

K822500 CDL LYSING REAGENTSep 17, 1982
30 days to decisionK822500 · 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

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Device record: <https://www.510kdatabase.net/k822500/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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