

**K790310 HEMOGLOBIN CONTROLS**Mar 21, 1979  
36 days to decisionK790310 · Product code: **GGM** · Hematology  
Source: <https://www.510kdatabase.net/k790310/>**SUBMISSION DETAILS**

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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**

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

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Device record: <https://www.510kdatabase.net/k790310/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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