

**WHAT WORKS CLEARINGHOUSE  
REGISTRY OF RANDOMIZED CONTROLLED TRIALS  
REGISTRATION FORM**

Thank you for choosing to register a study for the What Works Clearinghouse Registry of Randomized Controlled Trials (RCTs). Please note that some of the data items are mandatory and others are voluntary. If you have questions about filling out this form, or prefer to submit this information in paper format, please contact us at:

What Works Clearinghouse  
P.O. Box 2393  
Princeton, NJ 08543-2393  
Email: <http://ies.ed.gov/ncee/wwc/help/webmail/>  
Phone: (866) 503-6114  
Fax: (609) 799-0005

When your registration is complete, we will list you on the public site.

**Paperwork Burden Statement**

According to the Paperwork Reduction Act of 1995, no persons are required to respond to a collection of information unless such collection displays a valid Office of Management and Budget (OMB) control number. The valid OMB control number for this information collection is 1850-0788, which is valid through October 31, 2012. The time required to complete this information collection is estimated to average 30 minutes per response, including the time to review instructions, search existing data resources, gather the data needed, and complete and review the information collection. **If you have any comments concerning the accuracy of the time estimate(s) or suggestions for improving this form, please write to:** U.S. Department of Education, Washington, DC 20202-4651. If you have comments or concerns regarding the status of your individual submission of this form, contact the What Works Clearinghouse at <http://ies.ed.gov/ncee/wwc/help/webmail/>.

**Note: Required items are indicated with an asterisk (\*).**

**Contact Information:**

\*Researcher Name (include prefix): \_\_\_\_\_

\*Title: \_\_\_\_\_

\*Address 1: \_\_\_\_\_

Address 2: \_\_\_\_\_

\*City: \_\_\_\_\_ \*State: \_\_\_\_\_ \*ZIP: \_\_\_\_\_

Country: \_\_\_\_\_

\*Phone: (     )     -                      Fax (If None, write N/A): (     )     -

\*E-mail: \_\_\_\_\_

URL: \_\_\_\_\_

Organization 1 (if applicable): \_\_\_\_\_

Organization 2 (if applicable): \_\_\_\_\_

**\*Study Status:**

Funded and In Progress (complete pages 3–6)

Completed with Available Final Report (complete pages 7–10)

**RANDOMIZED CONTROLLED TRIALS DATA COLLECTION FORM  
FOR STUDIES FUNDED AND IN PROGRESS**

**Study Information:**

\*RCT Title: \_\_\_\_\_

\*Principal Investigator First Name: \_\_\_\_\_

\*Principal Investigator Last Name: \_\_\_\_\_

\*Prefix: \_\_\_\_\_

Principal Investigator Title: \_\_\_\_\_

Other Key Staff: \_\_\_\_\_

\*Start Date (MM/DD/YYYY):    /    /

\*Anticipated End Date (MM/DD/YYYY):    /    /

\*Sponsoring Organization (include grantee number where applicable): \_\_\_\_\_

\_\_\_\_\_

**Study Abstract Information:**

Related Publications (Citations in APA Format). *Include all publications related to this study that are publicly available. The final report should not be listed here.*

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\_\_\_\_\_

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\_\_\_\_\_

\*1. What is the study setting? *Include settings and locations where the data will be collected, as well as relevant eligibility criteria for participants.* \_\_\_\_\_

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\*2. What is the intervention? *Include intervention type, name(s), and description. Describe how the intervention is designed to be delivered to each group.* \_\_\_\_\_

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**Sample Characteristics:**

\*3. What is the randomized unit?

- Cluster
- Individual

4. Student Characteristics. *Below is a list of target populations identified by the What Works Clearinghouse. Select all that apply for the students for this RCT.*

**\*Gender**

- Female
- Male
- Not Applicable

**\*Student Race/Ethnicity**

- American Indian or Alaska Native
- Asian
- Black or African American
- White
- Hispanic or Latino
- Native Hawaiian or Other Pacific Islander
- Not Applicable

**\*Student Level(s) of Education**

- Preschool
- Kindergarten
- Elementary School
- Middle School
- High School
- Postsecondary Education
- Vocational/Career Education
- Adult/Continuing Education
- Not Applicable

**\*Student Disability**

- Autism
- Deaf-Blindness
- Emotional Disturbance (ED)
- Hearing Impairment including Deafness
- Mental Retardation
- Multiple Disabilities
- Orthopedic Impairment
- Other Health Impairment (OHI)
- Specific Learning Disability (LD)
- Speech or Language Impairment
- Traumatic Brain Injury
- Visual Impairment and Blindness
- Not Applicable

\*5. Additional Study Sample Information. *Include the expected size of the sample and any additional characteristics not described in question 4. Describe attrition that has occurred at this point, as well as any interim analyses.* \_\_\_\_\_

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**Design and Analysis**

\*6. What research design and methods will be used? *Include a description of any alterations to the sample after random assignment. Also note the statistical methods intended to compare compatibility of groups on primary outcome(s) and methods for additional analyses.* \_\_\_\_\_

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\*7. What is the counterfactual (control) condition? \_\_\_\_\_

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\*8. What is the data analytic strategy? *Include information about power level analysis and plans for estimating effects of the intervention, including calculation of effect sizes.* \_\_\_\_\_

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**Outcome and Summary**

\*9. What are the intended primary outcomes? *Include all relevant measures, subgroups, and time periods (e.g., Reading test scores for all female students in second grade one year after intervention), and list any planned subgroup analyses.* \_\_\_\_\_

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\*10. What are the intended secondary outcomes? *See question 9 for instructions.* \_\_\_\_\_

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\*11. Summary of the Study/Abstract. *State specific objectives and hypotheses.* \_\_\_\_\_

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**RANDOMIZED CONTROLLED TRIALS DATA COLLECTION FORM  
FOR STUDIES COMPLETED WITH AN AVAILABLE FINAL REPORT**

**Study Information:**

\*RCT Title: \_\_\_\_\_

\*Principal Investigator First Name: \_\_\_\_\_

\*Principal Investigator Last Name: \_\_\_\_\_

\*Prefix: \_\_\_\_\_

Principal Investigator Title: \_\_\_\_\_

Other Key Staff: \_\_\_\_\_

\*Start Date (MM/DD/YYYY):    /    /                      \*End Date (MM/DD/YYYY):    /    /

\*Sponsoring Organization (include grantee number where applicable): \_\_\_\_\_

**Study Abstract Information:**

Web Address for Report: \_\_\_\_\_

\*Published Report Citation (APA Format): \_\_\_\_\_

Related Publications (Citations in APA Format). *Include all publications related to this study that are publicly available. The final report should not be listed here.*

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\*1. What is the study setting? *Include settings and locations where the data was collected, as well as relevant eligibility criteria for participants.* \_\_\_\_\_

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\*2. What is the intervention? *Include intervention type, name(s), and description. Describe how and when the intervention was delivered to each group.* \_\_\_\_\_

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**Sample Characteristics:**

\*3. What was the randomized unit?

- Cluster
- Individual

4. Student Characteristics. *Below is a list of target populations identified by the What Works Clearinghouse. Select all that apply for the students for this RCT.*

**\*Gender**

- Female
- Male
- Not Applicable

**\*Student Race/Ethnicity**

- American Indian or Alaska Native
- Asian
- Black or African American
- White
- Hispanic or Latino
- Native Hawaiian or Other Pacific Islander
- Not Applicable

**\*Student Level(s) of Education**

- Preschool
- Kindergarten
- Elementary School
- Middle School
- High School
- Postsecondary Education
- Vocational/Career Education
- Adult/Continuing Education
- Not Applicable

4. Student Characteristics (continued).

**\*Student Disability**

- Autism
- Deaf-Blindness
- Emotional Disturbance (ED)
- Hearing Impairment including Deafness
- Mental Retardation
- Multiple Disabilities
- Orthopedic Impairment
- Other Health Impairment (OHI)
- Specific Learning Disability (LD)
- Speech or Language Impairment
- Traumatic Brain Injury
- Visual Impairment and Blindness
- Not Applicable

*\*5. Additional Study Sample Information. Include the sample size and any additional characteristics not described in question 4. Describe attrition that occurred, as well as any interim analyses.* \_\_\_\_\_

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**Design and Analysis**

*\*6. What research design and methods were used? Include a description of any alterations to the sample after random assignment. Also note the statistical methods used to compare compatibility of groups on primary outcome(s) and methods for additional analyses.* \_\_\_\_\_

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*\*7. What was the counterfactual (control) condition?* \_\_\_\_\_

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\*8. What was the data analytic strategy? *Include information about power level analysis and effect size.* \_\_\_\_\_

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### **Outcome and Summary**

\*9. What were the primary outcomes? *For each outcome, include a summary of results of each group and the estimated effect size and level of power analysis. Include all relevant measures, subgroups, and time periods, as well as the number of participants in each group included in each analysis.* \_\_\_\_\_

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\*10. What were the secondary outcomes? *See question 9 for instructions.* \_\_\_\_\_

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\*11. Summary of the Study/Abstract. *State specific objectives and hypotheses.* \_\_\_\_\_

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\*12. Interpretation of Results/Discussion. *Provide a brief discussion of findings. Clearly state any limitations and sources of bias.* \_\_\_\_\_

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