

**K842286 DIA T1, T2 & T3**Sep 11, 1984  
90 days to decisionK842286 · Product code: **JPK** · Hematology  
Source: <https://www.510kdatabase.net/k842286/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Mixture, Hematology Quality Control (JPK)
Date received	Jun 13, 1984
Decision date	Sep 11, 1984
Days to decision	90 days
Third-party review	No

**APPLICANT**

---

Company	<b>Diatech, Inc.</b>
Location	Mchenry, IL, US
510(k) history	17 submissions · 17 cleared · 1984-1988

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k842286/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated June 29, 2026