

**K963983 E. CAM PROFILE ATTENUATION CORRECTION**Mar 31, 1997  
178 days to decisionK963983 · Product code: **KPS** · Radiology  
Source: <https://www.510kdatabase.net/k963983/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Tomography, Computed, Emission (KPS)
Date received	Oct 4, 1996
Decision date	Mar 31, 1997
Days to decision	178 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Siemens Corp.</b>
Location	Mchenry, IL, US
Contact	PAUL G ORIS
Website	<a href="http://www.siemens.it/">http://www.siemens.it/</a>
510(k) history	66 submissions · 66 cleared · 1978-2010

Siemens Corp. is a global technology company headquartered in McHenry, US. The company develops medical imaging and diagnostic equipment for healthcare providers worldwide. Siemens has received FDA 510(k) clearances from total submissions. The company's regulatory focus centers on Radiology devices, which represent the dominant category of its cleared portfolio. FDA 510(k) clearances span from 1978 to 2010, establishing a significant historical record in medical device regulation. Recent cleared devices include advanced imaging systems such as CT scanners, MR systems, X-r...

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