The Nordic Alliance for Clinical Genomics

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NACG introduction slides

Updated 25. February 2020



NACG is an independent, non-governmental, not-for-profit Nordic association

We work together and learn from each other to lift our performance standards. We aim at responsible sharing of trustworthy data for improved diagnosis and treatment, and as a resource for research.







NACG goals



Facilitate the **responsible sharing** of genomic data, bioinformatics tools, sequencing methods and best practices for interpretation of genomic data.



Enhance quality of genomic data and processes, and explore methodologies to provide assurance.



Understand **legal barriers** to the implementation of personalized medicine and to engage with key stakeholders that influence these barriers



Develop **demonstration projects** that challenge perceived legal barriers that limit responsible and ethical sharing of genomic and health data.



Build bridges between research and clinical communities, technologies and practices to foster innovation





The NACG community





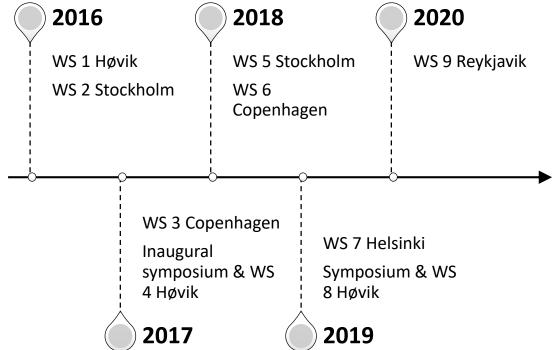


NACG organisation

Role	Name	Institution	Expiry of period
Steering Committee Chair	Dag E. Undlien	Oslo University Hospital, Dep of Medical Genetics	Nov 2022
Steering Committee Vice chair	Valtteri Wirta	Karolinska Institutet/ SciLifeLab	Nov 2020
Steering Committee Vice Chair	Morten Dunø	Rigshospitalet, Dep. of Clinical Genetics	Nov 2022
Steering Committee Member	Stephen McAdam	DNV GL	Nov 2020
Steering Committee Member	Jón Jóhannes Jónsson	Landspitali – National University Hospital	Nov 2020
Steering Committee Member	Janna Saarela	FIMM	Nov 2020
Steering Committee Member	Ane Yde Schmidt	Rigshospitalet, Center for Genomic Medicine	Nov 2022
Steering Committee Member	Kasper Thorsen	Danish Genome Centre	Nov 2022
Steering Committee Member	Gjertrud Bøhn Mageli	OUS legal, representative for Legal network	Nov 2022
Director of Secretariat	Guro Meldre Pedersen	DNV GL	



NACG development

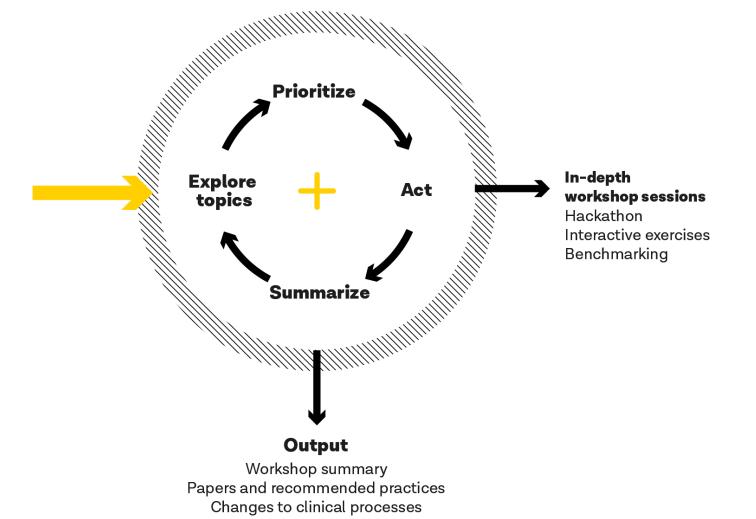


 Workshop reports and joint project reports available at <u>www.nordicclinicalgenomics.org/resources</u>





NACG - modus operandi



Nordic Alliance for Clinical Genomics



How to connect with NACG

Nordic Alliance for Clinical Genomics

About Events Resources Contact Us

The Nordic Alliance for Clinical Genomics (NACG)

The Nordic Alliance for Clinical Genomics is an association that gathers stakeholders in clinical genomics who collaborate to identify and address emerging challenges to the implementation of clinical genomics and precision medicine.

The Nordic Alliance for Clinical Genomics was named the Nordic Alliance for Sequencing and Precision Medicine (NASPM) until fall 2018

Find out more

How we work

Building on Nordic commonalities, advantages and shared challenges, NACG brings together professionals interested in sharing experiences, data and best practices for the implementation of precision medicine.



NACG website

- https://nordicclinicalgenomics.org/
- About
- Events
- Resources
 - NACG paper
 - NACG workshop reports
 - NACG governing documents
- How to join
 - Organisations
 - Individuals
- Contact us
 - post@nordicclinicalgenomics.org





How to join

- Open for organizational and individual members that will adhere to the guiding principles of NACG.
- Application forms available at https://nordicclinicalgenomics.org/
 - ..or contact us at post@nordicclinicalgenomics.org



Next workshop: Reykjavik 11-12 May 2020 More information to come at https://nordicclinicalgenomics.org (28. Feb 2020)

	Monday 11 May		Tuesday 12 May		-
A.S.M			Nordic consent framework and toolkit	Emerging technologies	No. of Contraction of
-			Preparing for IVDR	Collaborative software development	
	Lunch Welcome & keynote		Lunch		
			Preparing for IVDR	Collaborative software development	
	Cancer panel benchmarking	Variant interpretation and data sharing	Workshop closing & bus departure		
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