**Daily clinical screening**All patients fulfilling all inclusion criteria must be screened in the eCRF.

**Informed consent**It is legal to include inhabited participants in clinical trials if surrogate consentis obtained (Komitéloven §4).All patients intensive care patients is considered temporarily incompetent.

**Procedure**

1. Obtain informed consent from the first trial guardian before randomisation (oral consent by telephone is permitted).
2. Note date, time and name of first trial guardian in electronic medical journal.
3. Obtain written consent from the first trial guardian as soon as possible.

**Screen and include**

1. Log into the electronic medical journal
2. Screening of inclusion and exclusion criteria for new eligible patients

**Randomisation**When informed consent is obtained from first trial guardian AND the patient fulfil all inclusion criteria and none of the exclusion criteria THEN randomise the patient in the web based eCRF