FDA 510(k) Premarket Notification

K802917 MARK III ULTRASONIC SCANNING SYSTEM

CLEARED

Jan 16, 1981
58 days to decision

K802917 · Product code: **DXK** · Cardiovascular
Source: https://www.510kdatabase.net/k802917/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Echocardiograph (DXK)
Date received	Nov 19, 1980
Decision date	Jan 16, 1981
Days to decision	58 days
Third-party review	No

APPLICANT

Company	**Advanced Technology Laboratories, Inc.**
Location	Mchenry, IL, US
510(k) history	61 submissions · 59 cleared · 1980-2002