

Declaration of Conformity

Application of Council Directive(s): Medical Device Directive 93/42/EEC

Standards to which Conformity is Declared: EN ISO 13485:2003, EN ISO 14971:2007, EN ISO 11737-1:2006, EN ISO 10993:2003, ISO 15223:2007, EN 1041

Manufacturer's Name: Vycor Medical Inc

Manufacturer's Address: 80 Orville Drive, Suite 100, Bohemia, NY USA 11716

Community Representative: MediMark Europe, 11, rue Emile Zola, BP 2332
38033 Grenoble Cedex 2, France

Notified Body: Intertek Semko AB, Torshamnsgatan 43, Box 1103,
164 22 Kista, Sweden

Type of Equipment: Vycor ViewSite Surgical Access System

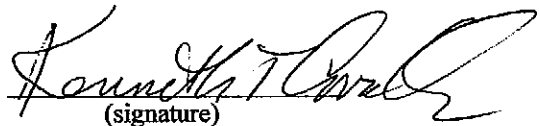
Device Classification per MDD 93/42/EEC: Class III, Rule 7

Model Numbers: TC120803, TC120805, TC120807, TC171103,
TC171105, TC171107, TC211503, TC211505,
TC211507, TC282003, TC282005, TC282007,
EC341403, EC341405, EC341407

Serial Numbers: N/A

I, the undersigned, hereby declare that the equipment specified above conforms to the above Directive (s) and Standard (s).

Place: Bohemia, NY USA


(signature)

Date: August 19, 2010

Kenneth T. Coviello
(full name)

President
(Position)



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