres

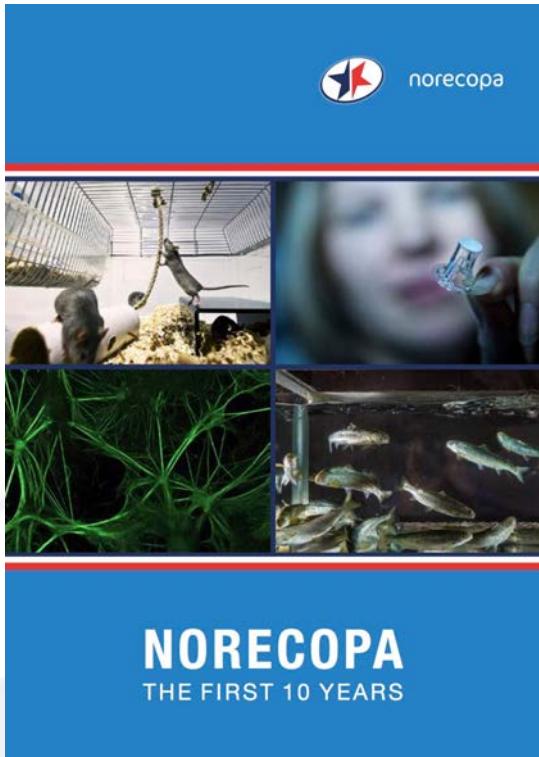
No common model for European/international 3R centres

Maybe with a special emphasis on knowledge dissemination about NAM's?



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2007-2017

<https://norecopa.no/about-norecopa/the-first-10-years>

Statutes (revised when necessary)

Activity Plan (revised yearly)

3R Prize (annual)

Web traffic

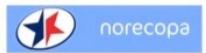
Annual Report with Milestones

Anniversary Reports



Planning guidelines: PREPARE

norecopa.no/PREPARE



PREPARATION

The PREPARE Guidelines Checklist

Recommendation		Definition of the study
1. Literature review	• Does a literature search, with primary and secondary sources, to identify relevant studies.	• Conduct the systematic review.
	• Identify gaps in the literature.	• Identify gaps in the literature of the question of interest. If the topic is unclear, identify the hypothesis associated with the research and its relevance.
2. Logical review	• Does a logical review of the evidence to answer the research question.	• Identify how the evidence is affected by relevant risk factors for patient selection and other risks, e.g. study design, incomplete health care, and study quality.
	• Identify the evidence that is relevant to the question of interest.	• Identify the evidence that is relevant to the question of interest.
3. Clinical benefit assessment and health impacts	• Does a clinical benefit assessment.	• Identify with other committees, consider whether the type of research has already been done.
	• Does a clinical benefit assessment, including risk and the risk versus great value, great availability.	• Assess the clinical benefit and the publication of negative results.
4. Evidence synthesis	• Does an evidence synthesis.	• Review the existing literature if the animal is used as a tool in testing processes.
	• Define study design, methodology, and analysis.	• Define study design, methodology, and analysis.
5. Peer review	• Does a peer review, for use and for end-point.	• Define the peer review process.
	• Define the peer review process.	• Define the peer review process.
6. Final report	• Does a final report.	• Define the final report.
	• Define the final report.	• Define the final report.
		(C) Quality control of the components in the study
7. Study design and methods	• Does a detailed description of the study design and methods.	• Define and describe all methods and procedures used at all stages of the study.
	• Define the study design and methods.	• Define the study design and methods.
8. Data collection and analysis	• Does a detailed description of the data collection and analysis.	• Relate the findings and conclusions of the study.
	• Define the data collection and analysis.	• Define the data collection and analysis.
9. Health risks, benefits, and costs/benefit-risk assessment and risk communication	• Does a detailed description of the health risks, benefits, and costs/benefit-risk assessment.	• Define a risk assessment, in accordance with the study design, for all persons and animals affected.
	• Define the health risks, benefits, and costs/benefit-risk assessment.	• Define the health risks, benefits, and costs/benefit-risk assessment.
10. Safety	• Does a detailed description of the safety.	• Define the safety.
	• Define the safety.	• Define the safety.
11. Peer review	• Does a detailed description of the peer review.	• Provide as much information as possible about the substances.
	• Define the peer review.	• Define the peer review.
12. Dissemination	• Does a detailed description of the dissemination of the results for the study to the public.	• Define the dissemination of the results.
	• Define the dissemination of the results.	• Define the dissemination of the results.
13. Sustainability	• Does a detailed description of the measures taken to prevent, assess and reduce, health monitoring and consequences for the person.	• Define the sustainability.
	• Define the sustainability.	• Define the sustainability.
14. Housing and husbandry	• Does a detailed description of the specific animal and rodent, in accordance with staff.	• Define the specific animal and rodent.
	• Define the specific animal and rodent.	• Define the specific animal and rodent.
15. Experimental procedures	• Does a detailed procedure for animal registration, handling, training, and exercise.	• Define the experimental procedures.
	• Define the experimental procedures for administration, sampling, analysis, and/or surgery and other techniques.	• Define the experimental procedures for administration, sampling, analysis, and/or surgery and other techniques.
16. Nutrition	• Does a detailed description of the nutrition.	• Define the nutrition.
	• Define the nutrition.	• Define the nutrition.
17. Veterinary care	• Does a detailed description of the veterinary care.	• Define the veterinary care.
	• Define the veterinary care.	• Define the veterinary care.
18. Monitoring	• Does a detailed description of the monitoring.	• Define the monitoring.
	• Define the monitoring.	• Define the monitoring.
		(D) Quality control of the components in the report
1. Report title	• Does a detailed description of the report title.	1.1. Report title: <i>Guidelines for the use of PRRSV in laboratory Animal Research and Testing</i>
	• Define the report title.	1.2. Report number: <i>SG-001</i>
2. Report date	• Does a detailed description of the date of the report.	2.1. Report date: <i>01/01/2018</i>
	• Define the date of the report.	2.2. Report version: <i>SG-001</i>
3. Report author	• Does a detailed description of the author.	3.1. Report author: <i>SG-001</i>
	• Define the author.	3.2. Report version: <i>SG-001</i>
4. Report publisher	• Does a detailed description of the publisher.	4.1. Report publisher: <i>SG-001</i>
	• Define the publisher.	4.2. Report version: <i>SG-001</i>
5. Report addressee	• Does a detailed description of the addressee.	5.1. Report addressee: <i>SG-001</i>
	• Define the addressee.	5.2. Report version: <i>SG-001</i>
6. Report purpose	• Does a detailed description of the purpose.	6.1. Report purpose: <i>SG-001</i>
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7. Report scope	• Does a detailed description of the scope.	7.1. Report scope: <i>SG-001</i>
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8. Report content	• Does a detailed description of the content.	8.1. Report content: <i>SG-001</i>
	• Define the content.	8.2. Report version: <i>SG-001</i>
9. Report structure	• Does a detailed description of the structure.	9.1. Report structure: <i>SG-001</i>
	• Define the structure.	9.2. Report version: <i>SG-001</i>
10. Report format	• Does a detailed description of the format.	10.1. Report format: <i>SG-001</i>
	• Define the format.	10.2. Report version: <i>SG-001</i>
11. Report language	• Does a detailed description of the language.	11.1. Report language: <i>SG-001</i>
	• Define the language.	11.2. Report version: <i>SG-001</i>
12. Report classification	• Does a detailed description of the classification.	12.1. Report classification: <i>SG-001</i>
	• Define the classification.	12.2. Report version: <i>SG-001</i>
13. Report distribution	• Does a detailed description of the distribution.	13.1. Report distribution: <i>SG-001</i>
	• Define the distribution.	13.2. Report version: <i>SG-001</i>
14. Report rights	• Does a detailed description of the rights.	14.1. Report rights: <i>SG-001</i>
	• Define the rights.	14.2. Report version: <i>SG-001</i>
15. Report obligations	• Does a detailed description of the obligations.	15.1. Report obligations: <i>SG-001</i>
	• Define the obligations.	15.2. Report version: <i>SG-001</i>
16. Report disclaimer	• Does a detailed description of the disclaimer.	16.1. Report disclaimer: <i>SG-001</i>
	• Define the disclaimer.	16.2. Report version: <i>SG-001</i>
17. Report signature	• Does a detailed description of the signature.	17.1. Report signature: <i>SG-001</i>
	• Define the signature.	17.2. Report version: <i>SG-001</i>
18. Report date	• Does a detailed description of the date.	18.1. Report date: <i>SG-001</i>
	• Define the date.	18.2. Report version: <i>SG-001</i>
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44. Report rights	• Does a detailed description of the rights.	44.1. Report rights: <i>SG-001</i>
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	• Define the rights.	74.2. Report version: <i>SG-001</i>
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	• Define the date.	78.2. Report version: <i>SG-001</i>
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	• Define the rights.	89.2. Report version: <i>SG-001</i>
90. Report obligations	• Does a detailed description of the obligations.	90.1. Report obligations: <i>SG-001</i>
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92. Report signature	• Does a detailed description of the signature.	92.1. Report signature: <i>SG-001</i>
	• Define the signature.	92.2. Report version: <i>SG-001</i>
93. Report date	• Does a detailed description of the date.	93.1. Report date: <i>SG-001</i>
	• Define the date.	93.2. Report version: <i>SG-001</i>
94. Report version	• Does a detailed description of the version.	94.1. Report version: <i>SG-001</i>
	• Define the version.	94.2. Report date: <i>SG-001</i>
95. Report addressee	• Does a detailed description of the addressee.	95.1. Report addressee: <i>SG-001</i>
	• Define the addressee.	95.2. Report version: <i>SG-001</i>
96. Report publisher	• Does a detailed description of the publisher.	96.1. Report publisher: <i>SG-001</i>
	• Define the publisher.	96.2. Report version: <i>SG-001</i>
97. Report date	• Does a detailed description of the date.	97.1. Report date: <i>SG-001</i>
	• Define the date.	97.2. Report version: <i>SG-001</i>
98. Report content	• Does a detailed description of the content.	98.1. Report content: <i>SG-001</i>
	• Define the content.	98.2. Report version: <i>SG-001</i>
99. Report structure	• Does a detailed description of the structure.	99.1. Report structure: <i>SG-001</i>



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*Looking forward to collaboration
Best wishes for the future!*

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