grafts that could be treated each year," said Peter Rule, president, CEO and chairman of PercuSurge.

The PercuSurge GuardWire Plus Temporary Occlusion and Aspiration system is designed to allow cardiologic specialists to capture embolic debris that might otherwise block downstream vessels and branches during in damage the heart. It consists of a balloon-tipped guidewire, which is inflated briefly to occlude blood flow an dislodged from the wall of the vessel during placement of a stent upstream. Captured material is then withdre PercuSurge Export aspiration catheter before the balloon of the GuardWire Plus is deflated and blood flow references.

The device has been used in over 5,000 procedures since its release in Europe during 1999 and was the first obe commercialized there. The product's first targeted indication is for the treatment of degenerated saphenor signs of disease following heart bypass surgery.

The GuardWire Plus is an investigational device in the United States, and clinical trials are already underway

Edited by Ursula Jones Managing Editor, Medical Design Online

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