

Permission no.: 561/2021/RHV

## DECISION on change of the permission to produce veterinary medicinal products

Institute for State Control of Veterinary Biologicals and Medicines, based in Brno (hereinafter the "Institute") as a relevant administrative body according to the § 16 paragraph 2 letter a) point 2 of the Act no. 378/2007 Coll., on Pharmaceuticals and on changes of some related acts (hereinafter "Act no. 378/2007 Coll., on Pharmaceuticals"), issues according to the § 63 of the Act no. 378/2007 Coll., on Pharmaceuticals, this

# PERMISSION to produce veterinary medicinal products

in the following range and under the following conditions:

1. Operator:

**DYNTEC spol. s r. o.**  
Company ID: 475 48 002

2. Registered office of the operator:

**Pražská 328  
411 55 Terezín**

3. Permission to produce no. 561/2021/RHV is granted for premises and equipment used by the manufacturer at the address:

**Pražská 328, 411 55 Terezín**

4. Range of permitted activity including activities provided under the contract in the area of manufacturing and quality control of veterinary medicinal products is stated in **Annex no. 1** to this permission. Annex no. 1 is an integral part of this permission.

5. According to the § 64 letter a) of the Act no. 378/2007 Coll., on Pharmaceuticals, the Qualified Person is:

**MVDr. Mgr. Ladislav Pažout**

**Ing. Pavla Žáková**

6. The manufacturer is obliged to follow the relevant provisions of the Act no. 378/2007 Coll., on Pharmaceuticals, Decree no. 229/2008 Coll., on Manufacturing and distribution of Pharmaceuticals (hereinafter "Implementing decree") and instructions issued by the Institute for State Control of Veterinary Biologicals and Medicines.

7. Holder of the permission is obliged, according to the § 64 letter i) of the Act no. 378/2007 Coll., on Pharmaceuticals, to allow the officers of the Institute to perform official supervision according to the § 16 paragraph 2 letter e) of the Act no. 378/2007 Coll., on Pharmaceuticals and is obliged to respect the imposed measures according to the § 16 paragraph 2 letter c) and d) of the Act no. 378/2007 Coll., on Pharmaceuticals.

8. The permission is granted for an indefinite period. It may be suspended or cancelled according to the § 63 paragraph 8 of the Act no. 378/2007 Coll., on Pharmaceuticals. Obligations may be imposed to the holder of the Permission to produce veterinary medicinal products even after the entry into the force of this decision. The original Permission to produce veterinary medicinal products issued by the Institute on 01.10.2018 with reg. no. 510/2018/RHV expires on the effective date of this Decision.

Logo	Hudcova 232/56a 621 00 Brno-Medlánky Czech Republic	+420 541 518 210 Data box: ra7aipu	uskvbl@uskvbl.cz www.uskvbl.cz	Date: 01/10/2021 Signature: <i>signature</i>
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9. The manufacturer is authorised to operate only with the equipment and in the premises approved by the inspection group during the inspection performed on 1.9. – 2.9.2021 and with respect to the conditions set in the Protocol on inspection of GMP ref. no. Ú S K V B L / 13218/2021/INS of 30.09.2021.

**Statement of the reasons:**

Based on the *Decision on change of permission to produce medicinal products* issued by the Institute on 01.10.2018 under reg. no. 510/2018/RHV,

*application for a change* delivered to the State Control of Veterinary Biologicals and Medicines on 25.08.2021, with ref. no. ÚSKVBL/11503/2021/POD, file number ÚSKVBL/11503/2021/POD submitted according to the § 63, paragraph 6 of the Act no. 378/2007 Coll., on Pharmaceuticals, and the implementing decree,

*payment of administrative fee* according to the Act no. 634/2004 Coll., on administrative fees, as amended (item 98),

*payment of the costs of professional actions performed on a request* according to the valid tariff of the State Control of Veterinary Biologicals and Medicines, item I-05 paid on 19.08.2021,

*changes of classification of production activities* published in “Compilation of Community Procedures” in accordance with Directive 2004/28/EC,

*submitted documentation* demonstrating compliance with the principles of good manufacturing practice under the provisions of implementing decree and

*result of the inspection* performed on 1.9. – 2.9.2021, according to the provisions of implementing decree

the Institute found that the applicant fulfilled the requirements for the production of veterinary medicinal products set by the Act no. 378/2007 Coll., on Pharmaceuticals and its implementing regulations and instructions of the Institute, and issued this Permission to produce veterinary medicinal products.

**Advice on appeal:**

It is possible to submit appeal against this decision within the limit of 15 days from the day of its delivery to the State Veterinary Administration of the Czech Republic through the Institute.

In Brno on 01.10.2021

Stamp

Signature  
MVDr. Jiří Bureš  
head of service office

**List of Annexes:**

Annex no. 1: Range of permitted activity (1 page)

Logo

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**Annex no. 1 to the Decision on change of permission to produce veterinary medicinal products no. 561/2021/RHV**

**A) Activities performed by the holder of permission to produce – DYNTEC spol. s r. o.**

<input checked="" type="checkbox"/> <b>Veterinary medicinal products</b>	
<b>Part 1 – Production operations</b>	
<b>1.1</b>	<b>Sterile products</b>
	<i>1.1.1 Prepared aseptically</i>
	1.1.1.1 Bulk liquid pharmaceutical forms
	1.1.1.2 Lyophilised
	1.1.1.4 Low-volume liquid pharmaceutical forms
	<i>1.1.2 Terminally sterilized</i>
	1.1.2.3 Low-volume liquid pharmaceutical forms
	<i>1.1.3 Batch release</i>
<b>1.3</b>	<b>Biological products</b>
	<i>1.3.1 Biological medicinal products</i>
	1.3.1.1 Blood derivatives
	1.3.1.2 Immunological products
	<i>1.3.2 Batch release</i>
	1.3.2.1 Blood derivatives
	1.3.2.2 Immunological products
<b>1.5</b>	<b>Packing</b>
	<i>1.5.2 Secondary packing</i>
<b>1.6</b>	<b>Quality control</b>
	<i>1.6.1 Microbiological – sterility testing</i>
	<i>1.6.2 Microbiological – non-sterile medicinal products</i>
	<i>1.6.3 Chemical/physical</i>
	<i>1.6.4 Biological</i>

Any restriction or clarification regarding the scope of manufacturing operations:

1.3.1.2. Immunological products: production of bacterial, viral, autogenous vaccines and immunosera

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**B) Activities provided under contract**

**ITEST Plus s.r.o., Kladská 1032, Hradec Králové Company ID: 620 61 828**

<b>1.6</b>	<b>Quality control</b>
	<i>1.6.4 Biological</i>

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**BIOPHARM, Research institute of biopharmacy and veterinary medicines  
 a.s., Jilové u Prahy, Pohří – Chotouň 90, Company ID: 463 56 606**

<b>1.6</b>	<b>Quality control</b>
	<i>1.6.3 Chemical/physical</i>

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