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Mr. Daniel Sterling is the founder and president of SterlingTech. With 25 years of experience in safety critical software development including all types of Class II and Class III systems, Mr. Sterling has led the organization in executing over 150 medical device projects.

Mr. Sterling is known as a medical device software expert having given numerous talks and workshops on software topics affecting medical device development.

Having worked with such prestigious organizations as St. Jude Medical, Medtronic, Boston Scientific, Johnson and Johnson, Teleflex Medical, Hitronics Design, Continua Health Alliance, and Advanced Neuromodulation Systems, Mr. Sterling has managed software projects spanning the medical device, aerospace, and telecommunication industries.

About SterlingTech:

SterlingTech provides development, testing, and quality systems services exclusively to the medical device industry.

Since its founding in 1998, SterlingTech has been recognized as a leader in medical device design providing innovative solutions for its clients. SterlingTech delivers the highest level of knowledge and commitment, optimizing hardware and software processes for their specific needs. Working from concept to delivery, maintaining cost efficiency, reducing risk, and keeping to schedule, SterlingTech brings software and hardware together to solve clients' needs.

The company has built integrated hardware and software systems leading to a proven track record of FDA/CE approvals for clients including multinational, midsize and startup medical device companies. The services provided include full life cycle design and development, independent test, quality system consulting, and prototype build capability.

SterlingTech's New York area development and testing laboratory was designed from the ground up to allow an unrivalled degree of collaboration with clients. Its quality system is ISO 13485 registered and ISO compliant to 60601, 45502 and 62304 standards.

SterlingTech's philosophy is to work diligently to get the job done – whether it is getting product to market or troubleshooting challenging technical issues. The company's dedicated staff does whatever it takes to get the job done right with the goal of meeting or exceeding FDA/CE expectations.

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