

**K800419 ADT PERCUTANEOUS TRANSHEPATIC DISP. CHOL**Mar 25, 1980  
28 days to decisionK800419 · Product code: **FCG** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k800419/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Biopsy Needle (FCG)
Date received	Feb 26, 1980
Decision date	Mar 25, 1980
Days to decision	28 days
Third-party review	No

**APPLICANT**

---

Company	<b>A.D.T. Lab Industries</b>
Location	Mchenry, IL, US
510(k) history	21 submissions · 21 cleared · 1980-1981

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k800419/> 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 July 4, 2026