FDA 510(k) Premarket Notification

K882232 DISCOVER(R)

CLEARED

Oct 25, 1988
151 days to decision

K882232 · Product code: **LCX** · Chemistry
Source: https://www.510kdatabase.net/k882232/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Kit, Test, Pregnancy, Hcg, Over The Counter (LCX)
Date received	May 27, 1988
Decision date	Oct 25, 1988
Days to decision	151 days
Third-party review	No

APPLICANT

Company	**Carter Products**
Location	Cranbury, NJ, US
Contact	STEPHEN KOLAKOWSKY
510(k) history	20 submissions · 20 cleared · 1988-1996