PROFORMA FOR REGISTRATION

Remarks against each of the column should be supported by suitable authorized documentation.

1. Name of the Firm.

2. Whether the firm is having registered office/Branch office. Give address & Tel. No. In case more than one manufacturing unit-details all to be inspected. In case marketing a product manufactured by other company. If 'Yes' that also to be inspected.

3. Whether firm is having 5years standing Marketing/Manufacturing of pharmaceutical products.

4. Whether valid drug license for each product exists from Drug controller (in case 'No' names of such product to be given)

5. Audited annual turn over figures for each product for the last 3 years.

6. Whether WHO GMP Certificate for production unit/s is there.

7. Company turn over figure for last 3years

8. R&D facilities with firm. Annual expenditure in R&D in last 3 years

9. Name of Original research products /formulations developed by firm

10. Names of products for which firm is original manufacturer

11. Whether any punitive action has been taken or contemplated by State/Central Institutional/Drug controller. If yes, give details.

Authorised Signatory