

K760917 LAS-R HUMAN HAPTOGLOBIN TESTDec 30, 1976
63 days to decisionK760917 · Product code: **DFT** · Pathology
Source: <https://www.510kdatabase.net/k760917/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Igm, Antigen, Antiserum, Control (DFT)
Date received	Oct 28, 1976
Decision date	Dec 30, 1976
Days to decision	63 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

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Device record: <https://www.510kdatabase.net/k760917/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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