carecentrix

Provider Newsflash
June 2026

FDA Recall – KayserBetten IDA Beds

Purpose of this communication:

We are writing to inform you that KayserBetten has issued a correction for KayserBett IDA pediatric care beds. This correction does not remove the beds from use or sale, but the FDA has identified it as the most serious type of recall because continued use without correction may cause serious injury or death. KayserBetten stated that the U.S. model has a similar risk of injury as a model sold outside the U.S. that was involved in a serious incident resulting in the death of a child.

If the hand control adjustment functions are not locked with the provided key when a child is unattended, there is an increased risk that the patient or other children could operate the hand control and become trapped under the bed frame or between the bed frame and the floor. This can result in serious injury or death.

What do I need to do?

- Please review the FDA recall notice.
- Notify impacted patients to facilitate hand control replacement by the manufacturer according to the guidelines issued by the manufacturer in the FDA notification.
- Confirm if you have any CareCentrix patients affected by this recall and notify the CareCentrix Quality Department by faxing information to: 919-792-6718 Attn: Quality Department.

Thank you in advance for your cooperation and continued partnership.

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