

**K832599 EARLY IN HOME PREGNANCY TEST**Sep 20, 1983  
49 days to decisionK832599 · Product code: **JHJ** · Chemistry  
Source: <https://www.510kdatabase.net/k832599/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Agglutination Method, Human Chorionic Gonadotropin (JHJ)
Date received	Aug 2, 1983
Decision date	Sep 20, 1983
Days to decision	49 days
Third-party review	No

**APPLICANT**

---

Company	<b>Warner-Lambert Co.</b>
Location	Mchenry, IL, US
510(k) history	50 submissions · 50 cleared · 1979-2003

---

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

Device record: <https://www.510kdatabase.net/k832599/> 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