Specification: <Title>

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| Reviewed by: | Job title: | Signature: | Date: |
| Quality representative signs to confirm document complies with quality management system | | | |
| Authorised by: | Job title: | Signature: | Date: |

# Product Profile

|  |  |
| --- | --- |
| Product: |  |
| Description: |  |
| Standard batch size: |  |
| How supplied: |  |
| Expiry dating: |  |
| Storage conditions: |  |
| Manufactured by: |  |
| Packed by |  |
| Originating company |  |

# Components

|  |  |
| --- | --- |
| Material Name | % volume |
|  |  |
|  |  |
|  |  |
|  |  |

# Certificate of analysis

The following information must be included in the supplier’s Certificate of Analysis.

|  |  |  |
| --- | --- | --- |
| Test | Test method | Limits |
|  |  |  |
|  |  |  |
|  |  |  |

# Stability

|  |  |  |
| --- | --- | --- |
| Test | Test method | Limits |
|  |  |  |
|  |  |  |
|  |  |  |

# Packaging

## Components

|  |  |
| --- | --- |
| Code | Description |
|  |  |
|  |  |
|  |  |
|  |  |

Document Information

| Revision History | | | |
| --- | --- | --- | --- |
| Revision | Modified by | Change Control No. | Description of Change |
| 01 |  |  |  |
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List all controlled procedural documents referenced in this document (for example, policies, procedures, forms, lists, work/operator instructions)

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DOCUMENT END