

PRODUCT MARKET WITHDRAWAL - Similasan Eye Drops 101823

Date of Issue – 2023-10-18

Recall # 101084

UNFI was notified by the manufacturer of a food safety event. Records indicate your location may have received impacted product. Refer to this notice and the supplier letter to identify and remove impacted product from the marketplace. Enter your findings into Recall Infolink.

Impacted UNFI Business Units: Natural & Conventional
Event Classification: Market Withdrawal

Supplier Name: EMERSON HEALTHCARE LLC

Supplier Contact: Jennifer Cameron, cameron@similasanusa.com, 303.539.4060 x108

Product Issue: Voluntary withdraw from Similasan as a result of the FDA warning letter sent out on 9/11/2023. Several concerns that the FDA had about the marketing and manufacturing of homeopathic eye drops and the safety of one of the preservatives used in their manufacture, silver sulfate.

Retail Disposition : Destruction

Brand Name & Description	Pack Size	Case UPC	Unit UPC	Impacted Best by Date(s)/ Lot Codes (MUST provide Best By Date)
Similasan – Styte Eye Relief 10ml	6	60094841300544	094841300542	All Best By Dates
Similasan – Dry Eye Relief 10ml	6	6009484130018	094841300146	
Similasan – Allergy Eye Relief 10ml	6	40094841300243	094841300245	
Similasan – Red Eye Relief 10ml	6	80094841300630	094841300634	
Similasan – Dry Eye Nighttime Gel 10ml	6	60094841300162	094841300160	
Similasan – Kids Allergy Eye Relief 10ml	6	60094841300261	094841300269	
Similasan – Kids Pink Eye Relief	6	60094841300353	094841300351	
Similasan – Pink Eye Nighttime Gel 10ml	6	60094841300360	094841300368	
Similasan – Pink Eye Relief 10ml	6	60094841300346	094841300344	
Similasan – Complete Eye Relief 10ml	6	60094841300605	094841300603	
Similasan – Computer Eye Relief 10ml	6	60094841300476	094841300474	
Similasan – Dry Eye Relief – Single Use 20ct	6	60094841300131	094841300139	
Similasan – Allergy Eye Relief – Single Use 20ct	6	60094841300230	094841300238	

**Cease distribution on the above product. Isolate the product in a secure location
Follow the below instructions to ensure proper disposition of the product.**

1. Remove items listed above from sales floor, displays and back stock.

2. Count the units of impacted product (**only date(s)/lot code(s) listed above**) and record in Recall Infolink.
3. Impacted product must not re-enter commerce or be made available for consumption by any means.
4. **For Credit:** Enter your impacted product count (**only date(s)/lot code(s) listed above**) through Recall Infolink
5. Complete disposition ASAP
6. Inspect inbound product for the next 48 hours to ensure no impacted product is received



September 28, 2023

To Our Valued Partners,

As you know, eight ophthalmic companies, including Similasan, recently received warning letters from the FDA which brought up several concerns that the FDA had about the marketing and manufacturing of homeopathic eye drops and the safety of one of the preservatives used in their manufacture, silver sulfite.

Each company has 15 business days to reply to the FDA to indicate how they intend to respond to their respective letters. Similasan's US leadership team has been in discussions with our retail partners, our manufacturing centers, and our shareholders as we develop an appropriate response.

Pending resolution of the FDA concerns, we have voluntarily suspended the sale and distribution of all Similasan eye drop products to US retailers. We are cooperating with the FDA to quickly resolve this matter.

Our ear drops and other ear relief products are not affected by the FDA's action, and we want to thank you for your continued support of them and of the Similasan brand as we work towards a solution with the FDA. We will continue to update you as we have more clarity.

Sincerely,

Dan Quail

President, North America

Similasan USA