

**K020056 DUET SUTURE ANCHOR**Jun 26, 2002  
169 days to decisionK020056 · Product code: **HWC** · Orthopedic  
Source: <https://www.510kdatabase.net/k020056/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Screw, Fixation, Bone (HWC)
Date received	Jan 8, 2002
Decision date	Jun 26, 2002
Days to decision	169 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Bionx Implants, Ltd.</b>
Location	Tampere, FI
Contact	TUIJA ANNALA
510(k) history	12 submissions · 12 cleared · 1999-2003

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k020056/> 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 20, 2026