

K811966 VISITEC NEEDLE RANGE (GENERAL HOSPITAL)Jul 23, 1981
13 days to decisionK811966 · Product code: **FMI** · General HospitalSource: <https://www.510kdatabase.net/k811966/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	Jul 10, 1981
Decision date	Jul 23, 1981
Days to decision	13 days
Third-party review	No

APPLICANT

Company	Visitec Co.
Location	Walker, MI, US
510(k) history	49 submissions · 49 cleared · 1979-1995

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Device record: <https://www.510kdatabase.net/k811966/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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