

**K801524 ZEST PLANT**Jul 21, 1980  
20 days to decisionK801524 · Product code: **DZE** · DentalSource: <https://www.510kdatabase.net/k801524/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implant, Endosseous, Root-form (DZE)
Date received	Jul 1, 1980
Decision date	Jul 21, 1980
Days to decision	20 days
Third-party review	No

**APPLICANT**

---

Company	<b>Zest Anchors, Inc.</b>
Location	Walker, MI, US
510(k) history	14 submissions · 14 cleared · 1980-2015

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k801524/> 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 June 4, 2026