

**K780240 PREDICTOR**Feb 27, 1978  
14 days to decisionK780240 · Product code: **LCX** · Chemistry  
Source: <https://www.510kdatabase.net/k780240/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Kit, Test, Pregnancy, Hcg, Over The Counter (LCX)
Date received	Feb 13, 1978
Decision date	Feb 27, 1978
Days to decision	14 days
Third-party review	No

**APPLICANT**

---

Company	<b>Organon, Inc.</b>
Location	Mchenry, IL, US
510(k) history	41 submissions · 41 cleared · 1977-1995

---

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

Device record: <https://www.510kdatabase.net/k780240/> 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