FDA 510(k) Premarket Notification

K902141 ANOLIFT - PORTABLE & ANOLIFT - GANTRY

CLEARED

Sep 12, 1990
120 days to decision

K902141 · Product code: **FSA** · General Hospital
Source: https://www.510kdatabase.net/k902141/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Lift, Patient, Non-ac-powered (FSA)
Date received	May 15, 1990
Decision date	Sep 12, 1990
Days to decision	120 days
Third-party review	No

APPLICANT

Company	**Anodyne Corp.**
Location	Brighton, MI, US
Contact	DONALD D MERRY
510(k) history	1 submissions · 1 cleared · 1990-1990