

**K894160 HAND TOOLS FOR ARTECH TYPE SS SUBMERGED  
SCREW IMPL**Sep 28, 1989  
106 days to decisionK894160 · Product code: **DZE** · Dental  
Source: <https://www.510kdatabase.net/k894160/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implant, Endosseous, Root-form (DZE)
Date received	Jun 14, 1989
Decision date	Sep 28, 1989
Days to decision	106 days
Third-party review	No

**APPLICANT**

---

Company	<b>Artech Corp.</b>
Location	Falls Church, VA, US
Contact	HENRY HAHN
510(k) history	8 submissions · 8 cleared · 1984-1992

---

**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k894160/>. 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 July 4, 2026