

K833849 ACTH RIAJan 30, 1984
87 days to decisionK833849 · Product code: **CKG** · Chemistry
Source: <https://www.510kdatabase.net/k833849/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Radioimmunoassay, Acth (CKG)
Date received	Nov 4, 1983
Decision date	Jan 30, 1984
Days to decision	87 days
Third-party review	No

APPLICANT

Company	Syncor Intl. Corp.
Location	Mchenry, IL, US
510(k) history	31 submissions · 31 cleared · 1983-1995

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Device record: <https://www.510kdatabase.net/k833849/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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