



PRoVENT-PED study

PRactice of VENTilation in Critically Ill PEdiatric Patients

Steering committee

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Objectives

- to describe practice of ventilatory support worldwide
- to identify potentially modifiable factors associated with outcome



Hypotheses

- practice of ventilatory support varies substantially
- potentially modifiable factors are associated with outcome



Endpoints

- key ventilator settings and ventilation parameters (primary)
- incidence of PARDS
- duration of ventilatory support
- LOS in ICU
- mortality rates



Methods

- data collection during 2 periods of 4 weeks each year
- an (optional) extra period in case of an epidemic or pandemic



Patients

Eligible:

- aged 0-18 years
- admitted to a participating PICU
- receiving ventilatory support > 12 hours

Exclusion:

- premature patients



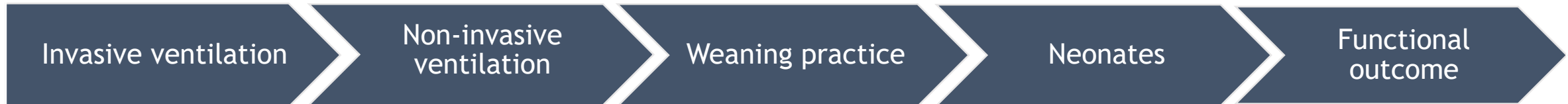
Data collection

- patient characteristics
- ventilator settings and parameters (at day 0-3)
- adjunctive respiratory therapies
- outcomes at ICU discharge (max 28 days)



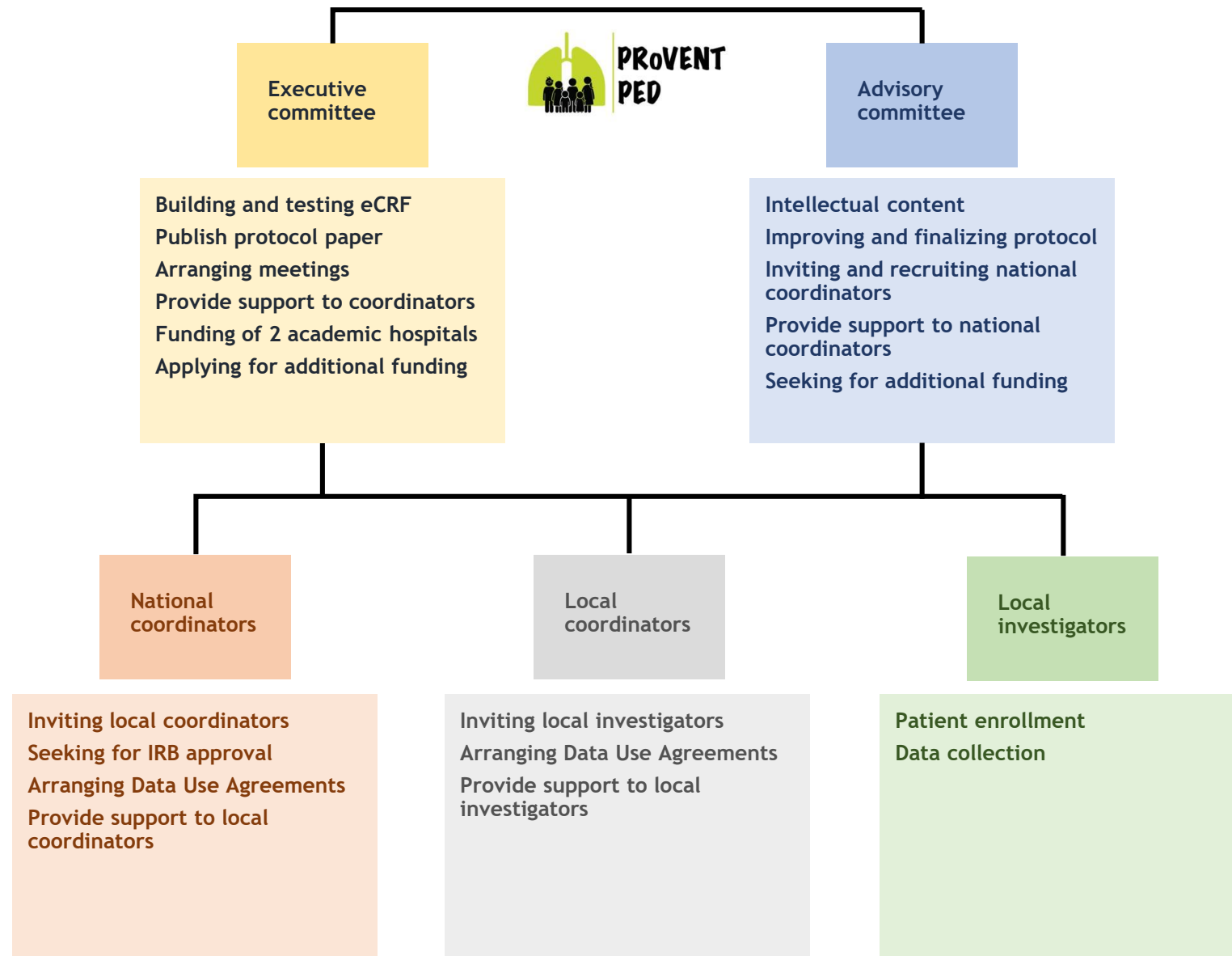
‘Adaptive’ study

- 10 years
- different focuses:

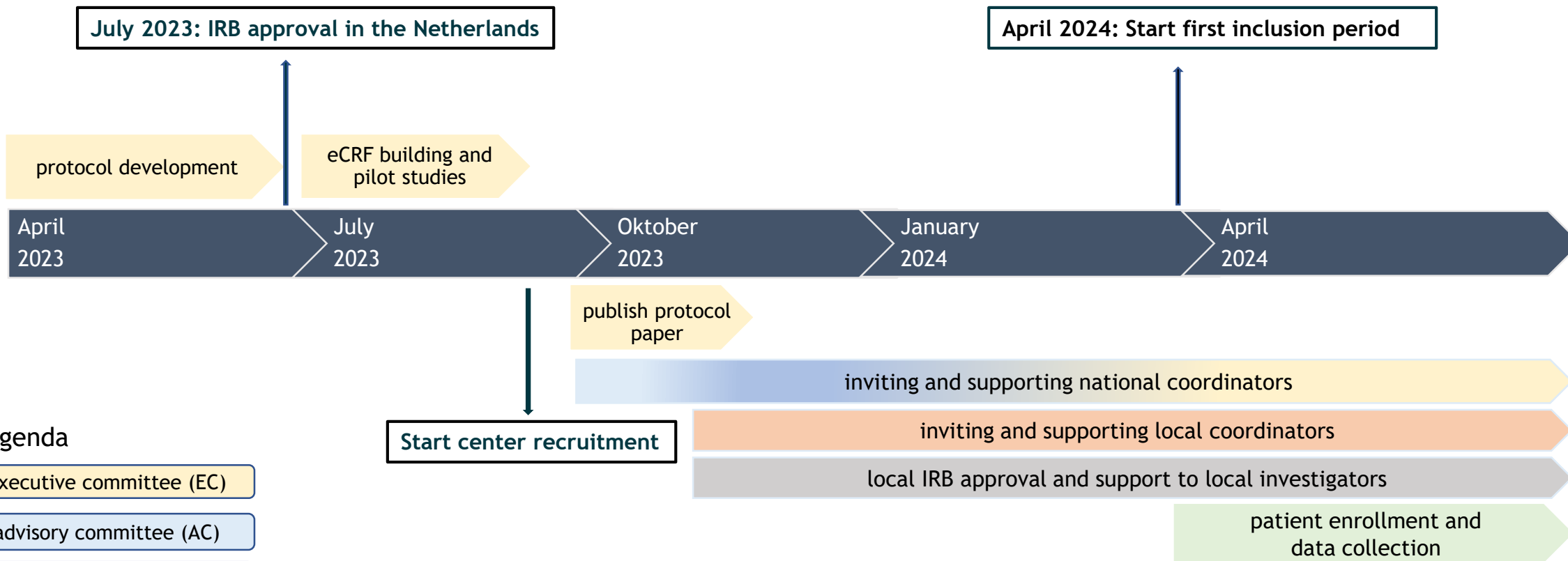


- centers have the option to opt out during certain time periods to make the study manageable for every participating country/center

Team



Timeline



Legenda

- executive committee (EC)
- advisory committee (AC)
- national coordinators
- local coordinators
- local investigators



Website

Please find the latest updates on the progress of the study and all documents on the PRoVENT-PED website.

[PRoVENT-PED | ICU Research](#)



Share ALL rewards

- all collaborators will participate with publication, also on every substudy
- participating centers are welcome to do additional studies on the data set
- the steering committee will coordinate these additional research questions and analysis



Please contact us for more information

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