

**K031018 AUTOGEL PAD**Dec 8, 2003  
252 days to decisionK031018 · Product code: **MUI** · Radiology  
Source: <https://www.510kdatabase.net/k031018/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Media, Coupling, Ultrasound (MUI)
Date received	Mar 31, 2003
Decision date	Dec 8, 2003
Days to decision	252 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

---

Company	<b>Rich-Mar Corp.</b>
Location	Mchenry, IL, US
Contact	DAVID RICHARDS
510(k) history	43 submissions · 41 cleared · 1976-2004

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k031018/> 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 18, 2026