

K812970 S/P CAPILLARY TUBES, HEPARINIZED (3A)Nov 6, 1981
15 days to decisionK812970 · Product code: **GIO** · Hematology
Source: <https://www.510kdatabase.net/k812970/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tube, Collection, Capillary Blood (GIO)
Date received	Oct 22, 1981
Decision date	Nov 6, 1981
Days to decision	15 days
Third-party review	No

APPLICANT

Company	American Dade
Location	Mchenry, IL, US
510(k) history	149 submissions · 149 cleared · 1980-1987

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Device record: <https://www.510kdatabase.net/k812970/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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