

LABORATORY SERVICE REQUEST (LSR) – Bioburden (ISO)

CLIENT INFORMATION

Instructions: Submit the form along with sample(s). Initiation of sample analysis will be delayed if form is not complete.

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| PBL Quote Number: | PO Number: <i>Credit Card to be used for payment</i> <input type="checkbox"/> |
| Sponsor (send report to): | Invoice To: <i>Check box if same as sponsor</i> <input type="checkbox"/> |
| Contact Name: | AP Contact Name: |
| Company Name: | Company Name: |
| Address: | Address: |
| City/State/Zip: | City/State/Zip: |
| Country: | Country: |
| Phone: | Invoice To Email: |
| Email: | AP Email: |

SERVICE INFORMATION

Regulatory Treatment: Non-regulatory cGMP GLP **Rush Service (will incur a surcharge):** YES NO

Suitability (required by GMP regulations):
 Suitability Completed*
 Suitability Declined (please specify testing parameters in the Special Instructions)
 Suitability to be conducted by Pacific BioLabs (Sterile samples required for Suitability test. Sample packaging will not be included unless specified in special instructions) Complete the suitability section on the second page.

DEVICE INFORMATION

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| Sample 1 Identification Please use the exact wording you want to appear in the final report. | Quantity of Samples Submitted Please indicate the number of units, volume, and/or weight of samples. |
| Lot Number: | Part Number: |
| Other Identifier: | Expiration Date: |
| Physical Description: <input type="checkbox"/> Device <input type="checkbox"/> Solid <input type="checkbox"/> Liquid <input type="checkbox"/> Powder <input type="checkbox"/> Gel <input type="checkbox"/> Other: | |
| *Suitability - PBL Report Number: | Guideline Number: |
| Sample 2 Identification Please use the exact wording you want to appear in the final report. | Quantity of Samples Submitted Please indicate the number of units, volume, and/or weight of samples. |
| Lot Number: | Part Number: |
| Other Identifier: | Expiration Date: |
| Physical Description: <input type="checkbox"/> Device <input type="checkbox"/> Solid <input type="checkbox"/> Liquid <input type="checkbox"/> Powder <input type="checkbox"/> Gel <input type="checkbox"/> Other: | |
| *Suitability - PBL Report Number: | Guideline Number: |
| Sample 3 Identification Please use the exact wording you want to appear in the final report. | Quantity of Samples Submitted Please indicate the number of units, volume, and/or weight of samples. |
| Lot Number: | Part Number: |
| Other Identifier: | Expiration Date: |
| Physical Description: <input type="checkbox"/> Device <input type="checkbox"/> Solid <input type="checkbox"/> Liquid <input type="checkbox"/> Powder <input type="checkbox"/> Gel <input type="checkbox"/> Other: | |
| *Suitability - PBL Report Number: | Guideline Number: |

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| Sample 4 Identification Please use the exact wording you want to appear in the final report. | Quantity of Samples Submitted Please indicate the number of units, volume, and/or weight of samples. |
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| Lot Number: | Part Number: |
| Other Identifier: | Expiration Date: |

Physical Description: Device Solid Liquid Powder Gel Other:

*Suitability - PBL Report Number: _____ Guideline Number: _____

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| Storage Condition <input type="checkbox"/> Room Temperature <input type="checkbox"/> 2 to 8C <input type="checkbox"/> -10 to -25C <input type="checkbox"/> -60 to -90C | Controlled Substance <input type="checkbox"/> No <input type="checkbox"/> Yes Schedule: NDC #: Concentration: | Hazardous <i>Include MSDS if samples are hazardous. A fee will apply for disposal of hazardous samples.</i> <input type="checkbox"/> Not Hazardous <input type="checkbox"/> Reactive <input type="checkbox"/> Biohazard <input type="checkbox"/> Toxic <input type="checkbox"/> Other: | Sample Disposition <i>Samples will be discarded per PBL SOP unless otherwise indicated.</i> <input type="checkbox"/> Return UNTESTED Samples to Client <input type="checkbox"/> Return TESTED Samples to Client <i>(must be preapproved by PBL)</i> <i>If return address is different than Client address above, indicate address in Special Instructions below.</i> Carrier: Account: Dispose in: <input type="checkbox"/> municipal waste <input type="checkbox"/> hazardous waste |
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TESTING REQUIRED

List part(s) of the Sample to be tested:

Final Intended Sterilization Method/Dose:

Final Intended Use/Application of Sample:

Test Article: Pharmaceutical Medical Device
Method: Pour Plate Membrane Filtration

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| Bioburden Validation (ISO/AAMI) <input type="checkbox"/> Aerobic Bacteria/Fungi <input type="checkbox"/> Spores (must indicate sterilization method above) <input type="checkbox"/> Anaerobes Liquids/Powders <input type="checkbox"/> Validation to be conducted by Pacific BioLabs Medical Devices <input type="checkbox"/> Validation to be conducted by Pacific BioLabs Method: <input type="checkbox"/> Spore Recovery Study (For sterile or near sterile samples) <input type="checkbox"/> Exhaustive Recovery (For samples with ≤300 and ≥100 CFU) | Bioburden Routine Testing (check all that apply) <input type="checkbox"/> Aerobic Bacteria/Fungi <input type="checkbox"/> Spores (must indicate sterilization method above) <input type="checkbox"/> Anaerobes <input type="checkbox"/> Verification Dose Calculation |
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Microbial Confirmation Tests (check all that apply, include quantity of each test)
 Bacterial ID, Qty. Yeast/Mold ID, Qty. Gram Stain/Colony Morphology, Qty.

OTHER TESTS/SPECIAL INSTRUCTIONS

Note that samples will be reported separately per lot unless otherwise indicated.

TESTING AUTHORIZED BY (please sign): _____ **DATE:** _____
(Signature and date, or electronic signature, is required for testing to begin; unsigned LSR forms will not be processed)