



MINISTRY OF AGRICULTURE AND FOOD
BULGARIAN FOOD SAFETY AGENCY

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Certificate No: 52/2015/GMP

CERTIFICATE OF GMP COMPLIANCE OF A MANUFACTURER
Part 1

Issued following an inspection in accordance with Art. 80(5) of Directive 2001/82/EC as amended

The competent authority of Republic of Bulgaria confirms the following:

The manufacturer: „NPP AVIVAC” Ltd.

Site address: **Russia, Leningradskaja province, Lomonosovski district, Gorbunki village**

Has been inspected under the inspection programme of the Bulgarian food safety agency in connection with manufacturing authorisation no. 19 of 05. 01. 2009, issued to "Mintech Co" EOOD company, (situated at the address: Sofia 1434, "Vitoscha", kv. "Simeonovo", 80 str., № 12), in accordance with Art. 44 of Directive 2001/82/EC/, transposed in the national legislation by: Veterinary Act, enforced on 2-nd May, 2006 and promulgated in the SG 87 on 1st November, 2005.

From the knowledge gained during inspection of this manufacturer, the latest of which was conducted on 08 - 10. 06. 2015, it is considered that it complies with the Good Manufacturing Practice requirements¹ referred to in the Agreement of Mutual Recognition between the European Community and [MRA partner]/The principles and guidelines of Good Manufacturing Practice laid down in 51 of Directive 2001/82/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. However, this period of validity may be reduced or extended using regulatory risk management principles by an entry in the Restrictions or Clarifying remarks field.

This certificate is valid only when presented with all pages and both Parts 1 and 2.

The authenticity of this certificate may be verified in EudraGMP. If it does not appear, please contact the issuing authority.

Done in Sofia on: 06.07. 2015

DR DAMYAN ILIEV
EXECUTIVE DIRECTOR
BULGARIAN FOOD SAFETY AGENCY
MINISTRY OF AGRICULTURE AND FOOD
SOFIA, BULGARIA

¹ The certificate referred to in paragraph 111(5) of Directive 2001/83/EC and 80(5) of Directive 2001/82/EC, is also applicable to importers.

² Guidance on the interpretation of this template can be found in the Help menu of EudraGMP database.

³ These requirements fulfil the GMP recommendations of WHO.

Part 2
Veterinary Medicinal Products

1 MANUFACTURING OPERATIONS *

- authorised manufacturing operations include total and partial manufacturing (including various processes of dividing up, packaging or presentation), batch release and certification, storage and distribution of specified dosage forms unless informed to the contrary.
- quality control testing and/or release and batch certification activities without manufacturing operations should be specified under the relevant items:
- if the company is engaged in manufacture of products with special requirements e.g. radiopharmaceuticals or products containing penicillin, sulphonamides, cytotoxics, cephalosporins, substances with hormonal activity or other or potentially hazardous active ingredients this should be stated under the relevant product type and dosage form.

1.3	Biological medicinal products
	<i>1.3.1 Biological medicinal products</i>
	1.3.1.2 Immunological products

Any restrictions or clarifying remarks related to the scope of this certificate:

This certificate reflects the status of the manufacturing of live lyophilized, as well as inactivated vaccines /oil-water emulsions and suspensions/, destined for immunization against viral and bacterial diseases in poultry, as listed in the Manufacturing Authorization No 19 of 05. 01. 2009, issued to "Mintech Co" EOOD, Sofia.

Done in Sofia on: 06.07. 2015

DR DAMYAN ILIEV
EXECUTIVE DIRECTOR
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