

The Board of Pharmacy has received notice of the following product recall:

Description	Lot # / Exp Date	NDC	UPC
THERAFLU COUGH RELIEF 30ML	19N1957675 05/31/21; 19N1793000 03/31/2 01M926004MA 05/31/21; M926003MA 05/31/21; M26002MA 05/31/21; M926001MA 05/31/21; K79C 03/31/21; M926006MA 05/31/21; 19N1958240 04/30/2021; M926007MA 05/31/21; 19N1952247 03/31/21; 19N1936583 03/31/21; 19N1936498 03/31/21; 19N1920630 04/30/21; 19N1895942 03/31/21; 19N1884668 03/31/21; K79B 03/31/21; M926203MB 02/28/21; Y23L05/31/21; XB2W 05/31/21; M926209MB 02/28/21; M926208MB 02/28/21; M926207MB 02/28/21; M926206MB 02/28/21; M926005MA 05/31/21; M926204MB 02/28/21; Y23V 05/31/21; M926202MB 02/28/21; M926202MA 05/31/21; M926201MB 02/28/21; M926201MA 05/31/21; M926009MA 05/31/21; M926008MA 05/31/21; M926205MB 02/28/21; K79A 3/31/2020	00676089030	30067116957

GlaxoSmithKline is recalling the above lots due to the absence of a warning statement “Ask your doctor before us if you have a sodium-restricted diet.” This recall is to the retail level. Affected product started shipping July 2019.

The Board of Pharmacy strongly encourages pharmacies to immediately review their quality assurance and recall policies and procedures to determine if any corrective action is required.