



The Intervention Centre - Clinical Test Facility at Oslo University Hospital

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Section for Method Development and Industry Collaboration

The Intervention Center



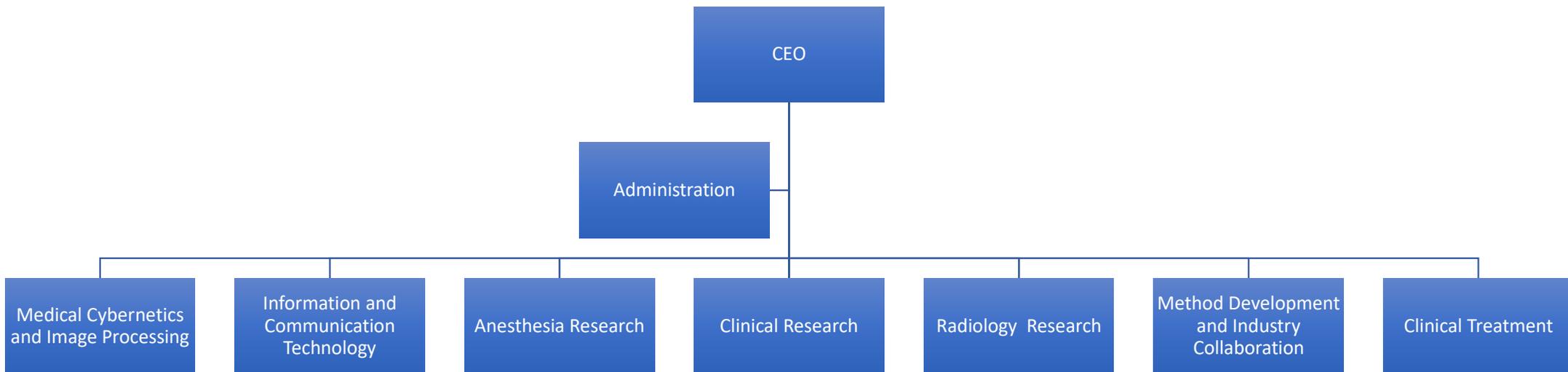
Mandate:

- Established in 1996 to introduce advanced technologies in medicine
 - Safe introduction of new technologies for the benefit of patients
- Department for Clinical Research and Development
- Image guided and minimally invasive therapies
- Support industry Collaboration:
- Non-production based funding

1. Cross Disciplinary Staff
2. Hybrid/ Multi Modal OR's
3. Backup to Gold Standard



Organization



Separation of R&D and production



Research Groups

1. Medical Robotics, visualization and navigation
2. Research Group for Industry Sponsored Studies
3. BigMed
4. Wireless Biomedical Sensor Network Research Group
5. Value-based and International Health Care Research
6. Clinical and experimental cardiovascular monitoring
7. MR guided High Intensity Focused Ultrasound treatment
8. Image guided general surgery and intervention



Infrastructure assets

Multi modal OR's:

- 2 Hybrid Angio OR's
- 1 CT on rails
- OR + Intraoperative CT
- Hybrid Videoscopic OR
- MRI Adjacent OR
- 2 MRI's
- Preclinical OR

-OR Lab

-IoT Lab

Formal Requirements



A screenshot of the Norwegian Medicines Act website (lovdata.no). The page displays the text of the Act, including the title 'Lov om medisinsk og helsefaglig forskning (helseforskningsloven)'. The text is organized into sections, with 'Kapittel 1. Lovens formål og virkeområde' being the most prominent. The page also features a search bar and a navigation menu on the left side.

A screenshot of the REK (Regional Committees for Medical Research Ethics) website. The page features the REK logo and the text 'REK REGIONAL COMMITTEES FOR MEDICAL RESEARCH ETHICS'. Below the logo, there are links for 'Deadlines and Forms' and 'Rules and Procedures'. The page also includes a section titled 'About Applying To REC' and a list of items that REK shall provide advance approval for.

About Applying To REC

REC shall provide advance approval for:

- Medical and health research projects
- General and thematic research biobanks
- Dispensation from professional secrecy requirements for other types of research

More information about applying to REC can be found under the specific menu option - see in particular **Rules and Procedures** and **Deadlines and Forms**.

Applications must be sent to REC in electronic format. You have to log in to get access to the forms.

Latest messages

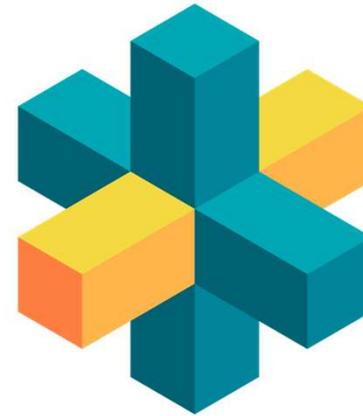
Change in publication of final reports from 12.4.2013
The submitted form for the final report to REC (Sluttmelding) is from 12.4.2013 onwards, made publicly available in its full text in the REC Public Records - Application Database. For final reports submitted before this date, the publication attached to the form is publicly available.

If you have not logged on to the portal since January 5th, you have to order a new password
Updating of the REC case portal January 6th, requires every user to order a new password. Before logging in, you must clear your browser cache and delete cookies. Please remember to edit your user profile in order to access your forms and applications. We apologize for any inconvenience. If you have problems logging in, please contact your local REC, or send an email to post@helseforskning.etikkom.no

A screenshot of the REK login page. The page features a 'Logg inn' button and a 'Create new user' link. Below the login fields, there are images of a microscope, a person, and a DNA helix. The page also includes a 'Forgotten/new password' link and a 'Change password' link.



Partners



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Competitive strenghts

- Preclinical Tials
- Human Trials
- Technology Trials

QoS

- Formalized collaboration with OUS
 - Nordic Proof
- High quality digital documentation
- Transparent public health system
 - Longitudinal patient follow up
- TestBed
 - Dedicated Infrastructure
 - Expertize: Clinical, Imaging, Technology, IT
 - “First in Man”/Early Phase Studies

