FDA 510(k) Premarket Notification

K831958 RADIAL IMMUNO-TEST FOR HUMAN C-REACTI

CLEARED

Jul 19, 1983
32 days to decision

K831958 · Product code: **DCK** · Chemistry
Source: https://www.510kdatabase.net/k831958/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	C-reactive Protein, Antigen, Antiserum, And Control (DCK)
Date received	Jun 17, 1983
Decision date	Jul 19, 1983
Days to decision	32 days
Third-party review	No

APPLICANT

Company	**Kent Laboratories, Inc.**
Location	Mchenry, IL, US
510(k) history	40 submissions · 37 cleared · 1977-1995