

**K790953 BORDISPO BRAND HYPODERMIC NEEDLE**Jun 27, 1979  
36 days to decisionK790953 · Product code: **FMI** · General HospitalSource: <https://www.510kdatabase.net/k790953/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	May 22, 1979
Decision date	Jun 27, 1979
Days to decision	36 days
Third-party review	No

**APPLICANT**

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Company	<b>Borda Products, Inc.</b>
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1979-1979

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k790953/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated July 10, 2026