# WHO PQT-VC Declaration of Manufacturing Sites

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| --- | --- |
| **Company:** | [Company name] |
| **Product Name:** | [Product name] |
| **PQ Ref #:** | [PQ Product Ref Number] (if not yet assigned, leave blank) |
| **Product Type[[1]](#footnote-2)** | [Product Type] |

**Instructions:**

Instructions and guidance for completing the Declaration of Manufacturing Sites (DMS) are included in this template in red text. All red text should be deleted from the DMS prior to submission.

Text in [square brackets] should be replaced by appropriate descriptive language.

Refer to the implementation guidance for Product Manufacturing Details for further guidance for ITNs.

Lines may be added to the tables if more space is needed.

The list of manufacturing sites for each prequalified product published on the WHO website is developed based on the information provided on this form. Please ensure that all information is accurate and complete.

## Identification of Manufacturing Sites

*The manufacturers and addresses for production of the AI source materials must be identified in the table below. The address provided must be the actual address of the manufacturing site. A non-manufacturing HQ/office address should not be provided.*

### Active Ingredients and Synergists

|  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- |
| **Active Ingredient/Synergist Name** | **Owner of Site** | **Name of Site** | **Address[[2]](#footnote-3) (including blocks/units)** | **Country** | **Supporting WHO Specification[[3]](#footnote-4)** |
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The manufacturers and addresses for production of the VCP must be identified in the table below. The address provided must be the actual address of the manufacturing site. A non-manufacturing HQ/office address should not be provided.

If a third-party manufacturing facility is contracted for use in the production of the VCP, the information provided must reflect the legal name of the contracted company. The company name of the applicant must not be provided in place of the name of the contracted company.

The information provided in the Activities field must rely on the established terminology found in the implementation guidance Terminology for Describing Activities of Manufacturing Sites for VCPs.

The information provided in the Supporting SMF field should reflect the file name of the submitted SMF. This may be the file name of a compressed file (e.g. .zip, .7z, or .rar) with compiled documents or the primary file in the SMF. The version number and effective date of the SMF should also be provided.

### End Use Vector Control Product

|  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- |
| **Owner of Site** | **Name of Site** | **Address (including blocks/units)** | **Country** | **Activities[[4]](#footnote-5)** | **Supporting SMF[[5]](#footnote-6)** |
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| **Name of Authorized Contact Person for the Manufacturer: \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_**  **Signature of Authorized Contact Person for the Manufacturer: \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_**  **Date:** |
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1. Examples: LLIN, IRS, Space Spray, Larvicide, Topical Repellent, Spatial Repellent, Molluscicide, Rodenticide, or other description. [↑](#footnote-ref-2)
2. Provide the actual address of manufacturer, not the HQ/Office address of the source provider. [↑](#footnote-ref-3)
3. Identify the WHO specification code (ex. 333/TC, 454/SC). [↑](#footnote-ref-4)
4. Rely on the established terminology found in the WHO PQT-VCP IG Terminology for Describing Activities of Manufacturing Sites for VCPs. [↑](#footnote-ref-5)
5. Identify the file name and version of the supporting SMF. [↑](#footnote-ref-6)