

K810136 URIC ACID REAGENT SYSTEMFeb 2, 1981
12 days to decisionK810136 · 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

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k810136/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 29, 2026