

K850410 GEN. DIAG. CHROMOSTRATE PLASMINOGEN ASSAYMay 8, 1985
93 days to decisionK850410 · Product code: **DDX** · Immunology
Source: <https://www.510kdatabase.net/k850410/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Plasminogen, Antigen, Antiserum, Control (DDX)
Date received	Feb 4, 1985
Decision date	May 8, 1985
Days to decision	93 days
Third-party review	No

APPLICANT

Company	Warner-Lambert Co.
Location	Mchenry, IL, US
Contact	LYNN DECARO
510(k) history	50 submissions · 50 cleared · 1979-2003

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k850410/>, 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 May 14, 2026