

Forsøksdyr: den skjulte kostnaden

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norecopa.no/kostnad



<https://norecopa.no>

peta.org



“Det du ikke vet, har de vondt av”



Thanks to animal research, they'll be able to protest 20.8 years longer.

 A group of protesters is shown holding signs that say "Stop Animal Research" and "DO NOT". One sign also says "NEEDS IT? TO MAKE STRONG!".

According to the US Department of Health and Human Services, animal research has helped extend our life expectancy by 20.8 years. Of course, how you choose to spend those extra years is up to you.

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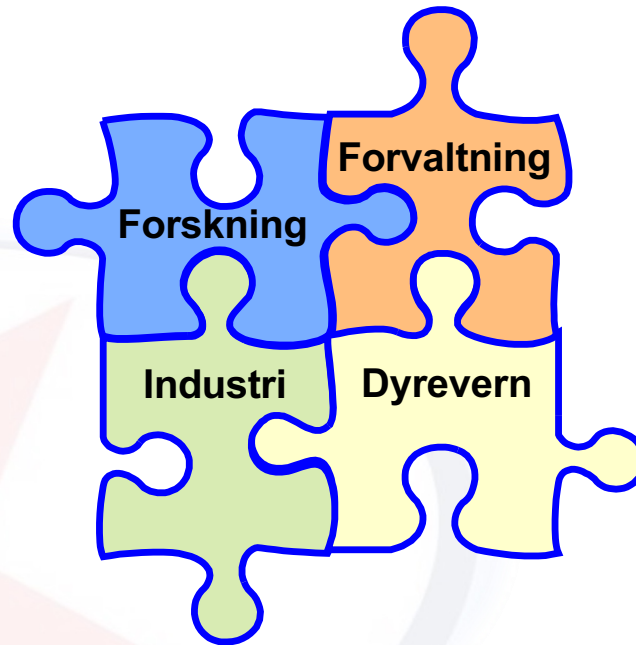
Norecopa: PREPARE for better Science

Norecopa er Norges nasjonale **konsensus**-plattform,

som skal *fremme alle de tre R'ene*:

Replacement, Reduction and Refinement

Styret representerer:

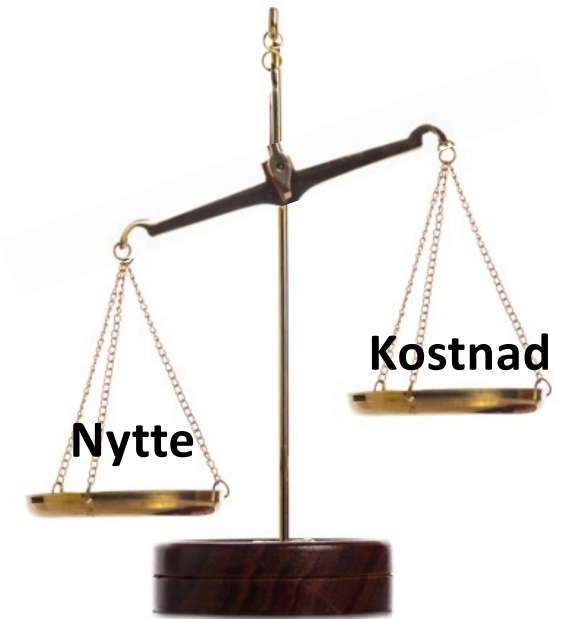


Stiftet i 2007

Hva er positivt med dyreforsøk?

- ✓ **Økt kunnskap** - i håp om nytte senere
- ✓ **Nytte for dyr**
 - ✓ bedre dyrevelferd
 - ✓ bedre sykdomsbehandling/profylakse
- ✓ **Nytte for mennesker**
 - ✓ bedre sykdomsbehandling/profylakse
 - ✓ HMS
 - ✓ utdannings- opplæringsøyemed
 - ✓ økonomisk gevinst

En viktig brikke i arbeidet med **Én Helse** (miljø/mennesker/dyr)



Det positive forutsetter:

- Klare og realistiske mål med forsøkene
- Korrekt eksperimentelt design – randomisering, blinding, tilstrekkelig statistisk styrke
- Korrekt statistisk analyse – ingen p-hacking!
- Ingen HARK-ing (Hypothesising after the Results are Known)
- Rapportering av **"alt"** (også negative og belastende funn) - for at ikke nye dyr skal brukes til det samme



norecopa.no/concerns

Vanlige kostnader ved alle dyreforsøk

- I motsetning til nytteverdien, som er *i fremtiden og usikker*: Belastningen på forsøksdyret er **NÅ og GARANTERT**
- Den eventuelle nytteverdien hjelper sjeldent selve forsøksdyret, men (noen ganger) sine artsfrender (som i husdyrforskning - dog ofte i menneskers interesser)
- Det kan være tvil om resultatene er pålitelige



Etter dyreforsøket

Det kan være vanskelig å vei ***kostnadene for enkeltdyrene*** opp mot ***nytteverdien for andre***

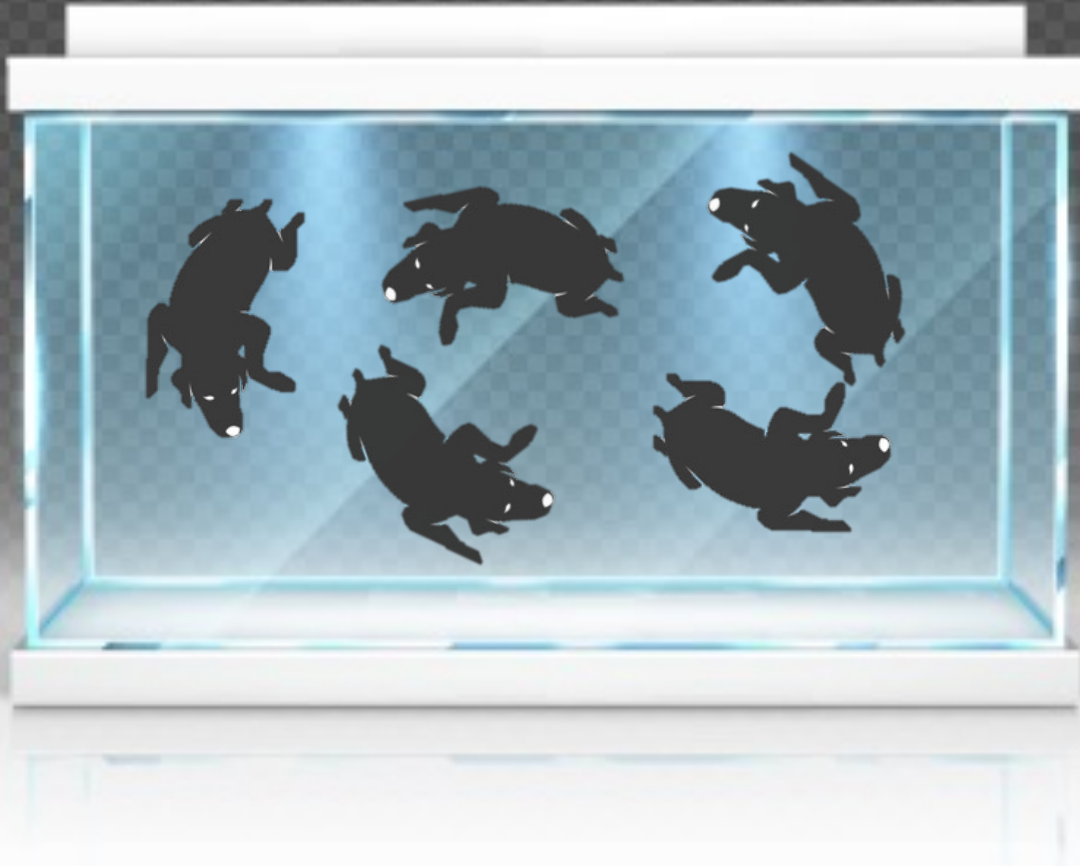


Forsøksdyret opplever ofte det stikk motsatte av “De 5 Frihetene” for dyrevelferd

- Frihet fra sult, tørste og feilernæring
- Frihet fra fysisk ubehag
- Frihet fra smerte, sykdom og skade
- Frihet til å utøve normal atferd
- Frihet fra frykt og stress

Fiskeforsøk har en del tilleggskostnader:

Selv enkel håndtering og enkle prosedyrer innebærer oftest å ta dem ut av sitt fysiologisk riktige element



Tilleggskostnader:



foto: T. Poppe



Fysiologiske endringer som de gjennomgår
Størrelsesvariasjoner mellom jevngamle dyr
Oppstallingsbehov – stim eller individuelt?
Bedøvelse og smertestillelse
m.m.

Vi kan påvirke fiskeforsøk i riktig retning

De 3 R'ene for å minimere belastningen:

- Erstatte de unødvendige forsøk
- Redusere antallet dyr pr. forsøk
- Forbedre forholdene for dyrene som må brukes

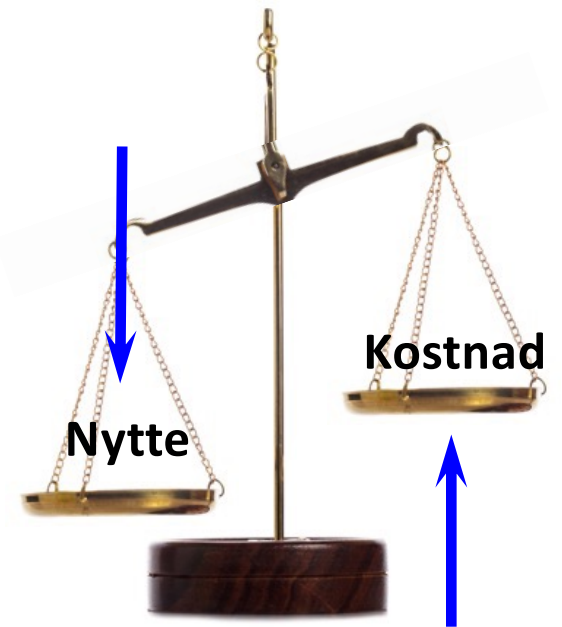
De 3 S'ene - det er lov å bruke både hue og hjertet

- Good Science
- Good Sense
- Good Sensibilities



De 3 V'ene for å øke gyldigheten av forsøket:

- Construct Validity (kan dyremodellen gi svaret på problemstillingen?)
- Internal Validity (er forsøket designet riktig?)
- External Validity (er resultatene overførbare til måldyrene/mennesker?)



norecopa.no/3R

norecopa.no/3S

norecopa.no/3V



Working Party Report

Guidance on the severity classification of scientific procedures involving fish: report of a Working Group appointed by the Norwegian Consensus-Platform for the Replacement, Reduction and Refinement of animal experiments (Norecopa)

P Hawkins (Convenor), N Dennison, G Goodman, S Hetherington, S Llywelyn-Jones, K Ryder and A J Smith

Research Animals Department, RSPCA, Welford Way, Southwark, West Sussex RH43 9RL, UK; Animals (Scientific Procedures) Inspectorate, Home Office, PO Box 4770, Dundee DD1 9BQ, UK; Biological Services, The University of Edinburgh, Charlotte Building, 48, Little France Crescent, Edinburgh EH8 4EB, UK; CERIAS, Paisley Road, Lowestoft, NR33 0HT, UK; King's College London, Biomedical Services Unit, 4th floor, Hodgkin Building, Guy's Campus, London SE1 1UL, UK; Norecopa, c/o Norwegian Veterinary Institute, PO Box 750 Sentrum, N-0108 Oslo, Norway

Abstract
The severity classification of procedures using animals is an important tool to help focus the implementation of refinement and to assist in reporting the application of the 3Rs (replacement, reduction and refinement). The recently revised Directive that regulates animal research and testing within the European Union requires Member States to ensure that all procedures are classified as 'non-recovery', 'mild', 'moderate' or 'severe', using assignment criteria set out by the European Commission (EC). However, these are focused upon terrestrial species, so are of limited relevance to fish users. A Working Group set up by the Norwegian Consensus-Platform for the 3Rs (Norecopa) has produced guidance on the classification of severity in scientific procedures involving fish, including examples of 'sub-threshold', 'mild', 'moderate', 'severe' and 'upper threshold' procedures. The aim is to complement the EC guidelines and help to ensure that suffering in fish is effectively predicted and minimized. Norecopa has established a website (www.norecopa.no/categories) where more information on severity classification for procedures using fish, including field research, will be made available.

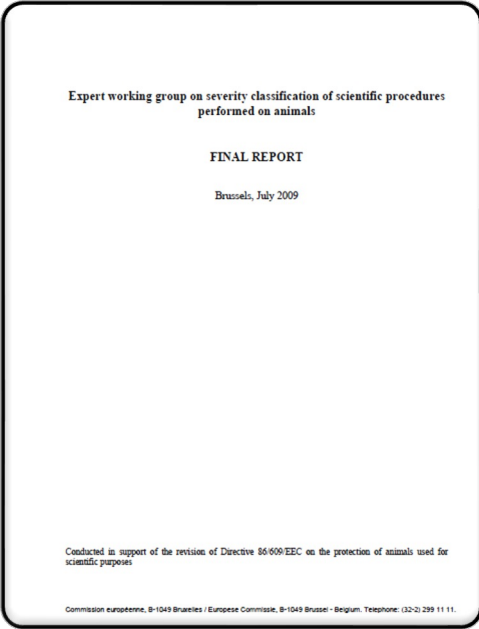
Keywords: Fish, harm-benefit assessment, humane endpoints, refinement, severity
Laboratory Animals 2011; 1-6. DOI: 10.1258/la.2011.010181

Background
An effective prediction of the effects of a research protocol on the animals concerned helps to ensure that any pain, suffering or distress they may experience will be effectively anticipated, recognized and alleviated. This is essential not only for animal welfare but also for scientific validity, because physiological and behavioural responses to suffering can significantly affect data quality. Severity classification is thus an important tool to help focus the implementation of refinement, including monitoring its progress, and to assist in reporting the application of the 3Rs (replacement, reduction and refinement) of 'Ranaid and Biach', which is now an integral part of the legislation on animal research and testing in many countries. Predictions of severity are also fundamental to the 'harm-benefit

assessment undertaken by bodies such as regulatory authorities and ethical committees when deciding whether or not a project should be licensed or funded. There may also be a legal requirement to predict and classify severity. For example, the new Directive regulating animal use within the European Union, which must be implemented within all Member States by January 2013, requires the severity of each procedure to be classified on the basis of the degree of pain, suffering, distress or lasting harm expected to be experienced by an individual animal during the course of the procedure, with the aim of enhancing transparency, facilitating the project authorization process and providing tools for monitoring compliance. Member States will have to ensure that all procedures are classified as 'non-recovery', 'mild', 'moderate' or 'severe' on a case-by-case basis, using the assignment

Guidance on the severity classification of procedures involving fish

Et supplement til EUs retningslinjer:



http://ec.europa.eu/environment/chemicals/lab_animals/pdf/report_ewg.pdf

P Hawkins, N Dennison, G Goodman, S Hetherington, S Llywelyn-Jones, K Ryder and AJ Smith (2011)

norecopa.no/categories

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PREPARE



The PREPARE Guidelines Checklist Planning Research and Experimental Procedures on Animals: Recommendations for Excellence

Adrian J. Smith¹, R. Eddie Clutton², Elliot Lilley³, Kristine E. Aa. Hansen⁴ & Tord Brattli⁵
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PREPARE consists of planning guidelines which are complementary to reporting guidelines such as ARRIVE¹. PREPARE covers the three broad areas which determine the quality of the preparation for animal studies:

1. Formulation of the study
2. Dialogue between scientists and the animal facility
3. Quality control of the components in the study

The topics will not always be addressed in the order in which they are presented here, and some topics overlap. The PREPARE checklist can be adapted to meet special needs, such as field studies. PREPARE includes guidance on the measurement of animal welfare, since in-house experiments are dependent upon their quality. The full www.norecopa.no website, with links to global resources, is available upon their quality. The full www.norecopa.no website, with links to global resources, is available upon their quality. The full www.norecopa.no website, with links to global resources, is available upon their quality.

Topic	Recommendation
(A) Formulation of the study	
1. Literature searches	<input type="checkbox"/> Form a clear hypothesis, with primary and secondary objectives. <input type="checkbox"/> Consider the use of systematic reviews. <input type="checkbox"/> Decide upon databases and information specialist services, and construct search terms. <input type="checkbox"/> Assess the relevance of the species to be used, its biology and suitability to answer the experimental questions with the least suffering, and its welfare needs. <input type="checkbox"/> Assess the reproducibility and translatability of the project.
2. Legal issues	<input type="checkbox"/> Consider how the research is affected by relevant legislation for animal research and other areas, e.g. animal transport, occupational health and safety. <input type="checkbox"/> Locate relevant guidance documents (e.g. EU guidance on project evaluation).
3. Ethical issues, harm-benefit assessment and humane endpoints	<input type="checkbox"/> Construct a lay summary. <input type="checkbox"/> In dialogue with ethics committees, consider whether statements about this type of research have already been produced. <input type="checkbox"/> Address the 3Rs (replacement, reduction, refinement) and the 3Ss (good science, good sense, good sensibilities). <input type="checkbox"/> Consider pre-registration and the publication of negative results. <input type="checkbox"/> Perform a harm-benefit assessment and justify any likely animal harm. <input type="checkbox"/> Discuss the learning objectives, if the animal use is for educational or training purposes. <input type="checkbox"/> Allocate a severity classification to the project. <input type="checkbox"/> Define objective, easily measurable and unequivocal humane endpoints. <input type="checkbox"/> Discuss the justification, if any, for death as an end-point.
4. Experimental design and statistical analysis	<input type="checkbox"/> Consider pilot studies, statistical power and significance levels. <input type="checkbox"/> Define the experimental unit and decide upon animal numbers. <input type="checkbox"/> Choose methods of randomisation, prevent observer bias, and decide upon inclusion and exclusion criteria.

En sjekkliste (+ tilhørende nettsider) for bedre forskning



Topic	Recommendation
(B) Dialogue between scientists and the animal facility	
5. Objectives and timescale, funding and division of labour	<input type="checkbox"/> Arrange meetings with all relevant staff when early plans for the project exist. <input type="checkbox"/> Construct an approximate timescale for the project, indicating the need for assistance with preparation, animal care, procedures and waste disposal/decontamination. <input type="checkbox"/> Discuss and disclose all expected and potential costs. <input type="checkbox"/> Construct a detailed plan for division of labour and expenses at all stages of the study.
6. Facility evaluation	<input type="checkbox"/> Conduct a physical inspection of the facilities, to evaluate building and equipment standards and needs. <input type="checkbox"/> Discuss staffing levels at times of extra risk.
7. Education and training	<input type="checkbox"/> Assess the current competence of staff members and the need for further education or training prior to the study.
8. Health risks	<input type="checkbox"/> Perform risk assessments for all persons and animals affected by the project. <input type="checkbox"/> Discuss the need for protective measures for all persons and animals affected by the project.
(C) Quality control of the components in the study	
<input type="checkbox"/> Provide as much information as possible about test substances, including the identity, source, and quality of the substances, and the suitability and validity of test procedures and the skills needed to perform them.	
<input type="checkbox"/> Decide upon the characteristics of the animals that are essential for the study and for reporting.	
<input type="checkbox"/> Avoid generation of surplus animals.	
11. Quarantine and health monitoring	<input type="checkbox"/> Discuss the animals' likely health status, any needs for transport, quarantine and isolation, health monitoring and consequences for the personnel.
12. Housing and husbandry	<input type="checkbox"/> Attend to the animals' specific instincts and needs, in collaboration with expert staff. <input type="checkbox"/> Discuss acclimatization, optimal housing conditions and procedures, environmental factors and any experimental limitations on these (e.g. food deprivation, solitary housing).
13. Experimental procedures	<input type="checkbox"/> Develop refined procedures for capture, immobilisation, marking, and release or rehoming. <input type="checkbox"/> Develop refined procedures for substance administration, sampling, sedation and anaesthesia, surgery and other techniques.
14. Humane killing, release, reuse or rehoming	<input type="checkbox"/> Consult relevant legislation and guidelines well in advance of the study. <input type="checkbox"/> Define primary and emergency methods for humane killing. <input type="checkbox"/> Assess the competence of those who may have to perform these tasks.
15. Necropsy	<input type="checkbox"/> Construct a systematic plan for all stages of necropsy, including location, and identification of all animals and samples.

References
 1. Smith AJ, Clutton RE, Lilley E, Hansen KEA & Brattli T. PREPARE: Guidelines for Planning Animal Research and Testing. *Laboratory Animals*, 2017, DOI: 10.1177/0023677217724923.
 2. Kilkenny C, Browne WJ, Cuthill IC et al. Improving Bioscience Research Reporting: The ARRIVE Guidelines for Reporting Animal Research. *PLoS Biology*, 2010, DOI: 10.1371/journal.pbio.1000412.

Further information
<https://norecopa.no/PREPARE> | post@norecopa.no | [@norecopa](https://twitter.com/norecopa)

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- 3-Ethical issues, harm-benefit assessment and humane endpoints
 - 3a Construct a lay summary.
 - 3b In dialogue with ethics committees, consider whether statements about this type of research have already been produced.
 - 3c Address the 3Rs (Replacement, Reduction, Refinement) and the 3Ss (Good Science, Good Sense, Good Sensibilities).
 - 3d Consider pre-registration and the publication of negative results.
 - 3e Perform a Harm-Benefit Assessment and justify any likely animal harm.
 - 3f Discuss the learning objectives, if the animal use is for educational or training purposes.
 - 3g Allocate a severity classification to the project.
 - 3h Define objective, easily measurable and unequivocal humane endpoints.
 - 3i Discuss the justification, if any, for death as an end-point.
- 4-Experimental design and statistical analysis

- 5. Have the experiments been carried out before, and is any repetition justifiable?
- 6. What [approaches to reduce distress](#) have been considered?

3a Construct a lay summary.

General principles For fish researchers

1. Have national or local research ethics committees already produced statements relevant to the research being planned? Consideration should also be paid to the broader context of the research. For example, research directed at increasing the productivity of farming at the expense of (or without improving) individual animal welfare, or wildlife research whose primary aim is population management.
2. Have the Three Rs ([Replacement, Reduction, Refinement](#)) been addressed, and will any advances in this area be mentioned in publications of the study (remembering that many databases only index the title and abstract of papers)? Which [non-animal alternatives](#) have been considered but rejected?
3. Have the Three S's ([Good Science, Good Sense and Good Sensibilities](#)) been addressed? Sufficient time should be allocated to this point, since two of the three S's are highly subjective, but equally important. The use of commonsense and critical anthropomorphism are justifiably part of the work to assess the impact of research on animals, not least when a scientific evidence base does not exist.
4. Does the proposed study have a clear rationale and scientific relevance, and what will be the next step if the hypothesis is supported or rejected?
5. Have the experiments been carried out before and is any repetition justifiable?
6. What [approaches to reduce distress](#) have been considered?
7. Will the project undergo [pre-registration](#) and will negative results be published, to avoid publication bias?

Many more [links to resources on ethics are available here](#).

Details about pre-registration of animal studies and reporting of critical incidents are to be found in the section on [Experimental Design and Statistical Analysis](#).

Harm-Benefit Assessment

Bruk av forsøksdyr (2021)



Antallet forsøksdyr er innrapporterte tall fra forskere og forsøksdyrvirksomheter.

Art	Antall
[A1] Mice (<i>Mus musculus</i>)	52 554
[A2] Rats (<i>Rattus norvegicus</i>)	4 498
[A3] Guinea-Pigs (<i>Cavia porcellus</i>)	120
[A4] Hamsters (Syrian) (<i>Mesocricetus auratus</i>)	30
[A7] Other Rodents (other Rodentia)	1 174
[A8] Rabbits (<i>Oryctolagus cuniculus</i>)	20
[A10] Dogs (<i>Canis familiaris</i>)	41
[A12] Other carnivores (other Carnivora)	139
[A13] Horses, donkeys & cross-breeds (Equidae)	92
[A14] Pigs (<i>Sus scrofa domesticus</i>)	705
[A15] Goats (<i>Capra aegagrus hircus</i>)	404
[A16] Sheep (<i>Ovis aries</i>)	569
[A17] Cattle (<i>Bos primigenius</i>)	294
[A27] Other Mammals (other Mammalia)	678
[A28] Domestic fowl (<i>Gallus gallus domesticus</i>)	867
[A29] Other birds (other Aves)	12 992
[A30] Reptiles (Reptilia)	10
[A33] Other Amphibians (other Amphibia)	500
[A34] Zebra fish (<i>Danio rerio</i>)	29 574
[A35] Other Fish (other Pisces)	2 186 386
SUM	2 291 647

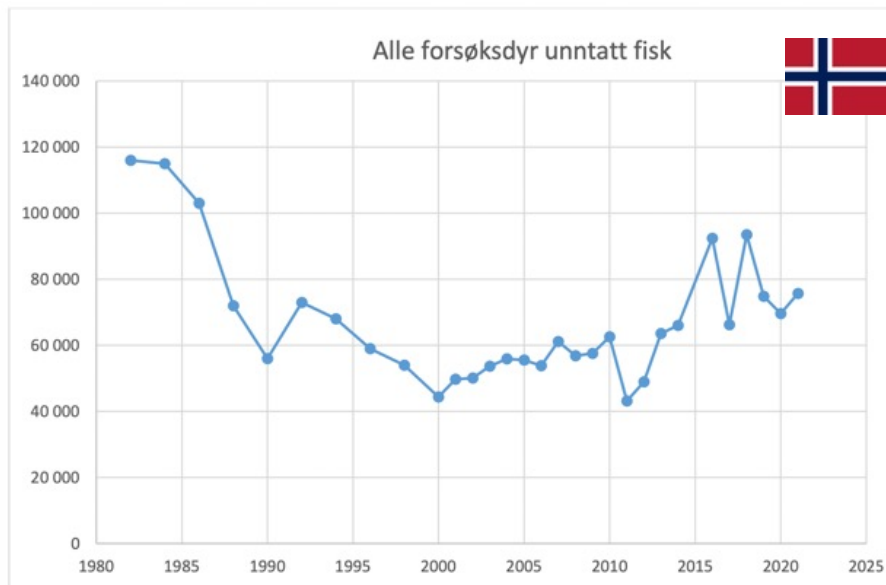


*Vaksineutvikling og testing
Utprøving av nye systemer for oppdrett
Forskning på dyrevelferd
Bekjempelse av lakselus*



Statistikken er lik for alle dyrearter:

- Belastningskategoriene er de samme
- Vanlig merking er ikke dyreforsøk
- Observasjoner er ikke dyreforsøk
- Det telles kun fisk som utsettes for prosedyrer



Norge og EU

	2015	2016	2017	2018 (EU-28 incl. NO) ²	2019 (EU-28 incl. NO) ²
Total	9,590,379	9,817,946	9,388,162	10,572,305	10,401,673

Table 1: Total numbers of animals used for the first time for research, testing, routine production and education purposes in the Union between 2015 and 2019 with the inclusion of data from Norway in 2018 and 2019

År	Sebrafisk	Alle fisk (inkl. sebrafisk)	Andre arter	Total
2003		796 497	53682	850 179
2004		267 375	55415	322 790
2005	2160	944 874	55552	1 000 426
2006	6049	670 235	53858	724 093
2007	4906	3 400 694	61170	3 461 864
2008	960	1 865 090	56862	1 921 952
2009	922	1 730 594	57579	1 788 173
2010	3355	1 357 795	62590	1 420 385
2011	1990	1 579 589	43125	1 622 714
2012	889	161 380	48986	210 366
2013	1551	5 458 434	63557	5 521 991
2014		4 823 202	65989	4 889 191
2015		1 140 975	89857	1 230 832
2016	8778	11 513 785	92383	11 606 168
2017	20724	1 093 413	66254	1 159 667
2018	38218	1 593 191	93467	1 686 658
2019	41148	1 206 789	74806	1 281 595
2020	38867	2 213 101	69609	2 282 710
2021	29574	2 215 960	75687	2 291 647
f.o.m. 2003		44 032 973	1240428	45 273 401

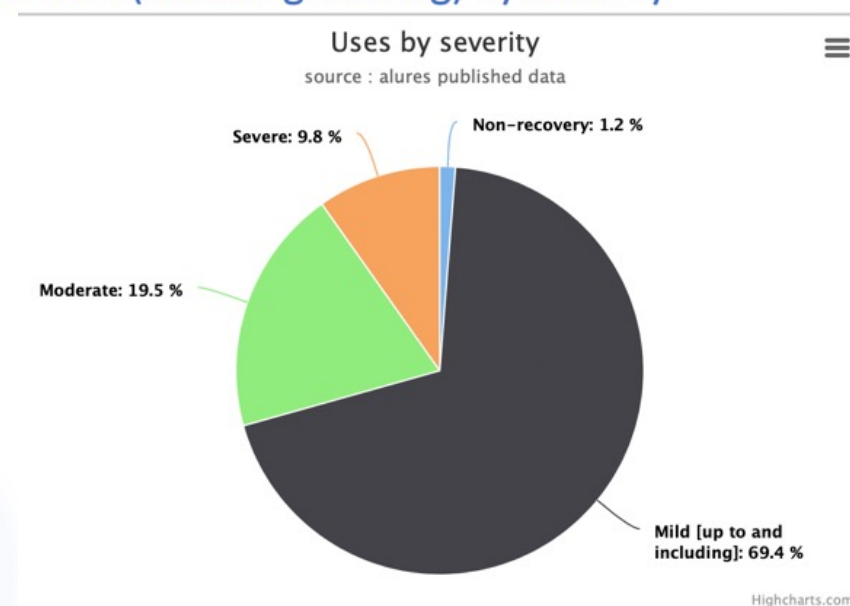
45 millioner forsøksdyr siden den forrige dyrevelferdsmeldingen:

- 44 millioner fisk
- 1,24 millioner landdyr

Norge: belastende dyreforsøk (tall fra 2019):

Uses of animals in research, testing, routine production and education (including training) by severity

Severity	Number of uses	Percentage
Non-recovery	36585	2.88%
Mild [up to and including]	866946	68.28%
Moderate	274767	21.64%
Severe	91434	7.2%
Total	1269732	100.00%



“Andre fisk” (ikke sebrafisk) i hele EU+Norge i 2019:
201.100 i betydelig belastende forsøk
400.000 i moderat belastende forsøk



Hvem vil ta kostnaden?

- Det norske forbruket av forsøksdyr er formidabelt. Vi må redusere denne bruken av levende dyr til eksperimenter som er strengt vitenskapelig nødvendig, sier landbruksminister Lars Sponheim til BT.

2003

Siden den forrige dyrevelferdsmeldingen...



2001-2005	Bondevik II	H, KrF, V
2005-2013	Stoltenberg II	A , Sp , SV
2013-2021	Solberg	H, FrP, V, KrF
2021-	Støre	A , Sp

Andre saker opptar stadig politikernes oppmerksomhet...

- *pelsdyr*
- *gris*
- *kyllinger*
- *veterinærvakten*

- *krigen i Ukraina*
- *energikrisen*
- *økningen i sosialutgifter*



...som fører til andre skjulte kostnader...

Arbeidstimer til:

- Lobbyvirksomhet
- Medieinnslag
- Leting etter støtte

Gratisarbeid



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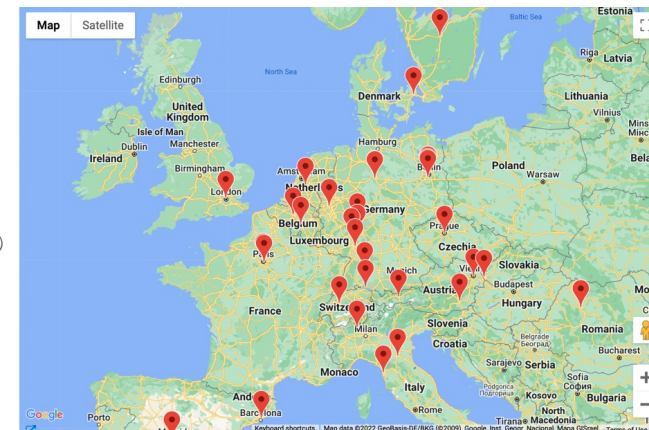
Det er bare fisk...?



...men vi bruker også relativt sett mange pattedyr, i et land uten stor farmasøytisk industri



Vi ser frem til en ny dyrevelferdsmelding i 2023



Den forrige meldingen (2003):

*Landbruksdepartementet vil derfor bidra til at **det opprettes en nasjonal plattform, et kompetansesenter, for alternativer til bruk av dyr i forskning slik det nå gjøres i en rekke europeiske land.***

*En slik plattform vil spre informasjon om alternativer nasjonalt og internasjonalt og **vil selv kunne initiere utvikling av alternative metoder og bedre oppstallingsystemer.***



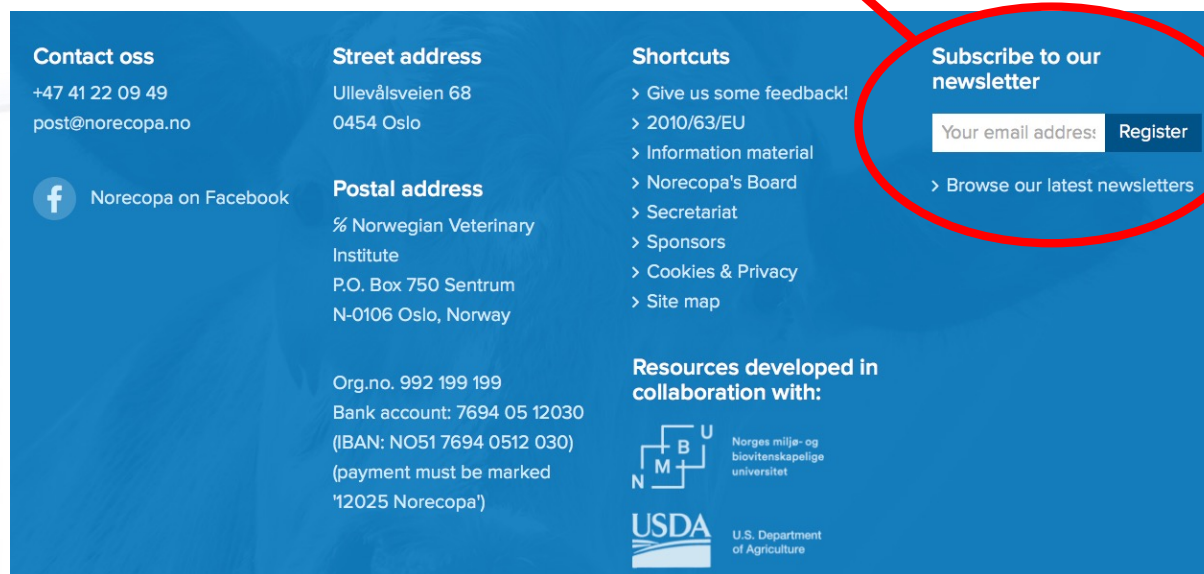
Mine konklusjoner

- 1. Det utvikles stadig nye 3R-alternativer til dyreforsøk, også i Norge*
- 2. Vi kan få ned antallet dyr, og belastningen, med en større innsats*
- 3. Vi trenger er **en solid norsk plattform, et 3R-senter**, som kan bidra til dette arbeidet.*

Takk for meg – og til forsøksdyrene

norecopa.no/kostnad

Nyhetsbrev - nederst på norecopa.no





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