L-citrulline to Improve Adverse Outcomes in Admitted Children (EChiLiBRiST, Clinical Trial 2, Inpatients)

NCT06426147

Status	RECRUITING
Phase	Not Applicable
Sponsor	Barcelona Institute for Global Health
Enrollment	2,200 participants

Key Eligibility Criteria

Inclusion (8)

- Enrolled in the initial prognostic screening component.
- Sick children with fever (axillary temperature\>37.5ºC) or a history of fever (within the preceding 72h) or with suspected severe disease.
- m-\<60 months of age.
- With an indication for admission, or having already been admitted to hospital due to their illness.
- With an sTREM-1 PoC result classifying their disease as of "moderate-high risk" ("yellow" or "red") upon study recruitment and within D3.

... and 3 more (see full listing online)

Exclusion (8)

- Admission to hospital for social reasons (and not on account of their disease).
- Children for which informed consent document has not been signed.
- Known allergy or contraindication to any of the study supplements including lactose intolerance or observing a lactose-free diet.
- Concurrent participation in any other clinical trial.
- Patient under NPO or "nothing by mouth" prescription .

... and 3 more (see full listing online)

Locations (2 total)

Hararghe Health Research, Harar, Ethiopia
Hospital Central de Maputo, Maputo, Mozambique