FDA 510(k) Premarket Notification

K802458 KEVORKIAN-YOUNGE BIOPSY CURETTE #90-6611

CLEARED

Jan 12, 1981
96 days to decision

K802458 · Product code: **HFB** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k802458/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Forceps, Biopsy, Gynecological (HFB)
Date received	Oct 8, 1980
Decision date	Jan 12, 1981
Days to decision	96 days
Third-party review	No

APPLICANT

Company	**J. Sklar Mfg. Co., Inc.**
Location	Mchenry, IL, US
510(k) history	18 submissions · 18 cleared · 1977-1984