



Sample Submission Form for Regulated Products

• Phone: (513) 471-1300

• Fax: (513) 471-5600

[Pharmaceutical (New Drug/Generic/API), Finished Product OTC (Antimicrobial, Sunscreen, Actives), Raw materials/components (Drugs), Dietary Supplements, Water/environmental samples (Drug or OTC product components)]

• www.QLABORATORIES.com

• Email: office@qlaboratories.com

<b>Customer Information</b>				<b>Date Submitted:</b>		
<b>Company Name:</b>				<b>Billing Contact (if different):</b>		
<b>Report Results To:</b>				<b>Billing Email:</b>		
<b>Street Address:</b>				<b>Street Address:</b>		
<b>City, State, Zip:</b>				<b>City, State, Zip:</b>		
<b>Phone #:</b>				<b>Phone #:</b>		
Please check option(s) for receiving results (electronic documents will be in Adobe Acrobat (PDF) format).				<b>Purchase Order # (if applicable):</b>		
<b>Fax to:</b>						
<b>Email to:</b>				* Conduct of regulated study? <input type="checkbox"/> GLP <input type="checkbox"/> GMP <input type="checkbox"/> Other _____		
<b>Q Labs Proposal # (if applicable):</b>				Is this product filed for NDA or ANDA (or IND)? <input type="checkbox"/> Yes <input type="checkbox"/> No		
<b>Turnaround time requested:</b> <input type="checkbox"/> Routine <input type="checkbox"/> Rush				Is this a Research and Development project? <input type="checkbox"/> Yes <input type="checkbox"/> No		
Sample #	Sample ID	Active	Test Name	Method (if client method, include version)	Has the method been validated? Yes or No	Specifications
1						
2						
3						
4						
5						
6						
7						
8						
9						
<b>Special Instructions:</b>				<b>Authorizing Signature</b>		
Completed and signed sample submission form indicates agreement with Q Laboratories terms and conditions and authorizes Q Laboratories to perform the requested tests. After testing is complete, samples will be placed into appropriate storage (e.g., refrigerator, freezer, dry storage) and held for a minimum of 30 days before discarded, unless otherwise dictated by the client.						
<b>Return sample(s) to client?</b> <input type="checkbox"/> Yes <input type="checkbox"/> No						

\* A minimum charge of \$295.00 may be assessed to conduct an Out of Specification/Microbiological Deviation investigation for each non-Research and Development product that does not meet specification.



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Sample #	Sample ID	Active	Test Name	Method (if client method, include version)	Has the method been validated? Yes or No	Specifications
10						
11						
12						
13						
14						
15						
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17						
18						
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20						
21						
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28						
29						
30						
31						

\* A minimum charge of \$295.00 may be assessed to conduct an Out of Specification/Microbiological Deviation investigation for each non-Research and Development product that does not meet specification.