FDA 510(k) Premarket Notification

K900735 MODEL 130 AUTOMATIC HEIGHT ADJUST (AHA) INCUBATOR

CLEARED

May 2, 1990
75 days to decision

K900735 · Product code: **FMZ** · General Hospital
Source: https://www.510kdatabase.net/k900735/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Incubator, Neonatal (FMZ)
Date received	Feb 16, 1990
Decision date	May 2, 1990
Days to decision	75 days
Third-party review	No

APPLICANT

Company	**Preemicare Corp.**
Location	Houston, TX, US
Contact	CHARLES S BOYD
510(k) history	9 submissions · 9 cleared · 1988-1991