



## Corrective and Preventive Action Form

(Ref. [SOP QMS-175](#))

Originator					
Subject					
Source of CAPA					
<input type="checkbox"/> Nonconformity from internal audit		<input type="checkbox"/> Complaint			
<input type="checkbox"/> Repeat deviations		<input type="checkbox"/> Suggestion for improvement			
<input type="checkbox"/> Product Review		<input type="checkbox"/> Other _____			
Description of issue requiring corrective action					
Name		Signature		Date	
Quality Assurance Review					
<input type="checkbox"/> Not approved					
Reason for not approving CAPA					
Name		Signature		Date	
<input type="checkbox"/> Approved      CAPA Plan Due Date: _____					
Immediate Action Required					
Assigned to:				CAPA #	
Name		Signature		Date	



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(Ref. SOP QMS-175)

Management Action					
Analysis of Root Cause: (if insufficient room please attach)					
If Risk Assessment is required please attach. If not required, please explain					
Corrective Action Plan			Due Date:		
Action		Responsibility		Due Date	
Manager		Signature		Date	
QA Approval		Signature		Date	
Quality Assurance Follow up / Verification					
List documented evidence of CAPA implementation					
Name		Signature		Date	
Effectiveness Review					
<input type="checkbox"/> CAPA Closed					
Name		Signature		Date	