

**K180183 Ritter Dental Implant System**Jan 24, 2019  
366 days to decisionK180183 · Product code: **DZE** · Dental  
Source: <https://www.510kdatabase.net/k180183/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implant, Endosseous, Root-form (DZE)
Date received	Jan 23, 2018
Decision date	Jan 24, 2019
Days to decision	366 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Ritter Implants GmbH CO KG</b>
Location	Monroe, CT, US
Contact	Wemer Schmitz
510(k) history	3 submissions · 3 cleared · 2014-2019

**REGULATORY CONSULTANT**

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Consulting firm	<b>Licensale, Inc.</b>
Contact	Raymond J Kelly

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: FDA accessdata.fda.gov

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k180183/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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