FDA 510(k) Premarket Notification

K890722 ADG BONE BIOPSY SYSTEM

CLEARED

Sep 21, 1989
220 days to decision

K890722 · Product code: **GDM** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k890722/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Aspiration And Injection, Reusable (GDM)
Date received	Feb 13, 1989
Decision date	Sep 21, 1989
Days to decision	220 days
Third-party review	No

APPLICANT

Company	**American Design Group, Inc.**
Location	Martinez, CA, US
Contact	DAVID W SCHLERF
510(k) history	3 submissions · 3 cleared · 1989-1991