Monthly Program Status Report - PROJECT

Reporting Period:	September 2015		
Contracting Agency:	Food and Drug Administration (FDA)		
FDA Project Manager:	Jingyee Kou, jingyee.kou@fda.hhs.gov, 301-796-9495		
FDA Subject Matter Expert:	Thomas Permutt, thomas.permutt@fda.hhs.gov, 301-796-1271		
FDA COTR:	Shaila Shaheed, Shaila.Shaheed@fda.hhs.gov,		
Contract / Order:	HHSF223201310230C		
Contractor PI:	Daniel Scharfstein, dscharf@jhu.edu, 410-955-2420		
Project Team:	Aidan McDermott (Computer Programmer)		
Description of Activity:	A recent FDA-sponsored National Research Council Report recommended that "examining sensitivity to the assumptions about the missing data mechanism should be a mandatory component of reporting." While the Report outlines a framework for conducting sensitivity analysis, there are two major problems with existing methods: (1) they have not been implemented in software packages and (2) they do not adequately address non-monotone missing data patterns (i.e., patients provide data irregularly). The objective of this project is to address these gaps by: 1) creating unified and coherent methods for global sensitivity analysis of clinical trials with monotone and non-monotone missing data, 2) developing free, open source and reproducible software in SAS and R to implement the methods, and 3) demonstrating the methods and software using real clinical trial data.		

Project Health Check						
Health ▶	Budget	Schedule	Resources	Deliverables		
Notes ►	Within Budget	On Schedule	Adequate	On Target		

Budget	Budget Tracking – (TOTAL CONTRACT CEILING)						
DOD	Ceiling Remaining	Cumulative Funding	Year Funding	Spent to Date	Year Funding Remaining	Month Invoice	Funding Covers
POP			(Year 1)				
Base	\$1,094,565	\$1,094,565	\$1,094,565	\$943,445.36 (\$306,047.69) committed)	\$151,119.64	\$562,466.07	Salary, fringe, other expenses, and indirect costs

Activity Summary and Highlights

We implemented a method for analyzing datasets with monotone and non-monotone missing data and are running an extensive simulation study. We worked on converting SAMON to SAS. We arranged FDA short course for November 30, 2015. We will present the SAS version of SAMON at the FDA short course.

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Key Accomplishments				
Current Reporting Period	Planned for Next Period			
 Implemented simulation study of intermittent missing data methods Worked on SAS version of SAMON 	 Develop SAS version of software Continue implementation of intermittent missing data methods 			

Issues and Risks					
Category	Prior ity	Status	Opened	Issue	Description
Contract (FDA)	1	Closed	9/30/13	Intellectual Property	Revision to contract regarding intellectual property language.
Dissemination (FDA)	2	Closed	2/15/14	Website	FDA Personnel cannot connect to www.missingdatamatters.org from their office computers.
Software (JHU)	1	Closed	3/15/14	Coverage of Confidence Intervals	Simulations indicate that standard procedures for constructing confidence intervals are not providing adequate coverage with typical sample sizes.
Computing (JHU)	1	Closed	4/21/14	Periods of slow performance of computing cluster	A new computing cluster was installed at Johns Hopkins. We are experiencing periods of slow performance on the cluster.
Personnel (JHU)	1	Closed	5/21/14	Re-Distribution of Effort	Starting April 1, Aidan McDermott has reduced his percent effort by 20%. Chenguang Wang joined the project starting July 15.
Invoicing (FDA)	1	Open	6/6/14	Payment of Invoices	Invoices have not been paid.
Computing (FDA)	1	Open	6/6/14	Software on FDA Cluster	Investigate the steps needed to run software on FDA cluster
Personnel (JHU)	1	Open	1/13/15	New Effort	Yi Lu joined the project to work on confidence intervals.

Other Activities

Attachments and References	