

**K022637 BD-3000 MICROKERATOME SYSTEM, MODEL K-3000  
MICROKERATOME**Oct 16, 2002  
69 days to decisionK022637 · Product code: **HMY** · Ophthalmic  
Source: <https://www.510kdatabase.net/k022637/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Keratome, Battery-powered (HMY)
Date received	Aug 8, 2002
Decision date	Oct 16, 2002
Days to decision	69 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Becton, Dickinson &amp; CO</b>
Location	Franklin Lakes, NJ, US
Contact	EILEEN T SCHWEIGHARDT
510(k) history	190 submissions · 190 cleared · 2001-2016

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k022637/>. Data sourced from FDA 510(k) public records (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. - Generated May 24, 2026