

# **DECLARATION OF CONFORMITY**

**We, TERUMO CORPORATION**

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

being the manufacturer of :

**TR Band**

**Product : Radial Artery Hemostasis Band (GMDN 10284)**

declare that the above products of **Class I sterile** are in conformity with the provisions of Annex I and Annex VII 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, 5 of the Directive, relating to the "EC Declaration of Conformity" set out in Annex VII, combined with the provisions set out in Annex V "Production Quality Assurance" limited to the aspects of the manufacture concerned with securing and maintaining sterile conditions, under the supervision of TÜV Rheinland Product Safety (Registration No.: HD 60012340 0001), as Notified Body authorized by the German Competent Authority and carrying the Notified Body No. 0197.

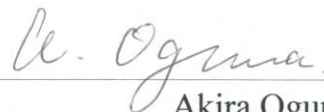
Authorized European Representative :

TERUMO EUROPE N.V.

Researchpark Zone 2 Haasrode, Interleuvenlaan 40  
3001 Leuven, Belgium

Tokyo, December 12, 2005

(place and date of issue)



Akira Oguma

Director, General Manager  
Quality Assurance Department  
TERUMO CORPORATION

