FDA 510(k) Premarket Notification

CLEARED

K020956 VIVOSONIC DIANOSTIC ULTRASOUND IMAGING COUPLING MEDIA

May 16, 2002
52 days to decision

K020956 · Product code: **MUI** · Radiology
Source: https://www.510kdatabase.net/k020956/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Media, Coupling, Ultrasound (MUI)
Date received	Mar 25, 2002
Decision date	May 16, 2002
Days to decision	52 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Microtek Medical, Inc.**
Location	Mchenry, IL, US
Contact	THOMAS BONNER
510(k) history	24 submissions · 24 cleared · 1984-2011