FDA 510(k) Premarket Notification

K802610 ALERT A,HDL-APOLIPOROTEIN A, TEST KIT

CLEARED

Jan 5, 1981
76 days to decision

K802610 · 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