

**K823693 MYELOGRAPHY NEEDLE**Feb 7, 1983  
61 days to decisionK823693 · Product code: **FMI** · General HospitalSource: <https://www.510kdatabase.net/k823693/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	Dec 8, 1982
Decision date	Feb 7, 1983
Days to decision	61 days
Third-party review	No

**APPLICANT**

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Company	<b>Ocean Medical Products, Ltd.</b>
Location	Mchenry, IL, US
510(k) history	9 submissions · 9 cleared · 1981-1988

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k823693/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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