



NATIONAL AGENCY FOR FOOD AND DRUG ADMINISTRATION AND CONTROL
NAFDAC
 Federal Ministry of Health
 Abuja, FCT, Nigeria
 Telephone: 0904 200 0000
 Website: www.nafdac.gov.ng

NAFDAC/REGISTRATION/001/2024

15th February 2024

Dear Sir/Madam,

Registration of New Drug

Reference is made to your application for the registration of the drug **Paracetamol Tablets**, **500mg**, manufactured by **ABC Pharmaceuticals Ltd**, Lagos, Nigeria.

- 1. The following information is required for the registration of the drug:
 - a. Certificate of Pharmaceutical Product (COPP) issued by the regulatory authority in the country of origin.
 - b. Certificate of Analysis (COA) for the drug, showing the results of the tests performed on the drug.
 - c. Certificate of Good Manufacturing Practice (CGMP) issued by the regulatory authority in the country of origin.
 - d. Certificate of Registration of the manufacturer issued by the regulatory authority in the country of origin.
 - e. Certificate of Registration of the drug issued by the regulatory authority in the country of origin.
 - f. Certificate of Registration of the drug issued by the regulatory authority in the country of origin.
 - g. Certificate of Registration of the drug issued by the regulatory authority in the country of origin.
 - h. Certificate of Registration of the drug issued by the regulatory authority in the country of origin.
 - i. Certificate of Registration of the drug issued by the regulatory authority in the country of origin.
 - j. Certificate of Registration of the drug issued by the regulatory authority in the country of origin.
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 - o. Certificate of Registration of the drug issued by the regulatory authority in the country of origin.
 - p. Certificate of Registration of the drug issued by the regulatory authority in the country of origin.
 - q. Certificate of Registration of the drug issued by the regulatory authority in the country of origin.
 - r. Certificate of Registration of the drug issued by the regulatory authority in the country of origin.
 - s. Certificate of Registration of the drug issued by the regulatory authority in the country of origin.
 - t. Certificate of Registration of the drug issued by the regulatory authority in the country of origin.
 - u. Certificate of Registration of the drug issued by the regulatory authority in the country of origin.
 - v. Certificate of Registration of the drug issued by the regulatory authority in the country of origin.
 - w. Certificate of Registration of the drug issued by the regulatory authority in the country of origin.
 - x. Certificate of Registration of the drug issued by the regulatory authority in the country of origin.
 - y. Certificate of Registration of the drug issued by the regulatory authority in the country of origin.
 - z. Certificate of Registration of the drug issued by the regulatory authority in the country of origin.

Yours faithfully,
Dr. [Name]
 Director General
 National Agency for Food and Drug Administration and Control