FDA 510(k) Premarket Notification

K792428 COE-SIL

CLEARED

Jan 23, 1980
57 days to decision

K792428 · Product code: **LDG** · Ear, Nose, Throat
Source: https://www.510kdatabase.net/k792428/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Kit, Earmold, Impression (LDG)
Date received	Nov 27, 1979
Decision date	Jan 23, 1980
Days to decision	57 days
Third-party review	No

APPLICANT

Company	**Coe Laboratories, Inc.**
Location	Mchenry, IL, US
510(k) history	20 submissions · 20 cleared · 1976-1990