

K792546 FETAL CHECKDec 18, 1979
8 days to decisionK792546 · Product code: **KJR** · Pathology
Source: <https://www.510kdatabase.net/k792546/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Erythrosin B (KJR)
Date received	Dec 10, 1979
Decision date	Dec 18, 1979
Days to decision	8 days
Third-party review	No

APPLICANT

Company	Alta Diagnostics, Inc.
Location	Mchenry, IL, US
510(k) history	5 submissions · 5 cleared · 1979-1980

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k792546/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 14, 2026