



Quick Facts about proDERM Pharma

To assure **Skin Tolerance**. To evaluate **Proof of Concept**. To document **Efficacy**.

Be it a dermatological or ophthalmological Medicinal Product, a Medical Device or a Transdermal Device; proDERM Pharma accomplishes your clinical studies according to ICH-GCP.

Study Protocols

- Local Tolerance

- 21 Day Cumulative Irritation Patch Test
- Human Repeat Insult Patch Test (HRIPT)
- Repeated Open Application Test (ROAT)
- Phototoxicity
- Photosensitization
- Scratch Patch Test
- Local Eye Tolerance

- Proof of Concept

- Psoriasis Plaque Test
- Vasoconstriction test / Blanching
- Suction Blister Model
- UV Erythema Model
- Acne
- Rosacea
- Actinic Keratosis
- Athlete's foot

proSERVICE

proSERVICE represents the Full-Service approach of proDERM and comprises all those services we offer combined with the clinical study:

- Consultancy
- Regulatory Affairs
- Recruitment
- Data Management
- Statistics
- Medical Writing
- Archiving

CRO Services:

In addition to Phase I/IIa trials, proDERM offers to be the responsible CRO for coordinating multicenter trials and is also available as an experienced study site. Our network of qualified study sites ensures a fast recruitment.

Indications

- Acne
- Atopic Dermatitis
- Psoriasis
- Allergy Type I & IV
- Rosacea
- Dry Eyes
- Inflammation / Wounds
- Hyperhidrosis
- Ophthalmology
- Herpes Simplex

Experience

- appr. 3,000 enrolled subjects since 2001
- Pilot studies, Proof of Concept, Phase I, Phase II / IIb

Recruitment

- Healthy volunteers
- Panels:
 - Acne
 - Atopic Dermatitis
 - Psoriasis
 - Rosacea

Equipment

- Tewameter
- Chromameter
- Corneometer
- Laser Doppler Blood Flow
- USR-CliP: Unit for Standardized and Reproducible Clinical Photography
- IR Thermography
- UVA/UVB Irradiation Sources
- Confocal Microscopy
- Ultrasound

Principal Investigator



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