FDA 510(k) Premarket Notification

K031018 AUTOGEL PAD

CLEARED

Dec 8, 2003
252 days to decision

K031018 · Product code: **MUI** · Radiology
Source: https://www.510kdatabase.net/k031018/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Media, Coupling, Ultrasound (MUI)
Date received	Mar 31, 2003
Decision date	Dec 8, 2003
Days to decision	252 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Rich-Mar Corp.**
Location	Mchenry, IL, US
Contact	DAVID RICHARDS
510(k) history	43 submissions · 41 cleared · 1976-2004