

K834129 MEDELA APGAR TIMERJan 10, 1984
41 days to decisionK834129 · Product code: **LHB** · General Hospital
Source: <https://www.510kdatabase.net/k834129/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Timer, Apgar (LHB)
Date received	Nov 30, 1983
Decision date	Jan 10, 1984
Days to decision	41 days
Third-party review	No

APPLICANT

Company	Medela, Inc.
Location	Mchenry, IL, US
510(k) history	19 submissions · 19 cleared · 1980-2017

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k834129/> 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 1, 2026