

K803157 AUTO-MATE CAMERAFeb 2, 1981
60 days to decisionK803157 · Product code: **IYO** · Radiology
Source: <https://www.510kdatabase.net/k803157/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Imaging, Pulsed Echo, Ultrasonic (IYO)
Date received	Dec 4, 1980
Decision date	Feb 2, 1981
Days to decision	60 days
Third-party review	No

APPLICANT

Company	Aptek , Ltd.
Location	Mchenry, IL, US
510(k) history	2 submissions · 2 cleared · 1980-1981

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Device record: <https://www.510kdatabase.net/k803157/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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