DEPARTMENT OF HEALTH AND HUMAN SERVICES

Agency for Healthcare Research and Quality

Supplemental Evidence and Data Request on Psychosocial and Pharmacologic

Interventions for Disruptive Behavior in Children and Adolescents

AGENCY: Agency for Healthcare Research and Quality (AHRQ), HHS.

ACTION: Request for Supplemental Evidence and Data Submissions

SUMMARY: The Agency for Healthcare Research and Quality (AHRQ) is seeking scientific information submissions from the public. Scientific information is being solicited to inform our review on *Psychosocial and Pharmacologic Interventions for Disruptive Behavior in Children and Adolescents*, which is currently being conducted by the AHRQ's Evidence-based Practice Centers (EPC) Program. Access to published and unpublished pertinent scientific information will improve the quality of this review.

DATES: Submission Deadline on or before [INSERT DATE 30 DAYS AFTER DATE OF PUBLICATION IN THE FEDERAL REGISTER].

ADDRESSES:

E-mail submissions: epc@ahrq.hhs.gov

Print submissions:

Mailing Address:

Center for Evidence and Practice Improvement

Agency for Healthcare Research and Quality

ATTN: EPC SEADs Coordinator

5600 Fishers Lane

Mail Stop 06E53A

Rockville, MD 20857

Shipping Address (FedEx, UPS, etc.):

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5600 Fishers Lane

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Rockville, MD 20857

FOR FURTHER INFORMATION CONTACT:

Kelly Carper, Telephone: 301-427-1656 or Email: epc@ahrq.hhs.gov.

SUPPLEMENTARY INFORMATION:

The Agency for Healthcare Research and Quality has commissioned the Evidence-based

Practice Centers (EPC) Program to complete a review of the evidence for *Psychosocial and*

Pharmacologic Interventions for Disruptive Behavior in Children and Adolescents. AHRQ is

conducting this systematic review pursuant to section 902 of the Public Health Service Act, 42

U.S.C. 299a.

The EPC Program is dedicated to identifying as many studies as possible that are relevant

to the questions for each of its reviews. In order to do so, we are supplementing the usual

manual and electronic database searches of the literature by requesting information from

the public (e.g., details of studies conducted). We are looking for studies that report on

Psychosocial and Pharmacologic Interventions for Disruptive Behavior in Children and

Adolescents, including those that describe adverse events. The entire research protocol is

available online at: https://effectivehealthcare.ahrq.gov/products/disruptive-

behavior/protocol

This is to notify the public that the EPC Program would find the following information on

Psychosocial and Pharmacologic Interventions for Disruptive Behavior in Children and

Adolescents helpful:

- A list of completed studies that your organization has sponsored for this indication. In the list, please *indicate whether results are available on* ClinicalTrials.gov along with the ClinicalTrials.gov trial number.
 - For completed studies that do not have results on ClinicalTrials.gov, a summary, including the following elements: study number, study period, design, methodology, indication and diagnosis, proper use instructions, inclusion and exclusion criteria, primary and secondary outcomes, baseline characteristics, number of patients screened /eligible /enrolled /lost to follow-up /withdrawn /analyzed, effectiveness/efficacy, and safety results.
- A list of ongoing studies that your organization has sponsored for this indication. In the list, please provide the ClinicalTrials.gov trial number or, if the trial is not registered, the protocol for the study including a study number, the study period, design, methodology, indication and diagnosis, proper use instructions, inclusion and exclusion criteria, and primary and secondary outcomes.
- Description of whether the above studies constitute ALL Phase II and above
 clinical trials sponsored by your organization for this indication and an index
 outlining the relevant information in each submitted file.

Your contribution is very beneficial to the Program. Materials submitted must be publicly available or able to be made public. Materials that are considered confidential; marketing materials; study types not included in the review; or information on indications not included in the review cannot be used by the EPC Program. This is a voluntary request for information, and all costs for complying with this request must be borne by the submitter.

The draft of this review will be posted on AHRQ's EPC Program website and available for public comment for a period of 4 weeks. If you would like to be notified when the draft is posted, please sign up for the e-mail list at: https://www.effectivehealthcare.ahrq.gov/email-updates.

The systematic review will answer the following questions. This information is provided as background. AHRQ is not requesting that the public provide answers to these questions.

Key Questions (KQ)

- **KQ 1:** In children under 18 years of age diagnosed with disruptive behaviors, which psychosocial interventions are more effective for improving short-term and long-term psychosocial outcomes compared to no treatment or other psychosocial interventions?
- **KQ 2:** In children under 18 years of age diagnosed with disruptive behaviors, which pharmacologic interventions are more effective for improving short-term and long-term psychosocial outcomes compared to placebo or other pharmacologic interventions?
- **KQ 3:** In children under 18 years of age diagnosed with disruptive behaviors, what is the relative effectiveness of psychosocial interventions alone compared with pharmacologic interventions alone for improving short-term and long-term psychosocial outcomes?
- **KQ 4:** In children under 18 years of age diagnosed with disruptive behaviors, are combined psychosocial and pharmacologic interventions more effective for improving short-term and long-term psychosocial outcomes compared to either psychosocial or pharmacologic interventions alone?

KQ 5: What are the harms associated with treating children under 18 years of age for disruptive behaviors with either psychosocial, pharmacologic or combined interventions?

KQ 6a: Do interventions for disruptive behaviors vary in effectiveness and harms based on patient characteristics, including gender, age (including pubertal changes and use of oral contraceptives), racial/ethnic minority, LGBTQ+ status, English proficiency, health literacy, socioeconomic status, insurance status, rural versus urban, developmental status or delays, family history of disruptive behavior disorders or other mental health disorders, prenatal use of alcohol and drugs (specifically methamphetamine), history of trauma or Adverse Childhood Experiences (ACEs), parental ACEs, access to social supports (neighborhood assets, family social support, worship community, etc.), personal and family beliefs about mental health (e.g. stigma around mental health), or other social determinants of health?

KQ 6b: Do interventions for disruptive behaviors vary in effectiveness and harms based on clinical characteristics or manifestations of the disorder, including specific disruptive behavior (e.g., stealing, fighting) or specific disruptive behavior disorder (e.g., oppositional defiant disorder, conduct disorder), co-occurring behavioral disorders (e.g., attention deficit hyperactivity disorder, autism spectrum disorder, internalizing disorders), related personality traits and symptom clusters, presence of non-behavioral comorbidities, age of onset, and duration?

KQ 6c: Do interventions for disruptive behaviors vary in effectiveness and harms based on treatment history of the patient?

KQ 6d: Do interventions for disruptive behaviors vary in effectiveness and harms based on characteristics of treatment, including setting (e.g., group homes, residential treatment, family setting), duration, delivery, timing, and dose?

Contextual Question 1. What are the disparities in the diagnosis of disruptive behavior disorders (based on characteristics such as gender, race/ethnicity, socioeconomic status, other social determinants of health, or other factors) in children and adolescents?

Contextual Question 2. What are the disparities in the treatment of disruptive behaviors or disruptive behavior disorders (based on characteristics such as gender, race/ethnicity, socioeconomic status, other social determinants of health, or other factors) in children and adolescents?

Contextual Question 3. How do disparities in the diagnosis and treatment of disruptive behaviors or disruptive behavior disorders affect behavioral and functional outcomes (e.g., compliance with teachers, contact with the juvenile justice system, substance abuse)?

Population, Intervention, Comparator, Outcome, Timing, Setting/Study Design (PICOTS)

PICOTS	Inclusion	Exclusion
Population	KQs 1-6. Children under 18 years of age who are being treated for disruptive behavior or a disruptive behavior disorder that includes oppositional defiant disorder, conduct disorder, and intermittent explosive disorder; children with a co-occurring diagnosis (e.g., ADHD, ASD) provided the disruptive behavior treated is due to a DBD will be included	Asymptomatic children At-risk children Treatment of disruptive behavior secondary to other conditions (e.g., substance abuse, developmental delay, intellectual disability, pediatric bipolar disorder, ADHD)
Interventions	KQs 1, 3-6. Psychosocial interventions for child, parents/family or both including: - Social skills training - Functional behavioral interventions - Parent training - Psychotherapy (e.g., cognitive behavior therapy, interpersonal psychotherapy, psychodynamic therapy, dialectical behavior therapy, equine-assisted psychotherapy with mental health provider) - Contingency management methods - Behavior management training KQs 2-6. Pharmacologic interventions that are FDA approved medications used on or off label, including the following class of drugs: - Alpha-agonists - Anticonvulsants - Second-generation (i.e., atypical) antipsychotics	 Preventive interventions for atrisk populations Preventive interventions for caregiver health Interventions that do not target disruptive behaviors Specialized diet or dietary supplements Speech, occupational, physical therapy Complimentary and Integrative Health interventions (e.g., acupuncture, herbal remedies) Exercise programs as the sole intervention Massage, chiropractic care

	- Beta-adrenergic blocking agents (i.e., beta-	- Invasive medical interventions
	blockers) - Central nervous system stimulants - First-generation antipsychotics	(e.g., surgery, deep brain stimulation)
	Selective serotonin reuptake inhibitors Selective norepinephrine reuptake inhibitors Mood stabilizers	
	- Antihistamines KQs 4-6. Combined psychosocial and	
Comparators	pharmacologic interventions included for KQs 1-3. - Other included psychosocial and/or	No comparison group, excluded
Comparators	pharmacologic interventions - Inactive treatment, including waitlist control, no	interventions
0.4	treatment and placebo KQs 1-4, 6. Behavioral outcomes:	Unvalidated outcomes measures
Outcomes	 Aggressive behavior Temper outbursts (not considered ageappropriate) Violent behavior 	Chyandated outcomes measures
	Delinquent behavior Fighting, property destruction, and rule violations	
	 Compliance with parents, teachers, and institutional rules Affective or mood elements of DBD 	
	- Treatment satisfaction - Other patient-centered outcomes	
	KQs 1-4, 6. Functional outcomes: - Family functioning/cohesion - School performance/attendance - Interpersonal/social function and competence/need for special accommodations - Interactions with legal/juvenile justice systems - Out of home placement - Health care system utilization - Substance abuse - Parenting stress - Logistical family outcomes (days of work lost, etc.) - Health-related quality of life (e.g., mental health, physical health) - Other patient-centered outcomes	
	 KQ 5. Adverse effects/harms: Metabolic effects: weight gain, hyperglycemia and diabetes, hyperlipidemia Extrapyramidal effects: parkinsonism, acute dystonia, akathisia, tardive dyskinesia Cardiac adverse effects: prolonged QT/arrhythmias, hypotension, cardiomyopathy Prolactin-related effects 	
	 Neutropenia as a potential adverse effect of atypical antipsychotics Allergic reaction Sleep disruption, fatigue Sudden death Suicide 	
	 Over-medication or inappropriate medication Negative effects on family dynamics Acne 	
	Stigma Harms/barriers to utilization of care related to psychosocial interventions (e.g., time investment, limited access to trained providers, and lower acceptability based on a misperception that family-focused psychosocial	

Timin	interventions carry implicit judgements about the quality of their parenting). - Study withdrawal due to medication adverse effects	
Timing	KQs 1-6. Any length of follow-up	
Setting	KQs 1-6. Clinical setting, including medical or psychosocial care that is delivered to individuals by clinical professionals (including telehealth), as well as individually focused programs to which clinicians refer their patients; may include classroom settings when intervention is directed to treat disruptive behavior(s) in a specific child (not the whole class) as part of that child's treatment plan	Exclude school wide or system wide settings (e.g., juvenile justice system) wherein interventions are targeted more widely
Study Design	Randomized controlled trials (no sample size limit), comparative nonrandomized controlled trials that adjust for confounding variables (N≥100), published in English on or after 1994.	Published before 1994

Abbreviations: ADHD=Attention-deficit/hyperactivity disorder; ASD=Autism Spectrum Disorder; DBD=Disruptive Behavior Disorders; FDA=U.S. Food and Drug Administration; KQ=Key Question

Dated: March 30, 2023.

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