**COMMITMENT**

I (Full Name) \_\_\_***\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_***

Certify that the product offered to UNICEF in reference to solicitation **No \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_**

Item description (Use INN name) **\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_**

UNICEF Material Number (Where applicable)\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

Registered trade/brand name if any\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

1. Is identical in all aspects of manufacturing and quality to that **previously approved by UNICEF** including formulation, method and site of manufacture, sources of active and excipient starting materials, quality control of the product and starting materials, packaging, shelf-life and product information. (Provide reference to previous approval or procurement)

**AND/OR**

1. Is identical in all aspects to that Prequalified by WHO, USFDA approved or Global Fund ERP

Name of Agency/Authority ***\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_***

Reference number ***\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_***

Valid until \_***\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_***

**AND/OR**

1. Is identical in all aspects to that registered and marketed in the following SRA[[1]](#footnote-1) country

Name of country ***\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_***

Name of regulatory authority***\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_***

Reference number ***\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_***

Valid until \_***\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_***

**AND**

1. Is identical in all aspects of manufacturing and quality to that registered and marketed in the following countries. (Attach separate list)

Describe any exceptions or changes to the product here (Use additional paper if necessary) \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

**Signature Date**

**Official stamp**

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|  |  |
| --- | --- |
| Name of company submitting BID |  |
| Physical address |  |
| Postal address |  |
| City, Country |  |
| Telephone, Fax |  |
| E-mail, website |  |

|  |  |
| --- | --- |
| Name of FPP manufacturer |  |
| Physical address of manufacturing site(s), including unit/block number |  |
| Postal address |  |
| City, Country |  |
| Telephone, Fax |  |
| E-mail, website |  |

|  |  |  |  |
| --- | --- | --- | --- |
| **Responsible person for** | **Name of contact person** | **Telephone and cell phone** | **E-mail** |
| Technical specifications & product quality |  | Tel:  Cell: |  |
| Regulatory & patent |  | Tel:  Cell: |  |
| Commercial/business |  | Tel:  Cell: |  |
| General enquiries |  | Tel:  Cell: |  |

**Signature Date**

**Official stamp**

1. Stringent Regulatory Authority <https://extranet.who.int/prequal/sites/default/files/documents/75%20SRA%20clarification_February2017_0.pdf> [↑](#footnote-ref-1)