FDA 510(k) Premarket Notification

K946050 LIQUISPEX(TM) LIQUID CRP CONTROL LEVEL 1 AND 2

CLEARED

Feb 10, 1995
60 days to decision

K946050 · Product code: **DCK** · Immunology
Source: https://www.510kdatabase.net/k946050/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	C-reactive Protein, Antigen, Antiserum, And Control (DCK)
Date received	Dec 12, 1994
Decision date	Feb 10, 1995
Days to decision	60 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Clinical Controls, Inc.**
Location	Grover Beach, CA, US
Contact	JAMES F GODREY
510(k) history	17 submissions · 17 cleared · 1994-1996