Terumo Medical Corporation
Corporate Headquarters
265 Davidson Avenue, Suite 320
Somerset, New Jersey 08873

Date: August 22, 2024

We, TERUMO Medical Corporation, being the manufacturer of the following products hereby declare:

LATEX DECLARATION

PINNACLE™ Introducer Sheath	GLIDESHEATH™ Introducer Sheath
PINNACLE™ TIF TIP™ Introducer Sheath	GLIDEACCESS™ Micro Accessing System
PINNACLE™ R/O II HiFlo Introducer Sheath	PINNACLE™ DESTINATION™ Guiding Sheath
PINNACLE™ R/O II Radiopaque Marker Introducer Sheath	R2P™ DESTINATION SLENDER™ Guiding Sheath
PINNACLE PRECISION ACCESS SYSTEM™ Sheath	ANGIO-SEAL™ VIP Vascular Closure Device
GLIDESHEATH SLENDER™ Introducer Sheath	FEMOSEAL™ Vascular Closure Device
GLIDESHEATH SLENDER™ Tibial Pedal Introducer Sheath	TR BAND™ Radial Compression Device
RADIFOCUS™ Introducer Sheath	RADIFOCUS™ R/O II Introducer Sheath
RADIFOCUS™ R/O II HiFlow Introducer Sheath	

- The above-mentioned devices and their packaging do not contain components made with natural rubber latex.
- During the manufacturing of the above-mentioned products, no natural rubber (latex) and no natural rubber (latex) containing parts have intentionally been used.
- During the manufacturing of the above-mentioned products, no natural rubber (latex) gloves have been worn.
- The possible presence of ubiquitous traces of natural rubber (latex) cannot be ruled out and may be present.

Sincerely,

Kimberly Feitl
Vice President, Quality