



# PROJECT AMENDMENT REQUEST

Applicant Name \_\_\_\_\_ Date \_\_\_\_\_  
 Address \_\_\_\_\_ Requester \_\_\_\_\_  
 \_\_\_\_\_ Phone # \_\_\_\_\_  
 \_\_\_\_\_ E-mail \_\_\_\_\_

REPORTS AFFECTED E #: \_\_\_\_\_  
 MET Report(s) #: \_\_\_\_\_  
 EN MET Project #: \_\_\_\_\_  
 CB MET US #: \_\_\_\_\_  
 Other: \_\_\_\_\_

PRODUCT(S) AFFECTED \_\_\_\_\_  
 \_\_\_\_\_

1) Description of file or product change: (ok to reference detailed attachments)  
 \_\_\_\_\_  
 \_\_\_\_\_

2) Manufacturer's drawings and/or other documents affected: (ok to reference detailed attachments)  
 \_\_\_\_\_  
 \_\_\_\_\_

3) Additional information: (please attach and reference any information that cannot be included here)  
 \_\_\_\_\_  
 \_\_\_\_\_

4) If this is a medical product for the US market, are the above changes considered significant changes in accordance with the guidance provided by the [FDA](#)?  
 \_\_\_\_\_  
 \_\_\_\_\_

To Be Completed by MET Personnel Only			
Reviewed By (Print) _____	Hold	Date	_____
(Signed) _____	Rejected	Date	_____
	Accepted	Date	_____
Comments/Instructions			
_____			
_____			
_____			
*If no response is received by the next follow-up inspection, the approval to apply the MET Mark may be revoked by the inspector upon the next follow-up inspection.			

Return this form to the Sales Representative at MET Laboratories, Inc. OR [info@metlabs.com](mailto:info@metlabs.com)