

DEVIATION REQUEST

ISF-QA-012

DATE		NUMBER	
REQUESTER PURCHASE / FACTORY CONTROL / PRODUCTION / OTHERS.			
Details of Deviation Requested (Product / Process)			
Part Number		Description	Qty/Period:
Existing Status (Indicate condition of Items (already in use) prior to this deviation):			
Status Expected after Deviation:			
Corrective Action / Target :			
Requester Signature			
Comments by production:			
Comments by Factory Control:			
Comments by Process Engineering:			
Deviation Category : A [] B []			
Comments by Quality Control:			

1. Are the final product characteristics impaired.	Yes <input type="checkbox"/>	No <input type="checkbox"/>
2. Any Updatons required in PPAP documents.	Yes <input type="checkbox"/>	No <input type="checkbox"/>
3. Any temporary work instruction / process sheets required.	Yes <input type="checkbox"/>	No <input type="checkbox"/>
4. Customer Approval required (As agreed with with customer)	Yes <input type="checkbox"/>	No <input type="checkbox"/>
5. Information required to customer (as agreed with customer)	Yes <input type="checkbox"/>	No <input type="checkbox"/>
6.any specific inspection / checks required.	Yes <input type="checkbox"/>	No <input type="checkbox"/>
7. Any specific records / check sheets to be maintained.	Yes <input type="checkbox"/>	No <input type="checkbox"/>
8. Any separate Identification requirements	Yes <input type="checkbox"/>	No <input type="checkbox"/>
If Yes , Details		
Any Another Remarks / Comments .		
Repeat Deviation		
Previous Dev . No.		Qty / Period
Previous Dev . No.		Qty. / Period
Final Conclusion :		
Deviation Accepted / Rejected for	Quantity	Period
Date :	Authorised :	
	HOD Process Engg.	Div. Head (For A category)
Deviation Monitoring		
Date		
Quantity		
Deviation Closure Date		