

**K952622 CRYO CHECK NORMAL HUMAN REFERENCE  
PLASMA**Oct 30, 1995  
144 days to decisionK952622 · Product code: **GGN** · Hematology  
Source: <https://www.510kdatabase.net/k952622/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Plasma, Coagulation Control (GGN)
Date received	Jun 8, 1995
Decision date	Oct 30, 1995
Days to decision	144 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Precision Biologicals, Inc.</b>
Location	Dartmouth, CA
Contact	STEPHEN L DUFF
510(k) history	20 submissions · 20 cleared · 1989-1999

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k952622/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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