FORM 39-A

[See sub-rule (f) of Rule 150-E]

Report of test or analysis by approved institution for an Individual or Organisation or Procurement agency

(1) Name of individual or organisation or procurement agency from whom sample is received.....

(2) Serial number and date of sender's memorandum.....

(3) Number of samples.....

(4) Date of receipt of the sample....

(5) Name of drug or cosmetics or raw material purporting to be contained in the sample.....

(6) Details of raw material or final product in bulk or final product in finished pack* as obtained by sender:

(a) Name and address of the Manufacturer and Licence number mentioned on the label……

(b) Name of original Manufacturer in the case of raw materials and re-packed drugs.....

(c) Batch number.....

(d) Date of manufacture, if any…..

(e) Date of expiry, if any…….

(7) Results of test or analysis with protocols of test or analysis applied.

In the opinion of the undersigned, the sample referred to above is *of standard quality/is not of standard quality as defined in the Act and the rules made thereunder for the reasons given below.

Date……..

Signature of Person-in-charge of testing

Note-Final product includes repacked material.

*Delete whichever is not applicable.]