FDA 510(k) Premarket Notification

K921418 ULTRA-CORE BIOPSY NEEDLES

CLEARED

Jun 19, 1992
87 days to decision

K921418 · Product code: **DWO** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k921418/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Biopsy, Cardiovascular (DWO)
Date received	Mar 24, 1992
Decision date	Jun 19, 1992
Days to decision	87 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Medical Device Technologies, Inc.**
Location	Gainsville, FL, US
Contact	PAUL BAKER
510(k) history	46 submissions · 46 cleared · 1992-2010