California Device Recall Information Sheet

Food and Drug Branch – Device Recalls

Baxter Healthcare Corporation Recalls Hillrom LikoScale and Q-Link 13 component

Recall Date	Product Description	Recalling Firm	Recall Reason
05/30/25	Hillrom LikoScale component, Product Code 3156232, used with the following adapter kits for patient lifts: 1) LikoScale 200 Accessory - Product Code 3156225; 2) LikoScale 350 Accessory - Product Code 3156228; 3) LikoScale 400 Accessory - Product Code 3156226. Hillrom Q-Link 13 component, Product Code 3156509, used with the following Mobile Patient Lifts: 1) Uno 102 EE Mobile lift - Product Code 2010004; 2) Viking L Mobile lift - Product Code 2040044; 3) Viking XL Mobile lift - Product Code 2040043; 4) Viking M Mobile lift - Product Code 2040045A; 5) Viking S Mobile Lift - Product Code 2040006; 6) Viking XS Mobile Lift - Product Code 2040007; 7) LikoLight Mobile lift - Product Code 2030001. .	Baxter Healthcare Corporation 1 Baxter Pkwy Deerfield, IL 60015-4625 United States	The Q-link 13 could allow for an improper attachment (false latching) of the Quick-Release Hook used on Sling Bars and other accessories. This could result in a critical injury from a patient fall as the false latched component may initially bear weight but can loosen from the Q-link resulting in a detachment and drop. The Q-link 13 could allow for an improper attachment (false latching) of the Quick-Release Hook used on Sling Bars and other accessories. This could result in a critical injury from a patient fall as the false latched component may

			initially bear weight but can loosen from the Q-link resulting in a detachment and drop.

Recall Class	Product Identification	Distribution	Affected Dates
I	All LikoScale Adapter Kits manufactured between 8/27/2013 to 2/27/2025 All Q-link 13 manufactured between 8/27/2013 to 2/27/2025: 1) Product Code 2010004, UDI/DI 00887761981775; 2) Product Code 2040044, UDI/DI 00887761981867; 3) Product Code 2040043, UDI/DI 00887761981850; 4) Product Code 2040045A, UDI/DI 00887761943193; 5) Product Code 2040006, UDI/DI N/A; 6) Product Code 2040007, UDI/DI N/A; 7) Product Code 2030001, UDI/DI N/A. .	Worldwide	May 2025 and prior

For additional information, please visit the FDA Website.

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