

## **Technical Service Bulletin**

## **Monitoring Products**

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**Bulletin Type:Information Only** 

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Date: 2011-01-24 SAP DIR #: 20012310

Subject: TSB - LXi Main PCB RoHS Compliance

CAR Number: N/A

**Distribution Scope:** WA Customers, ASP's, and Distributors

Define the intended

audience.

WA Technical Support, Product Service, &InternationalServiceCenters

Product(s) Referenced: Spot LXi Vital Signs Monitor

SW Version: ALL

Serial No. / Lot Code: ALL

Summary: Manufacturing process change for Spot LXi main PCB assembly

Issue: Improve 'environmentally-friendly' aspect of product by replacing main PCB

assembly (p/n 404940) that uses a lead-based solder to a main PCB assembly (p/n 403912) that utilizes a lead-free solder (ROHS Compliant).

**Action:** No specific action required. This is informational only.

Reference to • 21 CFR Part 820, ISO 13485, MPD SOP-0002

**Standards:** Add additional references as needed.

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Printed or electronic versions of this document not accessed directly from the designated Welch Allyn Controlled Quality Information System are For Reference Only. **Updates:** If required **Technical Manual** Repair Tool Service Plan Internet/Intranet **Procedures Price List Training Material** Other All service centers should return existing stock of 404940 to the **Service Strategy:** manufacturing facility, and replace stock with material 403912. N/A **Required Training:** N/A **Required Tools: Required Materials:** N/A N/A **Quality Process for** failed units or components: Recognition of the different versions of the Spot LXi main PCB assembly is **Procedure:** based on material number: The old board is labeled 404940. The new board is labeled 403912. **Quality Documents:** All service centers using SAP to record service transactions: For each monitor serviced, record the service activity in SAP. All other service centers and Field Service: For each monitor serviced, complete and file a service report and attach to the service DHR.

## **Notes:**

- 1. Contact the Welch Allyn Complaints Department to initiate or process a medical device complaint resulting from this or other issues.
- 2. Drawings, illustrations, and part numbers in this document are for reference purposes only and subject to change.

## **End of Bulletin**

Revision History					
Version	Description	Change #	Init	Release Date	Appr
A	Initial Release	n/a	NEA	2011-01-25	D*
D* - See SAP DIR for Change number, Approver Name and Date of Approval					

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