

# **DECLARATION OF CONFORMITY**

We, **TERUMO CORPORATION**

**44-1, 2-chome, Hatagaya, Shibuya-ku, Tokyo 151-0072, Japan**

being the manufacturer of :

## **Progreat** **Angiographic Catheter**

**Product : Angiographic Catheter (GMDN 10688)**

declare that the above products of **Class III** are in conformity with the provisions of Annex I and Annex II of the EC Council Directive 93/42/EEC of 14 June 1993 concerning medical devices, and have been subject to the following conformity assessment procedure laid down in the EC Council Directive 93/42/EEC Article 11, 1(a) relating to the "Full quality assurance" set out in Annex II, under the supervision of TÜV Rheinland (Registration No.: HD 60012340 0001 ID 60016788 0001), as Notified Body authorized by the German Competent Authority and carrying the Notified Body No. 0197.

Authorized European Representative :

**TERUMO EUROPE N.V.**

**Researchpark Zone 2 Haasrode, Interleuvenlaan 40**

**3001 Leuven, Belgium**

Tokyo, January 4, 2007

(place and date of issue)



**Akira Oguma**

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**Quality Assurance Department**  
**TERUMO CORPORATION**

