FDA 510(k) Premarket Notification

K812233 RATEMINDER

CLEARED

Aug 25, 1981
14 days to decision

K812233 · Product code: **LDR** · General Hospital
Source: https://www.510kdatabase.net/k812233/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Controller, Infusion, Intravascular, Electronic (LDR)
Date received	Aug 11, 1981
Decision date	Aug 25, 1981
Days to decision	14 days
Third-party review	No

APPLICANT

Company	**Anatros Corp.**
Location	Mchenry, IL, US
510(k) history	2 submissions · 2 cleared · 1981-1983