

K802610 ALERT A,HDL-APOLIPOROTEIN A, TEST KITJan 5, 1981
76 days to decisionK802610 · Product code: **DER** · Immunology
Source: <https://www.510kdatabase.net/k802610/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Alpha-1-lipoprotein, Antigen, Antiserum, Control (DER)
Date received	Oct 21, 1980
Decision date	Jan 5, 1981
Days to decision	76 days
Third-party review	No

APPLICANT

Company	Hyland Therapeutic Div., Travenol Laboratories
Location	Mchenry, IL, US
510(k) history	44 submissions · 44 cleared · 1976-1982

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k802610/> 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