Certificate

Quality Management System

ecm Zertifizierungsgesellschaft für Medizinprodukte in Europa mbH, Bismarckstr. 106, 52066 Aachen, Germany, hereby declares that an examination of the undermentioned quality assurance system has been carried out following the requirements of DIN EN ISO 13485:2016.

Through an audit performed on behalf of

VSY Biotechnology BV
Strawinskylaan 1143, 1077 XX Amsterdam, The Netherlands,

it could be demonstrated that a quality management system

according to **DIN EN ISO 13485:2016**
"Medical devices – Quality management systems – Requirements for regulatory purposes"

for the **design, production and sales activities of intraocular lenses and other ophthalmic medical devices and intra-articular visosupplementation medical devices**

has been established and implemented.

This certificate is only valid under the conditions stated in the audit report mentioned hereafter. Any substantial changes of the quality assurance have to be notified to ECM and are subject to a separate assessment.

Report Number	Registered under	Valid until
565-18-94	**Z/18/04340E**	**November 18th, 2021**

Valid as of: November 19th, 2018

Certification Body

Digitally signed by Cojuhari Alexandru
Date: 2020.11.18 09:26:21 EET
Reason: MoldSign Signature
Location: Moldova