

**K240796 myAblation Guide (VB80A)**Aug 6, 2024  
137 days to decisionK240796 · Product code: **QTZ** · Radiology  
Source: <https://www.510kdatabase.net/k240796/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Radiological Image Processing Software For Ablation Therapy Planning And Evaluation (QTZ)
Date received	Mar 22, 2024
Decision date	Aug 6, 2024
Days to decision	137 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Siemens Healthcare GmbH</b>
Location	Erlangen, DE
Contact	Allman Lynn
510(k) history	30 submissions · 30 cleared · 2016-2026

**REGULATORY CONSULTANT**

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Consulting firm	<b>Varian Medical Systems, Inc.</b>
Contact	Lynn Allman

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

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