

Press Release

SOURCE: CPC of America, Inc.

Med Enclosure, LLC – Market and Competitive Analysis: Vascular Closure

Sarasota, Fla., May 15, 2006 – CPC of America, Inc. (OTCBB: CPCF.PK) reports that it's subsidiary, Med Enclosure, LLC, which is developing the MedClose® Class III Medical Device which is a proprietary catheter based delivery system that uses an existing Food and Drug Administration ("FDA") licensed fibrin sealant to rapidly seal arterial puncture sites following angiography and angioplasty. This Vascular Closure System ("VCS") known as MedClose-Arterial® is not presently available for human use. For further details see the February 2005 Market and Competitive Analysis: Vascular Closure at <http://www.cpa2000.com/mcanalysis.htm>

This news release contains forward-looking statements within the meaning of the Private Securities Litigation Reform Act of 1995. Such statements are subject to certain risks and uncertainties, and actual circumstances, events or results may differ materially from those projected in such forward-looking statements. Factors that could cause or contribute to differences include, but are not limited to: the risk that the FDA may not approve the IDE application for MedClose® or, if approved, that any such investigation may be unfavorable; that that CPC may be unable to obtain FDA approval for commercial sale of MedClose®; and the risk that CPC may be unable to obtain capital as and when needed. For a discussion of these and other factors, which may cause actual events or results to differ from those projected, please refer to CPC's most recent annual report on Form 10-K/A and quarterly reports on Form 10-Q, as well as other subsequent filings with the Securities and Exchange Commission. CPC of America, Inc cautions readers not to place undue reliance on any forward-looking statements. CPC does not undertake, and specifically disclaims any obligation, to update or revise such statements to reflect new circumstances or unanticipated events as they occur.

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