

Specific safety evaluation programs follow International Organization for Standardization (ISO) 10993 standards and Food and Drug Administration (FDA) guidance (May 1, 1995). The table is based on ISO 10993-1 Evaluation and Testing, 2009 edition. While the table has been developed as a guideline for biocompatibility testing, it is essential that each device be evaluated based on its own unique characteristics.

Device Categories		Biological Effect										
			Cytotoxicity	Sensitization	Irritation/Intracutaneous	Acute Systemic Toxicity	Subchronic Toxicity (Subacute toxicity)	Genotoxicity	Implantation	Hemocompatibility	Chronic Toxicity	Carcinogenicity
<b>Body Contact</b>		<b>Contact Duration</b> A = Limited (≤24 Hours) B = Prolonged (24 Hours - 30 Days) C = Permanent (>30 Days)										
		Surface Devices										
		Externally Communicating Devices										
Surface Devices	Skin	A	x	x	x							
		B	x	x	x							
		C	x	x	x							
	Mucosal Membrane	A	x	x	x							
		B	x	x	x	o	o		o			
		C	x	x	x	o	x	x	o		o	
	Breached or Compromised Surfaces	A	x	x	x	o						
		B	x	x	x	o	o		o			
		C	x	x	x	o	x	x	o		o	
Externally Communicating Devices	Blood Path, Indirect	A	x	x	x	x				x		
		B	x	x	x	x	o			x		
		C	x	x	o	x	x	x	o	x	o	o
	Tissue/Bone/Dentin <sup>1</sup>	A	x	x	x	o						
		B	x	x	x	x	x	x	x			
		C	x	x	x	x	x	x	x			
	Circulating Blood	A	x	x	x	x		o <sup>2</sup>		x		
		B	x	x	x	x	x	x	x	x		
		C	x	x	x	x	x	x	x	x	o	o
Implant Devices	Tissue/Bone	A	x	x	x	o						
		B	x	x	x	x	x	x	x			
		C	x	x	x	x	x	x	x		o	o
	Blood	A	x	x	x	x	x		x	x		
		B	x	x	x	x	x	x	x	x		
		C	x	x	x	x	x	x	x	x	o	o

X = Tests per ISO 10993-1  
O = Additional tests that may be applicable in the U.S.

**Note<sup>1</sup>** – Tissue includes tissue fluid and subcutaneous spaces  
**Note<sup>2</sup>** – For all devices used in extracorporeal circuits

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Notes:

## Medical Device Development Process & The MRO<sup>®</sup> Approach

