

**K141787 SAGITTILT**Aug 7, 2014  
36 days to decisionK141787 · Product code: **IYE** · Radiology  
Source: <https://www.510kdatabase.net/k141787/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Accelerator, Linear, Medical (IYE)
Date received	Jul 2, 2014
Decision date	Aug 7, 2014
Days to decision	36 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

---

Company	<b>Orfit Industries NV</b>
Location	Wijnegem, BE
Contact	Lee Leichter
510(k) history	9 submissions · 9 cleared · 2013-2020

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k141787/> 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 15, 2026