**TO BE PREPARED ON YOUR OFFICIAL LETTERHEAD**

MANUFACTURER’S DECLARATION

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| **These conditions allow for the import of:**1. Polymerase Chain Reaction (PCR) diagnostic test kits.
2. Real-Time PCR or Quantitative PCR (qPCR) diagnostic test kits.
3. Reverse Transcriptase PCR (RT-PCR) diagnostic test kits.
4. Loop-Mediated Isothermal Amplification (LAMP) diagnostic test kits.

**Additional reagents, controls, calibrators etc. may also be imported:**1. when specifically designed for use with diagnostic test kits eligible for import under these conditions.
2. where they can meet all import conditions.
3. whether or not they are imported in the same consignment (separate to the diagnostic test kit) or in a separate consignment.
 |

Date

To Whom It May Concern

I hereby declare that the goods travelling on Air Waybill number: Obtain this number from your courier and include here

On Import Permit # 0005039155 are:

**Nucleic Acid Amplification (NAA) diagnostic test kits** *Commodity**4*

Product name: Enter name of each kit and each reagent, control, calibrator etc. (if imported separately to the kit) must be included on the manufacturer's declaration.

The goods are Nucleic Acid Amplification (NAA) diagnostic test kits only (or individual components specifically designed for use with kits eligible for import under these conditions).

The goods contain nucleic acid up to 1000 nucleotides, enzymes and chemical buffers only.

The goods are commercially manufactured and packaged.

The goods are for *in vitro* use only.

The following end uses will not be performed:

1. Culturing or isolating disease agents.

2. The synthesis of replication-competent disease agents or homologues.

3. Direct or indirect exposure to animals (including laboratory animals) or plants.

I declare all the information contained in this invoice to be certified true and correct.

Yours sincerely

Signature

Name, Position

Organisation