State of California—Health and Human Services Agency

California Department of Public Health

Food and Drug Branch – Device Recalls

CALIFORNIA DEVICE RECALL INFORMATION SHEET

Bard Access Systems Recalls Intraosseous Needle and Driver

Recall Date	Product Description	Recalling Firm	Recall Reason
6/20/2022	**Intraosseous needle and driver** Catalog/Device Name: D015151NK/BD Needle Kit for Powered Driver 15mm x 15Ga, D015151MK/ BD Manual Driver Needle Kit 15mm x 15Ga, D015251NK/ BD Needle Kit for Powered Driver 25mm x 15Ga, D015251MK/ BD Manual Driver Needle Kit 25mm x 15Ga, D015351NK/ BD Needle Kit for Powered Driver 35mm x 15Ga, D015351MK/ BD Manual Driver Needle Kit 35mm x 15Ga, D015451NK/ BD Needle Kit for Powered Driver 45mm x 15Ga, D015451MK/ …	**Bard Access Systems, Inc.** Salt Lake City, Utah	BD Intraosseous Needle Kits may exhibit: 1) Increased force required to remove the stylet from intraosseous needle during placement may cause removal of the entire needle assembly and loss of intraosseous access; 2) Stylet safety mechanism may not engage as the stylet is removed; 3) Metal discs in the powered driver may stick rendering the driver unusable.

Recall Class	Product Identification	Distribution	Affected Dates
I	Catalog/UDI/Lot: D015151NK/801741163586/, 121460 - D015551MK/80174 1163661/126834; D001001/008 01741163579/All Lots	Nationwide	June 2022 and prior

FOR ADDITIONAL INFORMATION, PLEASE VISIT THE FDA WEBSITE

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