

K203193 BD Intraosseous Infusion SystemMar 4, 2021
127 days to decisionK203193 · Product code: **MHC** · General Hospital
Source: <https://www.510kdatabase.net/k203193/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Interosseous (MHC)
Date received	Oct 28, 2020
Decision date	Mar 4, 2021
Days to decision	127 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Bard Access Systems, Inc(Bas)[Wholly-Owned Subsidiary of Bd]
Location	Salt Lake Ciy,, UT, US
Contact	Connor Dahl
510(k) history	1 submissions · 1 cleared · 2021-2021

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k203193/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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