FDA 510(k) Premarket Notification

K801898 URIC ACID DIAGNOSTIC REAGENT

CLEARED

Oct 10, 1980
63 days to decision

K801898 · 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