



Trauma-informed Research Partnerships

Practices for Creating Shared Knowledge

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recommendations expressed herein are those of the contributors and do not necessarily represent the official positions or policies of the U.S. Department of Justice.

Today

- 1. Introductions
- 2. Research and Partnerships
- 3. What is trauma informed research?
- 4. Reflections and Recommendations
- 5. Key takeaways

Introductions

Trauma-informed Care Principles

Safety

Trustworthiness & Transparency

Peer Support

Collaboration & Mutuality

Empowerment, Voice, & Choice

Cultural, Historical, & Gender Issues

Research Methods

Quantitative	Qualitative
Surveys	Interviews
Analysis of program or other administrative data	Focus groups
Experiments	Observations

Research Process

Planning & Design

Approval

Recruitment & Data collection

Analysis

Dissemination

Experiences with research and researchers

Research Partnerships

Benefits	Costs
Create new knowledge	Taking time away from services
Better understand programs	Research may be triggering for participants
Validation of models	Having to explain everything about a population to researchers
Help prepare for growth	Data is not used to help the organization
Communicate to funders	Unclear data ownership

What is trauma-informed research?

What training or background should researchers have?

What is the most appropriate method to use with individuals who have experienced trauma?

How should participants be honored for their time?

How can methods be adapted to minimize the potential for retraumatization?

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Community Resources

LGBTO* Service Providers

Central Illinois

Central Illinois Pride Health Center Bloomington, IL

https://cipridehealthcenter.or g/ (815) 893-7459 cipridehealthcenter@gmail.c om (LGBT parents and children support group, LGBT youth support group for ages 13 to

19)

Planned Parenthood-Bloomington Health Center

1319 N. Veterans Parkway Bloomington, IL 61704 www.ppil.org (309) 827-4014 (Transgender care: hormone therapy, Psycho social support services)

Planned Parenthood- Peoria Health Center

2709 N Knoxville Ave Peoria, IL 61604 www.ppil.org (309) 681-0350

Contact: Dana Garber (Pronouns: She, Her, Hers), Transgender Intake Coordinator and Community Educator danag@ppil.org OR 309-681-0350 Ext: 4002

("Informed Consent" Cross-hormone therapy for transgender clients, sexual health services, HIV counseling and treatment. Mental Health Counseling and Referral, Additional health services available)

The Phoenix Center

109 East Lawrence Avenue Springfield, IL 62704 http://www.phoenixcentersprin gfield.org/ (217) 528-5253 (Various support groups, STD testing, Educational training & programs, and transitional housing program)



Collar Counties

Community Alliance & Action Network Joliet. IL

http://www.caanmidwest.org/ (815) 726-7906 info@caanmidwest.org (Advocacy, support, and outreach services)

McHenry County Pride 5603 Bull Valley Rd.,

McHenry, IL 60050 http://mchenrycountypride.org McHenryCountyPride@gmail.com (Adult LGBT weekly support group)

Cook County

ACLU Illinois

150 N. Michigan Ave Suite 600 Chicago, IL 60601 https://www.aclu-il.org (312) 210-9740 acluofillinois@aclu-il.org (LGBTO and HIV advocacy)

Ann & Robert H. Lurie Children's Hospital of Chicago

225 E. Chicago Ave. Box #205
Chicago, IL 60611
https://www.luriechildrens.org
(312) 227-6117
Contact: Jennifer Leininger
JLeininger@luriechildrens.org
OR (773) 303-6056
(Gender development services,
comprehensive LGBTQ sexual health
drop-in services, Center for Gender,
Sexuality & HIV Prevention)

Center on Halsted 3656 N Halsted St

Chicago, IL 60613 http://www.centeronhalsted.org/ (773) 472-6469 (Programs: Anti-violence, behavioral health, community & cultural, HIV/AIDS & STD, senior, women, and youth)

Chicago Commission on Human Relations

Community Tensions & Hate Crimes

https://www.cityofchicago.org/city /en/depts/cchr.html (312) 744-2571 (Advocacy & assistance for hate crime victims, Educational workshops & presentations, community mediation, outreach services)

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Building rapport

BACKGROUND INFORMATION

First, I would like to get to know you a bit better before we talk about specific experiences you may have had. I will use this chart to help keep track of this information.

[Interviewer will present chart and record information in chart throughout interview.]

- 1. Thinking about right now, can you describe or tell me about:
 - Where you live?
 - Who you live with (e.g., parents/caregivers, children, siblings, other family members, peers)?
 - Dating partners or significant others you have?
 - How you spend most of your time? This may be at work, school, or home.
 - Anything else you would like me to know about you before we move on to what life was like for you as a child or youth?

Now I'd like to learn more about you as a child or youth.

- Can you describe or tell me about:
 - When and where you were born?
 - Where you lived, including the city or town?
 - How would you describe where you lived: mostly urban, suburban, or rural?
 - Who you lived with (e.g., parents/caregivers, children, siblings, other family | members, peers) growing up?

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Approval

What approvals are needed?

What type of identifying information will be collected and why?

Who will have access to participant identifying information and why?

How will data and identifying information be stored?

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Privacy & Confidentiality

What about privacy and confidentiality?

Only the research team will know that you are a research subject.

If you agree to be contacted for the survey you will be asked to provide contact information (i.e., name, phone number or email address, and/or mailing address). Your contact information will be separated from the survey responses and replaced with a unique code. Only research staff will have access to this master list that links a participant's identifying information to their unique code. This master list will be kept in a separate password protected file on password-protected computers.

The information collected in the study will be used to write a report. No potentially identifying information that can be linked directly to you will be included in the report or discussed with others.

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How do we reach marginalized populations?

How do we avoid "over-researching" a certain population?

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Safety

Out of a concern for the safety of yourself and others, if during the interview, you say you are going to cause serious harm to yourself or to someone else, the researcher may have to make a report. A description of different types of serious harm are below:

- a. Physical harm is any action that causes physical injury, pain, or leaves marks, such as hitting, shaking, or burning.
- b. Sexual abuse of a child includes any sexual contact with a child or youth. Sexual abuse of an adult is any unwanted sexual activity. Sexual abuse of an adult also includes any sexual activity with another person who is unable to understand or consent (e.g., are sleeping, have been using drugs or alcohol, have a mental disability).
- Emotional abuse is yelling, swearing, or criticizing another person repeatedly to make them feel bad about themselves.
- d. Neglect occurs when a parent or caregiver of a child or youth, or the caregiver of an elderly person or a person with disability, does not meet their basic needs (e.g., food, clothing, medical care) or keep them safe.

If you share about any current harm to someone who is a child, elderly, or disabled, the researcher may have to make a report. If the researcher needs to make a report, they may share your name and contact information, or information about the person causing the harm or being harmed, and a description of the harm to the appropriate agency such as the Department of Child and Family Services, Adult Protective Services, or local law enforcement.

What research questions will be answered by the analysis?

Who can provide input or ask questions of the data?

What can't the data tell us about?

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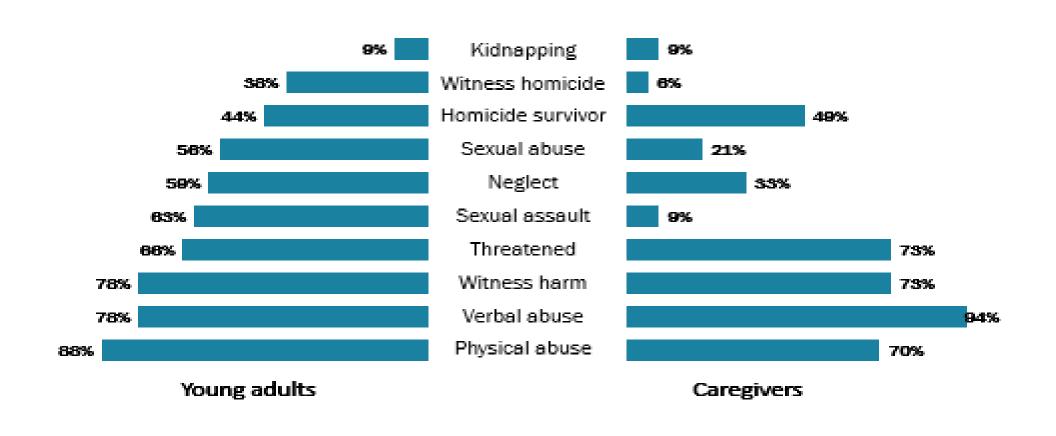
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What can't the data tell us about?

Sample

Young adults	Caregivers
N = 32	N = 33
59% female	64% female
M = 22 years old	M = 41 years old
75% heterosexual, 13% bisexual, 9% gay/lesbian	94% heterosexual
53% African American, 41% White, 13% Hispanic	76% African American, 30% White, 3% Hispanic
69% Cook County, 25% Southern region	76% Cook County, 24% Southern region

Interpreting results



What research questions will be answered by the analysis?

Who can provide input or ask questions of the data?

What can't the data tell us about?

Who is the intended audience?

Who will provide feedback on document drafts?

How can survivor voice be incorporated?

For researchers

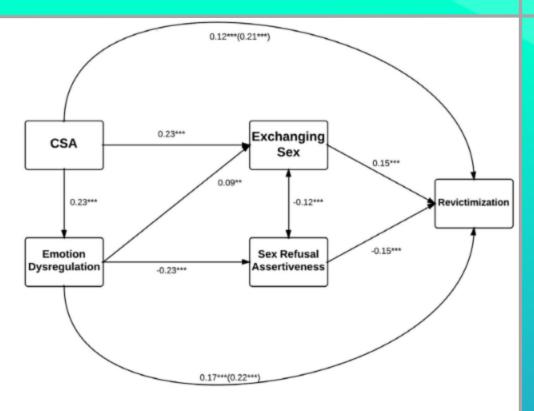


Figure 1.

Exchanging sex for money and sex refusal assertiveness as mediators of CSA and revictimization and CSA and emotion dysregulation

p<.01, *p<.001

For survivors

How Did Women React?

Everyone reacts differently to unwanted sexual experiences. However you react is normal for you. At first, many women are in shock. Later they may become confused, afraid, angry, or depressed. Women find many different ways to cope with their feelings.

The hardest time seems to be during the first years after an unwanted experience. When we compared women who experienced unwanted sex five years ago or more with women who experienced it one year ago, we found that women whose experiences were longer ago:

- Had fewer upsetting thoughts come into their head when they didn't want them to
- Were less emotionally upset when they were reminded of the experience
- Had fewer physical reactions when they were reminded of the experience
- · Were less likely to relive the experience

It is important to remember that women deal with their experiences in different ways and at different times.

Living with the Experience

One of the hardest things for some women to deal with is feeling like it was her fault

 In the first year after an unwanted experience women blamed themselves more. They often said they thought they should have been more cautious or that they didn't do enough to protect themselves.

Remember that no matter what a woman was wearing, where she was, who she was with, or what she did...It is not her fault. Women have the right to say "no."

Women who experienced the assault five or more years ago compared to women who had the experience in the past year seemed to have recovered more from the experience.

- Women felt there was more they could do to make themselves feel better.
- Women felt more spiritual. They said they had a stronger religious faith and had a better understanding of spiritual matters.
- Women felt more personal strength. They said they knew how to better handle difficulties and they discovered they were stronger than they thought they were.

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Journal articles

Original Articles

The Role of Victim Services for Individuals Who Have Experienced Serious Identity-Based Crime

ABSTRACT

Although much has been written about identity-based crimes, much less is known about victim services for identity-based crimes. This paper uses in-depth interview data collected from individuals who have both experienced a serious identity-based crime in the past 2 years and used services from the nonprofit Identity Theft Resource Center, an organization focused on helping victims deal with identity-based victimizations. In so doing, the study adds to the literature in several ways. First, the study echoes the existing knowledge about victim experiences, in that there are financial, emotional, physical, and psychological impacts for victims of serious identity-based crime that mirror the experiences of victims of other types of serious crimes. Second, it adds new knowledge and understanding about the actions taken by victims of different types of identity-based crime, including how they initially seek out services, are influenced by organizations and agencies other than law enforcement, and obtain knowledge about their perpetrators. Finally, the study includes recommendations made by interviewees about how identity-based crime victim services could be enhanced to better serve victims of serious identity-based crime.

KEYWORDS: White-collar crime, fraud, victimization



Get access

Conference Fees

National Training Institute Registration Fees VIRTUAL CONFERENCE ONLY

	Non-Member	Member
Early Bird	\$300	\$225
(February 24 – June 15,		
2020)		
Standard	\$400	\$300
(June 15 – September		
30, 2020)		
Late	\$450	\$350
(September 30 –		
November 13, 2020)		

Reflections & Recommendations

Key takeaways



Research as mutually beneficial to service providers and researchers



A trauma-informed approach to research can mitigate the potential for retraumatization



Each partner has rights and responsibilities that can be outlined in a formal agreement

Discussion & Questions

Presenters

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