FDA 510(k) Premarket Notification

K840174 LH QUIK RIA KIT

CLEARED

Mar 19, 1984
62 days to decision

K840174 · Product code: **CEP** · Chemistry
Source: https://www.510kdatabase.net/k840174/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Radioimmunoassay, Luteinizing Hormone (CEP)
Date received	Jan 17, 1984
Decision date	Mar 19, 1984
Days to decision	62 days
Third-party review	No

APPLICANT

Company	**Pacific Biotech, Inc.**
Location	Mchenry, IL, US
510(k) history	29 submissions · 29 cleared · 1983-1993