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	ORRECT	TON	SUBJECT:	Research Activities
DEPAR	TMENT OF	CORRECTIONS		
POLI	CIES AND	PROCEDURES		
RELATED ACA #1-H		C-3A-09(M), 5-ACI-1F-18	EFFECTIVE DA	ATE: July 15, 2023
STANDARDS:				
			SUPERSESSION	N: 02/07/2022
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DESCRIPTION:		REVIEW MONTH:	as	elly WOOKO
Admin. & Management -		July	K	XELLIE WASKO
Information Systems and				ARY OF CORRECTIONS
Research			SECKETA	INT OF CORRECTIONS

### I. POLICY

It is the policy of the South Dakota Department of Corrections (DOC) to support research activities relevant to its programs that assist in achieving department goals, objectives, or plans for the future; and that contribute to more effective and efficient supervision of offenders, use of DOC resources, and increased public safety.

#### II. PURPOSE

The purpose of this policy is to establish guidelines that govern voluntary offender participation in non-medical, non-pharmaceutical, and non-cosmetic research programs [5-ACI-1F-18] and to describe research procedures, responsibilities of administrators, and dissemination of information. The DOC will ensure the security of information and data collection systems, including verification of data, electronic transmission, and storage of data, and protection of the privacy of offenders, DOC employees, contract workers, and volunteers.

### III. DEFINITIONS

#### **DOC Research Activities:**

Research initiated by the DOC, either using DOC staff and/or contract researchers.

#### **Medical Research:**

Research activities that include, but are not limited to, medical experiments and pharmaceutical studies using offender subjects.

#### **Research Activities:**

The collection of information about a particular subject which is aimed at:

- 1. The discovery and/or interpretation of facts.
- 2. The revision of accepted theories in light of new facts; or
- 3. The practical application of new or revised theories.

#### **Social Science Research:**

Research activities that include, but are not limited to, the use of offender interviews, offender questionnaires, and reviews of offender case records. This does not include any research activities that will expose offenders to the possibility of physical, psychological, or other harm as a consequence of their participation.

#### IV. PROCEDURES

SECTION	SUBJECT	DOC POLICY	Page 2 of 5
Admin. & Management -	Research Activities	1.1.F.01	Effective:
Information Systems and			07/15/2023
Research			

### 1. General Research Activity Guidelines:

- A. The DOC may support and engage in internal research relevant to its programs, including approved research conducted by outside research professionals.
- B. Research activities other than those activities initiated by the DOC, which utilize the participation or records of DOC employees, interns, volunteers, or offenders which are held by the DOC, must be approved in writing by the secretary of corrections (SOC), deputy secretary of corrections (DSOC), or designee, prior to the start of the research.
- C. Those conducting research activities will be informed of, and shall adhere to, all DOC policies that are relative to the research project. All applicable security and safety procedures shall apply to the activities of the researcher and the research project.
- D. Research activities will comply with all state and federal law, including all laws and guidelines pertaining to the use and reporting of research findings. All approved research activities will conform to accepted professional standards and scientific ethics.

### 2. Application to Conduct Research:

- A. The principal researcher (excluding DOC research activities) must complete a *Research Application* (see attachment #1), prior to conducting any research within a DOC program, unit, or institution and prior to having contact with offenders or staff for research purposes. The completed application must be submitted to the SOC or designee.
- B. The DOC may require the researcher or sponsoring organization substantiate a professional standing in the field of corrections, criminal justice, education, or related field, subject to approval by the DOC.
- C. The applicant shall include an endorsement by a recognized research organization (e.g., university, college, private foundation, consulting firm, or public agency) that has a mandate to perform research, certifying that the research proposal is for valid scientific, educational, or other public purposes. If Human Subjects Reviews have been conducted and approved through the requesting research organization, a copy shall be included as part of the endorsement.
- D. The design of the research and information provided on the application shall be of sufficient quality to reasonably predict the results of the research will be reliable, valid, and reasonably relevant to the business of the DOC.
- E. Submitted applications or requests received by the DOC will generally be responded to within ten (10) working days of receipt by those with authority to consider the request.

#### 3. Obtaining Approval to Conduct Research:

- A. DOC staff may request additional documentation from the researcher concerning any aspect of the research project, prior to making a decision on the research request.
- B. If the research application is declined, a response will be sent in writing to the contact listed on the research application within fifteen (15) working days of receipt of the research application/request. All decisions are final. A copy of the response shall be retained on file in the DOC Central Office.

SECTION	SUBJECT	DOC POLICY	Page 3 of 5
Admin. & Management -	Research Activities	1.1.F.01	Effective:
Information Systems and			07/15/2023
Research			

- C. The *Research Agreement* (see attachment #2) must be signed and approved by the SOC or designee prior to initiation of the research.
  - 1. The secretary or designee will maintain the original signed research agreement with the research request and any other responses or documentation related to the research request.
  - 2. A copy of the respective signed research agreement form will be provided to the principal researcher.
  - 3. The researcher shall not modify, revise, or otherwise change the agreement or scope of research described and approved within the application, including extending the end date or broadening the number of research subjects without prior approval from the approver.

### 4. Conduct of the Researcher(s):

- A. The researcher(s) and/or designated DOC staff will obtain a signed *Research Consent Form* (see attachment #3) from all offenders, staff, interns, or volunteers (hereafter also referred to as "research subjects") selected to participate in social research, prior to conducting any research activity that directly involves the research subject. The applicant may propose an alternative informed consent form by including it as an attachment to the application.
  - 1. If the social research involves research subjects under the age of eighteen (18), signed consent forms are required from the juvenile's parent or guardian.
  - 2. Consent forms are not required when the research does not involve direct contact with a research subject under the authority of the DOC; or does not include publication of personally identifiable information.
  - 3. Research data identifying individual offenders is subject to the same confidentiality and security standards required for case records and personnel files.
- B. The researcher is responsible for ensuring the subject's rights and wellbeing are not compromised through participation or association with the research.
- C. At least one (1) DOC staff member from the Office and Planning and Analysis will be assigned to monitor the research project and oversee the researcher's compliance with all applicable DOC policies and rules, including preservation of protected, privileged information or confidential material provided or acquired through the research.
- D. The principal researcher shall provide an explanation of the research activity, goals, expectations, and purpose to all research subjects prior to participation in the research activity or signing the consent form.
  - 1. A written summary will be prepared to facilitate this explanation. A copy of the explanation shall be provided to each research subject. If the subject is an offender, a copy of the explanation will be included in the offender's institutional record.
  - 2. The explanation will make it clear to all research subjects that participation in the research activity is voluntary and participating or not participating in the research project will in no way impact the subject's incarceration or supervision (offenders) or employment with the DOC (DOC staff members).
  - 3. All fees and costs associated with translation of the explanation and documents/forms for those research subjects with communication disabilities, or those whose primary language is not English, shall be the responsibility of the researcher. No participant may be discriminated against on the basis of a disability.
- E. The principal researcher will maintain or have access to, adequate records showing the current status of the research project, which shall be made available to the SOC or designee upon request.

SECTION	SUBJECT	DOC POLICY	Page 4 of 5
Admin. & Management -	Research Activities	1.1.F.01	Effective:
Information Systems and			07/15/2023
Research			

- F. Data collected during the course of the research activity will only be used in the manner described to the research subject(s), or in a manner subsequently and specifically permitted by the research subject, and as approved by the DOC through acceptance of the application and agreement.
- G. No research subject will receive direct or indirect compensation or special favors for participation in the research or derive any benefit from such participation not expressly described by the researched and approved by the DOC, unless approved by the SOC or designee.
- H. Research data which identifies research subjects shall be used only for specified and approved research or statistical purposes and shall not be revealed for any purpose other than those specified. The privacy and identity of all research subjects who have not provided consent shall be maintained.
  - 1. Identifying information shall not be included in research reports or publications unless specifically agreed to by the research subject(s) and must serve a legitimate purpose relevant to the research.
  - 2. Identifying information shall be maintained under physically secure conditions. Upon termination of the research project, personally identifying information, such as names or ID numbers of the research subject(s) shall be destroyed or otherwise separated from the data.
- I. Researchers visiting DOC facilities or having direct contact with offenders shall be required to submit to a criminal records background check, which shall be conducted by authorized DOC staff prior to the researcher's admission to any DOC facility or having direct contact with DOC offenders. The results shall be reviewed by designated DOC staff and are subject to approval.

### 5. Follow-Up Reports on Research:

- A. A draft copy of the preliminary or final findings/results of the research project may be required by the SOC or designee prior to further dissemination of such, to the public or otherwise.
- B. When the research project is complete, a final report of the findings/results will be provided by the principal researcher to the SOC and/or designated DOC staff for review.
- C. At the discretion of the SOC or designee, the researcher may be required to verify the accuracy of research data collected prior to publication.
- D. The DOC shall be granted access to the data collected though the research project upon request by the SOC or designee.

### 6. Violations of Research Regulations:

- A. Permission to conduct research utilizing DOC employees, interns, volunteers, or offenders may be rescinded, suspended, terminated, or denied if there is reason to believe the research violates state or federal law, DOC policy, agency standards, or the research or associated processes become detrimental to offenders or staff, or compromise the safe, secure, and efficient operation of the facility, or conflicts with the legitimate penological interests of the DOC.
- B. Violations regarding the release of offender record information may subject the violator to civil or criminal penalty (See SDCL §§ 24-2-20, 24-15-1 and 26-7A-120).

#### 7. Medical Research:

SECTION	SUBJECT	DOC POLICY	Page 5 of 5
Admin. & Management -	Research Activities	1.1.F.01	Effective:
Information Systems and			07/15/2023
Research			

A. The use of offenders for medical, pharmaceutical, or cosmetic experiments is prohibited. This does not preclude voluntary offender participation in clinical trials that are approved by clinical services based on the offender's need for specific medical intervention. The institutions permit offender participation in medical or pharmaceutical research. Facilities electing to perform such biomedical research shall be in compliance with all state and federal guidelines and laws [ACA #1-HC-3A-09 (M)].

### 8. Surveys:

- A. Designated DOC staff shall respond in a timely manner to surveys sent to the DOC by an academic or professional organization seeking statistical data.
- B. Completed surveys/responses shall be forwarded to the DOC public information officer (PIO) or designee for inclusion in the department's correspondence database.
- C. Surveys received by the DOC from private, for-profit businesses for commercial purposes shall be responded to by designated DOC staff as time and duties allow. Staff may consider any clear or perceived benefit to the DOC from participation in the survey when prioritizing a response.

#### V. RESPONSIBILITY

The directors of Juvenile Services and Finance and Administration are responsible for the annual review and maintenance of this policy.

#### VI. AUTHORITY

A. SDCL §§ 24-2-20, 24-15-1, 26-7A-120

#### VII. HISTORY

July 2023

January 2022

December 2019

December 2018

December 2017

D 1 2017

December 2016

December 2015 December 2014

December 2013

December 2012

March 2012

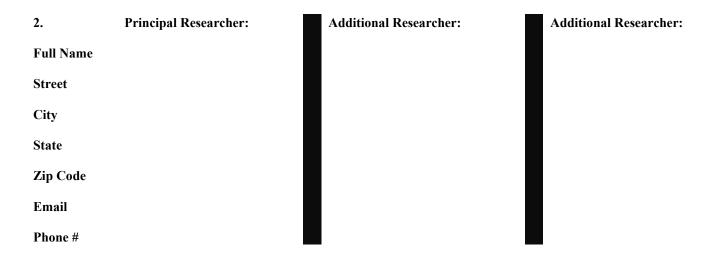
### **ATTACHMENTS** (\*Indicates document opens externally)

- 1. Research Application\*
- 2. Research Agreement\*
- 3. Research Consent Form\*
- 4. DOC Policy Implementation / Adjustments

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# RESEARCH APPLICATION

1. Title of Research Activity:



If additional researchers are needed, please provide their contact information on a separate sheet of paper in the provided format above.

- 3. Research Endorsed By:
- 4. Summary of the goals of the Research Activity:

5. The justification of the Research Activity:

- 6. Details of Research Design:
  - a. DOC resources/personnel needed:

Revised: 07/20/2022
Page 1 of 2

Effective: 07/15/2023

Distribution: Public

- b. Sampling procedures for selecting offender subjects or offender records for this research and any criteria that will be used to determine the sample selection:
- c. Procedures used for data collection and copies of research instruments to be used, including interview schedules, questionnaires, data collection forms and tests.
- d. The security procedures to be followed to protect the privacy and confidentiality of participants:
- e. Details of compensation, if any to be paid:
- f. What are the plans for dissemination of research findings:

All research requests will normally be answered within ten (10) working days of being received by the Secretary of Corrections, Deputy Secretary of Corrections, or designee.

All research will be conducted in accordance with DOC policy 1.1.F.01 Research Activities.

Effective: 07/15/2023

Revised: 07/20/2022 Page 2 of 2

# RESEARCH AGREEMENT

Name of Principal Researcher *					Date	
Address:						
Street or P.O. Bo	ΟX					
City					State	Zip Code
·	(	`				
Home Telephone Number:						
Work Telephone Number:		)				
Cell Phone Number:	_(	)				
E-Mail Address:						
I agree to abide by the following	guide	lines re	egarding my	research act	ivities:	
To follow all policies an	d dire	ctives t	hat apply to	the conducti	ng of research.	
To not change any part of end date, or broadening						
The use and dissemination member) requires a sign						
<ul> <li>To permit the DOC to m any research activity if detrimental to offenders</li> </ul>	there	is rea	son to belie	ve the proj	ect violates po	olicy, law, or becomes
To include in any public DOC's participation in t						
<ul> <li>To submit a plan discu approval prior to the sta project prior to dissemin the DOC upon completion</li> </ul>	art of nation	the res to the I	earch and to OOC. A copy	forward a	draft copy of	the completed research
Signature of Principal Researche	er				Date	
Signature of Department of Corn	rection	s Repr	esentative		Date	

\* This form must also be completed by each member of the research staff.

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## RESEARCH CONSENT FORM

I freely and knowingly agree to participate in the research project entitled				
I fully rea choosing and I agree not to hold the South Dakota Department of liable for any physical or psychological harm that may happen t DOC related to my voluntary participation in the research project.	o me, and I release any claim against the			
Offender I understand my decision to participate or not participate in this research project will have no impact on the terms of my incarceration or supervision and there is no penalty for not participating in the research project.				
DOC Staff Member I understand my decision to participate or not participate in this re employment with the DOC and there is no penalty for not particip				
☐ I consent to having my identity revealed in the research projec	t and any reports.			
☐I DO <u>NOT</u> consent to having my identity revealed in the resear	rch project or any reports.			
I understand that if the DOC is not the sponsor of the research project, the decision as to whether my ide will be protected is up to the researcher and is not under the control of the DOC				
I affirm this research project has been satisfactorily explained t answered. I understand that my participation if voluntary and of m to discontinue participation at any time.				
Printed Name of Offender/Staff	Title (if applicable)			
Signature of Offender/Staff	Date			
Signature of Staff Witness	Date			

Revised: 10/24/2022 Effective: 07/15/2023