

K803110 PROTEOLYTIC ENZYME CONTROLFeb 17, 1981
69 days to decisionK803110 · Product code: **LGP** · Chemistry
Source: <https://www.510kdatabase.net/k803110/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Assay, Alpha-2-antiplasmin (LGP)
Date received	Dec 10, 1980
Decision date	Feb 17, 1981
Days to decision	69 days
Third-party review	No

APPLICANT

Company	American Dade
Location	Mchenry, IL, US
510(k) history	149 submissions · 149 cleared · 1980-1987

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Device record: <https://www.510kdatabase.net/k803110/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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