FDA 510(k) Premarket Notification

K960066 IMMULITE ACTH

CLEARED

Feb 16, 1996
42 days to decision

K960066 · 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