

Phase II Medical Manufacturing Names Christine Beverly as New Quality Assurance Manager

(ROCHESTER, NH) June 3, 2010 -- Phase II Medical Manufacturing Inc., the innovative leader in the assembly, testing, and packaging of single use disposable medical devices, announced today that Christine Beverly has joined Phase II as its Quality Assurance Manager.

Ms. Beverly brings more than 22 years of experience in quality engineering and quality management in regulated industries. Prior to joining Phase II, Ms. Beverly spent two years as the Division Quality Assurance Manager at Lydall Corp. Prior to Lydall Corp, Ms. Beverly held positions in Quality Assurance Management and Quality Engineering at Parker Hannifin, Micro Med, NESLAB Instruments, and North Atlantic Energy Service Corporation. Ms. Beverly began her career as a mechanical engineer with International Paper Company.

Ms. Beverly has a Masters of Business Administration from Boston University and a Masters of Science in Mechanical Engineering from Northeastern University. She has a Bachelor of Science in Mechanical Engineering from The University of Vermont. She is ASQ (American Society for Quality) certified as a Manager of Quality/Organizational Excellence.

“Christine brings proven leadership skills in quality that have been honed in highly regulated industries including nuclear power and medical device manufacturing,” said Adam Prime, President, Phase II Medical Manufacturing, Inc. “Her focus on data driven structured problem solving and her vast experience in Lean Manufacturing will be an asset to Phase II Medical and our customer base.”

About Phase II Medical Manufacturing

Phase II Medical Manufacturing Inc., based in Rochester, NH, is an innovative leader and turn key contract manufacturer of single use disposable medical devices. Phase II Medical Manufacturing Inc., provides material procurement, device assembly, testing, packaging, shipping, sterilization, and warehousing and fulfillment services to its customers. Phase II Medical Manufacturing Inc., is FDA registered and licensed in Canada, as well as ISO 9001:2000 and ISO 13485:2003 certified. For more information, visit www.phaseiimed.com or call 603-332-8900.

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