



For Immediate Release:

October 3, 2005

QA EDGE, INC., WELCOMES VARNER AS GENERAL MANAGER, SOUTHERN REGION

WILMINGTON, DE – QA Edge, Inc., provider of a range of industry-proven Computer Validation and 21 CFR Part 11 training and services, is pleased to announce the addition of Barry L. Varner to its team of talented professionals as General Manager, Southern Region.

Varner brings an extremely strong background in providing Computer Validation and 21 CFR Part 11 solutions to QA Edge. In more than 10 years in the field, he has contributed to validation efforts at such pharmaceutical industry leaders as Amgen, Eli Lilly, Aventis, Bayer, McNeil, Wyeth, Merck, and Johnson & Johnson. Trained in Electrical Engineering at North Carolina State University, Varner worked in the automation, control, and electrical engineering fields before beginning a career as a Validation Engineer and Project Manager in 1992. He has extensive experience in the development of validation Master Plans, Standard Operating Procedures, Design Requirements, and User Requirements, in addition to extensive Project Management experience.

As General Manager, Southern Region, Varner will bring his wealth of expertise to QA Edge to help expand its services and clientele base in Maryland, Virginia, the Carolinas, and nearby locations.

About QA Edge

QA Edge, Inc., headquartered in Wilmington, DE, provides its customers with real-world strategies to avoid costly mistakes through a diverse range of Computer Validation and Part 11 training and services – with a “Computer Compliance Made Easy” methodology. The company’s knowledgeable and experienced professionals provide clients the advantage needed in meeting today’s regulatory requirements. Recognized in 2005 with a coveted Philadelphia 100 Award as one of the fastest-growing, privately-held companies in the Philadelphia region, the individualized solutions crafted for clients by QA Edge carry the industry’s only ironclad corporate guarantee: documents authored by QA Edge staff will pass **any** FDA inspection. In the unlikely event FDA has an issue, the document will be corrected immediately at no charge. More about QA Edge and its products and services can be found on the Web at <http://www.QAedge.com>.

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