REPUBLIC OF MACEDONIA
AGENCY FOR MEDICINES AND MEDICAL DEVICES

MALMED
MACEDONIAN AGENCY FOR MEDICINES AND MEDICAL DEVICES

Certificate No: 18-4276/3

CERTIFICATE OF GMP COMPLIANCE OF A MANUFACTURER

Part 1

Issued following an inspection in accordance with Art. 111(5) of Directive 2001/83/EC

The competent authority AGENCY FOR MEDICINES AND MEDICAL DEVICES OF THE REPUBLIC OF MACEDONIA confirms the following:

The manufacturer **Pharmaceutical Industry INTER- EVROGENEKS DOOEL**

Site address: Str.Goce Delcev No.12, 2434 Novo Selo, R.of Macedonia

Has been inspected under the national inspection programme in connection with manufacturing authorization and in accordance with Art.40 of Directive 2001/83/ECtransposed in the national legislation Law on medicines and medical devices (Official gazette of the RM No. 106/07,88/2010, 36/11, 53/11, 136/11, 11/12, 147/13, 27/14, 43/14, 88/15, 154/15, 228/15, 7/16, 53/16).

From the knowledge gainedduring inspection of this manufacturer, the latest of which was conducted on 10.05.2016 it is considered thatit complies with:

- The principles and guidelines of Good Manufacturing Practice requirements laid down in Directive 2003/94/EC.

This certificate reflects the status of the manufacturing site at the time of the inspection noted above and should not be relied upon to reflect the compliance status if more than three years have elapsed since the date of that inspection, after which time the issuing authority should be consulted. This certificate is valid only when presented with all pages and both Parts 1 and 2. The authenticity of this certificate may be verified with the issuing authority.

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Part 2

HUMAN MEDICINAL PRODUCTS

1. Manufacturing operations

1.1	***Non-sterile products***
1.1.1	Manufacturing and packing of **Herbal products and tradicional herbal products** in solid and liquid dosage forms

Agency for medicines and medical devices
Director,
Mr.ph.spec. Marija Darkovska –Serafimovska

Certificate No: 18-4276/3
Date: 11 -05- 2016
Skopje