

K840526 AMETHERMApr 30, 1984
83 days to decisionK840526 · Product code: **FMZ** · General Hospital
Source: <https://www.510kdatabase.net/k840526/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Incubator, Neonatal (FMZ)
Date received	Feb 7, 1984
Decision date	Apr 30, 1984
Days to decision	83 days
Third-party review	No

APPLICANT

Company	Egnell, Inc.
Location	Mchenry, IL, US
510(k) history	10 submissions · 8 cleared · 1981-1985

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k840526/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 30, 2026