MEDICAL PRODUCT ALERT

DRAP ALERT NO. N° II/S/10-21-06

SUBSTANDARD BLOMOX FORTE SUSPENSION

(MANUFACTURED BY M/S. BLOOM PHARMACEUTICALS (PVT.) LTD. HATTAR)

Date: 21st October, 2021.

Target Audience:

- Healthcare Professionals- Physicians, Pharmacists, and Nurses.
- General Public

Alert Summary:

CDL Karachi has declared the batch No. PF-045 of product "Blomox Forte Suspension" as substandard drug product. Details of the product are given as under:

Sr.	Product Name	Active Ingredient	Mfg. & Exp. date	Batch No.	Test/Analysis result of CDL
1	**Blomox Forte Suspension**	**Amoxicillin**	**Mfg: 03-2021** **Exp: 02-2023**	**PF-045**	**Substandard**

Identification of product:

Off white which reconstitute into off white suspension on mixing with water.

Action to be taken/ Advice for Healthcare Professionals and general public: -

DRAP requests increased vigilance within the supply chains of institutions/pharmacies/healthcare facilities likely to be affected by this **substandard product** batch.

If anyone is in possession of the above stated product batch, please do not use. If anyone has used this **substandard product** batch, or if anyone suffer an adverse reaction/event having used this product batch, it is advised to seek immediate medical advice from a qualified healthcare professional, and to report the incident to Drug Regulatory Authority of Pakistan/National Pharmacovigilance Centre.

All medical products must be obtained from licensed, authentic and reliable sources. Their authenticity and condition should be carefully checked. Seek advice from a healthcare professional in case of any doubt.

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