Certificate BE18/819943012.01

The management system of

RedPharma Productie B.V.

Draaibrugweg 29-39
1332 AB Almere, The Netherlands

has been assessed and certified as meeting the requirements of

ISO 13485:2016
EN ISO 13485:2016

For the following activities

Contract manufacture of non-sterile liquid and semi-solid medical devices for dermatological, nasal, ear, nail, oral, intimate zones, foot, anal and gastro-intestinal use.
Design & Development related to validation and transfer of production processes for non-sterile liquid and semi-solid medical devices for dermatological, nasal, ear, nail, oral, intimate zones, foot, anal and gastro-intestinal use.
Packaging (including primary packaging) and warehousing of non-sterile liquid and semi-solid medical devices for dermatological, nasal, ear, nail, oral, intimate zones, foot, anal and gastro-intestinal use.
Commissioning, qualification, validation (including verification), installation and maintenance of production equipment and production related installations for RedPharma Productie B.V. and Kolibrie-NL B.V.

This certificate is valid from 4 July 2021 until 3 July 2024
and remains valid subject to satisfactory surveillance audits.
Issue 4. Certified since 4 July 2018.
Re certification audit due before 13 March 2024.
Multiple certificates have been issued for this scope.
The main certificate is numbered BE18/819943012.00.

Authorised by

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Accreditation Number
005-QMS
EN ISO/IEC 17021-1:2015

SGS Belgium 13485-2 0308