

K800003 CORTEX DOUBLE HELIX SAMPLERFeb 1, 1980
30 days to decisionK800003 · Product code: **GLE** · Hematology
Source: <https://www.510kdatabase.net/k800003/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Mixer, Blood Tube (GLE)
Date received	Jan 2, 1980
Decision date	Feb 1, 1980
Days to decision	30 days
Third-party review	No

APPLICANT

Company	Cortex Research Corp.
Location	Walker, MI, US
510(k) history	1 submissions · 1 cleared · 1980-1980

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Device record: <https://www.510kdatabase.net/k800003/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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