The data for the CSE will come from three sources: the CSE portal (implementation and systems outcomes data), annual stakeholder interviews (implementation and systems outcome data), and findings from the annual End of Year Evaluation Reports.

Procedure for Data Extraction and Coding

We propose to extract both implementation and outcome findings from the End of Year Evaluation Reports using systematic procedures. This Procedures Guide describes which data elements will be extracted from the evaluation reports on provider, parent and child outcomes and how they will be coded. One component of the data extraction involves rating the strength of evidence of each study finding. Effects generated from quasi-experimental designs will be rated using the What Works Clearinghouse (WWC) Procedures and Handbook.¹ Outcomes generated from pre-post studies will be rated using The R-SEED (Review of Studies with Emergent Evidence Designs) Procedures and Handbook (in preparation).

The CSE final report will include data on effects on providers, families/parents, and children in two ways. First, for each service strand, there will be tables of outcome findings for each sample (providers, parents, and children). Although individual findings will be shown, the findings will not be identified with the name of the LAUNCH site. For each of these findings, we will attach a rating of strength of evidence of that finding. Second, if we have multiple findings for any of these three samples within a service strand, we will calculate a weighted average effect. This average effect will be calculated separately for findings within each of the major levels of strength of evidence, e.g., findings rated as being strong evidence (QEDs with baseline equivalence and a measure of known psychometric adequacy), findings rated as representing intermediate strength of evidence, and findings rated as having limited strength of evidence. Findings that do not meet standards for limited strength of evidence will not be used in the calculation of overall LAUNCH effects.

The coding guide is organized following the 7 sections of the LAUNCH CSE Outcome Findings Data Extraction Guide. These sections include:

- Tab 1: Services description
- Tab 2: Provider outcomes (includes assessment of strength of evidence)
- Tab 3: Provider outcome data
- Tab 4: Parent outcomes (includes assessment of strength of evidence)
- Tab 5: Parent outcome data
- Tab 6: Child outcomes (includes assessment of strength of evidence)
- Tab 7: Child outcome data

¹ The current version of the WWC standards is contained in the September 2011 What Works Clearinghouse Procedures and Standards Handbook (Version 2.1), U.S. Department of Education, Institute for Education Sciences. Version 3.0 (February 2013) is the most recent update and is currently available for public comment.

Surveys to Obtain Incomplete Data

Following the data extraction from the End of Year Evaluation Reports, surveys will be sent to the local evaluators to request any data elements that were not in the reports. These surveys will match the data elements in the 7 sections of the Extraction Guide. The surveys will be pre-populated with the available data and evaluators will be asked to provide data for elements that, on the survey, are shown to be missing. This process implies that the surveys to evaluators will be individually tailored to request only the missing data elements.

The Paperwork Reduction Act Burden Statement: This collection of information is voluntary and will be used to evaluate implementation and outcomes of the Project LAUNCH program. Public reporting burden for this collection of information is estimated to average 480 minutes per response, including the time for reviewing instructions, gathering and maintaining the data needed, and reviewing the collection of information. An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number. The OMB control number for this collection is 0970-XXXX and it expires XX/XX/XXXX.

TAB1: S	SERVICE (intervention/program/service being evaluated)		
Row	Data element	Definition	Comments
1	Project LAUNCH Grantee	LAUNCH Grantee (Cohorts 1,2,4= State/tribe; Cohort 3 = community)	Pre-populated on remaining tabs
2	Local Evaluator	Last name of lead evaluator	Pre-populated on remaining tabs
3	Grant Year of EOY Evaluation Report (One-Five)	Year of grant represented in report (1 – 5)	Pre-populated on remaining tabs
4	Date of EOY Evaluation Report	Report date	Pre-populated on remaining tabs
6	Description of LAUNCH-supported Service		
7	Strand (Home visiting (HV), Family support (FS), mental health consultation in preK (MHC-ECE), mental health consultation in school (MHC-ELEM), integration of behavior health in primary care (IBH-PC), developmental screening (DS)	Enter abbreviation for SAMHSA strand	Pre-populated on remaining tabs
3	Other strands/types of services (mental health consultation in other settings (MHC-OTH), early childhood education (ECE)	Enter abbreviation or specify type of service if not listed	Pre-populated on remaining tabs
)	Name of service/model	Specify	Pre-populated on remaining tabs
LO	Initiated or enhanced	Enter "yes" if initiated by LAUNCH; enter "no" if LAUNCH is enhancing an existing service	
L2	LAUNCH-supported enhancements		
L3	Program expansion (additional staff, slots)	Enter yes or no	
14	Workforce enhancement		
L5	Itraining on mental health & development	Enter yes or no	
L6	Itraining on screening/assessment	Enter yes or no	
17	Mental health consultation	Enter yes or no	
18	New component added to model (specify component)	Enter yes or no	
19	Other	Enter yes or no	

TAB 2:	PROVIDER OUTCOMES		
Row	Data element	Definition	Comments
1	Project LAUNCH Grantee	LAUNCH Grantee (Cohorts 1,2,4= State/tribe; Cohort 3 = community)	Pre-populated from Service tab
2	Local Evaluator	Last name of lead evaluator	Pre-populated from Service tab
3	Grant Year of EOY Evaluation Report (One-Five)	Year of grant represented in report (1 – 5)	Pre-populated from Service tab
4	Date of EOY Evaluation Report	Report date	Pre-populated from Service tab
6	Description of LAUNCH-supported Service		
7	Strand (Home visiting (HV), Family support (FS), mental health consultation in preK (MHC-ECE), mental health consultation in school (MHC-ELEM), integration of behavior health in primary care (IBH-PC), developmental screening (DS)	Enter abbreviation for SAMHSA strand	Pre-populated from Service tab
8	Other strands/types of services (mental health consultation in other settings (MHC-OTH), early childhood education (ECE)	Enter abbreviation or specify type of service if not listed	Pre-populated from Service tab
9	Name of service/model	Specify	Pre-populated from Service tab
10	Provider outcome measures		
12A	Measure name	Specify	E.g., "LAUNCH Provider Survey," Provider Stress Checklist
12B	Measure #		
12C	Domain	Specify overall domain being addressed	E.g., provider knowledge, provider attitudes, provider practices
12D	Construct	Indicate construct being measures	E.g., provider stress, provider use of standardized assessments
12E	Reference	Citation for measure	Full published name or reference in which measure was described. Enter NA is measure is site-developed.
12F	Type of measure	Specify type of measure	E.g., interview, survey, observation, test
12G	Scoring	Specify how measure is score	E.g., binary, ordinal(ordered), categorical/nominal (unordered), continuous
12H	Outcome is valid for implementation (matches program objectives, enhancements)	Enter yes or no	Outcome must appear to measure the domain into which it is classified; outcome must align with program theory of change
121	Over-aligned	Enter yes or no	Measure must not be designed or administered in ways that are specifically aligned with the program intervention model
12J	Measure reliability: standardized tests	If test is standardized, enter "yes"; otherwise, enter "no"	Outcomes should have test-retest reliability = .40 or higher (for scale measures based on survey items) or inter-rater reliability = .50 higher for data based on observation measures. Standardized tests are assumed to satisfy the reliability criterion. Other measures exempt from the reliability criterion include health indicators such as immunizations.
12K	Reliability of non-standardized measure	Enter test-retest reliability, internal consistency, or inter- rater reliability	Enter reliability statistic appropriate to type of measure

TAB 2:	PROVIDER OUTCOMES		
Row	Data element	Definition	Comments
13A- 13K	Repeat 13A-13K for 2 nd provider outcome		
14A- 14K	Repeat 13A-13K for 3 rd provider outcome		
15A- 15K	Repeat 13A-13K for 4 th provider outcome		
16	Design for measurement of outcome		
18A	Measure name		Pre-populated from 12a
18B	Measure #		Pre-populated from 12b
18C	Type of design: QED	Enter "yes" if design includes comparison group; otherwise, enter "no"	Tre-populated from 225
18D	Type of design: ITS	Enter "yes" if design is ITS with longitudinal pre and/or post data; otherwise, enter "no"	
18E	ITS: # of baseline data points	Enter # data points	
18F	ITS: # of data points during program implementation	Enter # data points	
18G	Type of design: pre-post	Enter "yes" if design includes one pre- and one post-test data point; otherwise enter "no"	2 measurement points
18H	Type of design: pre vs. norm	Enter "yes" if design includes one pre- test data point and test is normed so that standardization sample can be used as comparison; otherwise enter "no"	
181	Type of design: retrospective pre-post	Enter "yes" if design includes post-test that measures change since an assumed baseline; otherwise, enter "no"	
18J	Type of design: post-only	Enter "yes" if design includes post-test measurement only and no pre-test; otherwise, enter "no"	
19A-J	Repeat 18A-18J for 2 nd provider outcome		
20A-J	Repeat 18A-18J for 3 rd provider outcome		
21A-J	Repeat 18A-18J for 4th provider outcome		
	Counterfactual explanation		
23A	Measure name		Pre-populated from 12a
23B	Measure #		Pre-populated from 12b
23C	Data on outcome collected in comparable ways: same timing of data collection for all sample members (e.g., data should be collected at the same time point in a family's participation for all families), and the same data collection procedures for all sample members, (e.g., if measure is to be implemented as a self-administered questionnaire, this same procedure should be used with all sample members).	Enter "yes" if data collection is comparable; otherwise, enter "no"	Data for outcome defined and collected in a way that ensures the outcome measure is comparable for all groups being compared
23D	Measure defined consistently	Enter "yes" if data collection is comparable; otherwise, enter "no"	Outcome is defined in same way for all groups being compared
23E	Low likelihood of growth in intervention time period in absence of intervention	Enter "yes" if similar growth is unlikely in absence of the intervention; otherwise, enter "no"	

TAB 2: F	PROVIDER OUTCOMES		
Row	Data element	Definition	Comments
23F	Lack of other interventions/interruptions likely causes of growth	Enter "yes" if are no other interventions likely to create similar growth in absence of the intervention; otherwise, enter "no"	
24A-F	Repeat 24a-24f for 2 nd provider outcome		
25A-F	Repeat 25a-25f for 3 rd provider outcome		
26A-F	Repeat 26a-26f for 4 th provider outcome		
27	Evidence rating		
29A	Measure name		Pre-populated from 12a
29B	Measure #		Pre-populated from 12b
29C	Design level	Enter design number from R-SEED rating system	Based on R-SEED design rating system
29D	Outcome meets standards	Enter "yes" if outcomes meets all WWC standards; otherwise enter "no"	Must meet standards for validity, over-alignment, and reliability. Measures will pass the screen for overalignment unless there is clear evidence otherwise. Over-alignment arises when an outcome measures concepts or is administered in ways that are specifically aligned to the content of the intervention. For example, if an intervention includes repeatedly having parents practice reading specific texts and responding to questions from the interventionist and the outcome is an assessment that asks parents to read the same text and respond to the same (or very similar) questions, that assessment would be over-aligned.
29E	Study does not have a serious confound	If study is a QED, enter "yes" if there is no serious confound (i.e., an n=1 confound); otherwise enter "no." If study is a pre-post, enter NA.	A serious confound occurs when there is a factor that has a separate effect on the outcome that cannot be eliminated by the study design, and which will bias the estimated effect of the intervention. An n=1 confound is a serious confound that occurs when an effect is estimated on the basis of comparing one unit (e.g., one teacher, one class, one school) to one or more such entities.
29F	Study can establish baseline equivalence of the analytic sample	If study is a QED, enter "yes" if baseline equivalence has been established; otherwise enter "no." If study has no comparison group, enter NA.	The study must provide evidence that the groups being contrasted as equivalence at baseline on a pre-intervention measure of the outcome.
29G	The study design has a well-justified counterfactual explanation	If study is a pre-post, enter "yes" if study has well- justified counterfactual explanation (rows	
30A-G	Repeat 30A-30G for 2 nd provider outcome		
31A-G	Repeat 31A-31G for 3 rd provider outcome		
32A-G	Repeat 32A-32G for 4th provider outcome		

TAB 3: I	TAB 3: PROVIDER OUTCOME DATA			
ROW	Data element	Definition	Comments	
1	Project LAUNCH Grantee	LAUNCH Grantee	Pre-populated from Service tab	
		(Cohorts 1,2,4= State/tribe; Cohort 3 = community)		
2	Local Evaluator	Last name of lead evaluator	Pre-populated from Service tab	
3	Grant Year of EOY Evaluation Report (One-Five)	Year of grant represented in report (1 – 5)	Pre-populated from Service tab	
4	Date of EOY Evaluation Report	Report date	Pre-populated from Service tab	
6	Description of LAUNCH-supported Service			
7	Strand (Home visiting (HV), Family support (FS), mental	Enter abbreviation for SAMHSA strand	Pre-populated from Service tab	
	health consultation in preK (MHC-ECE), mental health			

TAB 3:	PROVIDER OUTCOME DATA		
ROW	Data element	Definition	Comments
	consultation in school (MHC-ELEM), integration of behavior health in primary care (IBH-PC), developmental screening (DS)		
8	Other strands/types of services (mental health consultation in other settings (MHC-OTH), early childhood education (ECE)	Enter abbreviation or specify type of service if not listed	Pre-populated from Service tab
9	Name of service/model	Specify	Pre-populated from Service tab
	Eligible outcomes		
19A	1st eligible provider outcome	Specify name of outcome from Provider Outcome tag, A29-32)	Eligible outcomes are those whose evidence rating is Limited, Intermediate, or Strong
20A	Repeat 19A for 2 nd eligible outcome		
21A	Repeat 19A for 3 rd eligible outcome		
22A	Repeat 19A for 4th eligible outcome		
	Samples		
19C	Baseline sample (time 1)	Describe sample participants at time 1	
19D	Posttest sample (time 2)	Describe sample participants at time 2	
19E	Matched sample	Enter "yes" if samples for two timepoints are matched; otherwise enter "no."	
19F	Pre-post time period	Indicate number of months between time 1 and 2 and amount of exposure	
20C-F	Repeat 19C-F for 2 nd eligible outcome		
21C-F	Repeat 19C-F for 3 rd eligible outcome		
22C-F	Repeat 19C-F for 4th eligible outcome		
	Sample sizes		
19W	Eligible sample time 1	Indicate # of units in baseline or time 1 eligible sample	# of all eligible units in sample at time 1
19X	Analysis sample time 1	Indicate # of units in baseline or time 1 analysis sample	# of units with data in analysis sample at time 1
19Z	Eligible sample time 2	Indicate # of units in posttest or time 2 eligible sample	# of all eligible units in sample at time 1
19AA	Analysis sample time 2	Indicate # of units in posttest or time 2 analysis sample	# of units with data in analysis sample at time 1
20W- AA	Repeat 19W-AA for 2 nd eligible outcome		
21W- AA	Repeat 19W-AA for 3 rd eligible outcome		
22W- AAF	Repeat 19W-AA for 4th eligible outcome		
	Findings		
19AV	Method used to calculate significance of pre-post difference	Select from drop-down menu	
19AW	Mean outcome for analysis sample at time 2	Enter mean or proportion	
19AY	Standard deviation of outcome for analysis sample at time 2	Enter standard deviation (if applicable)	
10BC	Mean outcome for analysis sample at time 1	Enter mean or proportion	

TAB 3: I	TAB 3: PROVIDER OUTCOME DATA			
ROW	Data element	Definition	Comments	
19BE	Standard deviation of outcome for analysis sample at time 1	Enter standard deviation (if applicable)		
19BL	T statistic	Enter t-statistic from t-test of pre-post difference on outcome		
19BO	P-value	Enter p-value for t-statistic		
19BP	Significance	Enter "yes" if finding is statistically significant; otherwise, enter "no"		
20AV- BP	Repeat 19AV-BP for 2 nd eligible provider outcome			
21AV- BP	Repeat 19AV-BP for 3 rd eligible provider outcome			
22AV- BP	Repeat 19AV-BP for 4 th eligible provider outcome			

TAB 4: F	TAB 4: PARENT OUTCOMES				
Row	Data element	Definition	Comments		
1	Project LAUNCH Grantee	LAUNCH Grantee	Pre-populated from Service tab		
		(Cohorts 1,2,4= State/tribe; Cohort 3 = community)			
2	Local Evaluator	Last name of lead evaluator	Pre-populated from Service tab		
3	Grant Year of EOY Evaluation Report (One-Five)	Year of grant represented in report (1 – 5)	Pre-populated from Service tab		
4	Date of EOY Evaluation Report	Report date	Pre-populated from Service tab		
6	Description of LAUNCH-supported Service				
7	Strand (Home visiting (HV), Family support (FS), mental	Enter abbreviation for SAMHSA strand	Pre-populated from Service tab		
	health consultation in preK (MHC-ECE), mental health				
	consultation in school (MHC-ELEM), integration of				
	behavior health in primary care (IBH-PC), developmental				

TAB 4:	PARENT OUTCOMES		
Row	Data element	Definition	Comments
	screening (DS)		
8	Other strands/types of services (mental health consultation in other settings (MHC-OTH), early childhood education (ECE)	Enter abbreviation or specify type of service if not listed	Pre-populated from Service tab
9	Name of service/model	Specify	Pre-populated from Service tab
	_		
10	Parent outcome measures		
12A	Measure name	Specify	E.g., Parenting Stress Index
12B	Measure #	0	
12C	Domain	Specify overall domain being addressed	E.g., parent knowledge, parent attitudes, parent practices
12D	Construct	Indicate construct being measures	E.g., parent stress, parent use of standardized assessments
12E	Reference	Citation for measure	Full published name or reference in which measure was described. Enter NA is measure is site-developed.
12F	Type of measure	Specify type of measure	E.g., interview, survey, observation, test
12G	Scoring	Specify how measure is score	E.g., binary, ordinal(ordered), categorical/nominal (unordered), continuous
12H	Outcome is valid for implementation (matches program objectives, enhancements)	Enter yes or no	Outcome must appear to measure the domain into which it is classified; outcome must align with program theory of change
121	Over-aligned	Enter yes or no	Measure must not be designed or administered in ways that are specifically aligned with the program intervention model
12J	Measure reliability: standardized tests	If test is standardized, enter "yes"; otherwise, enter "no"	Outcomes should have test-retest reliability = .40 or higher (for scale measures based on survey items) or inter-rater reliability = .50 higher for data based on observation measures. Standardized tests are assumed to satisfy the reliability criterion. Other measures exempt from the reliability criterion include health indicators such as immunizations.
12K	Reliability of non-standardized measure	Enter test-retest reliability, internal consistency, or inter- rater reliability	Enter reliability statistic appropriate to type of measure
13A- 13K	Repeat 13A-13K for 2 nd parent outcome		
14A- 14K	Repeat 13A-13K for 3 rd parent outcome		
15A- 15K	Repeat 13A-13K for 4 th parent outcome		
16	Design for measurement of outcome		
18A	Measure name		Pre-populated from 12a
18B	Measure #		Pre-populated from 12b
18C	Type of design: QED	Enter "yes" if design includes comparison group; otherwise, enter "no"	
18D	Type of design: ITS	Enter "yes" if design is ITS with	

TAB 4: I	PARENT OUTCOMES		
Row	Data element	Definition	Comments
		longitudinal pre and/or post data; otherwise, enter "no"	
18E	ITS: # of baseline data points	Enter # data points	
18F	ITS: # of data points during program implementation	Enter # data points	
18G	Type of design: pre-post	Enter "yes" if design includes one pre- and one post-test data point; otherwise enter "no"	2 measurement points
18H	Type of design: pre vs. norm	Enter "yes" if design includes one pre- test data point and test is normed so that standardization sample can be used as comparison; otherwise enter "no"	
181	Type of design: retrospective pre-post	Enter "yes" if design includes post-test that measures change since an assumed baseline; otherwise, enter "no"	
18J	Type of design: post-only	Enter "yes" if design includes post-test measurement only and no pre-test; otherwise, enter "no"	
19A-J	Repeat 18A-18J for 2 nd parent outcome		
20A-J	Repeat 18A-18J for 3 rd parent outcome		
21A-J	Repeat 18A-18J for 4 th parent outcome		
	Counterfactual explanation		
23A	Measure name		Pre-populated from 12a
23B	Measure #		Pre-populated from 12b
23C	Data on outcome collected in comparable ways	Enter "yes" if data collection is comparable; otherwise, enter "no"	Data for outcome defined and collected in a way that ensures the outcome measure is comparable for all groups being compared
23D	Measure defined consistently	Enter "yes" if data collection is comparable; otherwise, enter "no"	Outcome is defined in same way for all groups being compared
23E	Low likelihood of growth in intervention time period in absence of intervention	Enter "yes" if similar growth is unlikely in absence of the intervention; otherwise, enter "no"	
23F	Lack of other interventions/interruptions likely causes of growth	Enter "yes" if are no other interventions likely to create similar growth in absence of the intervention; otherwise, enter "no"	
24A-F	Repeat 24a-24f for 2 nd parent outcome		
25A-F	Repeat 25a-25f for 3rd parent outcome		
26A-F	Repeat 26a-26f for 4th parent outcome		
27	Evidence rating		
29A	Measure name		Pre-populated from 12a
29B	Measure #		Pre-populated from 12b
29C	Design level	Enter design number from R-SEED rating system	Based on R-SEED design rating system
29D	Outcome meets standards	Enter "yes" if outcomes meets all WWC standards; otherwise enter "no"	Must meet standards for validity, over-alignment, reliability
29E	Study does not have a serious confound	If study is a QED, enter "yes" if there is no serious confound (i.e., an n=1 confound); otherwise enter "no." If	A serious confound occurs when there is a factor that has a separate effect on the outcome that cannot be

TAB 4: I	TAB 4: PARENT OUTCOMES			
Row	Data element	Definition	Comments	
		study is a pre-post, enter NA.	eliminated by the study design, and which will bias the estimated effect of the intervention. An n=1 confound is a serious confound that occurs when an effect is estimated on the basis of comparing one unit (e.g., one teacher, one class, one school) to one or more such entities.	
29F	Study can establish baseline equivalence of the analytic sample	If study is a QED, enter "yes" if baseline equivalence has been established; otherwise enter "no." If study has no comparison group, enter NA.	The study must provide evidence that the groups being contrasted as equivalence at baseline on a pre-intervention measure of the outcome.	
29G	The study design has a well-justified counterfactual explanation	If study is a pre-post, enter "yes" if study has well- justified counterfactual explanation (rows		
30A-G	Repeat 30A-30G for 2 nd parent outcome			
31A-G	Repeat 31A-31G for 3 rd parent outcome			
32A-G	Repeat 32A-32G for 4th parent outcome			

TAB 5: I	PARENT OUTCOME DATA		
ROW	Data element	Definition	Comments
1	Project LAUNCH Grantee	LAUNCH Grantee (Cohorts 1,2,4= State/tribe; Cohort 3 = community)	Pre-populated from Service tab
2	Local Evaluator	Last name of lead evaluator	Pre-populated from Service tab
3	Grant Year of EOY Evaluation Report (One-Five)	Year of grant represented in report (1 – 5)	Pre-populated from Service tab
4	Date of EOY Evaluation Report	Report date	Pre-populated from Service tab
6	Description of LAUNCH-supported Service		
7	Strand (Home visiting (HV), Family support (FS), mental health consultation in preK (MHC-ECE), mental health consultation in school (MHC-ELEM), integration of behavior health in primary care (IBH-PC), developmental screening (DS)	Enter abbreviation for SAMHSA strand	Pre-populated from Service tab
8	Other strands/types of services (mental health consultation in other settings (MHC-OTH), early childhood education (ECE)	Enter abbreviation or specify type of service if not listed	Pre-populated from Service tab
9	Name of service/model	Specify	Pre-populated from Service tab
	Eligible outcomes		
19A	1 st eligible parent outcome	Specify name of outcome from Parent Outcome tag, A29-32)	Eligible outcomes are those whose evidence rating is Limited, Intermediate, or Strong
20A	Repeat 19A for 2 nd eligible outcome		
21A	Repeat 19A for 3 rd eligible outcome		
22A	Repeat 19A for 4th eligible outcome		
	Samples		
19C	Baseline sample (time 1)	Describe sample participants at time 1	E.g., "new mothers at time of entry into program"
19D	Posttest sample (time 2)	Describe sample participants at time 2	E.g., "mothers at one year of participation in program"
19E	Matched sample	Enter "yes" if samples for two timepoints are matched; otherwise enter "no."	
19F	Pre-post time period	Indicate number of months between time 1 and 2 and amount of exposure	E.g., average of 9 months pre-post during mothers' 1st year of participation
20C-F	Repeat 19C-F for 2 nd eligible outcome		
21C-F	Repeat 19C-F for 3 rd eligible outcome		
22C-F	Repeat 19C-F for 4th eligible outcome		
	Sample sizes		
19W	Eligible sample time 1	Indicate # of units in baseline or time 1 eligible sample	# of all eligible units in sample at time 1
19X	Analysis sample time 1	Indicate # of units in baseline or time 1 analysis sample	# of units with data in analysis sample at time 1
19Z	Eligible sample time 2	Indicate # of units in posttest or time 2 eligible sample	# of all eligible units in sample at time 1
19AA	Analysis sample time 2	Indicate # of units in posttest or time 2 analysis sample	# of units with data in analysis sample at time 1
20W- AA	Repeat 19W-AA for 2 nd eligible outcome		
21W-	Repeat 19W-AA for 3 rd eligible outcome		

TAB 5: I	PARENT OUTCOME DATA		
ROW	Data element	Definition	Comments
22W- AAF	Repeat 19W-AA for 4th eligible outcome		
	Findings		
19AV	Method used to calculate significance of pre-post difference	Select from drop-down menu	
19AW	Mean outcome for analysis sample at time 2	Enter mean or proportion	
19AY	Standard deviation of outcome for analysis sample at time 2	Enter standard deviation (if applicable)	
10BC	Mean outcome for analysis sample at time 1	Enter mean or proportion	
19BE	Standard deviation of outcome for analysis sample at time 1	Enter standard deviation (if applicable)	
19BL	T statistic	Enter t-statistic from t-test of pre-post difference on outcome	
19BO	P-value	Enter p-value for t-statistic	
19BP	Significance	Enter "yes" if finding is statistically significant; otherwise, enter "no"	
20AV- BP	Repeat 19AV-BP for 2 nd eligible parent outcome		
21AV- BP	Repeat 19AV-BP for 3 rd eligible parent outcome		
22AV- BP	Repeat 19AV-BP for 4 th eligible parent outcome		

TAB 6:	TAB 6: CHILD OUTCOMES			
Row	Data element	Definition	Comments	
1	Project LAUNCH Grantee	LAUNCH Grantee (Cohorts 1,2,4= State/tribe; Cohort 3 = community)	Pre-populated from Service tab	
2	Local Evaluator	Last name of lead evaluator	Pre-populated from Service tab	
3	Grant Year of EOY Evaluation Report (One-Five)	Year of grant represented in report (1 – 5)	Pre-populated from Service tab	
4	Date of EOY Evaluation Report	Report date	Pre-populated from Service tab	
6	Description of LAUNCH-supported Service			
7	Strand (Home visiting (HV), Family support (FS), mental health consultation in preK (MHC-ECE), mental health consultation in school (MHC-ELEM), integration of behavior health in primary care (IBH-PC), developmental screening (DS)	Enter abbreviation for SAMHSA strand	Pre-populated from Service tab	
8	Other strands/types of services (mental health consultation in other settings (MHC-OTH), early childhood education (ECE)	Enter abbreviation or specify type of service if not listed	Pre-populated from Service tab	
9	Name of service/model	Specify	Pre-populated from Service tab	
10	Child outcome measures			
12A	Measure name	Specify	E.g., CBCL	
12B	Measure #			
12C	Domain	Specify overall domain being addressed	E.g., child knowledge, child attitudes, child practices	
12D	Construct	Indicate construct being measures	E.g., child stress, child use of standardized assessments	
12E	Reference	Citation for measure	Full published name or reference in which measure was described. Enter NA is measure is site-developed.	
12F	Type of measure	Specify type of measure	E.g., interview, survey, observation, test	
12G	Scoring	Specify how measure is score	E.g., binary, ordinal(ordered), categorical/nominal (unordered), continuous	
12H	Outcome is valid for implementation (matches program objectives, enhancements)	Enter yes or no	Outcome must appear to measure the domain into which it is classified; outcome must align with program theory of change	
121	Over-aligned	Enter yes or no	Measure must not be designed or administered in ways that are specifically aligned with the program intervention model	
12 J	Measure reliability: standardized tests	If test is standardized, enter "yes"; otherwise, enter "no"	Outcomes should have test-retest reliability = .40 or higher (for scale measures based on survey items) or inter-rater reliability = .50 higher for data based on observation measures. Standardized tests are assumed to satisfy the reliability criterion. Other measures exempt from the reliability criterion include health indicators such as immunizations.	
12K	Reliability of non-standardized measure	Enter test-retest reliability, internal consistency, or inter- rater reliability	Enter reliability statistic appropriate to type of measure	
13A- 13K	Repeat 13A-13K for 2 nd child outcome			
14A-	Repeat 13A-13K for 3 rd child outcome			

TAB 6: 0	AB 6: CHILD OUTCOMES			
Row	Data element	Definition	Comments	
14K				
15A-	Repeat 13A-13K for 4th child outcome			
15K				
16	Design for measurement of outcome			
18A	Measure name		Pre-populated from 12a	
18B	Measure #		Pre-populated from 12b	
18C	Type of design: QED	Enter "yes" if design includes comparison group; otherwise, enter "no"		
18D	Type of design: ITS	Enter "yes" if design is ITS with		
		longitudinal pre and/or post data; otherwise, enter "no"		
18E	ITS: # of baseline data points	Enter # data points		
18F	ITS: # of data points during program implementation	Enter # data points		
18G	Type of design: pre-post	Enter "yes" if design includes one pre- and one post-test data point; otherwise enter "no"	2 measurement points	
18H	Type of design: pre vs. norm	Enter "yes" if design includes one pre- test data point		
		and test is normed so that standardization sample can be		
401		used as comparison; otherwise enter "no"		
18 I	Type of design: retrospective pre-post	Enter "yes" if design includes post-test that measures		
		change since an assumed baseline; otherwise, enter "no"		
18J	Type of design: post-only	Enter "yes" if design includes post-test measurement		
103	Type of design. post-only	only and no pre-test; otherwise, enter "no"		
19A-J	Repeat 18A-18J for 2 nd child outcome			
20A-J	Repeat 18A-18J for 3rd child outcome			
21A-J	Repeat 18A-18J for 4th child outcome			
	Counterfactual explanation			
23A	Measure name		Pre-populated from 12a	
23B	Measure #		Pre-populated from 12b	
23C	Data on outcome collected in comparable ways	Enter "yes" if data collection is comparable; otherwise, enter "no"	Data for outcome defined and collected in a way that ensures the outcome measure is comparable for all groups being compared	
23D	Measure defined consistently	Enter "yes" if data collection is comparable; otherwise, enter "no"	Outcome is defined in same way for all groups being compared	
23E	Low likelihood of growth in intervention time period in absence of intervention	Enter "yes" if similar growth is unlikely in absence of the intervention; otherwise, enter "no"		
23F	Lack of other interventions/interruptions likely causes of growth	Enter "yes" if are no other interventions likely to create similar growth in absence of the intervention; otherwise, enter "no"		
24A-F	Repeat 24a-24f for 2 nd child outcome			
25A-F	Repeat 25a-25f for 3rd child outcome			

confound (i.e., an n=1 confound); otherwise enter "no." If study is a pre-post, enter NA. As a separate effect on the outcome that cannot be eliminated by the study design, and which will bias estimated effect of the intervention. An n=1 confound a serious confound that occurs when an effect is estimated on the basis of comparing one unit (e.g., teacher, one class, one school) to one or more such entities. Study can establish baseline equivalence of the analytic sample	TAB 6: 0	TAB 6: CHILD OUTCOMES			
Evidence rating Pre-populated from 12a	Row	Data element	Definition	Comments	
Measure name Pre-populated from 12a	26A-F	Repeat 26a-26f for 4 th child outcome			
Measure name Pre-populated from 12a	27	Fuidana vatina			
Pre-populated from 12b				B	
Design level Enter design number from R-SEED rating system Enter "yes" if outcomes meets all WWC standards; otherwise enter "no" Study does not have a serious confound If study is a QED, enter "yes" if there is no serious confound (i.e., an n=1 confound); otherwise enter "no." If study is a pre-post, enter NA. Study can establish baseline equivalence of the analytic sample The study design has a well-justified counterfactual explanation The study design has a well-justified counterfactual explanation Repeat 30A-30G for 2nd child outcome Enter "yes" if outcomes meets all WWC standards; otherwise enter "no." If study is a QED, enter "yes" if baseline equivalence has been established; otherwise enter "no." If study has no comparison group, enter NA. Enter "yes" if outcomes meets all WWC standards; must meet standards for validity, over-alignment, reliability A serious confound occurs when there is a factor that a same separate effect on the outcome that cannot be eliminated by the study design, and which will bias estimated of the basis of comparing one unit (e.g., teacher, one class, one school) to one or more such entities. If study is a QED, enter "yes" if baseline equivalence has been established; otherwise enter "no." If study has no comparison group, enter NA. If study is a pre-post, enter "yes" if study has well-justified counterfactual explanation (rows If study is a pre-post, enter "yes" if study has well-justified outcome Repeat 30A-30G for 2nd child outcome					
29E Study does not have a serious confound 29E Study can establish baseline equivalence of the analytic sample 29F Study can establish baseline equivalence of the analytic sample 29G The study design has a well-justified counterfactual explanation 29G Repeat 30A-30G for 2 nd child outcome 31A-G Repeat 31A-31G for 3 nd child outcome 29E Study does not have a serious confound occurs when there is a factor the nucleon of the explanation of the study is a QED, enter "yes" if there is no serious confound occurs when there is a factor thas a separate effect on the outcome that cannot be eliminated by the study design, and which will bias estimated on the basis of comparing one unit (e.g., teacher, one class, one school) to one or more such entities. 29F Study can establish baseline equivalence of the analytic sample enter "no." If study has no comparison group, enter "yes" if study has no comparison group, enter when there is a factor that a separate effect on the outcome and the initiate of the intervention. An n=1 confound, in the set of the intervention. An n=1 confound a serious confound that occurs when there is a factor that as a separate effect on the outcome that cannot be eliminated by the study design, and which will bias estimated on the basis of comparing one unit (e.g., teacher, one class, one school) to one or more such entities. The study must provide evidence that the groups be contrasted as equivalence at baseline on a pre-intervention measure of the outcome. 19 Study can establish baseline equivalence at baseline on a pre-intervention measure of the outcome.	-				
Otherwise enter "no" Study does not have a serious confound If study is a QED, enter "yes" if there is no serious confound (i.e., an n=1 confound); otherwise enter "no." If study is a pre-post, enter NA. Study can establish baseline equivalence of the analytic sample Study design has a well-justified counterfactual explanation The study design has a well-justified counterfactual explanation Otherwise enter "no" If study is a QED, enter "yes" if there is no serious confound occurs when there is a factor that a serious confound that occurs when an effect is estimated effect of the intervention. An n=1 confound a serious confound that occurs when an effect is estimated on the basis of comparing one unit (e.g., teacher, one class, one school) to one or more such entities. If study is a QED, enter "yes" if baseline equivalence has been established; otherwise enter "no." If study has no comparison group, enter NA. The study must provide evidence that the groups be contrasted as equivalence at baseline on a preintervention measure of the outcome. If study is a pre-post, enter "yes" if study has well-justified counterfactual explanation (rows 30A-G Repeat 30A-30G for 2nd child outcome Repeat 31A-31G for 3nd child outcome	29C	Design level		Based on R-SEED design rating system	
confound (i.e., an n=1 confound); otherwise enter "no." If study is a pre-post, enter NA. As a separate effect on the outcome that cannot be eliminated by the study design, and which will bias estimated effect of the intervention. An n=1 confound a serious confound that occurs when an effect is estimated on the basis of comparing one unit (e.g., teacher, one class, one school) to one or more such entities. Study can establish baseline equivalence of the analytic sample	29D	Outcome meets standards			
sample been established; otherwise enter "no." If study has no contrasted as equivalence at baseline on a pre-intervention measure of the outcome. 29G The study design has a well-justified counterfactual explanation (rows If study has well-justified counterfactual explanation	29E		confound (i.e., an n=1 confound); otherwise enter "no." If	estimated on the basis of comparing one unit (e.g., one teacher, one class, one school) to one or more such entities.	
explanation justified counterfactual explanation (rows 30A-G Repeat 30A-30G for 2 nd child outcome 31A-G Repeat 31A-31G for 3 nd child outcome	29F	sample	been established; otherwise enter "no." If study has no comparison group, enter NA.		
30A-G Repeat 30A-30G for 2 nd child outcome 31A-G Repeat 31A-31G for 3 nd child outcome	29G				
	30A-G	Repeat 30A-30G for 2 nd child outcome			
32A-G Repeat 32A-32G for 4th child outcome	31A-G	Repeat 31A-31G for 3rd child outcome			
	32A-G	Repeat 32A-32G for 4th child outcome			

TAB 7:	CHILD OUTCOME DATA		
ROW	Data element	Definition	Comments
1	Project LAUNCH Grantee	LAUNCH Grantee	Pre-populated from Service tab
		(Cohorts 1,2,4= State/tribe; Cohort 3 = community)	
2	Local Evaluator	Last name of lead evaluator	Pre-populated from Service tab
3	Grant Year of EOY Evaluation Report (One-Five)	Year of grant represented in report (1 – 5)	Pre-populated from Service tab
4	Date of EOY Evaluation Report	Report date	Pre-populated from Service tab
	·	·	
6	Description of LAUNCH-supported Service		
7	Strand (Home visiting (HV), Family support (FS), mental health consultation in preK (MHC-ECE), mental health	Enter abbreviation for SAMHSA strand	Pre-populated from Service tab
	consultation in school (MHC-ELEM), integration of		
	behavior health in primary care (IBH-PC), developmental		
	screening (DS)		
8	Other strands/types of services (mental health	Enter abbreviation or specify type of service if not listed	Pre-populated from Service tab
	consultation in other settings (MHC-OTH), early		
	childhood education (ECE)		
9	Name of service/model	Specify	Pre-populated from Service tab
	Eligible outcomes		
19A	1 st eligible child outcome	Specify name of outcome from Child Outcome tag, A29-	Eligible outcomes are those whose evidence rating is
		32)	Limited, Intermediate, or Strong
20A	Repeat 19A for 2 nd eligible outcome		
21A	Repeat 19A for 3 rd eligible outcome		
22A	Repeat 19A for 4th eligible outcome		
400	Samples		
19C	Baseline sample (time 1)	Describe sample participants at time 1	E.g., "Children at time of entry into program"
19D	Posttest sample (time 2)	Describe sample participants at time 2	E.g., "children at one year of participation in program"
19E	Matched sample	Enter "yes" if samples for two timepoints are matched; otherwise enter "no."	
19F	Pre-post time period	Indicate number of months between time 1 and 2 and	E.g., average of 9 months pre-post during children's' 1st
		amount of exposure	year of participation
20C-F	Repeat 19C-F for 2 nd eligible outcome		
21C-F	Repeat 19C-F for 3 rd eligible outcome		
22C-F	Repeat 19C-F for 4th eligible outcome		
	Sample sizes		
19W	Eligible sample time 1	Indicate # of units in baseline or time 1 eligible sample	# of all eligible units in sample at time 1
19X	Analysis sample time 1	Indicate # of units in baseline or time 1 analysis sample	# of units with data in analysis sample at time 1
19Z	Eligible sample time 2	Indicate # of units in posttest or time 2 eligible sample	# of all eligible units in sample at time 1
19AA	Analysis sample time 2	Indicate # of units in posttest or time 2 analysis sample	# of units with data in analysis sample at time 1
20W-	Repeat 19W-AA for 2 nd eligible outcome		
AA			
21W-	Repeat 19W-AA for 3 rd eligible outcome		
AA	Damas 400M A A San Atharias Island		
22W-	Repeat 19W-AA for 4th eligible outcome		
AAF			

TAB 7: CHILD OUTCOME DATA				
ROW	Data element	Definition	Comments	
	Findings			
19AV	Method used to calculate significance of pre-post difference	Select from drop-down menu		
19AW	Mean outcome for analysis sample at time 2	Enter mean or proportion		
19AY	Standard deviation of outcome for analysis sample at time 2	Enter standard deviation (if applicable)		
10BC	Mean outcome for analysis sample at time 1	Enter mean or proportion		
19BE	Standard deviation of outcome for analysis sample at time 1	Enter standard deviation (if applicable)		
19BL	T statistic	Enter t-statistic from t-test of pre-post difference on outcome		
19BO	P-value	Enter p-value for t-statistic		
19BP	Significance	Enter "yes" if finding is statistically significant; otherwise, enter "no"		
20AV- BP	Repeat 19AV-BP for 2 nd eligible child outcome			
21AV- BP	Repeat 19AV-BP for 3 rd eligible child outcome			
22AV- BP	Repeat 19AV-BP for 4 th eligible child outcome			