FDA 510(k) Premarket Notification

K834129 MEDELA APGAR TIMER

CLEARED

Jan 10, 1984
41 days to decision

K834129 · 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