FDA 510(k) Premarket Notification

K896019 MODIFIED DISP. CHIBA-TYPE NEEDLE DESIGNATED ASPI.

CLEARED

Jan 3, 1990
79 days to decision

K896019 · Product code: **GAA** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k896019/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Aspiration And Injection, Disposable (GAA)
Date received	Oct 16, 1989
Decision date	Jan 3, 1990
Days to decision	79 days
Third-party review	No

APPLICANT

Company	**Dlp, Inc.**
Location	Mchenry, IL, US
Contact	CHET VANHOF
510(k) history	56 submissions · 56 cleared · 1979-1997