

Motivating Persons with Schizophrenia Spectrum Disorders to Exercise: Rationale and Design

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Introduction

Schizophrenia spectrum disorders (SSDs) include schizophrenia, schizoaffective disorder and schizophreniform disorder. The diagnostic overlap between the three is well known (1); recent research indicates that persons with these disorders have similar psychiatric symptoms (2) and cognitive deficits (3, 4). Another commonality is the increased risk for adverse physical health outcomes shared by persons with these illnesses.

Death rates from diabetes, respiratory/cardiovascular, and other obesity-related illnesses are significantly higher among the over two million Americans with SSDs than in the general population (5). The most effective medications for managing SSDs (second-generation antipsychotics such as clozapine and olanzapine) are associated with weight gain, glucose dysregulation and diabetes (6-8). Despite the well-known benefits of exercise and the health dangers associated with obesity, persons with SSDs rarely adhere to exercise regimens and few exercise adherence interventions have been tested in this group.

Few published studies have examined the effects of exercise in persons with SSDs (9-15) and only two addressed exercise motivation (16, 17). All of the studies examining psychiatric outcomes reported symptom improvement (9-12); three of four studies addressing physical health outcomes reported positive changes (12-15). Dropout rates ranged from 5 to 38% over periods from 8–16 weeks in length. Attendance at exercise sessions ranged from 23 to 91% with attendance being positively correlated with degree of improvement. Archie et al. (16) provided free access to a fitness facility to twenty outpatients with schizophrenia for six months and monitored their exercise behavior. Dropout rates were 40% after four months, 70% after five months and 90% after six months. These rates compare unfavorably with exercise cessation in the general population, which is approximately 50% after six months (24). The most common reason given for poor attendance at the exercise facility was lack of motivation (16). Menza and colleagues (17) tested a one-year weight control program in thirty-one outpatients with SSDs. The program incorporated exercise as well as behavioral interventions, and compared BMI and weight with a control group of twenty nonintervention patients. Weight and BMI decreased significantly in the intervention group, with 20 of 31 persons completing the study, for a one-year retention rate of 66% (69% of sessions were attended). The study intervention consisted of nutritional counseling, exercise, and behavioral interventions incorporating motivational counseling techniques. Motivational counseling consisted of professional support, encouragement, and strategies to improve exercise habits. This study shows that interventions can increase exercise adherence in persons with SSDs. However, because the control group had no opportunity to participate in an exercise program, it is

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not possible to separate the motivational effects of the intervention from the motivational effects of exercise alone. Our study addresses this limitation by offering an exercise group to every participant.

Methods

This paper reviews the rationale and design of our study. The purpose of this study is to test the effects of the **Walk, Address sensations, Learn about exercise, Cue exercise behavior** for persons with SSDs (WALC-S) motivational intervention on the following three aspects of exercise behavior:

- Attendance—the ratio of the total number of exercise sessions attended to the total number of exercise sessions offered;
- Persistence—the number of consecutive weeks that the participant attends at least one exercise session; and,
- Compliance—the performance of activities during exercise sessions.

We hypothesize that experimental participants will have higher: 1) attendance rates; 2) persistence rates; and, 3) compliance rates at the exercise program than controls.

Design

This experimental study is testing the WALC-S motivational intervention by comparing the attendance, persistence and compliance of an experimental group of persons with SSDs (who receive the WALC-S intervention) with the exercise attendance, persistence and compliance of a time-and-attention control group (not receiving the WALC-S

intervention). The WALC-S and the time-and-attention control will be provided prior to a sixteen-week walking group that will be offered to all participants (see Figure 1).

Motivational Intervention

The **Walk, Address sensations, Learn about exercise, Cue exercise behavior** (WALC) was originally designed to motivate community-dwelling older adults to exercise. The WALC is based upon self-efficacy theory and has been theoretically and empirically validated (18-23). Persons with SSDs face many similar exercise barriers as elderly persons. Research has shown that both elderly persons and those with SSDs lack exercise motivation (16, 23, 24). Further, elderly persons perceive the following barriers to exercise: fear of pain (18), problems with goal setting (25), and lack of knowledge regarding the positive benefits of exercise (15, 24-26). Problems with goal setting may be partially related to cognitive deficits associated with aging (27, 28). The literature supports the presence of similar exercise barriers among persons with SSDs. Persons with SSDs have documented cognitive deficits that lead to difficulties setting goals and carrying out goal-directed plans (29). In one study, over 33% of persons cited fear of pain as a barrier to exercise participation and participants reported that health benefits experienced were unexpected (12), highlighting the similarities with elderly persons who lack knowledge as to the strong relationship of health benefits and exercise (25). We believe these similarities make the use of the WALC a reasonable starting point in our search for interventions to increase exercise behavior in persons with SSDs.

Figure 1 Study Design

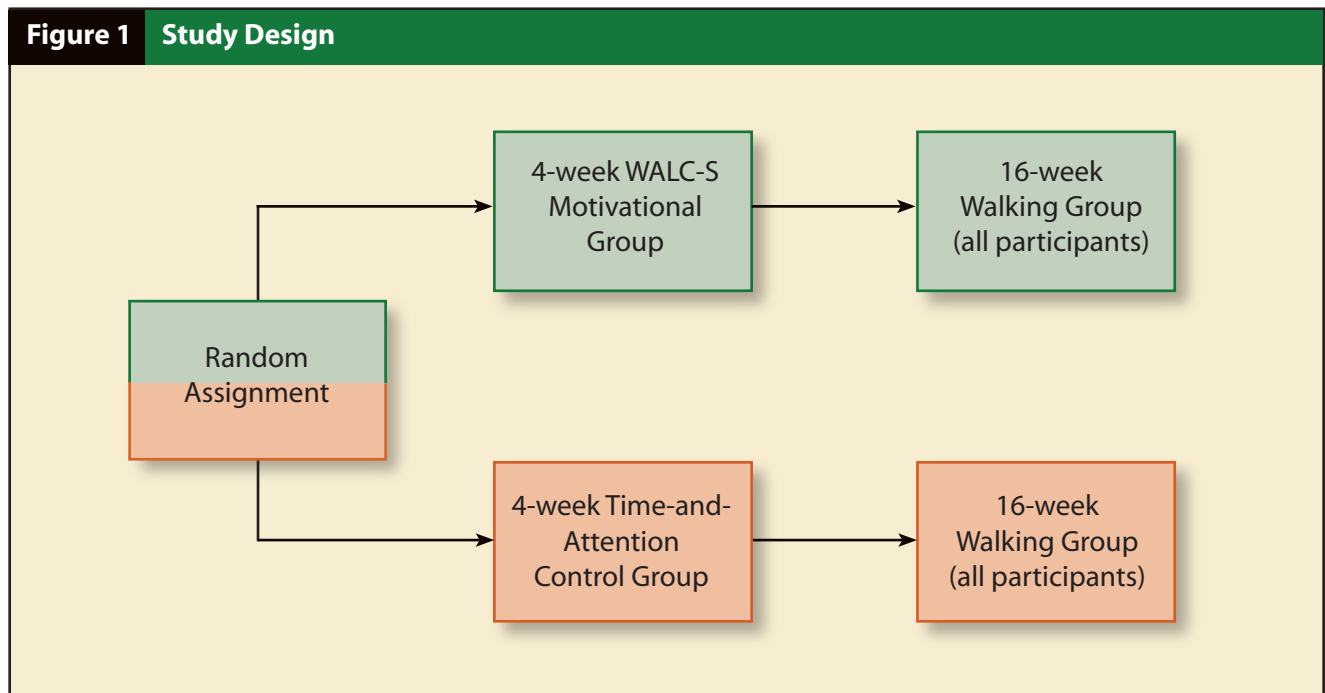
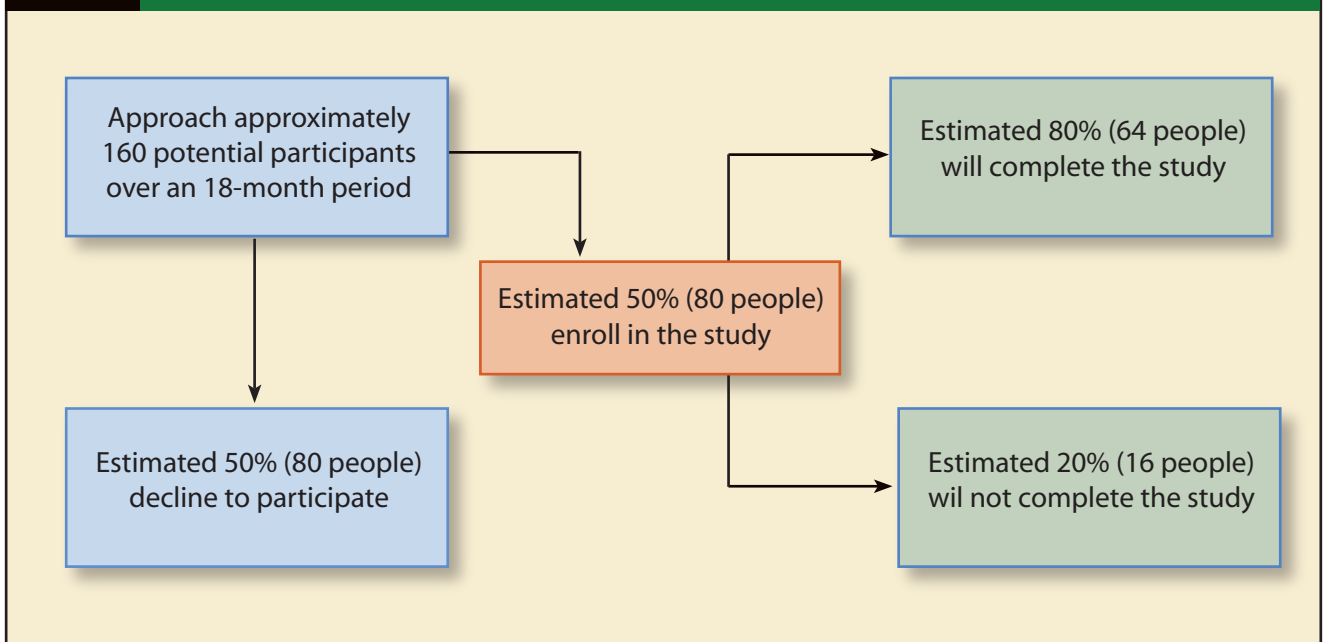


Figure 2 Estimated Study Enrollment and Completion Rates

Enrollment Criteria

The sample will be selected from persons with SSDs aged 18–68 years, receiving outpatient care at a community agency and meeting the following criteria: 1) a chart diagnosis of schizophrenia (any subtype), schizoaffective disorder or schizophreniform disorder, according to the criteria described in the *Diagnostic and Statistical Manual for Mental Disorders-IV, Text Revision* (1); 2) English speaking; 3) stable medication regimen (defined as no medication changes within the last month); and, 4) medical clearance for moderate exercise in writing from primary care provider.

Exclusion criteria include chart documentation of mental retardation, developmental delay, uncorrected visual, hearing impairments or any physiological condition preventing safe participation. Persons with the following physical conditions per chart documentation are excluded: 1) hospitalization within the past twelve months for angina pectoris, myocardial infarction, or cardiac surgery of any kind; 2) congestive heart failure; 3) cardiac pacemaker; 4) heart rate >100 or <50 at rest; 5) uncontrolled hypertension defined as a blood pressure exceeding 140/90 on three consecutive readings despite adequate treatment; 6) history of spinal or hip fractures or hip or knee arthroplasty; 7) neuromuscular or orthopedic limitations to normal, unassisted ambulation; and, 8) other medical conditions that, in the opinion of the primary care provider or investigators, involve risk exceeding the potential benefit of participation.

Our enrollment target is 80 participants. Allowing for a 20% attrition rate, it is estimated that approximately 64 participants (32 per group) will complete the study (see Fig-

ure 2). We use a computer-generated random number table to assign patients to either the control or the intervention group in a 1:1 ratio. The randomization process is expected to result in equal distribution of important sociodemographic and illness variables between the two groups. The sample will not be stratified by gender, ethnicity, or other variables, since no data are available at this time to indicate differential response to the intervention based upon differences between participants.

Study Interventions

Upon study entry, participants are randomly assigned to either the experimental or control group. Experimental participants (8–9 persons per group) attend the one-hour WALC-S motivational group weekly for four weeks prior to their sixteen-week walking group. The WALC-S was designed to address cognitive deficits common to persons with SSDs. While variability exists between individuals, cognitive impairments are present to some degree in virtually every SSD (30). Specific deficits include mild-to-moderate impairment in attention and memory, and severe impairment in executive functioning (31, 32). Executive functioning deficits involve frontal lobe dysfunction in the areas of planning, organization, and mental flexibility (30) that impede problem-solving and decision-making processes (33). We used a number of strategies to offset these difficulties. For example, we addressed memory deficits (34) through repetition of information during WALC-S sessions, and by providing written and verbal reminders of WALC-S groups. To assist participants to sustain attention during WALC-S

Table 1 Activities and Materials for the WALC-S

Week	Activities and Materials
One	<ul style="list-style-type: none"> • Introductions of research staff and group members. • Review section of booklet on basics of exercising safely. • Perform stretches: Poster with photograph of each stretch, demonstration by research staff and return demonstration by group members.
Two	<ul style="list-style-type: none"> • Introductions and check-in with each member. • Review section of booklet on basics of exercising safely. • Review section of booklet on exercise benefits. • Perform stretches: Poster with photograph of each stretch, demonstration by research staff and return demonstration by group members.
Three	<ul style="list-style-type: none"> • Introductions and check-in with each member. • Review section of booklet on basics of exercising safely. • Review section of booklet on exercise benefits. • Barriers and solutions: Discuss and write suggestions for dealing with common exercise barriers on blackboard. • Perform stretches: Poster with photograph of each stretch, demonstration by research staff and return demonstration by group members.
Four	<ul style="list-style-type: none"> • Introductions and check-in with each member. • Review section of booklet on basics of exercising safely. • Review section of booklet on exercise benefits. • Barriers and solutions: Discuss and write suggestions for dealing with common exercise barriers on blackboard. • Set individual goals. • Provide reminder calendars to cue exercise. • Perform stretches: Poster with photograph of each stretch, demonstration by research staff and return demonstration by group members.

groups, information was presented in a variety of formats: first verbally, then pictorially with visual aids, and finally demonstrated in person. Research staff assisted with goal setting and identifying ways to overcome exercise barriers in order to offset difficulties with problem solving and decision making (35). Finally, because participants in prior studies cited the importance of the social aspects of exercise (36), we designed the WALC-S as a one-hour group intervention (8–9 participants per group) that met weekly for four weeks.

The WALC-S intervention is *not* a walking group; rather, it is a forum for discussion of the basics of walking for exercise supplemented by a booklet provided to every experimental participant. The group practices a series of basic stretches at each meeting. Study personnel assist with goal setting, a process that is difficult for persons with schizophrenia due to limitations in executive functioning (35, 37).

Study personnel provide suggestions on reducing common exercise discomforts, such as using heat or massage for muscular soreness. Study personnel provide instruction on potential exercise benefits and overcoming barriers to exercise. Participants are provided with reminder calendars to track WALC-S attendance and any other exercises performed independently. Activities for each weekly group are listed in Table 1.

Control participants (8–9 persons per group) attend the one-hour time-and-attention control group weekly for four weeks prior to their sixteen-week walking group. Control groups are conducted by the same research staff as the WALC-S groups, and meet on the same day in the same location for an identical period of time. Control groups focus on healthy behaviors such as medication adherence, smoking cessation and progressive muscle relaxation.

Sixteen-Week Walking Group (All Participants)

After the four-week experimental or control group is completed, all participants participate in a sixteen-week walking group. Walking groups are conducted by different research personnel than those who led the WALC-S group or the time-and-attention control group to minimize the effects of relationship with group leader alone upon outcomes. Each walking group begins with a series of warm-up stretches (38). Warm up is followed by walking, beginning with five minutes and gradually increasing to thirty minutes over the first four weeks. Interaction during the walking groups is social rather than therapeutic (e.g., discussion of daily activities, family members, recreational plans, and the like). Walking group research personnel do not initiate any discussions of WALC-S content during walking groups. Should any participant initiate discussion or questions regarding WALC-S concepts, group leaders respond with appropriate education, goal setting or reinforcement. For example, if a participant remarks that he or she is experiencing less depression since beginning the walking group, the group leader reinforces information regarding the benefits of exercise on mood, but refrains *from initiating* any such discussions. Research personnel note that this has occurred on only one occasion; thus, we estimate that the likelihood of the blind being broken as a result of discussion of WALC-S training during the walking groups appears to be minimal. Each walking session concludes with a series of cool down stretches. Walking sessions occur outdoors (weather permitting) at the recruitment facility on Monday, Wednesday, and Friday, either in the morning or afternoon. Control and experimental participants do not attend the same walking group, in order to prevent cross-contamination.

Treatment Fidelity

Two different research personnel blinded to treatment group are providing walking groups to experimental and control participants. The first author conducted an eight-hour training session before any walking groups; both personnel scored 100% on a knowledge test after the training. The first author provides at least two hours of supervision each week, and attends 50% of the walking groups. A measure of concurrence with the walking group training is generated by completion of a fidelity form once a month. The form provides an objective measure of the provision of each aspect of the walking group, ranging from 1 (none of the time) to 5 (all of the time). The first author provides remediation for any area that scores below 4 (most of the time). Remediation is organized accordingly: 1) identification of specific concerns or questions; 2) item-by-item discussion of walking group procedures; 3) provision of feedback and clarification; and, 4) immediate correction of deficiencies. These activities ensure that the walking groups are implemented as stated, increasing uniformity and reducing alternative explanations based upon differences in walking group procedures.

Outcomes

Walking group attendance will be measured as a ratio of number of exercise sessions attended to total number of exercise sessions offered. For example, a person attending twenty-four exercise group sessions of the total forty-eight sessions offered would have an attendance rate of 50%. Walking group persistence will be measured as the number of consecutive weeks that the participant attends at least one walking group. Walking group compliance is measured by pedometer. The number of steps walked per group will be used to generate an ordinal measure of compliance as follows: Full—the number of steps equals or exceeds that of the previous walking group; Partial—the number of steps is not more than 10% less than that of the previous walking group; and, None—the number of steps is more than 10% less than that of the previous walking group.

Data collection will occur over a two-year period. Data regarding sociodemographic characteristics, living arrangements, prescribed medications and medication and/or dosage changes will be collected on all participants at study entry. Attendance, persistence, and compliance are documented at each walking group. Sixty-four participants have been recruited; this partial sample consists mostly of female (N=36, 56.3%) Caucasians (N=42, 65.6%) from 24–68 years with a mean age of 47.88 (SD±8.85). The majority live with family members (N=30, 46.9%). Forty-eight participants have completed all study activities including descriptive measures, the WALC-S or the time-and-attention control group, and the 16-week walking group with its

associated measures of attendance, persistence and compliance.

Data Analysis Plan

The Statistical Package for the Social Sciences (SPSS) and the Statistical Analysis System (SAS) will be used to analyze the data. Data analysis will begin with data plots and basic descriptive statistics appropriate for the level of measurement. Groups will be compared for pretreatment equivalence on sociodemographic characteristics, as well as type and dosage of prescribed medications. Expected mean squares will be calculated and the appropriate combination will be used for hypothesis tests with specific functions of the repeated measures. General linear model analyses in SAS (GLM and MIXED procedures) will be used to examine the effects of time, treatment, and time by treatment interaction. For hypotheses 1 and 2, a sample size of 32 in each group will have 80% power to detect an effect size of 0.629 using a two-group t-test with a 0.05 one-sided significance level. For hypothesis 3, a sample size of 32 in each group will have 80% power to detect a probability of 0.679 that an observation in the control group is less than an observation in the experimental group using a Wilcoxon rank-sum test with a 0.05 one-sided significance level.

Summary

In addition to hypothesis testing, this study will provide information to estimate effect sizes to calculate power and determine appropriate sample sizes for future inquiries. This work is the first step toward our ultimate goal of developing and empirically evaluating exercise interventions to enhance health for persons with SSDs.

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