262 Buffalo Avenue
Paterson, NJ 07503
Phone 973.278.6500
Fax 973.278.2228
www.ppi-timezero.com

News Release

For Immediate Release

Contact | Francis R. (Frank) Simons | Executive Vice President
Phone | 973.278.6500 | fsimons@ppi-timezero.com

FDA Quality Audit Successful for PPI-Time Zero

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Frank Simons, Executive Vice President, Sales and Marketing announced this week that PPI-Time Zero, Inc, has successfully concluded a Food and Drug Administration (FDA) Quality Audit.

SQA, a professional services company that specializes in supplier quality management and operates across a broad industry spectrum, including medical devices that are highly regulated, high precision or high value. In the audit process SQA represented PPI-Time Zero customer Stryker/Osteonics a broadly based, global leader in medical technology, and one of the largest players in the worldwide orthopedic market. SQA conducted the FDA Audit which stated in part that "PPI-Time Zero has a Quality System that is in compliance with ISO 9001:2000."

As the result of the SQA Audit Report it was concluded that no FDA-483 observations (deficiencies in a quality system and potential non-compliance issues) were noted. Additional auditor comments included "PPI-Time Zero showed evidence of numerous areas in which they demonstrated continuous improvements as part of a day-to-day-quality management effort," said Frank Simons.

Since 1971, PPI-Time Zero has produced over 20 million electronic assemblies for the high-reliability military, aerospace, medical and industrial controls markets. They manufacture sub assemblies, circuit card assemblies, burn-in boards and complete systems integration using the most advanced design automation technology for a wide range of specialized applications.

PPI-Time Zero is an ISO 9001:2000 Registered and HUBZone Certified manufacturer. Their 56,000 square foot facility, in Paterson, NJ, complies with the requirements of high-performance, high-reliability, industrial and military workmanship and performance standards.

For more information about the services of PPI-Time Zero, Inc., visit www.ppi-timezero.com

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