DOSIsoft

Job Announcement

Cachan, France – November 24, 2026

Post-Market Clinical Application Specialist – Radiotherapy Patient-Specific QA software products

DOSIsoft Radiotherapy develops software medical devices for External Beam Radiation Therapy, in particular the patient specific quality assurance (PSQA) solution, called ThinkQA. www.dosisoft.com.ThinkQA is distributed worldwide and complies with CE/MDR, US/FDA and other national regulatory obligations in the 60+ countries where it is marketed.

Responsibilities

The Post-Market Clinical Application Specialist helps ensure that clinical evidence, product performance, user needs and customer experience are consistently captured, assessed and translated into product and business improvements. The Post-Market Clinical Application Specialist helps also put in place practices, and perform activities, necessary to fulfill regulatory obligations regarding post-market surveillance. The Post-Market Clinical Application Specialist contributes to collaboration with international clinical centers.

Main Activities

The activities and primary responsibilities of the Post-Market Clinical Application Specialist include:
- Contribute to the definition of the clinical evaluation strategy for our medical device
- Contribute to the requested regulatory documents: Clinical Evaluation Plan, Clinical Evaluation Report
- Contribute to the Post-Market Clinical Follow-up activities, plans and reports
- Collect and synthesize relevant clinical literature
- Collect, analyze and summarize user feedback, clinical data, product-use information, complaints, incidents and emerging clinical trends.
- Work with Product Management, R&D, Quality and Regulatory Affairs to translate product features and intended purposes into consistent clinical claims, risks and performance statements
- Share clinical insights with Marketing, Product Management and Sales; support product demonstrations, customer meetings and congresses, identify sales opportunities and develop clinical materials with reference customers. Study, report and disseminate clinical uses of the Products
- Provide clinical input to CE/FDA submissions and regulatory audits.
- Document and manage customer claims and Product-related safety issues together with DOSIsoft Quality and Regulatory Affairs team
- Develop, support, contribute to customer pilot activities and critical initial deployments together with the R&D/Product Management and Customer Services teams
- Develop general or specific Customer satisfaction surveys
- Contribute to collaboration projects regarding clinical use of products with selected customers

Post-Market Clinical Application Specialist reports to the QA Clinical Product Manager, works closely with the Customer Services team, the Scientific / R&D / Product Management teams, the Marketing Manager, the Sales team, and with the Quality and Regulatory Affairs team.

Skills and knowledge

Medical Physics master degree or a relevant equivalent combination of experience and education in the healthcare domain
Knowledge and good understanding of EU MRD and US FDA regulatory framework and obligations

Experiences

- 1-3 years proven initial clinical experience in oncology / radiotherapy / nuclear medicine
- Ability to speak and write fluently in both French and English.
- Customer oriented: ability to develop strong relationships with customers
- Business oriented: as for every member of a small and medium company
- Ability to work both independently and collaboratively across multidisciplinary teams.
- Excellent verbal and written communication skills
- Technical proficiency with computers and software
- Resourceful and proactive problem-solver, adaptable and quick to learn

Located at DOSIsoft Radiotherapy headquarters in Cachan, France
Partial remote working possible, Occasional international travels to congresses, partners, and customers
Salary according to experience Conditions as per “Syntec” collective agreement or more advanced company conditions
Send CV + letter to: jobs@dosisoft.fr