

**K834026 RDW PARAMETER FOR THE TECHNICON H6000**Jan 30, 1984  
69 days to decisionK834026 · Product code: **GKL** · Hematology  
Source: <https://www.510kdatabase.net/k834026/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Counter, Cell, Automated (particle Counter) (GKL)
Date received	Nov 22, 1983
Decision date	Jan 30, 1984
Days to decision	69 days
Third-party review	No

**APPLICANT**

---

Company	<b>Technicon Instruments Corp.</b>
Location	Mchenry, IL, US
510(k) history	157 submissions · 156 cleared · 1976-1991

---

**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k834026/> 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 May 14, 2026