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For Immediate Release:

GlobalSubmit Extends Signature Partnership Program

Philadelphia, PA— October 6, 2009 - GlobalSubmit Inc., (www.globalsubmit.com) a leading developer of software designed exclusively for the review and validation of electronic Common Technical Document (eCTD) global submissions, announces that it is extending its Signature Partners Program to include OEMs. As a continuing commitment to its partners, GlobalSubmit confirms that its REVIEW™ and VALIDATE™ software solutions are available exclusively to GlobalSubmit partners and clients. While competitive electronic submission solutions providers claim to have access to REVIEW and VALIDATE, and to be able to train customers in the use of same, these claims are unequivocally untrue.

According to GlobalSubmit CEO Rahul Mistry, “The GlobalSubmit Signature Partners program has, to date, enabled GlobalSubmit to work with outstanding eCTD consultants including e-SubmissionSolutions.com and with elite clinical research organizations such as [CATO Research](http://CATO_Research). Aligning with these partners allows our organization to offer products and services that, taken together, provide an almost universal eCTD submission solution. Expanding our network of Signature Partners to include OEMs is the next logical step in making our offerings even more comprehensive.”

The expansion of the GlobalSubmit Signature Partners program comes at a time when GlobalSubmit is on the verge of releasing a new version of its solution suite and introducing a new product that will provide customers an even more complete solution set at an extremely competitive price.

For more information, please contact Rahul Mistry at 215-253-7471.

About GlobalSubmit

GlobalSubmit is a products and services company that provides transparency in regulated healthcare products. The U.S. Food & Drug Administration and leading Life Sciences companies use our flagship software applications, REVIEW™ and VALIDATE™, to review and validate electronic submissions. GlobalSubmit’s thought leaders lead international efforts, constantly working with the industry and with government agencies to standardize product and study information. The company is headquartered in Philadelphia, Pennsylvania.

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OCTOBER 5, 2009