

MHRA

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United Kingdom

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DR. P Punia
PHARMEXON CONSULTING SRO
PITTEROVA 2855/7
PRAGUE
CZ-130 00
CZECH REPUBLIC

05/04/2018

Dear DR. Punia,

GRANT / RENEWAL OF MARKETING AUTHORISATION

Our Reference: PL 04465/0017 - 0001
Your Reference: 4465
Product: Ghemaxan 8,000 IU (80 mg)/0.8 mL solution for injection in pre-filled syringes
Type of Procedure: Decentralised
Submission Type: Initial
Submission Category: Abridged
EU Procedure Number (if applicable): UK/H/5798/006/DC

The Licensing Authority agrees to the grant or renewal of the marketing authorisation for the above submission on the basis of the data provided. This includes any replacement and amendment of the original dossier.

In line with Article 23a of Directive 2001/83/EC as amended, the Marketing Authorisation Holder should submit notification of the actual date of marketing of the product to the Competent Authority.

This notification should be provided by email to the following address: sunsetclause@mhra.gov.uk.

The formal documents are enclosed. These constitute evidence of authorisation. If you consider them to contain information that is incorrect or not in accordance with the dossier, please return immediately indicating any errors.

All Marketing Authorisations are subject to standard provisions contained in current medicines regulations full details of which are published on the MHRA website:

<http://www.mhra.gov.uk/Howweregulate/Medicines/Licensingofmedicines/Informationforlicenceapplicants/Provisionstowhichthemarketingauthorisationisgranted/index.htm>

Yours sincerely,

Ahsana Israel

The Human Medicines Regulations SI 2012/1916, as amended.

GRANT / RENEWAL OF MARKETING AUTHORISATION

Product: Ghemaxan 8,000 IU (80 mg)/0.8 mL solution for injection in pre-filled syringes
Submission Type: Initial

Granted to: CHEMI SPA
VIA DEI LAVORATORI 54
CINISELLO BALSAMO (MI)
I-20092
ITALY

This Marketing Authorisation, under the above reference number is hereby granted / renewed in respect of the product named above. The Summary of Product Characteristics of the product is set out in the attached document.

The application is subject to the further provisions set out or referred to in the above Regulations.

This Marketing Authorisation, as now granted / renewed, unless previously revoked, will continue in force until the expiry date (if applicable) given below.

Grant Date: 05/04/2018

Date of Expiry: 07/03/2023

Ahsana Israel

A person authorised to sign on behalf of the Secretary of State for Health