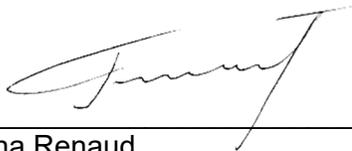


CE DECLARATION OF CONFORMITY

We, SmartHealth, Inc. (dba SmartPractice) of 3400 East McDowell Road, Phoenix, Arizona, USA declare under our sole responsibility that the following Class I Finn Chamber Products for skin testing described below are in conformity with the applicable provisions of Medical Devices Directive 93/42/EEC of 14 June 1993 concerning the medical devices and its Annex VII as amended as per 2007/47/EU.

- Finn Chambers on ScanPor packaged as (patch test strip/box, chamber size)
25 (5x2), 8 mm; 100 (5x2), 8 mm; 50 (5x1), 8 mm; 200 (5x1), 8 mm; 150 (1x1), 8 mm;
200 (1x1), 12 mm (Large); 100 (1x1), 18 mm (Extra Large)
- Finn Chambers OPEN packaged as (patch test strip/box, chamber size)
50 (5x2), 8 mm
- Finn Chambers AQUA packaged as (patch test strip/box, chamber size)
100 (5x2), 8 mm
- Finn Chambers, loose, packaged as (package size, chamber size)
1000 chambers, 8 mm; 200 chambers, 12 mm (Large); 100 chambers, 18 mm (Extra Large)
- Filter Paper Discs packaged as (package size, disc size)
1000 discs, 8 mm; 200 discs, 8 mm; 200 discs, 12 mm (Large); 100 discs, 18 mm (Extra Large)
- Finn Chamber ATOPY packaged as (patch test strip/box, chamber size)
50 (5x1), 12 mm

Phoenix, Arizona



Elena Renaud
Director of Regulatory Affairs
SmartHealth, Inc.
3400 East McDowell road
Phoenix, AZ, 85008 USA
Phone: +1 (602) 225-0595 x 7290
e-mail: erenaud@smarthealth.com

Date: May 7, 2020

Expiration date: May 25, 2021

Note: Our Authorized Representation within EU is Dr. Ebeling & Assoc. GmbH Isestrasse 5 · 20144 Hamburg Germany