FDA 510(k) Premarket Notification

K923872 BONE MARROW BIOPSY SYSTEM

CLEARED

Dec 15, 1992
134 days to decision

K923872 · Product code: **GDM** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k923872/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Aspiration And Injection, Reusable (GDM)
Date received	Aug 3, 1992
Decision date	Dec 15, 1992
Days to decision	134 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**The Straumann Co.**
Location	Cambridge, MA, US
Contact	CAROLYN BITETTI
510(k) history	14 submissions · 14 cleared · 1992-2004