FDA 510(k) Premarket Notification

K792032 LEIF CENTRIFUGAL CYTOLOGY BUCKET

CLEARED

Nov 30, 1979
51 days to decision

K792032 · Product code: **IFB** · Pathology
Source: https://www.510kdatabase.net/k792032/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cytocentrifuge (IFB)
Date received	Oct 10, 1979
Decision date	Nov 30, 1979
Days to decision	51 days
Third-party review	No

APPLICANT

Company	**Robert C. Leif, Ph.D.**
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1979-1979