



A service of the National Library of Medicine
and the National Institutes of Health

My NCBI [?](#)
[\[Sign In\]](#) [\[Register\]](#)

All Databases

PubMed

Nucleotide

Protein

Genome

Structure

OMIM

PMC

Journals

Books

Search

PubMed



for

Go

Clear

Limits

Preview/Index

History

Clipboard

Details

[About Entrez](#)Display | [Abstract](#)

▼ Show | 20

▼ Sort by

▼ Send to

Text Version

All: 1 Review: 0 **Entrez PubMed**[Overview](#)
[Help | FAQ](#)
[Tutorial](#)
[New/Noteworthy](#)
[E-Utilities](#)**PubMed Services**[Journals Database](#)
[MeSH Database](#)
[Single Citation Matcher](#)
[Batch Citation Matcher](#)
[Clinical Queries](#)
[Special Queries](#)
[LinkOut](#)
[My NCBI](#)**Related Resources**[Order Documents](#)
[NLM Mobile](#)
[NLM Catalog](#)
[NLM Gateway](#)
[TOXNET](#)
[Consumer Health](#)
[Clinical Alerts](#)
[ClinicalTrials.gov](#)
[PubMed Central](#) 1: [Skin Pharmacol Appl Skin Physiol.](#) 2002;15 Suppl 1:31-9.[Related Articles, Links](#)**Improvement of the experimental setup to assess cutaneous bioavailability on human skin models: dynamic protocol.**[Dreher F](#), [Patouillet C](#), [Fouchard F](#), [Zanini M](#), [Messenger A](#), [Roguet R](#), [Cottin M](#), [Leclaire J](#), [Benech-Kieffer F](#).

L'Oreal Research, Aulnay-sous-Bois, France. fdreher@recherche.loreal.com

Human skin models, such as EpiDerm and Episkin, are not easily mounted into static or dynamic diffusion cells that are commonly used to perform bioavailability studies with human skin ex vivo. For various reasons, such as fragility, small sample size, and other morphological constraints, skin absorption studies with human skin models are often carried out on the delimited skin surface obtained by gluing a ring onto the reconstituted epidermis and manually exchanging the receptor solution. However, such an experimental setup is prone to artifacts. Discontinuous removal of the receptor fluid leads to alternating sink conditions, and an area of application smaller than the area in contact with the receptor fluid, as well as imperfect seal of the glued ring, may result in inaccurate penetration rates. Human skin models were shown to be relatively easily mounted into In-Line cells (PermeGear Inc.), vertical diffusion cells which appear to be appropriately designed for such a purpose. In-Line cells allowed accurate determination of solute penetration as well as automated sampling of receptor fluid. Excised human skin can be mounted into these cells as well, making it possible to compare penetration rates through different types of skin samples under identical conditions. Using mannitol as a reference compound, penetration profiles and epidermal distribution similar to those obtained with human skin ex vivo were obtained both with EpiDerm and Episkin. Under the present conditions, human skin models were more permeable to mannitol than excised human skin, which was only slightly permeable to mannitol. Due to these experimental innovations and to the good agreement with the absorption characteristics through human skin ex vivo, EpiDerm and Episkin seem to be promising human skin models for testing the cutaneous bioavailability of topical products in vitro. Copyright 2002 S. Karger AG, Basel

PMID: 12476007 [PubMed - indexed for MEDLINE]

Display | [Abstract](#)

▼ Show | 20

▼ Sort by

▼ Send to

[Write to the Help Desk](#)[NCBI](#) | [NLM](#) | [NIH](#)[Department of Health & Human Services](#)[Privacy Statement](#) | [Freedom of Information Act](#) | [Disclaimer](#)

Nov 29 2005 11:19:28