

K801898 URIC ACID DIAGNOSTIC REAGENTOct 10, 1980
63 days to decisionK801898 · Product code: **KNK** · Chemistry
Source: <https://www.510kdatabase.net/k801898/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Acid, Uric, Uricase (colorimetric) (KNK)
Date received	Aug 8, 1980
Decision date	Oct 10, 1980
Days to decision	63 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/k801898/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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