

K811931 NEEDLE GUIDE ATTACHMENTAug 18, 1981
43 days to decisionK811931 · Product code: **IYO** · Radiology
Source: <https://www.510kdatabase.net/k811931/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Imaging, Pulsed Echo, Ultrasonic (IYO)
Date received	Jul 6, 1981
Decision date	Aug 18, 1981
Days to decision	43 days
Third-party review	No

APPLICANT

Company	Technicare Corp.
Location	Mchenry, IL, US
510(k) history	32 submissions · 32 cleared · 1980-1985

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Device record: <https://www.510kdatabase.net/k811931/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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