

**Generic Clearance for CDC/ATSDR  
Formative Research and Tool Development  
OMB# 0920-1154**

**Title: Caring Adults Perceptions on Teen Mental Health**

Supporting Statement A

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### List of Attachments

Attachment 1: Email

Attachment 2: Screening Survey

Attachment 3: Consent Form

Attachment 4: Interview Discussion Guide

Attachment 5: Focus Group Discussion Guide

Attachment 6: Notice of IRB Approval

Attachment 7: Privacy Forms

## Summary

**Goal of the study:** This project is intended to collect data from adults that mentor or support girls and nonbinary adolescents (ages 13-17) seeking and accessing mental health care in rural America, with a focus on identifying trusted resources and building resiliency for teens through social support systems, coping skills, and mental health literacy. This project is the second component of the “Adolescent Mental Health *Journey Mapping Project*.”

**Intended use of the resulting data:** The information collected for this qualitative formative testing will be used to inform CDC’s journey map(s) and inventory of recommended prevention strategies. The data collected through this work and outputs of this project will be for CDC internal use only and is not intended for external publication.

**Methods to be used to collect:** Data will be collected through a combination of virtual and in-person discussions.

**The subpopulation to be studied:** The population of focus for the adult data collection component of this work are adults that mentor or support girls and nonbinary adolescents (ages 13-17) in rural areas of Montana, New Mexico, and North Carolina. We plan to recruit coaches, teachers, youth leaders, and/or other adult mentors who work with girls and/or nonbinary adolescents in rural communities.

**How data will be analyzed:** We will analyze and identify major themes by conducting thematic analysis, a combination of notes, as well as the transcripts will be analyzed. The research team will analyze themes from across the data sources as well and across subgroups and respondent groups to compare and contrast findings.

## A. JUSTIFICATION

### 1. Circumstances Making the Collection of Information Necessary

The Centers for Disease Control and Prevention (CDC), National Center for Injury Prevention and Control (NCIPC), Office of Policy and Partnerships (OPP) requests OMB approval of a new generic information collection (GenIC) under the generic information collection entitled *Generic Clearance for CDC/ATSDR Formative Research and Tool Development* for a period of 10 months as the conclusion date of this project is late September 2024.

According to the US Centers for Disease Control and Prevention (CDC)'s 2021 Youth Risk Behavior Survey, 29% of high school students experienced poor mental health during the past 30 days, and female students were more likely than male students to experience mental health challenges. Additionally, those living in rural areas experience disparities in mental health outcomes compared to those living in metropolitan and suburban areas. As a response to this data and the national emergency for children and adolescent mental health declared in 2022, CDC's National Center for Injury Prevention and Control's (NCIPC) Office of Policy and Partnerships (OPP) is looking to better understand the mental health experiences of adolescent youth (ages 13-17) identifying as girls/female or nonbinary living in rural areas. We will not explicitly be asking about any traumatic experiences. This project is formative in nature and intended to guide CDC's programmatic planning to address adolescent mental health needs in the US.

### 2. Purpose and Use of Information Collection

This project is the second component of the "Adolescent Mental Health Journey Mapping Project." The information collected for this qualitative formative testing will be used to inform CDC's journey map(s) and inventory of recommended prevention strategies. The final report and journey map(s) will be used to inform and support the future of CDC's work in adolescent mental health. The data collected through this work and outputs of this project will be for CDC internal use only and is not intended for external publication. We will not be sharing any identifiable data with partners. However, we intend to share synthesized, summarized findings from across all qualitative data collection to share learning with project partners. The primary goals of this low-burden and low-cost project are to better understand the experiences of girls and nonbinary adolescents (ages 13-17) seeking and accessing mental health care in rural America, and to inform opportunities for this population for public health interventions and upstream prevention at CDC.

To meet the project goals, we will conduct primary data collection using in-person and virtual interviews and focus groups with adults that mentor girls and nonbinary adolescents. The project will include these questions:

- How can we build resilience and foster positive mental health outcomes for adolescent girls and nonbinary youth by identifying key protective and risk factors?
- How could we bring awareness to the trusted resources that adolescent girls and nonbinary youth turn to when experiencing mental health challenges?
- How might we amplify cultural or community related strengths that bolster protective factors for adolescent girl and nonbinary youth mental health?
- How can we improve identification of mental health challenges and expedite the path to support for adolescent girls and nonbinary youth in rural areas?

- Are there non-traditional ways for adolescent girls and nonbinary youth in rural areas to access mental health support?
- Can we better understand the meaning of “mental health” to adolescent girls and nonbinary youth and the ways they invest in their own mental health?

A maximum of 10 adult mentor interviews and 5 adult mentor focus groups will be conducted via an interactive in-person or virtual session between the participant(s) and the facilitator(s) using a communication interface consistent with Zoom Video Communications Inc. (Zoom) or at a community-based organization facility.

The semi-structured discussions with participants will use discussion guides. The discussion guides include questions/topics that will elicit feedback from the participants about their experiences with adolescent mental health, with a focus on social support systems, coping skills, and mental health literacy.

Interviews will be 60 minutes while focus groups will be 90 minutes. Tables 1 and 2 detail the general structure and the estimated time allocation by stage of the adult mentor interviews and focus groups, respectively.

**Table 1: General Structure of Adult Mentor Semi-Structured Interview Sessions (60-minute discussion)**

Estimated Time	Stage	Description
~3 minutes	Introduction	Facilitator(s) will introduce topic and obtain verbal consent for the session to be audio recorded.
~3 minutes	Rapport building	Short introductions to create cohesion and build rapport with the interview participant (no last names) and an ice breaker question/activity if appropriate.
~53 minutes	In-depth conversation	Facilitator(s) will initiate the major discussion topics and rely on the interview guide to ensure all topics are addressed.
~1-2 minutes	Closure	The facilitator(s) will thank the interviewee for attending, answer any last questions, and let them know that the gift-card will be emailed to them within 1-2 weeks of the session.

**Table 2: General Structure of Adult Mentor Focus Group Sessions (90-minute discussion)**

Estimated Time	Stage	Description
~3 minutes	Introduction	Facilitators will introduce topic and obtain verbal consent for the session to be audio recorded.
~10 minutes	Rapport building	Short introductions to create group cohesion (no last names) and an ice breaker question/activity if appropriate.
~75 minutes	In-depth investigation	Facilitator will initiate the major discussion topics and rely on the discussion guide to ensure all topics are addressed.
~1-2 minutes	Closure	The facilitator will thank participants for attending, answer any last questions, and let them know that the gift-card will be emailed to them within 1-2 weeks of the session.

### **3. Use of Information Technology and Burden Reduction**

The screening surveys for the session will be conducted online via an online survey platform to limit burden on both the participants and the project team. Individuals will be selected via a convenience sampling approach, through meeting potential participants where they are with advertisements in community-based organization facilities and newsletters. The survey platform has built-in analytics, which will allow the research team to quickly analyze data. The team will use a visual guide/worksheet on the screen for participants to follow along and annotate during the virtual sessions. In person, the team will use printed copies of the visual guide available for participants to use for notetaking, drawing, and interactive engagement purposes. This is intended to provide an alternative avenue to encourage engagement throughout the session audience.

### **4. Efforts to Identify Duplication and Use of Similar Information**

A literature review found no instances of similar information to be available. This information collection request represents a new effort – the first of its kind – to collect qualitative data from girls and nonbinary adolescents (ages 13-17) in an intend to map their experiences seeking and accessing mental health care in rural America. This study represents the first attempt to systematically assess this topic. There is no known information available that can substitute data collection.

### **5. Impact on Small Businesses or Other Small Entities**

This data collection will not involve small businesses. Some data collection involves from community-based organizations, including YMCA of Southeastern North Carolina, YMCA Mountain States Alliance, Family YMCA of Los Alamos, North Carolina Native American Youth Organization, and Time Out Youth; however, these data collections will not have a significant impact on these non-profit organizations. The project team will be partnering with these local non-profit, community-based organizations that serve adolescent girls and nonbinary youth in rural communities to support a convenience sampling approach to recruitment.

### **6. Consequences of Collecting the Information Less Frequently**

This request is for a one-time data collection.

### **7. Special Circumstances Relating to the Guidelines of 5 CFR 1320.5**

This request fully complies with the regulation 5 CFR 1320.5

### **8. Comments in Response to the Federal Register Notice and Efforts to Consult Outside the Agency**

A. A 60-day Federal Register Notice has already been published for the Generic Clearance. No Federal Register Notice is required for this GenIC submission.

B. A federal technology consultancy is the contractor for the data collection.

### **9. Explanation of Any Payment or Gift to Respondents**

Krueger and Casey (2009) note that the gift helps emphasize to participants that the assessment is important, which in turn is intended to make them more inclined to make time to participate.<sup>1</sup> To encourage and improve response rates for this project, each session participant will be given \$50 per session in the form of a Visa gift card for their time within 1-2 weeks of the session. This can help minimize bias in the perspectives, improve the validity and reliability of the data, and ensure equitable treatment of project participants, all of which are of utmost importance in this project.

### **10. Protection of the Privacy and Confidentiality of Information Provided by Respondents**

This submission has been reviewed by the NCIPC's Information Systems Security Officer, who has determined that the Privacy Act does not apply (**Attachment 6**). PII will only be used to screen for, and to contact, respondents who match the specified respondent profile for the session. PII will be collected via a web-based survey application as part of the prescreening process. PII for each selected individual will be destroyed before the conclusion of this effort in September 2024. Individuals not selected for a session will not be contacted and their information will immediately be deleted. PII will not be retained or used to link respondents' individual comments during the session or during the analysis. All findings will be reported in the aggregate. PII will also not be used to describe the respondents in the room or to provide context to the analysis. Data are treated in a private manner, unless otherwise compelled by law. No paper documents with PII will be collected. Data containing PII will be disposed of as soon as session participants have been contacted and scheduled for sessions. Emails will be disposed of after the incentive has been sent to the participants. Participants will have the option to turn their Zoom camera on, but it will not be mandatory during any of the virtual sessions and only audio will be recorded for all sessions. Names of participants entering the Zoom group discussion will be changed to their first names only (last names will be removed) to protect the privacy of the individual.

Audio recordings and transcriptions will be stored on a secure, password-protected cloud storage. Audio will be recorded via Zoom and recordings will be encrypted and transcribed using Zoom's transcription mechanism which will only be used by the project team to validate the transcribed audio. Transcripts of the audio recordings will be de-identified prior and during analysis. Access to audio and de-identified transcript files is limited to authorized project team personnel only. Session audio files will be destroyed once the session has been transcribed including deleting files from the password-protected cloud and any local file storage. All staff and contractors working on the project agree to safeguard the data and not to make unauthorized disclosures. Responses in published reports are presented in aggregate form and no individuals are identified by name.

All consent forms include appropriate information about privacy, including purpose for collecting the data, with whom identifiable information will be shared, the voluntary nature of the information collection and the effect upon the respondent for not participating.

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<sup>1</sup> Krueger, R. A., & Casey, M. A. (2009). Focus groups: A practical guide for applied research (4th ed.). San Francisco: Sage

## 11. Institutional Review Board (IRB) and Justification for Sensitive Questions

### *IRB Approval*

CDC has received IRB approval (**Attachment 5**) through Solutions IRB (FWA00021831).

### *Sensitive Questions*

This data collection asks sensitive questions around teen mental health and experiences adult mentors have had in supporting teens with their mental health. Adult mentors will receive a recruitment email explaining the purpose of the study invitation (**Attachment 1**) and be directed to the weblink of the screening survey (**Attachment 2**) if they are interested in participating. Once the list of potential participants is compiled, the project team will contact the adult mentor by email to provide consent and select which session dates/times work with their schedule (**Attachment 3**).

## 12. Estimates of Annualized Burden Hours and Costs

Recruitment for the interviews and focus will be conducted via convenience sampling through partnerships with local community-based organizations. Interviews will take 60 minutes to complete while focus groups will take 90 minutes to complete. There is no cost to participants beyond the participation burden time. The table below provides the burden estimates for this study.

**Table 1. Estimated Annualized Burden Hours**

Category of Respondent	Form Name	No. of Respondents	Participation Time (Hours)	Burden (Hours)
Adult Mentor Participants	Emails ( <b>Attachment 1</b> )	20	15/60	5
	Screening Survey ( <b>Attachment 2</b> )	20	10/60	3
	Adult Mentor Consent Form ( <b>Attachment 3</b> )	10	10/60	2
	Interview Discussion Guide ( <b>Attachment 4</b> )	10	1	10
	Focus Group Session Discussion Guide ( <b>Attachment 5</b> )	5	1.5	8
<b>Totals</b>				<b>28</b>

**12b.** The estimates of the annualized cost to adult mentors for the burden hours for the collection of information is derived from the November 2023 average hourly wage of \$34.23 across all occupations, per the [US Bureau of Labor Statistics website](#).

**Table 2. Estimated Annualized Burden Costs**

Type of Respondents	Form Name	Total Burden Hours	Hourly Wage Rate	Total Respondent Costs
Adult Mentor Participants	Emails ( <b>Attachment 1</b> )	5	\$34.23	\$171.15
	Screening Survey ( <b>Attachment 2</b> )	3	\$34.23	\$102.7
	Adult Mentor Consent Form ( <b>Attachment 3</b> )	2	\$34.23	\$68.5
	Interview Discussion Guide ( <b>Attachment 4</b> )	10	\$34.23	\$342.3
	Focus Group Session Discussion Guide ( <b>Attachment 5</b> )	8	\$34.23	\$273.8
Total				<b>\$958.5</b>

**13. Estimates of Other Total Annual Cost Burden to Respondents or Record Keepers**

This data collection does not involve other annual cost burdens to respondents or record keepers.

**14. Annualized Cost to the Government****Table 3. Annualized Costs to the Government**

Description of Services	Estimated Annualized Cost
Contractor costs for labor, adult mentor data collection, and other overhead costs, per 3 months of the contract year + hard costs (travel, participant compensation, and independent IRB review)	<b>\$15,760</b>

**15. Explanation for Program Changes or Adjustments**

This is a new data/information collection.

**16. Plans for Tabulation and Publication and Project Time Schedule**

All activities for the project are expected to be completed within 10 months after IRB approval.

**17. Reason(s) Display of OMB Expiration is Inappropriate**

None; the display of the OMB expiration date is not inappropriate.

### **18. Exceptions to Certification for Paperwork Reduction Act Submissions**

None; there are no exceptions to the certification.

#### **References**

Krueger, R. A., & Casey, M. A. (2009). *Focus groups: A practical guide for applied research* (4th ed.). San Francisco: Sage