

# **New information and consent forms ECSS and DCC**

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# Why new forms?

- Previous forms too long and too hard to understand.
- No uniformity across Radboud University
- RU Workgroup with privacy, data, and ethics experts



- **More concise information and consent forms**
- **Understandable for a wider audience**
- **Part of the privacy information to website**
- **Basis for all faculties**

# What is new for IC?



- Create a study-specific information and consent form for each study
- Based on a template which is available via Ethics Committee Social Sciences (ECSS) website



- All collected data presented in a table



- Part of the information - RU website
- Approval participant



- Lab studies: name and signature
- Online studies: press / *agree* + time stamp



- Online studies:
  - Also here study-specific forms
  - Templates for Limesurvey and JSPsych under construction, available before end of year
  - Consent no personal data, can be collected together with research data
  - After acquisition - separate consent answers from research data

**NEW**

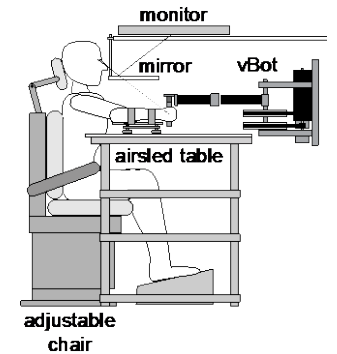
# What is new for method forms?

All available via [ECSS website](#)

Simplified texts

Method forms

- EEG, Sled, robot arm
- Still to be finalized TUS, MRI



# What is new for screening forms?

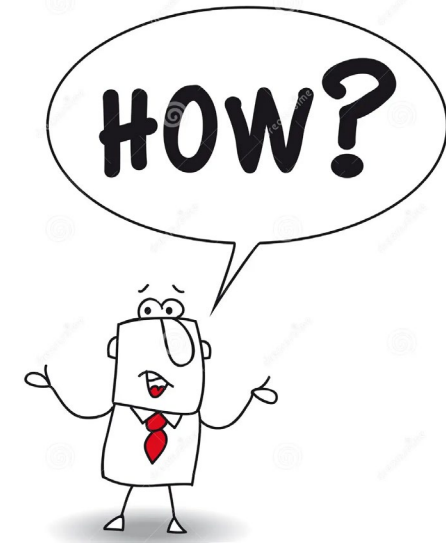


- All available via [ECSS website](#)
- Simplified texts
- Not always, but only screening form when important for data quality or safety
  - EEG, sled
  - still to be finalized: TUS, TMS, TES, MRI
- No screening on age anymore
- Changed protocol Adverse Events
  - In case of Adverse Event: Send mail to [dcc-researchintegrity-officer@donders.ru.nl](mailto:dcc-researchintegrity-officer@donders.ru.nl), within 24 hours
  - Note on [DCC Research Admin Statement](#) whether there have been any Adverse Events



# How does it work?

- Download **ECSS informed consent template** and the **template table personal data (docx, 64 kB)**
- Generate your study specific consent forms by following the instructions within the template documents

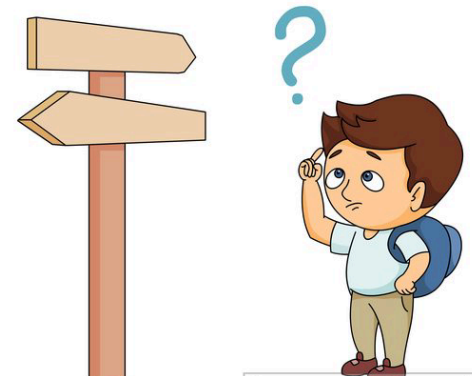


# Where can I find the new templates?

*ECSS informed consent template and template table data* [English](#) or [Dutch](#)

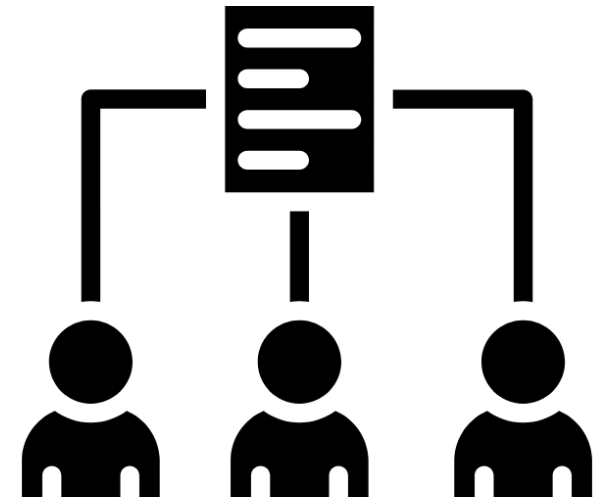
Website additional information privacy: [English](#) / [Dutch](#)

[Updated DCC intranet](#)



# How to share info with SONA participants?

- Must: provide the information >24 hours before experiment
- Within SONA Participant system no attachments
- Put information document(s) on [Surfdrive](#) and create public link.
- Check out [step-by-step manual](#) on how to create a public link.
- Share this public link with your participant via the Direct Message option of SONA.





# Who should start using new forms, and when?

- All research that has been or is going to be approved by Ethical Committee Social Sciences
- For ongoing research transition period. You can choose continue to work with current forms. However, when starting a new study you should use the new forms
- In case of doubt/questions, contact Miriam
- Btw, for WMO research new 'PIF', one with DCC instructions will be put on intranet before end of 2025.



# Feedback DCC researchers

*Although the new forms require a bit more study specific information, I found it useful to write the information up and believe the experiment and procedure will be clearer to participants with these forms.*

*A suggestion would be to make some sample forms available. This would make the "administrative hurdle" even less.*

*I'm very happy with the new forms. Adapting them to our experiment took us a couple of hours at most. My information brochure went from ~10 pages to 2 pages. This is much clearer for both the participant and the researchers!*



# QUESTIONS?