Manufacturing summary sheet

This form should be completed for each new site master file (SMF) revision submitted. Revised versions of the SMF should be submitted in PDF format with the changes tracked.

1. Applicant details

|  |  |
| --- | --- |
| Name and address of authorisation/licence holder |  |
| Address of manufacturing premises |  |
| Companies registration office Number |  |
| Licence/authorisation number |  |
| Last issued endorsement number  (see footer of current authorisation) |  |

1. Manufacturer’s licence/authorisation details

Please indicate if each of the following parts of your manufacturer’s licence/authorisation are currently up-to-date:

|  |  |
| --- | --- |
| Address of authorisation holder | Yes  No  If ‘no’, please give brief details: |
| Address of manufacturing site(s) | Yes  No  If ‘no’, please give brief details: |
| Manufacturing operations (see Annex 1) to this document.) | Yes  No  If ‘no’, please give brief details: |
| Product categories (see Annex 1) | Yes  No  If ‘no’, please give brief details: |
| List of imported products (see Annex 1) | Yes  No  If ‘no’, please give brief details: |
| Contract manufacturers/storage sites | Yes  No  If ‘no’, please give brief details: |
| Contract laboratories | Yes  No  If ‘no’, please give brief details: |
| Named personnel | Yes  No  If ‘no’, please give brief details: |

1. Qualified Person details

|  |  |
| --- | --- |
| Name |  |
| Telephone number |  |
| Fax number |  |
| E-mail |  |

1. Number of Personnel working at the Authorised Site

If additional space is required, please complete on a separate sheet.

|  |  |
| --- | --- |
| Department | No. of personnel |
|  |  |
|  |  |
|  |  |
|  |  |
|  |  |
|  |  |
|  |  |
|  |  |
|  |  |
|  |  |
|  |  |
|  |  |
|  |  |
|  |  |
|  |  |

1. declaration

|  |  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- | --- |
| Current site master file revision number  Copy provided to the HPRA  Yes  No  Any other relevant information  I hereby declare that, to the best of my knowledge and belief, all the particulars given in this site master file reflect the facilities, operations and organisational structures at the site   |  |  | | --- | --- | | Signature (primary QP): | Date: | |  |  | | Print name: | Title/position: |     *Notes:*  Applications must bear the signature of the applicant. Where the application is on behalf of a limited company, the declaration must be signed by a director or the secretary and, in the case of a partnership, by a partner. |

1. return address

Send to:

Licensing Section

Compliance Department

Health Products Regulatory Authority

Kevin O’Malley House

Earlsfort Centre

Earlsfort Terrace

Dublin 2

Tel: + 353 1 676 4971

Fax: + 353 1 676 7836

E-mail: [compliance@hpra.ie](mailto:compliance@hpra.ie)

Annex 1

Please include a full list of product names and dosage form for the following activities.

|  |  |  |
| --- | --- | --- |
|  | PRODUCT NAME(S) | DOSAGE FORM |
| Bulk manufactured on site |  |  |
| Primary packaged on site |  |  |
| Secondary packaged on site |  |  |
| QC tested on site |  |  |
| Batch released to market |  |  |
| Site of importation only |  |  |