Precautionary batch recall
Maltofer drops, 30 mL up to patient level

In consultation with the Swiss Agency for Therapeutic Products, Swissmedic, Vifor (International) Inc. is recalling the following batches of Maltofer drops 50 mg/1 mL, 30 mL, as a precautionary measure up to the patient level.

This recall is carried out as a precautionary measure because a plastic particle from the dropper was found in a batch of Maltofer drops. Further clarifications have shown that the defect is due to the packaging process. The following batches are affected:

Product	Batch	Expiration date
Maltofer drops 30 mL	AAK34801	06.2024
Maltofer drops 30 mL	AAL56402	09.2024
Maltofer drops 30 mL	AAQ03801	02.2025
Maltofer drops 30 mL	AAW72205	11.2025
Maltofer drops 30 mL	AAR88201	06.2025
Maltofer drops 30 mL	AAW72103	11.2025
Maltofer drops 30 mL	AAZ24904	01.2026
Maltofer drops 30 mL	NAA14003	05.2026
Maltofer drops 30 mL	NAA69701	09.2026
Maltofer drops 30 mL	NAA69701	03.2027

So far, there are no reports of adverse drug reactions that could be related to the deviation. If you use or have a package of Maltofer drops with one of the batch numbers listed above, do not use them anymore and please return them to your place of purchase (pharmacy, doctor's office or drugstore). The packages will be refunded or exchanged.

All other batches delivered in Switzerland as well as other Maltofer products (syrup) are not affected by this recall and can continue to be used. The safety and quality of our medicinal products are our top priority. That in mind, we would like to thank you in advance for your understanding and apologize for any inconvenience you may have experienced.

If you have any questions, please contact our Customer Service department:
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