

**K160610 Kobold Fletcher-model Tandem and Ovoid Applicator Set, Kobold Henschke-model Tandem and Ovoid Applicator Set, Kobold Vaginal Cylinder Applicator Set, Kobold Miami Cylinder Applicator Set**Feb 13, 2017  
347 days to decisionK160610 · Product code: **JAQ** · Radiology  
Source: <https://www.510kdatabase.net/k160610/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Applicator, Radionuclide, Remote-controlled (JAQ)
Date received	Mar 3, 2016
Decision date	Feb 13, 2017
Days to decision	347 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Kobold, LLC</b>
Location	Liberty Lake, WA, US
Contact	Brian Stoddart
510(k) history	3 submissions · 3 cleared · 2015-2017

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k160610/> 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 16, 2026