# Contact

## Contact Pixmeo

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**Subject**

**Message (in English or French)**

Send

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