GOVERNMENT OF MAHARASHTRA
FOOD AND DRUGS ADMINISTRATION

Press Note

<u>Public Alert – Stop Use Notice</u>

Stop Use Notice Regarding Coldrif Syrup (Batch No. SR-13) Due to

Toxic Adulteration

Date: 05-10-2025

The Food and Drugs Administration, Maharashtra, has been alerted to

reports of the tragic deaths of children in Madhya Pradesh and Rajasthan, Coldrif

Syrup (Phenylephrine Hydrochloride, Chlorpheniramine Maleate Syrup), Batch

No. SR-13, Mfg. Dt. May-2025, Exp. Dt. April-2027, manufactured by Sresan

Pharma, Sunguvarchathiram, Kancheepuram District, Tamil Nadu, which has been

allegedly adulterated with **Diethylene Glycol (DEG)**, a toxic substance.

In view of this, all the licensee and public hereby instructed to **immediately**

stop sale/distribution/use of Coldrif Syrup, Batch No. SR-13, if anybody is in

possession, and report it to the local Drugs Control Authorities without delay.

The public may also report possession of the said drug directly to the Food

and Drugs Administration, Maharashtra, through the toll-free number 1800222365

or by email to jchq.fda-mah@nic.in or ph- 9892832289

FDA Maharashtra officials are coordinating with the DCA authorities of Tamil

Nadu, where the manufacturer **Sresan Pharma** is located, to track the distribution

of the said product batch to Maharashtra. All Drugs Inspectors and Assistant

Commissioners have been instructed to immediately alert retailers, wholesalers,

and hospitals to freeze any stocks of the said product batch if available in the

market.

The Food and Drugs Administration, Maharashtra is taking necessary

measures in this regard. The public is urged to exercise caution to prevent any

further risk to the life

Date: 05-10-2025

D. R. Gahane
Drug Contoller,

FDA Maharashtra