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Effectiveness of Motivational Interviewing-Enhanced Behavior Therapy for Adolescents With Attention-Deficit/Hyperactivity Disorder: A Randomized Community-Based Trial

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Abstract

Objective: This study tests the effectiveness of parent-teen psychotherapy for adolescent ADHD (Supporting Teens' Autonomy Daily; STAND) versus Usual Care (UC) in four community clinics.

Method: A randomized clinical trial was conducted with double randomization of adolescents and therapists to STAND vs. UC. Participants were 278 culturally-diverse adolescents diagnosed with *DSM-5* ADHD at baseline and 82 community therapists. Seven primary outcomes were assessed at baseline (BL), post-treatment (PT; $M=5.11$ months post-BL, $SD=2.26$), and follow-up (FU; $M=9.81$ months post-BL, $SD=2.50$): inattention (IN; parent/teacher-rated), academics (parent-rated/official records), family functioning (parent/adolescent-rated), and disciplinary records. Treatment engagement indicated consumer fit (e.g., number or sessions received, percentage of sessions attended by parent, satisfaction). The impact of treatment on concurrent

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Disclosure:

Dr. Sibley has received royalties from Guilford Press for the treatment manual described in this paper. She has received royalties from Vimeo, Inc. for STAND training videos and has provided consultation to Takeda Pharmaceuticals. Drs. Graziano, Coxé, Bickman and Mr. Martin have reported no biomedical financial interests or potential conflicts of interest.

medication use was also examined. Service delivery features were examined as moderators of outcome.

Results: Intent to treat ($N=278$) analyses indicated no significant group $\times$ time effects. STAND only led to superior outcomes when therapists were licensed (22% of sample) vs. unlicensed (parent-rated IN: $p<.001$, $d=1.08$; parent-rated academic impairment: $p=.010$, $d=1.17$). Compared to UC, STAND was associated with greater parent participation ($p<.001$, $d=.88$) and higher scores on certain indices of parent satisfaction. STAND also was associated with superior medication engagement over time compared to UC ($OR=7.18$).

Conclusion: Evidence-based psychosocial treatment for adolescent ADHD did not outperform UC on outcome trajectories despite improving some indices of treatment engagement. STAND requires additional adaptation for community contexts.

Clinical trial registration information: STAND Community Trial (STAND); clinicaltrials.gov; [NCT02694939](https://clinicaltrials.gov/ct2/show/study/NCT02694939).

Keywords

ADHD; community-based treatment; psychotherapy; adolescence

Introduction

Practice parameters recommend medication as first-line treatment for adolescent Attention-deficit/hyperactivity disorder (ADHD) with psychosocial treatment encouraged as needed^{1,2}. Despite the efficacy of both approaches, adolescents with ADHD access far fewer services than children in their communities^{3–5}. Medication is the cornerstone childhood ADHD treatment; however, adolescents often discontinue ADHD medication perceiving adverse effects, stigma, and ineffectiveness^{6–7}. Adolescents with ADHD appear willing to engage in psychosocial treatments^{8–9}. However, evidence-based practices (EBPs) are costlier and more burdensome to implement than medication and usual care psychological services^{10,11}; as a result, they are offered in few community clinics¹². To bridge this gap, research should adapt and refine adolescent ADHD EBPs to fit the constraints of community contexts, evaluating their implementation and effectiveness. In this trial, we utilize a Hybrid Type 1 implementation-effectiveness design¹³ to evaluate an EBP for adolescent ADHD (Supporting Teen's Autonomy Daily; STAND)^{14–16} compared to usual care at four community clinics. Implementation outcomes are previously reported.¹⁷

STAND

STAND is a ten-session manualized EBP that targets ADHD symptoms and related impairments (i.e., family conflict, organization skills, homework problems) and is delivered individually to parent-teen dyads¹⁸. Like other EBPs for adolescent ADHD, STAND directly teaches skills (e.g., organization and time management, communication) and engages adults in autonomy-supportive roles—not merely supervisors of behavior modification^{18,19}. Because patient barriers often prompt premature disengagement from adolescent ADHD treatment²⁰, STAND includes engagement-focused components: (1) parent-teen collaboration, (2) Motivational Interviewing (MI)²¹, (3) modular treatment²², and (4)

strength-based, autonomy-supportive activities. STAND is divided into engagement, skills, and planning sessions. Population-relevant process issues are addressed through MI (i.e., motivation deficits, inconsistent family routines, intrusive parenting, regulating electronics, skepticism about behavioral techniques)²³.

Three randomized clinical trials (RCTs) of STAND demonstrate efficacy in university settings^{14–16}. In a pilot RCT ($N=36$), STAND was delivered with high fidelity (i.e., 93%-100% content fidelity across sessions) and was highly acceptable to families. Compared to treatment as usual, large effects were present for ADHD and ODD symptoms ($d=.82$ to 1.42), academic indices ($d=1.30$ to 5.15), and parent-teen conflict ($d=.65$) from baseline to post-treatment¹⁴. A larger RCT ($N=128$) demonstrated full or partial maintenance of similar effects at six-month follow-up ($d=.63$ to 1.01) and detected effects on parental outcomes including parenting stress ($d=.60$) and use of behavioral strategies ($d=.49$ to 1.07)¹⁵. A third RCT ($N=123$) compared STAND to standard group behavioral parent training and teen organization skills training and found superior effects for STAND when parents possessed elevated ADHD ($d=.50$) or depression symptoms ($d=.64$) or dyads displayed high parent-teen conflict ($d=.51$)¹⁶.

Community-Based STAND Implementation Outcomes

Recently, we reported implementation outcomes for this trial¹⁷. Adolescents with ADHD ($N=278$) were randomly assigned to receive psychosocial treatment from agency therapists who also were randomized either to delivery usual care psychotherapy or receive training and supervision in STAND. Approximately 96% of therapists assigned to STAND completed the three-day training and, on average, attended 87% of scheduled weekly supervision sessions. Although therapists rated STAND as highly acceptable, relevant, and lower burden than UC practices ($d=.40$ -.86), fidelity was poorer than in university-based trials^{14–16}. Average community-based fidelity ranged from 85% in skills sessions to 24% in planning sessions¹⁷. Therapists delivered STAND at a slower pace and lower intensity than the manual intended. However, they also demonstrated significantly higher MI integrity scores compared to UC therapists ($d=.21$ to $.79$) and delivered STAND with high fidelity in office-based and early sessions. Thus, implementation of STAND in community contexts was promising but requires refinement to improve aspects of fidelity.

Aim 1: Effect of Community-Based STAND on Symptoms and Impairment

The current study's primary aim is to test whether STAND improves patient outcome trajectories (i.e., ADHD symptoms, academic, family, and behavioral functioning) compared to UC. Due to previously documented fidelity disruptions²², we hypothesize that effectiveness will decrease when STAND is delivered in community contexts (versus past university trials) but will exceed UC. Although standard EBPs for youth psychiatric disorders often do not outperform UC,^{24–25} STAND includes elements that promote engagement and effectiveness in community settings^{21–22,26}. STAND also outperforms standard behavior therapy when families are clinically complex, which is characteristic of community contexts^{27–28}. Thus, even a diluted dose of STAND may outperform UC.

Aim 2: Therapeutic Engagement

We also examined whether STAND increases therapeutic engagement (i.e., improved retention in services, parent participation, stakeholder satisfaction), including concurrent medication utilization. Because of its engagement-focused approach, we hypothesized that community-delivered STAND would be associated with higher retention in services, greater parent participation, and higher levels of stakeholder satisfaction across targeted domains of treatment. We also hypothesized that engagement in concurrent medication treatment would be enhanced by STAND because STAND targets facilitators of adherence (i.e., oppositional behavior, consistency of daily routine, teen motivation to succeed).²⁹

Aim 3: Service-Delivery Features that Moderate of Effectiveness

If STAND demonstrates reduced effectiveness compared to university trials, identification of service delivery features (i.e., therapist characteristics, treatment setting, whether treatment is combined with medication) that moderate effectiveness can guide refinement of STAND's community implementation strategy. For example, if clinician years of experience influences outcomes, future implementation efforts might focus on skill development. If indices of therapist autonomy (e.g., achievement of clinical licensure) predict effectiveness, therapist empowerment or motivational strategies might increase engagement in STAND delivery³⁰. Given this study's culturally diverse context, we also hypothesized that therapist-patient ethnicity match might enhance treatment outcomes—signaling a need to pair adolescents with therapist of similar cultural backgrounds³¹.

Childhood EBPs for ADHD demonstrate reduced patient engagement when delivered subsequent to ADHD medication³². If the same finding stands in adolescence, STAND might be maximally effective when implemented with unmedicated teens. However, if ADHD medication use is associated with stronger treatment response, combined treatment might be optimal for adolescents in community settings. Additionally, if patient engagement indices are significant treatment moderators, implementation efforts might enhance STAND's engagement strategies. Finally, if therapy is more effective in office-based (vs. home-based) sessions, future efforts might address barriers to effective home-based care.

Method

All procedures were approved by the Florida International University Institutional Review Board. All parents, therapists, and adolescents signed consent/assent documents prior to participating.

Participants

Adolescents.—Adolescents ($N=278$; ages 11–17) were incoming patients at four community agencies in a large pan-Latinx and pan-Caribbean U.S. city (see Figure 1). They were required to meet full DSM-5 ADHD criteria (see Supplement 1, available online). Autism spectrum disorder and intellectual disability ($IQ<70$) were exclusionary. Adolescents were randomly assigned to STAND or UC using a stratified randomization procedure within agency. Randomization occurred after agency and study intake and before initiation of

treatment at the agency (see Supplement 1, available online). Table 1 presents sample demographic characteristics. There were no significant group differences on any variable.

Therapists.—Therapists ($N=82$) were mental health professionals employed at four agencies. Therapists self-identified as 19.8% non-Hispanic White ($n=16$), 14.8% Black or African-American ($n=12$), 64.2% Hispanic ($n=53$), and 1.2% Other ($n=1$). 86.6% were female therapists ($n=71$), with 61.0% ($n=50$) offering treatment in both Spanish and English. 22.0% of therapists ($n=18$) were licensed and 86.6% ($n=71$) held a master's degree [7.3% held a doctorate ($n=6$) and 6.1% were bachelor's level interns ($n=5$)]. On average, clinicians reported 5.24 years delivering therapy ($SD=5.00$). STAND ($n=44$) and UC therapists ($n=38$) did not differ on any of the background variables noted above.

Procedures

Recruitment and Intake.—At agency intake, agency staff provided study information to parents of 6th-12th grade students with attention, organization, motivation, or behavior problems. Parents signed a permission to contact form and study staff administered an eligibility screen by phone that queried ADHD symptoms, impairment, exclusionary criteria, and treatment priority²⁵. If another presenting problem (e.g., anxiety, substance use) took priority over ADHD, the teen was not eligible. Students with at least four inattention (IN) or hyperactivity/impulsivity (H/I) symptoms according to the screen attended a full diagnostic assessment to evaluate inclusion criteria. The study intake included an IQ screener (Wechsler Abbreviated Scale of Intelligence-2nd Edition)³³ and parent-administered Diagnostic Interview Schedule for Children (DISC)³⁴.

Therapist Recruitment.—Detailed information about therapist recruitment can be found in Supplement 1, available online. All therapists were randomly assigned to STAND or UC at baseline.

Intervention Content.—STAND is a manualized engagement-focused psychosocial treatment for adolescent ADHD. STAND consists of 10 weekly 60-minute sessions attended by the adolescent and parent¹⁶. Skill instruction is blended with MI and guided parent-teen behavioral contracting¹⁷. Treatment targets family, behavioral, and academic impairment. Treatment is modular to promote flexibility and treatment tailoring. In the engagement phase, MI increases awareness of personal values and goals, identifies strengths, and recognizes ways to achieve personal goals and act consistently with values. The skills phase teaches parent-teen communication, parent behavioral strategies, and organization, time management and planning skills applied to homework, school, and chores. Planning sessions teach families to integrate skills into a daily routine, transfer new habits to school settings, and build a final parent-teen contract. MI in the final session promotes maintenance of change.

Therapist Procedures.—Therapy was delivered across three years. Duration of treatment varied naturalistically to avoid built-in between-group dose differences. Participating therapists treated an average of 2.74 study participants (range: 0 to 14). Study interventions were provided by agency employees using typical billing procedures. Therapists randomized

to STAND were offered a three-day training and 30-minutes of weekly supervision while treating study participants. Every 12 months, a four-hour booster training was provided. STAND therapists were provided with a treatment manual and a family workbook for each participant. Therapists in both groups were instructed to utilize usual care procedures for termination, allowing STAND therapists to continue treatment after completing STAND manualized content. UC therapists were instructed to treat study participants using usual procedures in the agency and the treatments they believed would be most effective for the youth. They received weekly supervision for study participants from agency supervisors according to typical agency practices. UC therapists were offered STAND training at study conclusion.

Treatment Differentiation.—Therapists in both groups were asked to provide sample audio-recorded sessions for each participant (available for 70.2% of treated sample). Fidelity data for STAND are reported in detail elsewhere¹⁷. Coding of 78 available UC audio tapes using STAND fidelity checklists indicated high treatment differentiation (53.8% of items were not present on any UC recordings). Seven items were present on greater than 5% of recordings and typically represented non-specific therapy activities: discussion of presenting problems (43.8%), cognitive behavioral strategies for emotion regulation (8.2%), reviewing therapy homework (8.2%), reviewing progress on goals (5.5%), instruction in daily planner utilization (9.6%), instruction in time management skills (6.8%), and recognizing positive client changes (5.5%).

Data Collection.—Participants were permitted to utilize naturalistic stimulant medication during the study; all medications were monitored and controlled for in analyses. Because therapy duration was allowed to vary naturalistically, PT assessments were scheduled for 16 weeks after the participant's first session at the agency, which provided ample time for families to complete the 10 session STAND protocol with assumed cancellations. On average, PT assessments occurred 5.11 months after BL ($SD=2.26$). FU assessments were attempted at approximately 12 weeks after PT. On average, FU assessments occurred approximately 4.70 months after PT ($SD=2.50$). Retention was 99.3% ($n=276$) at PT and 97.5% ($n=271$) at FU (data provided by at least one informant). Academic records and teacher ratings were obtained directly from schools. Electronic health records were accessed directly. Parent ratings were available in Spanish or English. Teachers and therapists received \$20 and families received \$100 for each assessment.

Primary Outcome Measures

ADHD Symptoms.—Parent and teacher reports of IN and HI symptom severity were measured on the Conners 3 Parent Short Form Rating Scale (C3RS)³⁵ and parent and teacher DSM-5 ADHD checklist³⁶. Respondents rated symptoms on both scales as 0 (*not at all*) to 3 (*very much*). Symptom severity scale scores was the mean level (0–3) of subscale items. For the C3RS, scale scores were converted into T-scores based on age and gender for IN (five items) and HI (six items). The DSM-5 checklist contains nine items per subscale that correspond with DSM-5 items. Psychometric properties of both measures are very good, with empirical support for an internally consistent subscales^{36–37}. In this sample, ADHD subscale alphas ranged from .86–.92.

Academic Impairment.—Two indices of academic impairment were Grade Point Average (GPA) and parent report of academic organization, time management, and planning (OTP) problems. Report cards were obtained directly from the school district. GPA for each academic quarter was calculated by converting class grades (e.g., English, Math) to a 5-point scale (i.e., 4.0=A to 0.0=F). At each assessment, GPA was calculated for the immediately preceding academic quarter. The 24-item parent *Adolescent Academic Problems Checklist* (AAPC) measures observable secondary-school specific OTP problems and is validated in samples of adolescents with ADHD³⁸. The AAPC possesses two distinct factors and a total score, with strong internal reliability and concurrent validity³⁸. In this study, total score was used ($\alpha=.91$).

Family Impairment.—For family impairment, the parent and adolescent versions of the *Conflict Behavior Questionnaire-20* (CBQ-20) assessed parent-teen conflict³⁹. Informants rated statements about the parent-teen relationship on a five-point scale from 1-*strongly agree* to 5-*strongly disagree*. In this study, alpha ranged from .92 to .93.

Disciplinary Incidents.—The school district provided records of student disciplinary incidents. Counts of all disciplinary incidents (e.g., detention, in-school suspension) were calculated for the academic quarter immediately preceding each assessment.

Measures of Treatment Engagement

Psychosocial Treatment Engagement.—Electronic health records at each agency were accessed for all participants for a 12-month period beginning with the baseline assessment. Information was collected about the dates and durations of all therapy sessions (including STAND sessions), who attended, who provided therapy, dates of cancellations, and the location that treatment was provided. Parent treatment participation was calculated by dividing the total number of sessions the parent attended by the total of number of sessions the youth received.

Stakeholder Satisfaction.—At PT, parents and adolescents completed 20-item treatment utility scales designed to measure stakeholder satisfaction (i.e., perceptions of *how* ADHD treatment was helpful)¹⁶. Parents and adolescents indicated level of agreement with statements on a 5-point scale (1=strongly disagree to 5=strongly agree). As in previous trials, items were analyzed individually to identify stakeholder perspectives of specific treatment features that differentiated STAND from active control¹⁶. In the current trial, alpha for the parent and youth scales were excellent ($\alpha=.95-.97$).

Medication Utilization.—ADHD medication use (stimulant or non-stimulant) was naturalistic. At each assessment, parents and teens completed a medication interview used extensively in past trials of adolescent ADHD treatment⁴⁰. Respondents indicated current medications received, doses, administration schedules, settings taken, changes made since the last assessment, reasons for changes, and information about frequency of medication visits. Data were screened for discordant parent-teen reports, which were resolved by discussion. Based on these combined reports, current medication utilization was coded for each time point.

Measurement of Moderators

Service Delivery Moderators.—Receipt of combined treatment, number of sessions received, therapist ethnicity match, therapist years of experience, therapist licensure, percentage of office-based sessions, and percentage of sessions attended by a parent were moderators. Medication status was derived from the medication interview; number of sessions received over twelve months, percentage office-based sessions, and percentage of sessions attended by parent were collected from the electronic health record. Therapists provided information about their ethnicity, years of experience, and licensure status at BL.

Analytic Plan

Initially planned sample size was 300 with 15% attrition (i.e., $N=255$); however, at the end of the three-year planned recruitment period, power analyses were conducted for the sample size to date ($N=278$) with true, rather than estimated, design effects. The originally estimated design effects due to clustering between were between 1.3 and 1.6; however, true design effects were between 1.1 and 1.3, based on Intraclass Correlation (ICC) values at BL and PT for primary outcomes. Based on these new estimates of design effect, 202 participants achieved 80% power for expected effects ($d=.3$ to $.4$). The variable with the most missing data possessed 239 participants at FU (14% attrition). All participants possessed complete baseline data. ICC and design effects for adolescent-level outcomes revealed that all design effects were < 2 . As a result, we elected not to include clustering in the analysis based on the recommendation of Muthen and Satorra⁴¹. A false-discovery rate correction was applied within outcome domain for all analyses⁴².

Primary Outcomes

Linear mixed models (LMMs) with random intercepts were conducted in SPSS 25. We first conducted intent to treat analyses including all randomized participants ($N=278$). Separate Linear Mixed Models were conducted for each outcome. In LMMs, dummy codes were specified for group (UC=0, STAND=1). To model agency-specific effects, we included three dummy codes with agency 1 (largest) serving as the reference group. We tested various time curves and found linear time to possess the best fit. Time was coded as a continuous, subject-specific measure reflecting months since BL (BL time=0). Data were assumed missing at random (MAR)⁴³. A full information robust Maximum Likelihood estimator was employed. For each outcome, the following specifications were evaluated. The linear effects of time and group $\times$ time were the effects of interest to test aim 1 hypotheses.

$$\text{Level 1: } Y_{ij} = \pi_{0i} + \pi_{1i}(\text{time}) + e_{ij}$$

$$\begin{aligned} \text{Level 2: } \pi_{0i} &= \beta_{00} + \beta_{01}(\text{agency 2}) + \beta_{02}(\text{agency 3}) + \beta_{03}(\text{agency 4}) + \beta_{04}(\text{group}) + r_{0i} \\ \pi_{1i} &= \beta_{10} + \beta_{14}(\text{group}) \end{aligned}$$

$$\text{Combined: } Y_{ij} = \beta_{00} + \beta_{01}(\text{agency 2}) + \beta_{02}(\text{agency 3}) + \beta_{03}(\text{agency 4}) + \beta_{04}(\text{group}) + \beta_{14}(\text{group} * \text{time}) + r_{0i} + e_{ij}$$

We then conducted per protocol (PP) analyses that included only participants who initiated treatment ($n=225$). This supplemental analysis isolates the upper limit on true effects of STAND in community settings, given that ITT effects may be deflated by randomized participants who never engaged in services after agency intake⁴⁴. Compared to those who

initiated treatment, those who did not ($n=53$) did not differ on demographic or clinical variables listed in Table 1.

Treatment Engagement

Linear regression tested group differences on cross-sectional continuous outcomes and logistic regression tested group differences on cross-sectional categorical outcomes. A generalized estimating equation using a binomial probability distribution, logit link function, and maximum likelihood estimator was used to test group differences on categorical longitudinal outcomes (medication use). Cohen's d effect size was calculated for continuous outcomes and relative risk odds ratios were calculated for categorical outcomes.

Service Delivery Moderators

Per protocol LMMs tested service delivery moderators (non-initiators had no values for these variables). For moderator analyses, the list of outcomes was trimmed to include one variable from each of the four domains. Three-way interactions of moderator $\times$ group $\times$ time indicated whether the group's effect on outcome over time varied by the moderator level. False discovery rate corrections were applied within domain.

Results

Primary Outcomes: Intent to Treat (ITT) Analyses

Results (see Table 2) indicated no group $\times$ time difference on any outcome. The full sample demonstrated significant improvements over time on all IN, HI, and academic impairment measures.

Primary Outcomes: Per Protocol Analyses

The full sample demonstrated significant improvement over time on all IN, HI, and academic impairment measures (see Table S1, available online). Group $\times$ time effects were significant for parent-rated DSM-5 IN, such that UC showed greater decreases in IN from BL to FU than STAND ($b=.02$, $SE=.01$, $p=.009$, $d=.37$, 95% CI: .01 to .04). Results indicate that null findings in ITT analyses were not solely accounted for by participants who did not initiate treatment.

Treatment Engagement

There were no group differences in treatment initiation after agency intake (STAND=82.6%, UC=79.3%; $b=-.22$, $SE=.31$, $p=.481$, $OR=.81$) or number of sessions that initiating participants received over 12 months [STAND $M=13.99$, $SD=13.80$, UC $M=17.38$, $SD=15.26$; $F(1,223)=3.43$, $p=.065$, $d=-.23$]. Among initiating participants, parents of STAND participants attended a greater percentage of sessions than UC parents [STAND $M=74.2\%$, $SD=32.0\%$, UC $M=45.7\%$, $SD=32.8\%$; $F(1, 223)=43.72$, $p<.001$, $d=.88$]. There were no group differences in cancellations ($p=.296$).

After applying false discovery rate corrections (see Table S2, available online), parents who received STAND had greater satisfaction than UC parents in 3 of 20 domains: (1) receiving new parenting techniques as a result of treatment ($p<.001$, $d=.53$), (2) gaining greater

awareness of how their habits influence the adolescent's behavior ($p=.001$, $d=.47$), and (3) learning valuable lessons from practicing skills at home ($p=.006$, $d=.39$). There were no group differences in adolescent satisfaction (see Table S3, available online).

For medication utilization, both the linear ($b=.20$, $SE=.05$, $p=.002$, 95% CI: .08 to .33) and quadratic ($b=-.02$, $SE=.01$, $p=.001$, 95% CI: $-.03$ to $-.01$) group $\times$ time effects were significant. Results (see Figure 3) indicated that participants in STAND were significantly more likely to utilize medication over time compared to UC. At BL, the odds of using medication in the STAND group were 1.43 times the odds of using medication in the UC group; this effect grew to 3.99 at PT and 10.27 at FU. Compared to the BL odds ratio, the FU between-group odds ratio is 7.18 times larger. Between BL and PT, 13.0% of STAND participants ($n=18$) either started medication for the first time ($n=8$) or resumed ($n=10$) a pre-BL prescription (UC=5.0%; $n=7$ new prescriptions, $n=0$ resumptions) and 26.1% of STAND group ($n=36$) sustained an ongoing prescription (UC=17.9%; $n=25$). Between PT and FU, an additional 3.8% of STAND participants ($n=5$) started ($n=1$) or resumed ($n=4$) a prescription (UC=6.7%; $n=4$ new prescriptions; $n=5$ resumptions) and 30.3% ($n=42$) sustained an ongoing prescription (UC=7.8%; $n=11$).

Service Delivery Moderators

See Table S4, available online. Significant three-way interactions (see Figure 3) indicated that STAND (vs. UC) led to greater reductions in parent-rated IN over time ($b=.04$, $SE=.01$, $p<.001$, 95% CI: .02 to .06) and parent-rated academic problems over time ($b=.02$, $SE=.01$, $p=.017$, 95% CI: .00 to .03) when the therapist was licensed. The standardized difference scores for these three-way interactions (time $\times$ group $\times$ licensure) was $d=1.08$ for IN and $d=1.13$ for academic problems, indicating large differences in the impact of group over time for therapists who are licensed vs. unlicensed. Group $\times$ Time $\times$ Moderator and Time $\times$ Moderator effects were non-significant for medication status, number of sessions received, billing source, therapist ethnicity match, therapist years of experience, percentage of office-based sessions, and percentage of sessions attended by a parent.

Discussion

Study findings were as follows. First, there were no significant group $\times$ time effects in ITT analyses and PP demonstrated a significant group $\times$ time on one of eleven outcomes (indicating that UC outperformed STAND on parent-rated DSM-5 IN; $d=.37$). Thus, overall, community-based STAND did not outperform UC on any primary outcome. Second, significant improvements in ADHD symptom severity and academic impairment were demonstrated for both STAND ($d=.33$ to .70) and UC ($d=.23$ to .89); however, these within-group effect sizes were notably lower than those reported for similar indices in university trials of STAND ($d=.71$ – 1.93)^{15,16}. Despite reduced effectiveness in the community setting, STAND outperformed UC on some indices of treatment engagement (i.e., parent participation in session, parent satisfaction indices, concurrent medication engagement). In addition, STAND outperformed UC on primary outcomes when therapists were licensed (22% of therapists), but not when they were unlicensed (78% of therapists).

Regarding clinical outcomes, both groups demonstrated average parent-rated IN symptoms within the C3RS³⁵ “Very Elevated” range at BL and “Elevated” range by FU. Parent-rated C3RS HI symptom averages were in the “Elevated” range at BL and “High Average” range at FU. Parent and teacher ratings of DSM-5 ADHD symptom were of similar magnitudes. Given the chronicity of ADHD, full remission of symptoms after discontinuation of ADHD treatment is not expected.⁴⁵ However, unlike in past trials of STAND, symptoms remained in the clinical (rather than subclinical) range^{15,16}. Similarly, average school grades in this trial improved from a D to a C- average, which indicates continued impairment following treatment for both groups. No improvements in family impairment of disciplinary incidents were present.

Among adolescents who attended at least one therapy session, retention in agency services was strong in both groups. Compared to UC, STAND parents showed greater participation in treatment ($d=.88$) and higher levels of satisfaction with the impact of treatment on their parenting ($d=.39-.53$). STAND also outperformed UC on concurrent medication engagement, including both sustainment of ongoing medication, resuming stopped medication, and initiating new medication. This finding is not surprising because several STAND elements are known to improve medication utilization in other populations (i.e., psychoeducation, MI, goal-setting, increasing self-awareness, and parent-teen behavioral contracting)⁴⁶. However, despite these promising effects, treatment engagement in the STAND group was lower than in university trials¹⁵⁻¹⁶ did not lead to superior outcomes than UC at approximately 10 months post-treatment (see Table 2).

STAND was more effective when delivered by licensed (versus unlicensed) therapists, suggesting that therapist engagement may be an important target. In this trial, therapist licensure was unrelated competence or fidelity¹⁷; however, literature review suggests that licensed therapists demonstrate higher levels of autonomy and self-efficacy than unlicensed therapists³⁰. Perhaps therapist engagement and empowerment efforts may improve STAND implementation beyond basic fidelity metrics. Because the majority of community therapists are unlicensed, these initiatives may have a meaningful effect on outcomes⁴⁷. Future work should identify strategies to increase therapist engagement within the constraints of community clinics.

There were no significant moderation or prediction effects for patient attendance, concurrent medication use, therapist ethnicity match, therapist years of experience, or setting of treatment, indicating that these factors did not influence treatment effectiveness (see Table S1, available online). The null attendance finding is common⁴⁸⁻⁴⁹ when youth characteristics that dampen treatment outcomes (i.e., adolescent symptom severity) also increase motivation to engage in treatment. Our results also fail to replicate Pelham and colleagues’ finding that receiving medication prior to psychosocial treatment reduces treatment response in younger children³². Similar to university trials, community-based STAND was equally effective when delivered as monotherapy or adjunctive to medication—despite improved medication use in the STAND group¹⁴. Future analyses are planned to examine the extent to which STAND’s effectiveness was impacted by reduced implementation outcomes (i.e., therapist knowledge and competence, fidelity and MI

integrity, parent use of behavioral strategies during treatment, pace of delivery). These analyses will further guide a refined community-based implementation strategy.

Therapist participation in the study was voluntary; thus, we may have oversampled therapists with openness to new interventions. Given the nature of the treatment, it was not possible to mask therapists and participants to study group, though they were masked to study hypotheses (teachers and coders were masked). Therapist to client ratio was low in this trial (i.e., 1:2.74) due to high turnover in community contexts and fine-grained patient-therapist matching that required consideration of agency, parent language, therapist catchment area, and insurance type. As a result, we did not cluster within therapist in analyses (though we covaried for agency). We did not assess the number of potential participants (therapist or adolescent) who were present at agencies but elected not to enroll in the trial. We did not assess psychological service utilization outside of the agencies. It is not clear which therapies were administered in UC; future work is needed to develop measures that can detect common UC practices for adolescent ADHD.

In sum, STAND implemented by community practitioners demonstrated overall effectiveness that was no different than UC, despite improving some aspects of therapeutic engagement. Only licensed community therapists delivered STAND in a manner that outperformed standard services. Additional work with this sample will pursue a revised implementation strategy by: (1) investigating relationships between implementation outcomes and patient outcomes and (2) querying stakeholder perspectives (e.g., therapists, supervisors, agency leadership, parents, adolescents) about barriers and facilitators to community-based STAND implementation. Future iterations of community-based treatment for adolescent ADHD will address key barriers directed during the course of this trial.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

Acknowledgments

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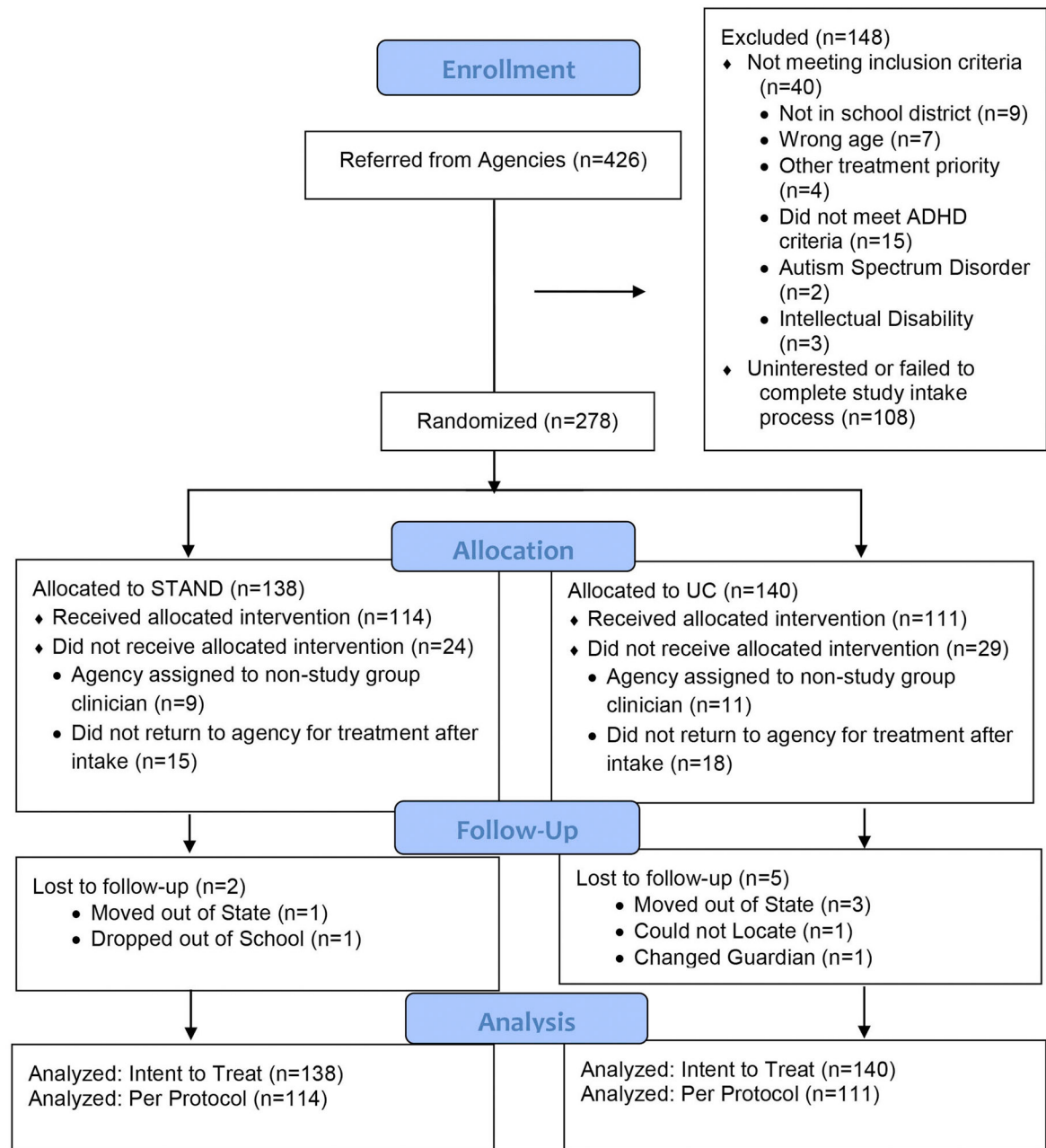


Figure 1.
CONSORT Flow Diagram: Adolescent Participants

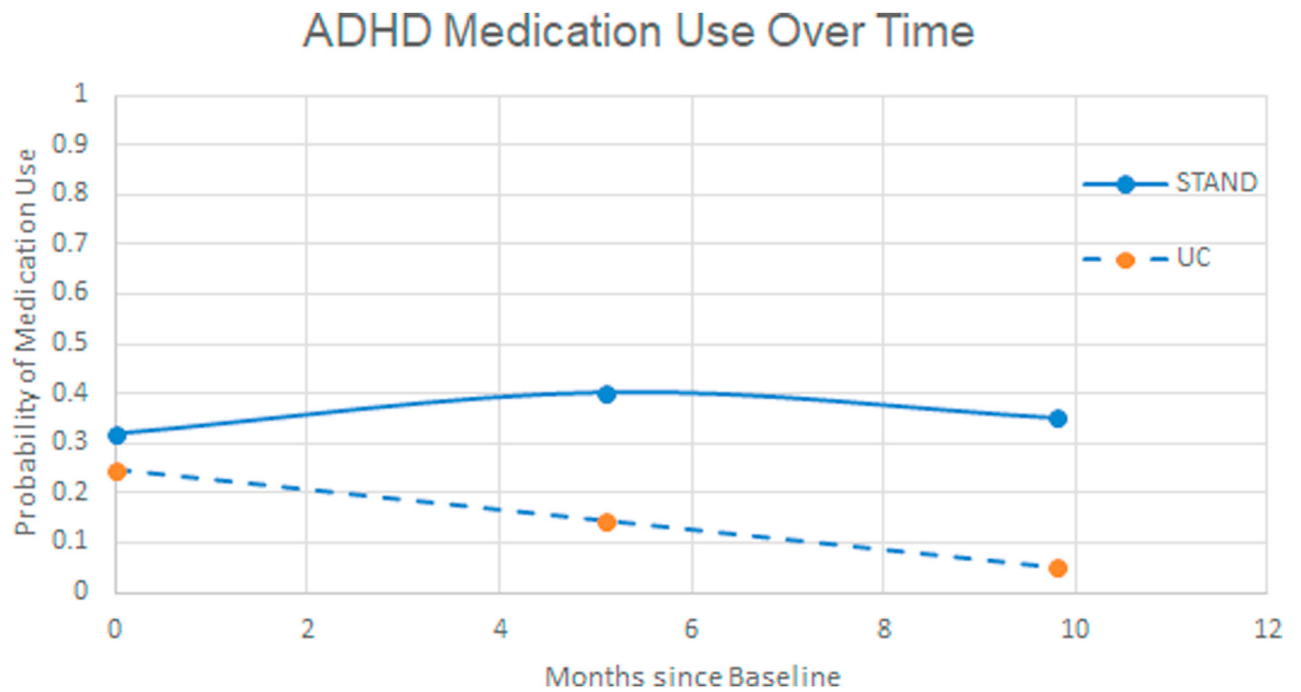


Figure 2.
Attention-Deficit/Hyperactivity Disorder (ADHD) Medication Use over Time
Note. Figure represents marginal probabilities derived from the generalized estimating equation.

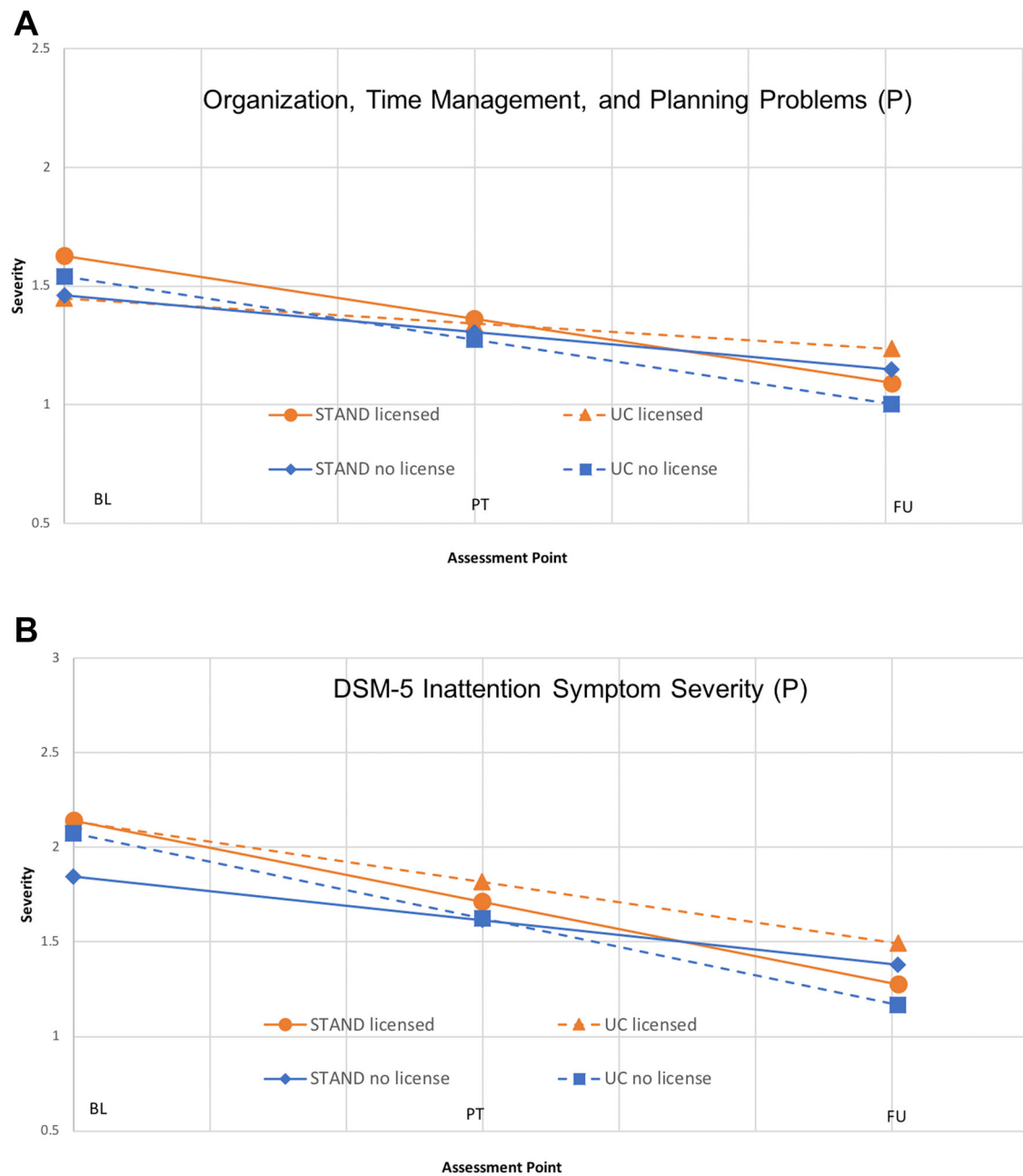


Figure 3.

The Effect of Licensure Status and Group on Effectiveness in Community Clinics

Note. Post-treatment (PT) and follow-up (FU) represent mean functioning for each group at the mean number of months since baseline (BL) that PT and FU assessments occurred.

STAND = Supporting Teens' Autonomy Daily; UC = Usual Care.

Table 1.

Baseline Characteristics of Adolescent Sample

Diagnostic Variables	STAND (N = 138)	UC (N = 140)
WASI estimated Full-Scale IQ <i>M(SD)</i>	94.15(14.07)	96.81(13.20)
ADHD Presentation		
ADHD-Predominantly Inattentive (%)	50.0	54.3
ADHD-Combined (%)	50.0	45.7
ODD/CD	50.7	47.1
Current ADHD Medication (%)	31.2	23.6
Demographic Variables		
Age <i>M(SD)</i>	13.97(1.51)	14.08(1.50)
Male Patients (%)	70.3	70.7
Race/Ethnicity (%)		
White Non-Hispanic	5.1	3.6
Black Non-Hispanic	16.7	10.0
Hispanic Any Race	77.5	85.7
Other	0.7	0.7
Single Parent (%)	35.5	36.4
Limited Parent English Proficiency (%)	36.2	46.4
Billing Source (%)		
Medicaid	57.0	55.0
State/County Subsidy	12.2	14.4
Sliding Scale	29.8	28.8
Pro Bono	0.0	1.8
Private Insurance	0.9	0.0
Parent Education Level		
High School Grad or less (%)	23.9	27.3
Part College or Specialized Training (%)	30.4	30.2
College or University Graduate (%)	33.3	33.1
Graduate Professional Training (%)	12.3	9.4

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Note. ADHD = attention-deficit/hyperactivity disorder; CD = conduct disorder; *M* = mean; ODD = oppositional defiant disorder; STAND = Supporting Teens Autonomy Daily; UC = usual care; WASI = Wechsler Abbreviated Scale of Intelligence.

Table 2.

Intent to Treat (ITT) Analyses: Results From Linear Mixed Models

	ITT STAND (N=138)				ITT UC (N=140)				ITT Time				ITT Group × Time			
	BL M (SD)	PT M (SD)	FU M (SD)	d	BL M (SD)	PT M (SD)	FU M (SD)	d	b (SE)	p	95% CI	b (SE)	p	d ^a	95% CI	
Inattention																
Parent C3RS	75.51(13.32)	69.54(13.54)	66.54(14.67)	.52	73.74(12.66)	69.02(13.15)	64.68(13.84)	.72	-.92(.11)	<.001	-1.15 to -.70	.21(.16)	.171	-.16	-.09 to .52	
Parent DSM	1.92(.69)	1.69(.70)	1.44(.65)	.70	2.00(.66)	1.70(.67)	1.42(.74)	.89	-.06(.01)	<.001	.00 to .03	.01(.01)	.162	-.16	.00 to .03	
Teacher DSM	1.84(.66)	1.66(.76)	1.49(.83)	.53	1.63(.65)	1.48(.80)	1.33(.80)	.47	-.03(.01)	<.001	-.05 to -.02	.00(.01)	.634	.07	-.03 to .02	
H/I																
Parent C3RS	65.37(17.19)	62.44(15.82)	59.74(16.71)	.33	63.02(16.71)	60.99(14.75)	59.12(15.33)	.23	-.40(.11)	<.001	-.61 to -.19	.18(.15)	.236	.10	-.46 to .11	
Parent DSM	1.02(.79)	.86(.71)	.71(.69)	.39	1.06(.78)	.94(.73)	.83(.78)	.30	-.02(.00)	<.001	-.03 to -.01	.01(.01)	.272	.09	-.02 to .01	
Teacher DSM	.99(.87)	.78(.70)	.56(.70)	.50	.90(.81)	.69(.73)	.50(.72)	.50	-.04(.01)	<.001	-.06 to -.03	.00(.01)	.801	-.03	-.02 to .02	
Academic Impairment																
School	1.60(.84)	1.71(.80)	1.80(.91)	.24	1.83(.75)	1.93(.82)	2.02(.78)	.26	.02(.01)	.002	.01 to .03	.00(.01)	.904	-.01	-.02 to .02	
Parent	1.53(.56)	1.36(.53)	1.21(.53)	.57	1.52(.50)	1.34(.53)	1.16(.55)	.72	-.04(.00)	<.001	-.05 to -.03	.00(.01)	.466	-.08	-.01 to .02	
Family Impairment																
Parent	2.95(.78)	2.85(.80)	2.75(.84)	.26	2.85(.78)	2.82(.81)	2.79(.82)	.08	-.01(.01)	.286	-.02 to .01	.01(.01)	.082	.18	-.03 to .00	
Adolescent	2.42(.69)	2.46(.78)	2.50(.76)	-.10	2.32(.75)	2.31(.67)	2.31(.74)	.01	.00(.01)	.869	-.01 to .01	.01(.01)	.258	-.11	-.01 to .02	
Disciplinary Incidents	6.21(15.62)	4.89(9.90)	3.66(8.04)	.16	3.03(5.03)	2.78(6.40)	2.55(4.53)	-.10	-.05(.008)	.549	-.21 to .11	.21(.11)	.063	.20	-.43 to .01	

Note. Means are marginal estimates controlling for agency. Cohen's *d* within groups is difference between baseline (BL) and follow-up (FU) divided by baseline pooled standard deviation. Boldface type indicates significant *p* values. *alpha*=0.05. C3RS = Conners 3 Rating Scale; H/I = Hyperactivity/Impulsivity; PT = post-treatment.

^a difference between group change scores divided by baseline pooled standard deviation.