

K810355 ALT (UV) TESTFeb 23, 1981
13 days to decisionK810355 · Product code: **CKA** · Chemistry
Source: <https://www.510kdatabase.net/k810355/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Nadh Oxidation/nad Reduction, Alt/sgpt (CKA)
Date received	Feb 10, 1981
Decision date	Feb 23, 1981
Days to decision	13 days
Third-party review	No

APPLICANT

Company	Boehringer Mannheim Corp.
Location	Mchenry, IL, US
510(k) history	340 submissions · 340 cleared · 1976-1999

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Device record: <https://www.510kdatabase.net/k810355/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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