FDA 510(k) Premarket Notification

K780240 PREDICTOR

CLEARED

Feb 27, 1978
14 days to decision

K780240 · Product code: **LCX** · Chemistry
Source: https://www.510kdatabase.net/k780240/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Kit, Test, Pregnancy, Hcg, Over The Counter (LCX)
Date received	Feb 13, 1978
Decision date	Feb 27, 1978
Days to decision	14 days
Third-party review	No

APPLICANT

Company	**Organon, Inc.**
Location	Mchenry, IL, US
510(k) history	41 submissions · 41 cleared · 1977-1995