FDA 510(k) Premarket Notification

K772397 COTTLE COLUMELLA FORCEP

CLEARED

Jan 17, 1978
21 days to decision

K772397 · Product code: **KAE** · Ear, Nose, Throat
Source: https://www.510kdatabase.net/k772397/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Forceps, Ent (KAE)
Date received	Dec 27, 1977
Decision date	Jan 17, 1978
Days to decision	21 days
Third-party review	No

APPLICANT

Company	**Edward Weck, Inc.**
Location	Mchenry, IL, US
510(k) history	140 submissions · 140 cleared · 1976-1992