FDA 510(k) Premarket Notification

K782159 DIAGNOSTIC KIT, FERRITAB-FE-59

CLEARED

Feb 1, 1979
41 days to decision

K782159 · Product code: **JJA** · Chemistry
Source: https://www.510kdatabase.net/k782159/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Radio-labeled Iron Method, Iron (non-heme) (JJA)
Date received	Dec 22, 1978
Decision date	Feb 1, 1979
Days to decision	41 days
Third-party review	No

APPLICANT

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