FDA 510(k) Premarket Notification

K842861 PRO-PAR CLEARANT

CLEARED

Aug 21, 1984
29 days to decision

K842861 · Product code: **KEM** · Pathology
Source: https://www.510kdatabase.net/k842861/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Agent, Clearing (KEM)
Date received	Jul 23, 1984
Decision date	Aug 21, 1984
Days to decision	29 days
Third-party review	No

APPLICANT

Company	**Anatech, Ltd.**
Location	Mchenry, IL, US
510(k) history	14 submissions · 14 cleared · 1984-1988