

## APPENDIX I: Method Development and Validation Project Record

<b>MDVP REQUEST PROJECT RECORD **</b>		<b>1. PROJECT NUMBER</b>	
<b>5. TITLE</b>		<b>2. LABORATORY</b>	
		3. FISCAL YEAR(S)	
		4.	
6. SCIENTIST(S), NAME, SIGNATURE AND DATE			
7. CONTINUING RESEARCH		8. PROJECT PLAN	
<input type="checkbox"/> YES (If yes explain in item 10) <input type="checkbox"/> NO		STARTING DATE	
		COMPLETION DATE	
		FISCAL YEAR	HOURS
		TOTAL HOURS	0
10. COMMENTS <b>PURPOSE</b>  <b>MILESTONE(S)</b> <b>TASKS IDENTIFIED TO REACH MILESTONE(S)</b> Identify approximate timeframes for each task  <b>PROBLEM THE PROJECT INTENDS TO SOLVE</b>  <b>REGULATORY SIGNIFICANCE AND RELEVANCE TO FDA MISSION (PUBLIC HEALTH IMPACT)</b>  <b>EXPECTED COMPLETION DATE</b>  <b>EXPERIMENTAL PROTOCOL/VALIDATION PROTOCOL</b>			
11. NAME OF SCIENCE ADVISOR (Date)	12. NAME OF SUPERVISOR (Date)	13. NAME OF APPROVING SUPERVISOR	

FDA MDVP Template, Version 1.0 (07/07/2011)

\*\* All new projects submissions must be submitted between October and June of that Fiscal Year. Project submitted after this time will be review for the following fiscal year. Projects which will continue into the next fiscal year must be resubmitted for hour approvals.