FDA 510(k) Premarket Notification

K810662 LEECO LH-QUANT DIAGNOSTIC KIT

CLEARED

Mar 31, 1981
20 days to decision

K810662 · Product code: **CEP** · Chemistry
Source: https://www.510kdatabase.net/k810662/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Radioimmunoassay, Luteinizing Hormone (CEP)
Date received	Mar 11, 1981
Decision date	Mar 31, 1981
Days to decision	20 days
Third-party review	No

APPLICANT

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