FDA 510(k) Premarket Notification

K800023 LUNG AEROSOL UNIT

CLEARED

Jan 1, 1985
1821 days to decision

K800023 · Product code: **CAF** · General Hospital
Source: https://www.510kdatabase.net/k800023/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Nebulizer (direct Patient Interface) (CAF)
Date received	Jan 7, 1980
Decision date	Jan 1, 1985
Days to decision	1821 days
Third-party review	No

APPLICANT

Company	**Cadema Medical Products, Inc.**
Location	Walker, MI, US
510(k) history	6 submissions · 6 cleared · 1985-1990