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 URF TYPE P5
4. Name of product	OES URETERO-RENO FIBERSCOPE
5. Serial or Lot No.	from 2600001 to
6. Classification	Class IIa

7. Authorized representatives in EU

■ Name: Olympus Europa Holding GmbH
Address: Wendenstr. 14-18 20097 Hamburg,Germany

We herewith declare that the above mentioned product complies with the requirements of EC Directive 93 / 42 / EEC (MDD).

This declaration is based on : MDD, Annex II

8. Certification of a quality system : Issued by TÜV Rheinland LGA Products GmbH (0197)

Place	2951 Ishikawa-cho, Hachioji-shi, Tokyo, Japan
Signature	
Name	Seiya Raiju
Title	General Manager, Regulatory Affairs & Quality Assurance Department
Date	2010/04/30(yyyy.mm.dd)

[N-OIS D28001 Appendix 3]
Revision 2 :2010/04/30 (yyyy.mm.dd)