ATTACHMENT 2a: NCCDPHP DMP Template

Form Approved OMB NO: 0920-xxxx Exp. Date: X/XX/XXXX

NCCDPHP Data Management Plan (DMP)

for use by applicants and awardees of contracts and notices of funding opportunity (NOFOs)

Background: The DMP should be developed during the project planning phase prior to the initiation of collecting or generating **public health data** and regularly updated as plans evolve. The DMP will be evaluated by CDC for completeness and quality at the time of application submission, award, or submission of the evaluation plan; at least annually thereafter; and when the project approaches termination. **Public health data** means digitally recorded factual material commonly accepted in the scientific community as a basis for public health findings, conclusions, and implementation. It does <u>not</u> include grantee progress reports, administrative data, preliminary analyses, drafts of scientific papers, plans for future research, reports, communications with colleagues, or physical objects, such as laboratory notebooks or specimens. In most cases, acquisition of secondary data does not require a DMP. For projects in which CDC aggregates, analyzes, and disseminates awardees' data, CDC may choose to develop the DMP. If the applicant or awardee believes that their project does not meet the criteria for submission of a DMP, the applicant/awardee must provide a justification.

Data Management Plan Requirement: Required Not Required	
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Reason not required: ______

Phase of project (check one):
New Application
New Award
Evaluation Plan
Continuation
Final

Date:

NOFO or Contract name:

NOFO or Contract (solicitation or award) Number:

Awardee Name or Number:

Description of the Data

In the following table, identify the data to be generated or collected for the public health dataset. If the project involves more than one, repeat this table for each public health dataset. All cells should be filled with brief answers. Expand cells as needed. Where necessary, state "n/a" or "plans pending"; however, the final DMP must have substantive responses for all cells

Dataset Title	Study Type/Design	Frequency of Data Collection	Data Collection Timeframe	Where Data Will Be Maintained During Study Period	Responsible Person / Contact Information
Describe the content (top	ics, variables) of the data				
	ndards for Collecting	g Data			
Describe the data collect	ion/generation methods				
Describe measures to en	sure data quality				
Providing Access t					
access) of data will be pr	ccess (free public access, res rovided and when it will be ne-off collection; 12 months	made available (must be			
	nd how the data will be ave ess dataset, discoverability o ted dataset)	-			
If free public access to th	e data will not be provided	, give a justification			
	ased, describe procedures ((removal of PII, data use ag	•			
-	ndards Accompanyir	-			
	standards to be used to en e.g., ICD codes, CSV files, el	,			
Describe the documentation (e.g., population studied	tion that will be available re , response rate, etc.)	egarding data source			
Describe the documenta dictionary, sample code)	tion that will be available fo	or analysis (e.g., data			
	-Term Data Preserva				
-	g-term preservation (how l in be accessed, who has acc ation	-			
If applicable, name the p	lanned final location of the or governmental repository				

it can be accessed (including link or contact information for archived data)	