

FDA/CSD/CPE/PRS/25/011

THE FOOD AND DRUGS AUTHORITY (FDA) REVOKES MARKETING AUTHORIZATION FOR OMAMA HERBAL MIXTURE

The Food and Drugs Authority (FDA) in collaboration with the Ghana Police Service carried market surveillance and laboratory analysis of Omama Herbal Mixture.

The laboratory analysis revealed that Omama Herbal Mixture has been adulterated with Diazepam, Metronidazole, Paracetamol and Niacinamide, which are allopathic medicines.

It is important to note that Omama Herbal Mixture is registered as a herbal medicinal product indicated for Malaria and Loss of appetite and is not expected to contain any allopathic medicines.

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The medicines found in Omama Herbal Mixture are dangerous when taken without prescription and/or advise from a qualified health professional.

The public is, therefore, advised <u>not</u> to patronize Omama Herbal Mixture.

Meanwhile, the FDA is working with Omama Herbal Group Limited to recall all products on the market for safe disposal and appropriate regulatory actions including criminal prosecution initiated.

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The FDA wishes to assure the public that it will not relent on its effort of ensuring public health and safety.

The public is advised to give information on persons, pharmacies, and over-the-counter medicine sellers (OTCMS) offering Omama Herbal Mixture for sale to the nearest FDA Offices.

Your Well-being, Our Priority.

Signed

Ag. CHIEF EXECUTIVE OFFICER

Food and Drugs Authority

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