PROWESS

Prowess Inc. Announces FDA 510(k) Clearance for PUMA (OIS | R & V)

Concord, CA, June 2010 – Prowess Inc, a radiation oncology treatment-planning software company, has received FDA 510(k) clearance for Puma (Oncology Information System | Record and Verify). Offered at an enticing price, Puma is so affordable that medical professionals will be able to acquire a new OIS | R&V system for their oncology department without the typical anxiety over how to meet a tight budget.

Prowess created Puma to provide a comprehensive and highly reliable computerized Record and Verify System to help radiation therapists improve quality control and provide invaluable scheduling, record-keeping, and report-generating functionalities. Fully integrated, Puma enables a paperless office by importing radiation treatment plans in DICOM and RTP. The result is an intuitive system that expedites the clinical workflow and empowers the oncology team to focus on their highest priority – the patient.

Key Features and Benefits:

- Oncology specific EMR
- Diagnosis and Staging
- Radiation prescription monitoring
- Scheduling and Reports
- Simple, Intuitive Interface
- HIPAA compliance
- Multi-client access to central database

Contact Details

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Come visit us at the AAPM **Booth #142**, on July 18-22, 2010.

About Prowess Inc.

Prowess Inc., a leading innovator in Windows-based treatment planning systems for radiation therapy treatment and OIS software, is headquartered in Concord, California. Under the leadership of CEO John Nguyen, Prowess pioneered the DAO IMRT technology.

Prowess is in the process of launching **Panther Version 5.0,** the very first Windows 7 / 64 bit-based treatment planning system, with an intuitive, one-dimensional Ribbon UI and multiple-language capability.

Further information can be found by visiting http://www.prowess.com/