

Immunogenicity and Safety PCV-20 of the Vaccine Administered During an Acute Febrile Illness in Adults

NCT06822907

Status	RECRUITING
Phase	Phase 4
Sponsor	Centre Hospitalier Universitaire de Saint Etienne
Enrollment	1,052 participants

Key Eligibility Criteria

Inclusion (7)

- History of body temperature $\geq 38^{\circ}\text{C}$ measured at least twice prior to randomization (Randomization must be performed as soon as possible on a febrile patient or 72 hours after apyrexia at the latest)
 - Having at least one comorbidity that defines patients as medium or high risk for pneumococcal invasive infection:
 - Medium risk: Cyanogenic congenital heart disease; chronic heart failure; chronic respiratory failure; chronic obstructive pulmonary disease; emphysema; severe asthma under chronic treatment; chronic renal failure; chronic liver disease; diabetes mellitus treated; Osteo-meningeal leak or cochlear implant; Age ≥ 65 years old.
 - High risk : Hypo or asplenic people; hereditary immunodeficiency syndromes; people living with HIV; solid organ transplanted; People under immunosuppressors (corticosteroids, biotherapy) for an auto-immune or an inflammatory chronic disease; patients with nephrotic syndrome
 - Hospitalization for ≥ 24 hours long
- ... and 2 more (see full listing online)

Exclusion (18)

- Patient unable to give informed consent
 - Curators, wardship
 - History of previous vaccination with PCV-7 or PCV-13 or PCV-20
 - History of PPV-23 in the previous year
 - Patient having received another vaccination within one month prior to inclusion or planning another vaccination in the month after inclusion except for Influenza vaccine.
- ... and 13 more (see full listing online)

Locations (24 total)

CHU de Saint-Etienne, Saint-Etienne, France, France
Centre Hospitalier, Annecy, France
Centre Hospitalier Universitaire, Besançon, France
... and 21 more locations

<https://clinicaltrials.gov/study/NCT06822907>

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