DECLARATION OF CONFORMITY(MDD)

OLYMPUS®

1. Manufacturer	OLYMPUS MEDICAL SYSTEMS CORP.
2. Address	2951 Ishikawa-cho, Hachioji-shi, Tokyo 192-8507 Japan
3. Model	OLYMPUS BF-1TH190
4. Name of product	EVIS EXERA III BRONCHOVIDEOSCOPE
5. Serial or Lot No.	from 2400001 to
6. Classification	Class IIa

7. Authorized representative in EU

Name	Olympus Europa SE & Co. KG
Address	Wendenstr. 14-18 20097 Hamburg, Germany

We hereby declare that the above mentioned product complies with the requirements of EC Directive 93/42/EEC(MDD)under the sole responsibility as a legal manufacturer. This Declaration of Conformity is valid in manufactured devices with the above Serial/Lot number.

This declaration is based on : MDD, Annex II.3

8. Notified Body Approval

Issued by	TÜV Rheinland LGA Products GmbH (0197)
Address	Tillystrasse 2, D-90431 Nürnberg, Germany

9. Applied Standards — Refer to the Essential Requirements Checklist for above mentioned product.

Place	2951 Ishikawa-cho, Hachioji-shi, Tokyo, Japan
Signature	
Name	Hiroki Moriyama
Title	Deputy General Manager Regulatory Affairs & Quality Assurance Department Quality & Environment Division
Date	2014/03/12

[N-OIS D28001 Appendix 3]