FDA 510(k) Premarket Notification

K810232 LEECO P.A.P. - QUANT DIAGNOSTIC KIT

CLEARED

Feb 10, 1981
14 days to decision

K810232 · Product code: **JFH** · Chemistry
Source: https://www.510kdatabase.net/k810232/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Acid Phosphatase (prostatic), Tartrate Inhibited (JFH)
Date received	Jan 27, 1981
Decision date	Feb 10, 1981
Days to decision	14 days
Third-party review	No

APPLICANT

Company	**Leeco Diagnostics, Inc.**
Location	Mchenry, IL, US
510(k) history	49 submissions · 49 cleared · 1979-1989