FDA 510(k) Premarket Notification

K802999 BETA-TEC QUALITATIVE

CLEARED

Dec 22, 1980
27 days to decision

K802999 · Product code: **JHI** · Chemistry
Source: https://www.510kdatabase.net/k802999/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Visual, Pregnancy Hcg, Prescription Use (JHI)
Date received	Nov 25, 1980
Decision date	Dec 22, 1980
Days to decision	27 days
Third-party review	No

APPLICANT

Company	**Armkel, LLC**
Location	Mchenry, IL, US
510(k) history	68 submissions · 68 cleared · 1979-2004