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OSIRIX	SUPPORT	RESOURCES	ABOUT	My Accoun B uy OsiriX
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OsiriX	Support	Resources
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For Mac	Tutorials	PACS
For Mobile	Help Center	RIS Integration
Solutions	Premium Membership	DICOM Image Library
Institutions	FAQ	Plugins
Patients	OsiriX Cloud FAQ	Plugins Development
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OsiriX MD is cleared by the FDA, as a Class II Medical Device, for diagnostic imaging in medicine
OsiriX MD complies with European Directive 93/42/ EEC concerning medical devices. Under this directive, it is regarded as a
Class IIa product. Notified Body: DQS Medizinprodukte GmbH, 0297