Required documents for applying Sample Import Licence for Borderline Products
National Medicines Regulatory Authority

1. Classification form issued by NMRA with the approved formulation

2. Application for a Licence to import a limited Quantity of any Borderline Product(s) for Testing, Examination, Distribution, Sample Analysis or Clinical Trial

3. Copy of Business Registration Certificate

4. Copy of Letter of Authorization from the manufacturer which addressed to Director General, NMRA

5. Catalogue (If a device)

6. Copy of Free Sale Certificate /Copy of COPP/ or a copy of certificate to prove the registration status in country of origin

Application for a License to Import a Limited Quantity of any Borderline Product(s) for Testing, Examination, Distribution as Samples, Analysis or Clinical Trial

M/S.....(Name),of.....(Address) is hereby requesting to import from ..... the Borderline Product specified below for the purpose of testing, examination, distribution as samples, analysis or clinical trials.

Name(s) of Borderline Product(s) / Dosage Form / Type / Model and quantities which may be imported.

Name of the Borderline Device / Borderline Medicine / Borderline Cosmetic	Brand Name	Pack Size	Dosage Form / Type/ Model	Quantity

Signed:.....

Address:.....

Designation of applicant.....