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| General information Customer | Product information |
| Company name |       | Article name |       |
| Preparation type |       |
| Contact person |       | Batch number |       |
| Email address |       | Reference to order(PO number) |        |
| Telephone number |       | Quotation number |       [ ]  N/A |
| Add. Remarks CoA       |

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| **Sample and test information** |
| Method of analysis [ ]  EP [ ]  USP [ ]  In-house [ ]  Other (please specify)       |
| Routine tests (Acc. to validation report:                          ) |
|  [ ]  **Microbial quality of non-sterile product**[ ]  TAMC (<100 or 1000CFU/g/ml)[ ]  TYMC (<10 or 100CFU/g/ml)[ ]  *E.coli* (absence per 1 g/ml)[ ]  *S.aureus* (absence per 1 g/ml)[ ]  *P.aeruginosa* (absence per 1 g/ml)[ ]  *C.albicans* (absence per 1 g/ml)[ ]  Bile-tolerant gram-negative bacteria (absence per 1 g/ml)[ ]  *Salmonella* (absence per 10 g/ml)[ ]  Other       | [ ]  **Test for sterility*** Batch size
* Number of containers to be tested
 |
| [ ]  Test for bacterial endotoxins (BET), Method C (Turbidimetric kinetic method)**LIMIT**       |
| [ ]  Microbiological assay of antibiotics**LIMIT**       |
|  |
| Non-Routine tests |
| [ ]  Suitability of the method**, Microbial quality of non-sterile product** Please specify the required parameters above | [ ]  Validation Test for bacterial endotoxins (BET), Method C (Turbidimetric kinetic method)**LIMIT**       (3 batches required) |
| [ ]  Suitability of the method**,Test for sterility*** Batch size

Number of containers to be tested       | [ ]  Efficacy of antimicrobial preservation (challengetest) |
| Specifications Analysis (free text field):  |
|       |
|       |

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| **Sample Handling** | [ ]  Normal (10 working days target) | [ ]  Priority (Only on advance notice, confirmation by Synergy Health required) |

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| **Sample status** | **Sample size and/or container number** |
| [ ]  Final drug product; testing purpose: batch release [ ]  Final drug product; testing purpose: not for batch release (e.g., stability study, process validation)[ ]  Restricted substance **(OPIUM LAW)**[ ]  Final drug product, manufactured outside EU[ ]  Investigational medicinal product (IMP) |            | [ ]  gram [ ]  ml[ ]  pieces |
| **Sample storage conditions** |
| [ ]  -20°C ± 5°C [ ]  2 - 8°C [ ]  Ambient temperature[ ]  Protected from light |
| Customer is obliged to share a Safety Data Sheet /SDS when submitting **DANGEROUS** goods for the first time.  |

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| Signature Customer: |       | Date: |       |