

K830340 FITKIT I CHECKUP KITAug 16, 1983
195 days to decisionK830340 · Product code: **CDM** · Hematology
Source: <https://www.510kdatabase.net/k830340/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Diazonium Colorimetry, Urobilinogen (urinary, Non-quant.) (CDM)
Date received	Feb 2, 1983
Decision date	Aug 16, 1983
Days to decision	195 days
Third-party review	No

APPLICANT

Company	Boditestor
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1983-1983

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Device record: <https://www.510kdatabase.net/k830340/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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