Ref No	Rev No.	Issue Date

Design output Reviw

Medical Device name:xxxxxx

Stage	item	Outputs requirements	method of verification	design output	verify
	product basic requirements				
	raw materials, component parts, and sub-assemblies,				
	Biological/ Toxicity requirement	must be biocompatible per ISO 10993.	Biocompatible test according to the ISO-10993	Biocompatibility tests -All tests are passed AT folder	Done
	Label and IFU				

This is an example of some points only , so the manufacturer can increase or change this points regarding manufacturing process

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