

**K902032 O. B. PACK IV**Sep 28, 1990  
147 days to decisionK902032 · Product code: **HFC** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k902032/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Aspirator, Endocervical (HFC)
Date received	May 4, 1990
Decision date	Sep 28, 1990
Days to decision	147 days
Third-party review	No

**APPLICANT**

---

Company	<b>Customed, Inc.</b>
Location	Carolina Puerto Rico, US
Contact	FELIX B SANTOS
510(k) history	36 submissions · 26 cleared · 1990-1998

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k902032/>; 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 26, 2026