FDA 510(k) Premarket Notification

K021497 CYPRESS ULTRASOUND SYSTEM

CLEARED

Jul 9, 2002
61 days to decision

K021497 · Product code: **IYN** · Radiology
Source: https://www.510kdatabase.net/k021497/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	System, Imaging, Pulsed Doppler, Ultrasonic (IYN)
Date received	May 9, 2002
Decision date	Jul 9, 2002
Days to decision	61 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Acuson Corp.**
Location	Mountain View, CA, US
Contact	BOB LEIKER
510(k) history	37 submissions · 37 cleared · 1988-2002