

K811722 PARA-PAKJul 15, 1981
27 days to decisionK811722 · Product code: **HYI** · Toxicology
Source: <https://www.510kdatabase.net/k811722/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Iodine, Grams (HYI)
Date received	Jun 18, 1981
Decision date	Jul 15, 1981
Days to decision	27 days
Third-party review	No

APPLICANT

Company	Meridian Diagnostics, Inc.
Location	Walker, MI, US
510(k) history	92 submissions · 92 cleared · 1980-1999

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Device record: <https://www.510kdatabase.net/k811722/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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