The EMBARC registry now holds comprehensive clinical data for >20,000 individuals with bronchiectasis from 300 research sites across 27 EU and non-EU countries, including baseline data and data obtained from annual follow-up.

Clinical data collected within the EMBARC registry:

Demographics (age, sex, ethnicity, height, weight, Body Mass Index (BMI) etc)

Comorbidities

Respiratory and non-respiratory medications and treatment regimens including chest physiotherapy and pulmonary rehabilitation

Vaccinations

Bronchiectasis aetiologies and aetiological testing data

Microbiology (at stability and exacerbation)

Radiology

Spirometry

Breathlessness scores

QoL measurements

Disease history (e.g., exacerbations and hospitalisation in previous year)

Disease severity indices, including Bronchiectasis Severity Index (BSI) and FACED

Follow up data (e.g., exacerbations, hospitalisations, mortality)

EMBARC

The European Bronchiectasis Registry