FDA 510(k) Premarket Notification

K822500 CDL LYSING REAGENT

CLEARED

Sep 17, 1982
30 days to decision

K822500 · Product code: **GGK** · Hematology
Source: https://www.510kdatabase.net/k822500/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Products, Red-cell Lysing Products (GGK)
Date received	Aug 18, 1982
Decision date	Sep 17, 1982
Days to decision	30 days
Third-party review	No

APPLICANT

Company	**Connecticut Diagnostics, Ltd.**
Location	Mchenry, IL, US
510(k) history	14 submissions · 14 cleared · 1979-1991