Haleon – Recall of Robitussin® Honey CF Max products

- On January 24, 2024, Haleon announced a consumer level recall of eight lots of Robitussin Honey CF Max Day Adult and Robitussin Honey CF Max Nighttime Adult to the consumer level due to microbial contamination.
- This recall covers the following lots:

Product Description	Lot number (Exp Date)
Robitussin Honey CF Max Day Adult, 4 oz.	T10810 (10/31/2025)
Robitussin Honey CF Max Day Adult, 8 oz.	T08730 (5/31/2025); T08731 (5/31/2025); T08732 (5/31/2025); T08733 (5/31/2025); T10808 (9/30/2025)
Robitussin Honey CF Max NT Adult, 8 oz.	T08740 (6/30/2026); T08742 (6/30/2026)

- Robitussin Honey CF Max Day and Nighttime are cough syrups indicated for the temporary relief of symptoms occurring with cold or flu, hay fever, or other respiratory allergies.
- In immunocompromised individuals, the use of the affected product could potentially result in severe or life-threatening adverse events such as fungemia or disseminated fungal infection. In non-immunocompromised consumers, the population most likely to use the product, life-threatening infections are not likely to occur. However, the occurrence of an infection that may necessitate medical intervention cannot be completely ruled out.
- To date, Haleon has not received any reports of adverse events related to this recall.
- Consumers who purchased the recalled Robitussin should stop consumption immediately.
- Anyone with the affected products on hand should discontinue use, stop distribution and quarantine product immediately. Patients should contact their healthcare provider if they have experienced any problems that may be related to using the recalled products.
- Consumers may contact the Haleon Consumer Relation team at **1-800-245-1040** or by email at **mystory.us@haleon.com** for more information.