

K862641 U.S. DIAGNOSTIC INC. HEMOGLOBIN TEST (DRY PACK)

Aug 21, 1986
41 days to decision

K862641 · Product code: **KHG** · Hematology
Source: <https://www.510kdatabase.net/k862641/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Whole Blood Hemoglobin Determination (KHG)
Date received	Jul 11, 1986
Decision date	Aug 21, 1986
Days to decision	41 days
Third-party review	No

APPLICANT

Company	U. S. Diagnostics, Inc.
Location	San Luis Obispo, CA, US
Contact	ROY E SPECK
510(k) history	22 submissions · 22 cleared · 1986-2011

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Device record: <https://www.510kdatabase.net/k862641/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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