**Payment application**

|  |  |
| --- | --- |
| **Recipient of service**  (Name of marketing authorisation holder) | |
| Company registered address |  |
| Company/ VAT payer registration No. |  |
| Payment requisites |  |
| Contact person (name, surname, e-mail address) |  |
| **Fee Payer** (to be filled in if different from the recipient of the service) | |
| Company registered address |  |
| Company/ VAT payer registration No. |  |
| Payment requisites |  |
| Contact person (name, surname, e-mail address for invoice forwarding) |  |

|  |  |
| --- | --- |
| Intended date of application submission |  |

|  |  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- | --- |
| Tradename of medicines, strength/concentration, pharmaceutical form | Type of procedure/ role of LV1 | Procedure number or medicines authorisation number | Type of application2 | Legal basis/type of variations3 | Point in the pricelist4 | Amount EUR |
|  |  |  |  |  |  |  |

Explanations:

1 Indicate the type of procedure and Latvia’s role in it:

NP - national procedure;

MRP / CMS - mutual recognition procedure, if Latvia is a Concerned Member State;

DCP / CMS - decentralised procedure, if Latvia is a Concerned Member State;

MRP / RMS - mutual recognition procedure, if Latvia is a Reference Member State;

DCP / RMS - decentralised procedure, if Latvia is a Reference Member State.

If the application for variations (Latvia is a Reference Member State) is submitted, indicate IB/RMS or II/RMS.

2 Indicate the type of application, i.e. the purpose of submission: marketing authorization; renewal; variation.

3 Indicate the legal basis for an application for marketing authorization in accordance with the Directive 2001/83/EC:

Full application (Article 8(3) of Directive No 2001/83/EC),

Generic application (Article 10(1) of Directive No 2001/83/EC),

Hybrid application (Article 10(3) of Directive No 2001/83/EC),

Similar biological application (Article 10(4) of Directive No 2001/83/EC),

Well-established use application (Article 10a of Directive No 2001/83/EC),

Fixed combination application (Article 10b of Directive No 2001/83/EC),

Informed consent application (Article 10c of Directive No 2001/83/EC),

Simplified registration for homeopathic medicinal products (Article 14 of Directive No 2001/83/EC),

Homeopathic medicinal products (Article 16 of Directive No 2001/83/EC),

Traditional use registration for herbal medicinal product application (Article 16a of Directive No 2001/83/EC).

If the application for variations (Latvia is a Reference Member State) is submitted, indicate type of variations – IB or II.

4 Number of the service in the 10.12.2019 Cabinet of Ministers Regulation No 641 “The State Agency of Medicines Publicly Available Paid Service Price List” .