

K791462 HYPERDERMIC NEEDLE WITH SENSOR #514Sep 24, 1979
70 days to decisionK791462 · Product code: **FMI** · General HospitalSource: <https://www.510kdatabase.net/k791462/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	Jul 16, 1979
Decision date	Sep 24, 1979
Days to decision	70 days
Third-party review	No

APPLICANT

Company	Medtech Intl., Inc.
Location	Mchenry, IL, US
510(k) history	6 submissions · 6 cleared · 1978-1992

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Device record: <https://www.510kdatabase.net/k791462/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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