## SOP Reference #: QS016

Operation/Task:	Medtronic IFU Requirements			Equipment:	None
Owner:	Quality Manager	Date Created:  Revision History:	2/14/22022 See last page	Department:	Quality Systems
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ALERTS (see below): Critical Step ◆ Quality Check ✓ Tip ☺ Team Safety •

## Purpose: This SOP/work instruction describes specific requirements for producing work for customer Medtronic

Step #	Alerts	Step Description - "What to Do"	"How to Do it"	"Why to Do it"
1		Enhance inspection and QC techniques as required for customer Medtronic	Include additional production requirements for Medtronic IFU products	To achieve 100% conforming product
2	•	Offset press requirement for QC pulls is increased	Press requirement is double standard press pull frequency. Up to 5 per hour depending on press speed.	Additional QC pulls narrow field for non-conforming sheets.
3	•	Prepress requirement for bar code placement	Bar codes as close to press sheet gripper as possible and on first side impression when imposed.	Reduce possibility of bar code dot gain
4	•	Finish requirements	<ol> <li>Inspection of 3 sides of stitched material lifts</li> <li>Fan parts prior to loading stitcher pocket to inspect bar code.</li> <li>Test bar codes at packaging station</li> <li>Perform line clearance process for each version included in a production run</li> </ol>	Remove non-conformances at stitcher  Ensure bar codes are clean  Ensure bar codes can be read  Keep versions separate and product accurate.

5	•	Inspection requirement and trigger for 100% inspection	Inspection of finished product must follow the requirements of the established sampling technique. See QS017 for requirements.  If during the sampling process, a single nonconforming part is found, the sample must be doubled and inspected with no additional non-conforming parts found in order to pass	This technique is an accepted protocol for sampling and inspection.  The goal is 100% conformance so the standard is severe.
			inspection.  If more than one non-conforming part is found during sampling, either in the initial inspection or the expanded sample, a 100% inspection is triggered for the discovered non-conformance. This means all parts of the job must be inspected for conformance and a count kept of all parts discovered to be non-conforming.	The expansion of the sample is used to determine the breadth of the non-conformity
			If the 100% inspection is triggered, alert the department supervisor or quality manager. The supervisor or quality manager will supervise the inspection.	The supervisor or quality manager may determine that the inspection may be suspended based on finding no additional non-conforming parts after inspecting 20% of the total lot.
6	•	Project Management	Record customer requirements in MIS system database for customer Medtronic	Requirements are available to print on production documents specific to customer.

Notes:	lotes: This process is invoked upon behalf of customer.					

Definitions: QS017 is a quality system SOP defining the sampling technique deployed at ENPOINTE.

Revision History	Description of Changes	Requested by	Date
Original	First posting to intranet	Dean Milinkovich	3/7/2022
Rev. 1	Added instruction for 100% inspection requirement	Dean Milinkovich	6/22/2022