

**K852176 DUPONT ACA FIBRIN OGEN DEGRAD-PRODUCTS  
CALIBRATOR**Aug 12, 1985  
84 days to decisionK852176 · Product code: **GHH** · Hematology  
Source: <https://www.510kdatabase.net/k852176/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Fibrin Split Products (GHH)
Date received	May 20, 1985
Decision date	Aug 12, 1985
Days to decision	84 days
Third-party review	No

**APPLICANT**

---

Company	<b>E.I. Dupont DE Nemours &amp; Co., Inc.</b>
Location	Mchenry, IL, US
Contact	ROGER C BRIDEN
510(k) history	253 submissions · 252 cleared · 1976-1996

---

**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k852176/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 14, 2026