

K840018 DPC ACTH KITMar 23, 1984
78 days to decisionK840018 · Product code: **CKG** · Chemistry
Source: <https://www.510kdatabase.net/k840018/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Radioimmunoassay, Acth (CKG)
Date received	Jan 5, 1984
Decision date	Mar 23, 1984
Days to decision	78 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k840018/> 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 21, 2026