

Bulgarian Drug Agency

CERTIFICATE NUMBER: **BG/GMP/2025/304**

CERTIFICATE OF GMP COMPLIANCE OF A MANUFACTURER^{1, 2}

Part 1

Issued following an inspection in accordance with
Art. 111(5) of Directive 2001/83/EC as amended
Art. 63 of Regulation (EU) 536/2014 as amended

The competent authority of Bulgaria confirms the following:

The manufacturer: ***KeVaRo GROUP EOOD***

Site address: ***Floor 5, Ulitsa Tsaritsa Eleonora 9 Office 23, Sofia, 1618, Bulgaria***

OMS Organisation Id. / OMS Location Id.: ***ORG-100034005 / LOC-100053703***

Has been inspected under the national inspection programme in connection with manufacturing
authorisation no. ***BG/MIA-0499*** in accordance with Art. 61 of Regulation (EU) No 536/2014 and Art. 40 of
Directive 2001/83/EC.

From the knowledge gained during inspection of this manufacturer, the latest of which was conducted
on ***2025-02-04***, it is considered that it complies with:

- The principles and guidelines of Good Manufacturing Practice laid down in Directive (EU) 2017/1572
and/or Commission Delegated Regulation (EU) 2017/1569, as reflected by the product categories stated in
Part 2.³

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. Updates to restrictions or
clarifying remarks can be identified through the EudraGMDP website (<http://eudragmdp.ema.europa.eu/>).
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 EudraGMDP. If it does not appear, please contact the
issuing authority.

¹ The certificate referred to in paragraph Art. 111(5) of Directive 2001/83/EC and Art. 15 of Directive 2001/20/EC is also applicable to importers.

² Guidance on the interpretation of this template can be found in the Interpretation of the Union format for GMP certificate.

³ These requirements fulfil the GMP recommendations of WHO.

Part 2

Human Medicinal Products
Human Investigational Medicinal Products

1 MANUFACTURING OPERATIONS	
1.1	Sterile products
	1.1.3 Batch certification
1.2	Non-sterile products
	1.2.2 Batch certification
1.3	Biological medicinal products (list of product types)
	1.3.2 Batch Certification (list of product types) 1.3.2.5 Biotechnology products

2 IMPORTATION OF MEDICINAL PRODUCTS	
2.2	Batch certification of imported medicinal products
	2.2.1 Sterile products 2.2.1.1 Aseptically prepared 2.2.1.2 Terminally sterilised
	2.2.2 Non-sterile products
	2.2.3 Biological medicinal products 2.2.3.3 Cell therapy products 2.2.3.5 Biotechnology products
2.3	Other importation activities
	2.3.1 Site of physical importation

Clarifying remarks (for public users)

KeVaRo GROUP Ltd doesn't possess any storage premises. Medicinal products storage operations are performed under contract. The manufacturer is not authorized for following activities for Human Investigational Medicinal Products 1.1.Sterile products 1.1.3.Batch certification 1.2.Non-sterile products 1.2.2 Batch certification 1.3.Biological medicinal products 1.3.2.Batch certification 1.3.2.5 Biotechnology products. The following activity 2.2.3.3. Cell therapy products is only for Human Investigational Medicinal Products. KeVaRo GROUP Ltd doesn't possess any storage premises. Storage operations of investigational medicinal products are performed under contract at address: 425 Tsarigradsko Shose Blvd., Sofia 1137.

2025-03-25

Name and signature of the authorised person of the
Competent Authority of Bulgaria

Confidential
Bulgarian Drug Agency
Tel: ***Confidential***
Fax: ***Confidential***