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Sentinelle Medical Receives CE Mark Approval

Toronto, Canada: October 6th, 2008 – Sentinelle Medical Inc., a leading manufacturer of MRI coils and interventional software, today announced it has received CE Mark approval to begin marketing the Sentinelle Vanguard Breast MR Auxiliary Table[®] and other products in the European Economic Area (EEA).

CE Marking is a mandatory European requirement to indicate conformity with the essential health and safety requirements set out in the European Directives. CE Mark approval is another example of the international acceptance of Sentinelle's technologies. "This CE Mark approval is further evidence of Sentinelle's continued commitment to its customers to ensure quality and safety of its products and to increase its market presence," said Cameron Piron, Sentinelle's President and CEO. "We are pleased to receive the CE mark approval as it is a recognized symbol of quality."

About Sentinelle Medical Inc.

Sentinelle Medical is dedicated to improving patient care with early detection and accurate intervention. The company researches, develops, manufactures, sells and supports leading edge Magnetic Resonance (MR) technologies. The ground breaking Sentinelle Vanguard Breast MR Auxiliary Table[®] provides excellent quality MR images, complete and open access for intervention and award winning design that optimizes patient comfort and workflow. Sentinelle products are used in leading breast cancer sites throughout North America including Dartmouth-Hitchcock Medical Centre, H. Lee Moffitt Cancer Center and Research Institute (University of South Florida), Scripps Memorial Hospital, Sunnybrook Health Sciences Centre and University Health Network. Sentinelle Medical is a privately owned Canadian company.

For more information, visit www.sentinellemedical.com

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