

Master Formulation Record Pharmacy compounding of non-sterile preparations

| Name of compounded product | | Protocol number and version | | |
|------------------------------|-----------------|-----------------------------|---|--|
| Concentration | | Effective date | | |
| | | Day / Month / Year | | |
| Pharmaceutical form | _ | Developed by | _ | |
| | | | | |
| Route of administration | | Verfied by | | |
| | | | | |
| Formula | | | | |
| Ingredients | Quantities | Physical description | Other information (i.e DIN, unique identifier such as CAS, lot number, manufacturer, expiry date, expected yield) | |
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| Additional information about | the ingredients | | | |
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| Notes on calculations and measurements | | | | |
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| Required equipment, instruments, and materials | | | | |
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| Compounding method | | | | |
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| Quality controls | | | | |
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| Postonin a | | | | |
| Packaging | | | | |
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| Stability and storage | |
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| Labelling | Sample label |
| | |
| | Patient label |
| | Patient label |
| Training | |
| | |
| References consulted | |
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| Preparation date sheet history # | |
| Treparation date sheet history # | |

| Revised | | Revised by | | |
|-------------|--------------------|------------------------|-----|----|
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| | | | | |
| | Day / Month / Year | | | |
| Change made | | | | |
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| | | Version number changed | Yes | No |
| | | | | |
| Revised | _ | Revised by | | |
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| | | | | |
| | Day / Month / Year | | | |
| Change made | | | | |
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