Study to Evaluate the Maximal Use of Ruxolitinib Cream in Adult and Adolescent Participants With Hidradenitis Suppurativa

NCT07049575

Status	RECRUITING
Phase	Phase 1
Sponsor	Incyte Corporation
Enrollment	24 participants

Key Eligibility Criteria

Inclusion (5)

- Diagnosis of HS for at least 6 months before screening visit.
- Diagnosis of HS (Hurley Stage I, II, III) with a total abscess and inflammatory nodule count of at least 4 and affecting at least 3 distinct anatomical areas at screening and Day 1 visits.
- Total estimated treatment BSA \> 20% at screening and baseline.
- Agreement to not use topical or systemic antibiotics for treatment of HS during the Maximal Use Treatment Period.
- Willingness to avoid pregnancy or fathering children based on the criteria defined in the protocol.

Exclusion (2)

- Current or history of skin condition(s) other than HS that might confound the evaluation of HS; clinically uncontrolled cardiovascular disease; thrombosis; certain cancers; certain infections; severe anemia, thrombocytopenia, or neutropenia; other medical conditions at the discretion of the investigator.
- Laboratory values outside of the protocol-defined ranges.

Locations (23 total)

Saguaro Dermatology, Phoenix, Arizona, United States
First Oc Dermatology, Fountain Valley, California, United States
Amicis Research Center Valencia, Northridge, California, United States
... and 20 more locations