

K842861 PRO-PAR CLEARANTAug 21, 1984
29 days to decisionK842861 · 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

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k842861/> 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 May 14, 2026