FDA 510(k) Premarket Notification

K803157 AUTO-MATE CAMERA

CLEARED

Feb 2, 1981
60 days to decision

K803157 · 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