

**K895015 ARTROMOT(TM) CPM DEVICES**Sep 11, 1989  
34 days to decisionK895015 · Product code: **BXB** · Physical MedicineSource: <https://www.510kdatabase.net/k895015/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Exerciser, Powered (BXB)
Date received	Aug 8, 1989
Decision date	Sep 11, 1989
Days to decision	34 days
Third-party review	No

**APPLICANT**

---

Company	<b>Buckman Co., Inc.</b>
Location	Mchenry, IL, US
Contact	DAVID W SCHLERF
510(k) history	111 submissions · 104 cleared · 1983-1998

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k895015/>; 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 21, 2026