

**K900735 MODEL 130 AUTOMATIC HEIGHT ADJUST (AHA)
INCUBATOR**May 2, 1990
75 days to decisionK900735 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k900735/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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