

K960066 IMMULITE ACTHFeb 16, 1996
42 days to decisionK960066 · Product code: **CKG** · Chemistry
Source: <https://www.510kdatabase.net/k960066/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Radioimmunoassay, Acth (CKG)
Date received	Jan 5, 1996
Decision date	Feb 16, 1996
Days to decision	42 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Diagnostic Products Corp.
Location	Mchenry, IL, US
Contact	EDWARD M LEVINE, PHD.
510(k) history	321 submissions · 321 cleared · 1976-2006

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k960066/> 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 June 20, 2026