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January 25, 2020

ISSUES NATIONWIDE
VOLUNTARY PRODUCT RECALL OF SPECIFIC BATCH OF ENALAPRIL MALEATE 10 MG AND 20 MG TABLETS (PRILSEA 10 AND PRILSEA 20)

Purpose of the Press Release:

On August 9, 2019, Seapharma International Inc. initiated a nationwide Product recall of the following products and their pertinent details which were distributed from May 2017-July 2019.

PRODUCT INFO	REG. NO.	BATCH NO.	Mfg. Date	Exp. Date
Enalapril maleate 10 mg Tablet (Prilsea 10)	DRP-6090	ERT-1701	February 2017	January 2020
		ERT-1901	March 2019	February 2022
Enalapril maleate 20 mg Tablet (Prilsea 20)	DR-XY44331	EPFT-1701	February 2017	January 2020
		EPFT-1901	March 2019	February 2022

The products were found to be non-conforming with the required specifications.
It was verified that specific batches did not meet the approved specifications (90.0-110.0%) for Assay.

Consumers who have purchased these products should stop using them and return immediately the said products to the pharmacy from where it was bought. Pharmacies should then return the products to the wholesaler, and the wholesaler to the importer.

Any adverse drug reaction or quality problem experience with the use of this product must reported to FDA. OR may contact Seapharma International, Inc. Below are the contact details:

(Office) +632 8242 7175
Monday to Friday from 9:00 am to 5:00 pm
(Email) Pharmasurrey@yahoo.com

PARMINDER SEKHON
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