FDA 510(k) Premarket Notification

K922537 DURALIFE

CLEARED

Dec 4, 1992
190 days to decision

K922537 · Product code: **BZA** · Anesthesiology
Source: https://www.510kdatabase.net/k922537/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Connector, Airway (extension) (BZA)
Date received	May 28, 1992
Decision date	Dec 4, 1992
Days to decision	190 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Dhd Diemolding Healthcare Div.**
Location	Canastota, NY, US
Contact	JEAN WALLACE
510(k) history	11 submissions · 11 cleared · 1991-1998