FDA 510(k) Premarket Notification

K071851 HDC STERILE SPIDER SCREW

CLEARED

Oct 18, 2007
105 days to decision

K071851 · Product code: **DZE** · Dental
Source: https://www.510kdatabase.net/k071851/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implant, Endosseous, Root-form (DZE)
Date received	Jul 5, 2007
Decision date	Oct 18, 2007
Days to decision	105 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Hdc S.R.L.**
Location	Bethesda, MD, US
Contact	GUIDO BONAPACE
510(k) history	3 submissions · 3 cleared · 2005-2009