**FREE SALE AND GMP CERTIFICATE**

**POTRDILO O PRODAJI IN PROIZVODNJI**

**V SKLADU Z DOBRO PROIZVODNO**

**PRAKSO (DPP)**

No. of certificate: \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

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| **1. Ime izdelka:** | |  | | **1. Product name:** | |
| Kliknite tukaj, če želite vnesti besedilo. | |  | | Kliknite tukaj, če želite vnesti besedilo. | |
| **Farmacevtska oblika** | |  | | **Pharmaceutical form** | |
| Kliknite tukaj, če želite vnesti besedilo. | |  | | Kliknite tukaj, če želite vnesti besedilo. | |
| **Pakiranje** | |  | | **Supply** | |
| Kliknite tukaj, če želite vnesti besedilo. | |  | | Kliknite tukaj, če želite vnesti besedilo. | |
| **2. Ime in količina aktivnih sestavin:**  Kliknite tukaj, če želite vnesti besedilo. | |  | | **2. Name and amount of active ingredients:**  Kliknite tukaj, če želite vnesti besedilo. | |
| **3. Številka in datum izdaje odločbe / dovoljenja za promet:** | |  | | **3. Number of marketing authorisation and date of issue:** | |
| Kliknite tukaj, če želite vnesti besedilo. | |  | | Kliknite tukaj, če želite vnesti besedilo. | |
| **4. Proizvajalec:** | |  | | **4. Manufacturer:** | |
| Kliknite tukaj, če želite vnesti besedilo. | |  | | Kliknite tukaj, če želite vnesti besedilo. | |
| **5. Naslov:** | |  | | **5. Address:** | |
| Kliknite tukaj, če želite vnesti besedilo. | |  | | Kliknite tukaj, če želite vnesti besedilo. | |
| **6. Potrjujemo:**  a/ Da je proizvod odobren za uporabo v Izberite element. medicini in prodajo v deželi izvora. | |  | | **6. It is certified that:**  a/ This product has been approved for Izberite element. treatment only and sale in the country of origin. | |
| b/ je tovarna, v kateri se izdelek proizvaja, podvržena inšpekcijam v primernih razdobjih.  c/ proizvajalec ustreza zahtevam za ustrezno proizvodnjo in kontrolo kvalitete, kot priporoča Svetovna zdravstvena organizacija, kar zadeva izdelke, ki se prodajajo ali distribuirajo v deželi izvora ali se izvažajo. | |  | | b/ the manufacturing plant in which the product is produced is subject to Inspections at suitable intervals and  c/ the manufacturer meets the requirements for good practice in the manufacture and quality control, as recommended by the World Health Organization, in respect of products to be sold or distributed in the country of origin or to be exported. | |

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| Adress of Certifying authority |  | | **Agency for Medicinal Products and Medical Devices of the Republic of Slovenia, Slovenčeva ulica 22, 1000 Ljubljana, Slovenia** |
| Name of authorized person | | **Momir Radulović, MPharm**  **Director** | |

Signature:

Stamp and date: