FDA 510(k) Premarket Notification

K790310 HEMOGLOBIN CONTROLS

CLEARED

Mar 21, 1979
36 days to decision

K790310 · Product code: **GGM** · Hematology
Source: https://www.510kdatabase.net/k790310/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Control, Hemoglobin (GGM)
Date received	Feb 13, 1979
Decision date	Mar 21, 1979
Days to decision	36 days
Third-party review	No

APPLICANT

Company	**Electrophoresis Corp.**
Location	Mchenry, IL, US
510(k) history	4 submissions · 4 cleared · 1979-1980