

K772397 COTTLE COLUMELLA FORCEPJan 17, 1978
21 days to decisionK772397 · Product code: **KAE** · Ear, Nose, ThroatSource: <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

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k772397/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 23, 2026