FDA 510(k) Premarket Notification

K882659 AUGUSTINE INTUBATION GUIDE(TM)

CLEARED

Jul 21, 1988
23 days to decision

K882659 · Product code: **CCW** · Anesthesiology
Source: https://www.510kdatabase.net/k882659/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Laryngoscope, Rigid (CCW)
Date received	Jun 28, 1988
Decision date	Jul 21, 1988
Days to decision	23 days
Third-party review	No

APPLICANT

Company	**Augustine Medical, Inc.**
Location	Eden Prairie, MN, US
Contact	D AUGUSTINE
510(k) history	25 submissions · 24 cleared · 1987-2002