FDA 510(k) Premarket Notification

K840018 DPC ACTH KIT

CLEARED

Mar 23, 1984
78 days to decision

K840018 · 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