Device Schedule: Class III and Class IIb devices

Class III, Implantable	Intended purpose
Coated VICRYL™ Plus Antibacterial Suture	See MDR 730033
ETHIBOND EXCEL™ Suture	See MDR 730045
ETHICON PHYSIOMESH™ Open Flexible Composite Mesh	See MDR 730035
LAPRA-TY™ II Clips	See MDR 730059
MERSILENE™ Suture	See MDR 730030
MONOCRYL™ Suture	See MDR 730032
MONOCRYL™ Plus Antibacterial Suture	See MDR 730060
PDS™ II Suture	See MDR 730038
PDS™ Cord	See MDR 730058
PDS™ Plate	See MDR 730036
PDS™ Plus Antibacterial Suture	See MDR 730053
PERMA-HAND™ Braided Silk Non-Absorbable Suture	See MDR 730051
PROCEED™ Surgical Mesh	See MDR 730258
PROCEED™ Ventral Patch	See MDR 730040
ULTRAPRO™ Hernia System	See MDR 730049
ULTRAPRO™ Mesh and ULTRAPRO ADVANCED™ Mesh	See MDR 730047
ULTRAPRO™ Plug	See MDR 730048
VICRYL™ Mesh	See MDR 730031
VICRYL™ Suture	See MDR 730029
VICRYL RAPIDE™ Suture	See MDR 730044

Device Schedule: Class IIa, Custom-made and other devices

Device(s)	Risk Classification
Clipping Devices	Class Ir
For Class Ir devices (Class I re-usable surgical instruments), the Notified Body conformity assessment is limited to the aspects relating to the reuse of the device.	