11/11/2026

Dear Patient or Caregiver,

We are writing to inform you of an urgent medical device recall from McKesson on behalf of Intersurgical for their **22mm Smoothbore Breathing Systems (circuit).** The notice has been issued because the solvent used to bond the soft connectors onto the tube may have not been enough to secure connection. Reports from markets outside of the U.S. identified instances where the soft connectors disconnected from the tube.

If a connector disconnects from the Smoothbore tube during use, this could result in a loss of ventilation or compromised ventilation. **No adverse events or harm have been reported as a result of this issue.**

Refer to the table for a list of affected item(s) distributed by McKesson Medical-Surgical

MMS #	MFG Catalog #	Description	Affected Lot(s)
1182077	5191000	CIRCUIT, BREATH EXHALATION VALVE W/MONITIR 22MM 1.8M (25/CS)	32520670 32522190 32522802
1279862	5805000	BREATHING SYS, SMTHBORE BILEVEL W/MONIT LINE/SWIVL 22MM (35/	32516315 32517877
1279864	5191850	BREATHING SYS, SMTHBORE EXHALATION VLV W/MONIT LINE 22MM(25/	32521551 32521655
1228711	5183000	CIRCUIT, BREATHING UNIVERSAL ADLT 22MM (20/CS)	32527207
1189083	5804000	CIRCUIT, BOARD SMOOTH BILEVEL ADLT/PED 22MM 1.8M LNG (30/CS)	32508034 32509327 32510909 32516895 32518659 32519175 32525543

Immediate Actions you Should Take

- Inspect all products with matching item codes **(see table above)**. Compare to affected lots to determine if product is affected. If your product lot number does not match the lots in the notification, no further action is required.
- If affected product is identified, discard the product immediately and call PHS for a replacement.

If you have any questions or concerns related to any of this information, or would like a physical copy of this notice, please contact your local branch office.

Sincerely,
Samantha Hoy

National Product Safety Manager
Pediatric Home Service

22mm Smoothbore Breathing Systems Recall Letter