**LATEX DECLARATION**

- FINECROSS® MG Coronary Micro-Guide Catheter
- GLIDEATH® Hydrophilic Coated Catheter
- GLIDEATH® XP Hydrophilic Coated Catheter
- GLIDESHEATH SLENDER® Introducer Sheath
- GLIDEWIRE ADVANTAGE® Guidewire
- GLIDEWIRE® Endoscopic Hydrophilic Coated Guidewire
- GLIDEWIRE® Gold Hydrophilic Coated Guidewire
- GLIDEWIRE® GT Guidewire
- GLIDEWIRE® Hydrophilic Coated Guidewire
- GLIDEWIRE® Urologic Hydrophilic Coated Guidewire
- HEARTAIL® III Guiding Catheter
- MISAGO® RX Self-expanding Peripheral Stent
- NAVICROSS® Support Catheter
- Oburator
- OPTITORQUE® Diagnostic Catheter
- PROGREAT ALPHA™ Microcatheter
- PROGREAT® Coaxial Microcatheter System
- PROGREAT® Hydrophilic Coated Microcatheter
- R2P™ MISAGO® RX Self-expanding Peripheral Stent
- R2P™ SLENGUIDE™ Guiding Catheter
- RUNTHROUGH® NS Coronary Guidewires
- RUNTHROUGH® NS Extension Wire Guidewire
- RUNTHROUGH® NS HYPERCOAT™ Coronary Guidewire
- TORQUE™ Device
We, TERUMO CORPORATION, Japan, being the manufacturer of the following product hereby declare that:

- that above mentioned device and its packaging do not contain components made with natural rubber latex.

- that during the production of the above mentioned product no natural rubber (latex), and no natural rubber (latex) containing parts, have intentionally been used.

- that during the manufacturing process of the above mentioned product, no natural rubber (latex) gloves have been worn.

- that the possible presence of ubiquitous traces, of natural rubber (latex), can of course never be totally ruled out.

Sincerely,

Signature: [Signature]
Name: Tomoyuki Iida
Title: Deputy General Manager of Quality Assurance Department