FDA 510(k) Premarket Notification

K031894 SCANTEC PAD

CLEARED

Jul 18, 2003
29 days to decision

K031894 · Product code: **MUI** · Radiology
Source: https://www.510kdatabase.net/k031894/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Media, Coupling, Ultrasound (MUI)
Date received	Jun 19, 2003
Decision date	Jul 18, 2003
Days to decision	29 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Sonotech, Inc.**
Location	Bellingham, WA, US
Contact	MARGARET J LARSON
510(k) history	14 submissions · 14 cleared · 1995-2004