ForPatients

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Atopic Dermatitis

A Long Term Extension Study to Evaluate the Safety and Efficacy of Afimkibart (RO7790121) in Participants With Atopic Dermatitis

Trial Status Trial Runs In Trial Identifier
Not yet recruiting NCT07223697 2025-520771-59-00
CS45943

The information is taken directly from public registry websites such as ClinicalTrials.gov, EuClinicalTrials.eu, ISRCTN.com, etc., and has not been edited.

Official Title:

An Extension Study To Evaluate the Long-Term Safety and Efficacy of Afimkibart (RO7790121) in Patients With Atopic Dermatitis Who Participated in Previous Afimkibart Clinical Trials

Trial Summary:

This study will assess long-term safety and efficacy of Afimkibart (also known as RO7790121) in participants with Atopic Dermatitis (AD) who participated in previous afimkibart clinical trials.

Hoffmann-La Roche Sponsor		Phase 2 Phase	
NCT07223697 2025-5	CT07223697 2025-520771-59-00 CS45943 al Identifiers		
Eligibility Crite	ria:		
Gender All	Age #18 Years	Healthy Volunteers No	

Inclusion Criteria:

Ability and willingness to comply with all aspects of the protocol including completion of the efficacy
assessments (EASI and IGA), clinical outcome assessment instruments (DLQI, POEM), and safety and
PK sample collections for the duration of the study

Parent Clinical Trial-Specific Criteria:

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• Study CS45570 participants who continued to be evaluated at the Week 36 follow up visit and achieved => EASI50 response from study baseline

Exclusion Criteria:

- · Evidence of other skin conditions that would interfere with the assessment of AD
- Withdrawal of consent and/or premature discontinuation from parent study
- Any permanent discontinuation of study drug in parent study
- History of severe allergic reaction or anaphylactic reaction to any biologic agent or known hypersensitivity to any component of Afimkibart