

K812233 RATEMINDERAug 25, 1981
14 days to decisionK812233 · 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

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k812233/> 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 July 10, 2026