FDA 510(k) Premarket Notification

K841299 BIERMAN BIOPSY NEEDLE

CLEARED

May 2, 1984
33 days to decision

K841299 · Product code: **GDM** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k841299/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Aspiration And Injection, Reusable (GDM)
Date received	Mar 30, 1984
Decision date	May 2, 1984
Days to decision	33 days
Third-party review	No

APPLICANT

Company	**Popper & Sons, Inc.**
Location	Mchenry, IL, US
510(k) history	8 submissions · 8 cleared · 1984-1991