

**K810235 ACETAMINOPHEN/SALICYLATE CONTROL**Feb 10, 1981  
13 days to decisionK810235 · Product code: **DIF** · Toxicology  
Source: <https://www.510kdatabase.net/k810235/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Drug Mixture Control Materials (DIF)
Date received	Jan 28, 1981
Decision date	Feb 10, 1981
Days to decision	13 days
Third-party review	No

**APPLICANT**

---

Company	<b>Clin Tech Diagnostics Corp.</b>
Location	Mchenry, IL, US
510(k) history	5 submissions · 5 cleared · 1980-1982

---

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

Device record: <https://www.510kdatabase.net/k810235/> 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