FDA 510(k) Premarket Notification

K800003 CORTEX DOUBLE HELIX SAMPLER

CLEARED

Feb 1, 1980
30 days to decision

K800003 · 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