

DECLARATION OF CONFORMITY

Expiration 2028-12-31

MANUFACTURER: OASIS® Medical, Inc.
510-528 S. Vermont Ave.
Glendora, CA USA 91741

EUROPEAN REPRESENTATIVE: Donawa Lifescience Consulting Srl
Piazza Albania, 10
00153 Roma, Italy

PRODUCT: VISCO SHIELD® HPMC Viscoelastic (Injectable)
REF #: 51082, 5122

INTENDED USE: VISCO SHIELD® HPMC Viscoelastic is intended for use as a surgical aid by protecting the corneal endothelial cells and maintaining a deep anterior chamber during ophthalmic surgical procedures.

CLASSIFICATION: Class IIa, Rule 6 according to Annex IX of Directive 93/42/EEC

CONFORMITY ASSESSMENT ROUTE: Annex II of Directive 93/42/EEC

OASIS® MEDICAL, INC. DECLARES THAT THE ABOVE MENTIONED PRODUCTS CONFORM TO THE PROVISIONS OF THE COUNCIL DIRECTIVE 2023/607 AND MDR ARTICLE 120 FOR MEDICAL DEVICES. ALL TECHNICAL DOCUMENTATION IS RETAINED UNDER THE PREMISES OF THE MANUFACTURER.

TO THE BEST OF ITS KNOWLEDGE, INFORMATION AND BELIEF, OASIS® MEDICAL, INC. IS IN COMPLIANCE, IN ALL MATERIAL RESPECTS WITH ITS QUALITY MANAGEMENT SYSTEM ACCORDING TO EN ISO 13485:2016. THIS DECLARATION IS ISSUED UNDER THE SOLE RESPONSIBILITY OF OASIS® MEDICAL, INC.

NOTIFIED BODY: DQS 0297 – DQS Medizinprodukte GmbH
August-Schanz-Strasse 21
60433 Frankfurt a.M.
Germany

EC CERTIFICATE(S): EC Certificate Number: 288050 MR2
Issued: 2020-10-22
Valid Until: 2023-09-15

CONFIRMATION LETTER 2023-607: Issued 2023-06-23
(MDD CERTIFICATE EXTENSION GRANTED BY DQS)

QUALITY MANAGEMENT SYSTEM: EN ISO 13485:2016



Certificate Number: 288050 MP2016
Issued: 2024-09-04
Valid Until: 2027-09-15

 20 Dec 2024

Ishan Patil Date
Director of Regulatory Affairs, OASIS® Medical, Inc.