**Daily clinical screening**Clinicians identify all new eligible patients who fulfill all inclusion criteria. All patients fulfilling all inclusion criteria must be screened in the eCRF.

**Informed consent**Informed consent is obtaining according to national regulations. In Denmark, it is legal to include inhabited participants in clinical trials if surrogate consentis obtained (Komitéloven §4). All patients with COVID-19 and severe hypoxia are temporarily incompetent.

***Procedure for obtaining informed consent in Denmark***

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 by post or e-mail as soon as possible.

**Screen**

1. Log into the electronic medical journal
2. Screen inclusion and exclusion criteria for new eligible patients
3. Log into the web-based eCRF. All patients fulfilling all inclusion criteria must be screened in the web-based eCRF.

**Randomise**When informed consent is obtained from first trial guardian AND the patient fulfill all inclusion criteria and none of the exclusion criteria, THEN randomise the patient in the web-based eCRF.