



Health and Youth Care Inspectorate - Pharmaceutical Products

CERTIFICATE NUMBER: NL/H 20/2016993

CERTIFICATE OF GMP COMPLIANCE OF A MANUFACTURER

Part 1

Issued following an inspection in accordance with:

Art. 111(5) of Directive 2001/83/EC as amended

The competent authority of Netherlands confirms the following:

The manufacturer: DuSart Pharma B.V.

Site address: Nijverheidstraat 15, KATWIJK ZH, 2222AV, Netherlands

Has been inspected under the national inspection programme in connection with manufacturing authorisation no. 5304 F in accordance with Art. 40 of Directive 2001/83/EC transposed in the following national legislation:

Art. 100 of the Medicines Act

From the knowledge gained during inspection of this manufacturer, the latest of which was conducted on 2020-01-21, it is considered that it complies with:

The principles and guidelines of Good Manufacturing Practice laid down in Directive 2003/94/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 EudraGMDP. If it does not appear, please contact the issuing authority.

Signatory: Drs G. J. V. Bingen

The certificate referred to in paragraph 111(5) of Directive 2001/83/EC and 80(5) of Directive 2001/82/EC, shall also be required for imports coming from third countries into a Member State.

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

³ These requirements fulfil the GMP recommendations of WHO.



Part 2

Human Medicinal Products

1.2	Non-sterile products		
	1.2.1 Non-sterile products (processing operations for the following dosage forms)		
	1.2.1.6 Liquids for internal use		
1.4	Other products or manufacturing activity		
A 9 B			
	1.4.1 Manufacture of		

2020-01-22

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

Drs Gerrit Johannes van Ringen

Health and Youth Care Inspectorate - Pharmaceutical

Products

Tel: +31 88 1205000 Fax: +31 88 1205001



RELEASE CERTIFICATE

Concerning the following product:

Endoca CBD oil 10ml/flacon

Article number:

END-01001

Batch Number:

1852

Production date:

06/2020

Expiration date:

06/2022

The above mentioned product has been produced and tested in accordance with valid procedures and has been found to meet the required specifications.

Mr. C.M.J. de Groot, MSc Pharm D.'
Qualified Person

*Valid without signiature

DuSart Pharma by

Industriegebied 't Heen • Nijverheidstraat 15 • 2222 AV Katwijk ZH • P.O. Box 3219 • 2230 CE Katwijk ZH • The Netherlands T + 31 (0)71 4028564 • F + 31 (0)71 4028424 • E info@dspharma.nl • www.dusartpharma.com IBAN NL 14ING80578810133 • BIC INGBNL2A • KvK 28049143 • BTW nr A/AT no. NL 08838299881

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HOMEOPATHY * PHYTOTHERAPY * SPORTS FOOD SUPPLEMENTS * TABLETS * CAPSULES



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This is to certify that Henry Vincenty Staniulevicius today in my presence at the Notarial Office approved and signed the above document. No conspicuous corrections or addenda were found in the document.

Henry Vincenty Staniulevicius has proved his identity by showing passport.

The Court in Holstebro, Denmark, May 19, 2021

Ditte Aakjær Iversen Notary Public

SANMARKS DOMSTOLE

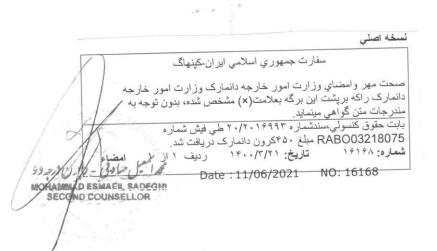
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2. has been signed by er underskrevet af	Ditte Aakjær I			
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4. bears the seal/stamp of er forsynet med segl/stempel af				
5. at	Copenhagen	6. the	25 May 2021	
i	København	den	25 maj 2021	
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-			Karl Johan Bech-Jessen	

This Apostille only certifies the authenticity of the signature and the capacity of the person who has signed the public document, and, where appropriate, the identity of the seal or stamp which the public document bears. This Apostille does not certify the content of the document for which it was issued.



To verify the issuance of this Apostille, scan the QR code or visit the following website:

https://e-register.um.dk



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