DECLARATION OF CONFORMITY

We, TERUMO CORPORATION
44-1, 2-chome, Hatagaya, Shibuya-ku, Tokyo 151-0072, Japan

being the manufacturer of :

Destination
Guiding Sheath

Product : Catheter, Intravascular, Guiding (GMDN 17846)

declare that the above products of **Class IIa** 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, 3(a) relating to the "Full quality assurance" set out in Annex II, under the supervision of TÜV Rheinland (Registration No.: HD 60012340 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, July 27, 2006
(place and date of issue)

A. Oguma.
Akira Oguma
Director, General Manager
Quality Assurance Department
TERUMO CORPORATION

TERUMO®