

# Alternative cast for distal radius fractures

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## 0. Administrative questions

### 1. Name of data management support staff consulted during the preparation of this plan.

Question not answered.

### 2. Date of consultation with support staff.

## I. Data description and collection or re-use of existing data

### 3. Provide a general description of the type of data you will be working with, including any re-used data:

Type of data	File format(s)	How will data be collected (for re-used data: source and terms of use)?	Purpose of processing	Storage location	Who will have access to the data
Anonymised data on age, gender and measures of the upper limb	.xlsx	Questionnaire	To understand relations between these data and comfort or fitting of the device	Project drive	The researcher and two supervisors
Subjective data related to comfort and experience	.xlsx	Questionnaire (and conversations during the testing)	The comfort should be evaluated, and the subjective questions with a likert scale can quantify this.	Project drive	The researcher and two supervisors
Quantitative data on the fitting of the device	.xlsx and .jpg	Measurements and photos	There will be evaluated if people fit in the device. Therefore there is recorded what settings the device is on	Project drive	The researcher and two supervisors
Subjective data on ease of application	.xlsx	Questionnaire (and conversations during the testing)	To get insights in ease of application useful for implementation in a hospital setting	Project drive	The researcher and two supervisors

### 4. How much data storage will you require during the project lifetime?

- < 250 GB

## II. Documentation and data quality

### 5. What documentation will accompany data?

- Methodology of data collection

### III. Storage and backup during research process

6. Where will the data (and code, if applicable) be stored and backed-up during the project lifetime?

- OneDrive

### IV. Legal and ethical requirements, codes of conduct

7. Does your research involve human subjects or 3rd party datasets collected from human participants?

- Yes

8A. Will you work with personal data? (information about an identified or identifiable natural person)

*If you are not sure which option to select, first ask your [Faculty Data Steward](#) for advice. You can also check with the [privacy website](#) . If you would like to contact the privacy team: [privacy-tud@tudelft.nl](mailto:privacy-tud@tudelft.nl), please bring your DMP.*

- Yes

Yes personal data will be collected but participants are not identifiable

8B. Will you work with any other types of confidential or classified data or code as listed below? (tick all that apply)

*If you are not sure which option to select, ask your [Faculty Data Steward](#) for advice.*

- No, I will not work with any confidential or classified data/code

9. How will ownership of the data and intellectual property rights to the data be managed?

*For projects involving commercially-sensitive research or research involving third parties, seek advice of your [Faculty Contract Manager](#) when answering this question. If this is not the case, you can use the example below.*

The datasets underlying the published papers will be publicly released following the TU Delft Research Data Framework Policy. During the active phase of research, the project leader from TU Delft will oversee the access rights to data (and other outputs), as well as any requests for access from external parties. They will be released publicly no later than at the time of publication of corresponding thesis.

10. Which personal data will you process? Tick all that apply

- Signed consent forms
- Photographs, video materials, performance appraisals or student results
- Gender, date of birth and/or age

11. Please list the categories of data subjects

Students

And possibly:  
Employees TU Delft  
Family

**12. Will you be sharing personal data with individuals/organisations outside of the EEA (European Economic Area)?**

- No

**15. What is the legal ground for personal data processing?**

- Informed consent

**16. Please describe the informed consent procedure you will follow:**

All study participants will be asked for their written consent for taking part in the study and for data processing before the start of test.

**17. Where will you store the signed consent forms?**

- Other - please explain below
- Same storage solutions as explained in question 6

The forms will be uploaded to onedrive and a paper copy will be stored and later destroyed.

**18. Does the processing of the personal data result in a high risk to the data subjects?**

If the processing of the personal data results in a high risk to the data subjects, it is required to perform [Data Protection Impact Assessment \(DPIA\)](#). In order to determine if there is a high risk for the data subjects, please check if any of the options below that are applicable to the processing of the personal data during your research (check all that apply).

If two or more of the options listed below apply, you will have to [complete the DPIA](#). Please get in touch with the privacy team: [privacy-tud@tudelft.nl](mailto:privacy-tud@tudelft.nl) to receive support with DPIA.

If only one of the options listed below applies, your project might need a DPIA. Please get in touch with the privacy team: [privacy-tud@tudelft.nl](mailto:privacy-tud@tudelft.nl) to get advice as to whether DPIA is necessary.

If you have any additional comments, please add them in the box below.

- None of the above applies

**22. What will happen with personal research data after the end of the research project?**

- Anonymised or aggregated data will be shared with others

**25. Will your study participants be asked for their consent for data sharing?**

- Yes, in consent form - please explain below what you will do with data from participants who did not consent to data sharing

## V. Data sharing and long-term preservation

### 27. Apart from personal data mentioned in question 22, will any other data be publicly shared?

- All other non-personal data (and code) produced in the project
- All other non-personal data (and code) underlying published articles / reports / theses

### 29. How will you share research data (and code), including the one mentioned in question 22?

- All anonymised or aggregated data, and/or all other non-personal data will be uploaded to 4TU.ResearchData with public access

### 30. How much of your data will be shared in a research data repository?

- < 100 GB

### 31. When will the data (or code) be shared?

- As soon as corresponding results (papers, theses, reports) are published

### 32. Under what licence will be the data/code released?

- CC0

## VI. Data management responsibilities and resources

### 33. Is TU Delft the lead institution for this project?

- Yes, the only institution involved

### 34. If you leave TU Delft (or are unavailable), who is going to be responsible for the data resulting from this project?

Karin Thomassen, PhD candidate, k.e.thomassen@tudelft.nl

Gerwin Smit, Assistant Professor biomechanical-engineering, g.smit@tudelft.nl

### 35. What resources (for example financial and time) will be dedicated to data management and ensuring that data will be FAIR (Findable, Accessible, Interoperable, Re-usable)?

Question not answered.