

# Procedure Incoming Nonconforming Product P8.3-2 Rev. G

**Approved By:** Mike Orsini, Quality Manager

**Purpose:** To document methods of identifying and dispositioning non-conforming material.

**Scope:** All non-conforming material received from suppliers or rejected in production lines.

**Authority:** The Quality Manager has the authority to change or modify this procedure.

## Reference Documents and Records

- Policy Element 8.3 Control of nonconforming product
- DMR
- Supplier Corrective and Preventive Action Procedure P8.5.2/3-2
- Quality Records Procedure P4.2.4

## Change History

Date	Change	Rev	By
12/10/96	Initial release after rewrite	C	MO
03/19/97	Changed to use new DMR form – step 2	D	MO
4/16/02	Removed ref. To DMR database – no longer used.	E	MO
10/29/2002	Updated to the ISO 9001:2000 requirements and flow charted.	F	MO
10/9/03	Updated procedure to reflect CPAP initiated as result of review.	G	MO

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