FDA 510(k) Premarket Notification

K810136 URIC ACID REAGENT SYSTEM

CLEARED

Feb 2, 1981
12 days to decision

K810136 · Product code: **KNK** · Chemistry
Source: https://www.510kdatabase.net/k810136/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Acid, Uric, Uricase (colorimetric) (KNK)
Date received	Jan 21, 1981
Decision date	Feb 2, 1981
Days to decision	12 days
Third-party review	No

APPLICANT

Company	**American Monitor Corp.**
Location	Mchenry, IL, US
510(k) history	73 submissions · 73 cleared · 1977-1990