Application Form for Imported Medicinal Materials Inspection

HS Code:

<table>
<tr><td rowspan="2">Name of Medicinal Material</td><td colspan="4">Chinese Name</td><td colspan="4">English Name</td></tr>
<tr><td colspan="4">Synonym</td><td colspan="4">Latin Name</td></tr>
<tr><td>Packaging Specification</td><td colspan="8"></td></tr>
<tr><td>Approval Document No.</td><td></td><td colspan="2">Contract No./Shipping Mark</td><td></td><td>Test Standards</td><td></td><td>Claim Period</td><td></td></tr>
<tr><td>Quantity of Goods</td><td></td><td>Unit</td><td></td><td>Value</td><td colspan="3"></td><td>Bill of Lading No.</td><td></td></tr>
<tr><td>Port (Place) of Shipment</td><td colspan="2"></td><td colspan="2">Date of Dispatch</td><td></td><td>Means of Transportation (flights/schedules)</td><td>Responsible Customs</td><td></td></tr>
<tr><td>Port (Place) of Arrival</td><td colspan="2"></td><td colspan="2">Date of Arrival</td><td></td><td colspan="3">Storage Location</td></tr>
<tr><td>Exporter</td><td colspan="4"></td><td>Country</td><td colspan="3"></td></tr>
<tr><td>Shipper</td><td colspan="4"></td><td>Country</td><td colspan="3"></td></tr>
<tr><td rowspan="4">Importer (inspection applicant)</td><td>Name</td><td colspan="3"></td><td rowspan="2">Number of Drug Operating License or Production License</td><td colspan="3" rowspan="2"></td></tr>
<tr><td>Social Credit Code</td><td colspan="3"></td></tr>
<tr><td>Address</td><td colspan="4"></td><td colspan="2" rowspan="2">(Official Seal)
DD/MM/YYYY</td></tr>
<tr><td>Contact Person</td><td colspan="2"></td><td>Telephone</td><td colspan="2"></td></tr>
<tr><td>Accompanying Documents</td><td colspan="8"></td></tr>
</table>

(Please pay attention to the "Notes" on the back) National Medical Products Administration

Precautions

1. This form shall be completed by the importer in duplicate. One copy shall be submitted to the port medical products administration for filing, and the other copy shall be submitted to the institute for drug control. The applicant shall affix its official seal in the designated "Official Seal" section.

2.When the imported medicinal material has been re-exported from other countries or regions, all of its purchase contracts, packing lists, bills of lading and shipping invoices of the country or region of origin and of every transit country or region shall be submitted at the same time.

3. "Test Standards" refer to quality specifications and their numbers specified in the *Approval Document for Imported Medicinal Material.*

4. "Quantity of Goods" refers to the total quantity of goods calculated in basic units based on the packaging specifications specified in the *Approval Document for Imported Medicinal Material*, such as the number of bottles, boxes, kilograms, etc.

5. "Storage Location" refers to the specific location where the goods will be stored after customs clearance procedures are completed, and shall indicate name and address.

6. Under the "Attached Documents" section, the numbers of all documents specified in *Approval Document for Imported Medicinal Material* shall be indicated item by item.

7. If space in this inspection application form is insufficient, additional sheets may be attached for supplementary explanations.