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EPA guidance on physical activity as a treatment for severe mental illness: a meta-review of the evidence and Position Statement from the European Psychiatric Association (EPA), supported by the International Organization of Physical Therapists in Mental Health (IOPTMH)

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Abstract (250/250)

Physical activity (PA) may be therapeutic for people with severe mental illness (SMI) who generally have low PA and experience numerous life style-related medical complications. We conducted a metareview of interventions and their impact on health outcomes for people with SMI, including schizophrenia-spectrum disorders, major depressive disorder (MDD) and bipolar disorder. We searched major electronic databases until January 2018 for systematic reviews with/without metaanalysis that investigated PA for any SMI. We rated the quality of studies with the AMSTAR tool, grading the quality of evidence, and identifying gaps, future research needs and clinical practice recommendations. For MDD, consistent evidence indicated that PA can improve depressive symptoms versus control conditions, with effects comparable to those of antidepressants and psychotherapy. PA can also improve cardiorespiratory fitness and quality of life in people with MDD, although the impact on physical health outcomes was limited. There were no differences in adverse events versus control conditions. For MDD, larger effect sizes were seen when PA was delivered at moderate-vigorous intensity and supervised by an exercise specialist. For schizophrenia-spectrum disorders, evidence indicates that aerobic PA can reduce psychiatric symptoms, improves cognition and various subdomains, cardiorespiratory fitness, whilst evidence for the impact on anthropometric measures was inconsistent. There was a paucity of studies investigating PA in bipolar disorder, precluding any definitive recommendations. No cost effectiveness analyses in any SMI condition were identified. We make multiple recommendations to fill existing research gaps and increase the use of PA in routine clinical care aimed at improving psychiatric and medical outcomes.

Key words: Physical activity, sedentary behaviour, exercise, psychosis, schizophrenia, severe mental illness, bipolar disorders, major depressive disorders

Introduction

Serious mental illness, physical co-morbidity and premature mortality

Severe mental illnesses (SMI), defined as schizophrenia-spectrum disorders, bipolar disorder (BD) and major depressive disorder (MDD), are leading causes of years lived with global disability and are of considerable public health importance ¹. In addition to the impact of the mental health symptoms and reduced daily life functioning, people with SMI are at increased risk of premature mortality by between 10 to 20 years compared to age- and sex-matched controls ²⁻⁵. While suicide accounts for a concerning portion of the early mortality ^{6,7}, there is increasing recognition that physical disorders account for approximately 70% of these premature deaths ^{3,8}. Of notable concern, cardiovascular and metabolic diseases appear to greatly increase the risk of early death in those with SMI⁹, which is of particular importance, given the high prevalence of these diseases in SMI ⁹⁻¹¹. People with SMI are also at increased risk of various other physical comorbidities, such as respiratory disease ^{12,13}, poor bone health ¹⁴ and physical multimorbidity ¹⁵. Moreover, people with SMI typically experience pronounced cognitive impairment, which often worsens over time ¹⁶⁻¹⁸ and for which treatment approaches remain limited ^{19,20}.

Current treatment for mental health symptoms and functioning largely revolves around psychotropic medication ^{21,22} and/or psychotherapeutic interventions ²³⁻²⁵. Whilst both of these dominant approaches, alone and in combination, have demonstrated treatment efficacy on mental health symptoms ²⁶, their impact on the rising physical health burden in this population is limited, and psychotropic medication may even have an adverse relationship with cardiometabolic/physical health ^{8,9}. In addition, antipsychotic medication has been associated with reduced grey matter volume in people with schizophrenia ²⁷ while psychotherapeutic interventions appear to have limited efficacy for cognitive impairment in this population ²⁸.

Established benefits of physical activity in the general population

In the general population, there is evidence that physical activity is equally effective as frontline pharmacological interventions, such as statins and beta-blockers, in preventing cardiovascular disease mortality ²⁹. Moreover, there is consistent evidence that physical activity and exercise can decrease the risk of developing cardiovascular and metabolic disease ³⁰⁻³² and reduce inflammatory parameters, such as C-reactive protein, ^{33,34} which are commonly raised in people with SMI ³⁵. Conversely, higher levels of sedentary behaviour (characterized by an energy expenditure ≤1.5 metabolic equivalents (METs), while in a sitting, reclining or lying posture during waking hours ³⁶) are independently associated with an increased risk of diabetes, cardiovascular disease and premature mortality ³⁷. In the general population, there is also evidence that lower levels of cardiorespiratory fitness are a more accurate determinant of premature death than body mass index (BMI) ³⁸. Moreover, there is evidence that aerobic exercise is effective in improving cognitive function in the general population ³⁹⁻⁴³ including potentially increasing hippocampal volume ⁴⁴. In addition, a recent global meta-analysis has demonstrated that higher levels of PA confers protection from the development of depressive symptoms and MDD⁴⁵.

Low levels of physical activity and fitness

Despite the aforementioned, there is evidence to suggest that less than half of people with SMI (schizophrenia ⁴⁶, bipolar disorder ⁴⁷ and major depression ⁴⁸ ⁴⁹) meet recommended physical activity levels of 150 minutes of moderate-vigorous physical activity per week ⁵⁰. Moreover, each of these populations engage in remarkably high levels of sedentary behaviour ⁴⁶⁻⁴⁸ and have low levels of cardiorespiratory fitness ⁵¹. People with SMI experience, a number of barriers from engaging in physical activity exist, such as side effects of medications, complications from obesity/poor physical health^{52,53}, lack of resources/professional support⁵⁴, various motivational factors,⁵⁵ which calls for targeted interventions in this highly sedentary population⁵⁶⁻⁵⁸.

In light of the above, we set out to conduct an evidence based guideline meta review and position statement on the use of physical activity and exercise among people with SMI.

Aims

The overall aims of this meta-review and position statement were as follows: First, to establish the benefits of physical activity / exercise across all categories of severe mental illness (SMI), using top-tier evidence from published systematic reviews and meta-analyses. Second, examine how the benefits of physical activity may differ across specific SMIs, including schizophrenia-spectrum disorders, BD and MDD. Finally, to use these findings to provide guidance for clinical practice, policy and future research.

Methods

Guidance development process

This guidance paper was performed in accordance with the PRISMA guidelines ⁵⁹ following a predetermined, published protocol (PROSPERO registration CRD42017068292). Moreover, the current guidance was conducted in accordance with the European Psychiatric Association (EPA) guidelines framework and wherever possible, adopted guidance based on the findings from systematic reviews and meta-analyses ⁶⁰.

Searches and study selection

Two independent authors searched from inception to 15th January 2018 Medline/ Pubmed, PsychInfo, EMBASE and the Cochrane database for systematic reviews (with and without meta-analyses) of studies investigating physical activity/ exercise among people with SMI, schizophrenia-spectrum disorders, BD or MDD. The search terms included (exercise or aerobic exercise or physical activity or resistance training) and (schizophrenia or psychosis or psychotic or major depression or depression or bipolar disorder or serious mental illness or serious mental disorder). The reference lists of included articles were also hand-searched.

Type of studies eligible for inclusion

We followed the European Psychiatric Association manual for identifying and conducting this review and evidence was considered in a hierarchical approach, going from systematic reviews to the highest level of meta-analyses. In the absence of these two data sources for particular outcomes, we sought to identify newer randomised/controlled trials.

The specific inclusion criteria were: 1) systematic reviews (with or without meta-analyses) that synthesised randomised, controlled clinical trials or randomised control trials (RCTs) or controlled clinical trials (CCTs); 2) physical activity/ exercise interventions, including aerobic, high intensity and resistance exercise as monotherapy or in conjunction with other treatment options, including psychotropic medication or psychological therapies; 3) systematic reviews of PA, which included

people with pooled SMI or schizophrenia-spectrum disorders, BD or MDD, confirmed through validated assessment measures (e.g. Diagnostic and Statistical Manual of Mental Disorders ⁶¹(DSM), International Classification of Diseases ⁶² (ICD) criteria), in any setting; 4) systematic reviews, which included a non-active/ non-exercise control group (e.g., does not include physical activity). We excluded mind-body physical activity interventions, such as yoga and tai-chi, since these activities are presumed to exert beneficial effects on mental health through additional factors distinct from the physical activity itself.

Definition of exercise and physical activity

We included systematic reviews investigating the benefits of exercise or physical activity in people with SMI. Exercise was defined as "planned, structured, and repetitive and has as a final or an intermediate objective the improvement or maintenance of physical fitness." Within this definition, we included aerobic exercise, high intensity exercise, resistance exercise and mixed exercise (i.e., aerobic and resistance exercise). Physical activity was defined as "any bodily movement produced by skeletal muscles that results in energy expenditure" 63. We considered exercise/physical activity studies used as monotherapy or in combination with other types of treatment, e.g., psychotropic medication or psychological interventions.

Outcomes

As indicated in the EPA guidelines manual ⁶⁰, as a first choice in the hierarchy of evidence, we drew evidence from systematic reviews of exercise/ physical activity interventions, including meta-analyses and/ or RCTs/ CCTs that considered the outcomes listed below. In accordance with our published protocol, we included data from the largest and/or most recent paper investigating each type of PA and each outcome in any population.

Primary outcomes

The primary outcomes focused on changes in the severity of symptoms, which characterise the included psychiatric disorders. For example, positive and negative symptoms in people with schizophrenia-spectrum disorders; and depressive symptoms in people with MDD or BD.

Secondary outcomes

We were interested in a range of secondary outcomes, including:

- Physical health factors, e.g., cardiovascular or metabolic parameter changes, anthropometric
 measures (e.g., BMI, waist circumference) or body composition measures (e.g., amount of
 intra-abdominal and cardiac adipose tissue).
- Cardiorespiratory fitness (expressed as percentage maximal or peak oxygen uptake), muscular fitness.
- Increasing physical activity levels or decreasing sedentary behaviour.
- Biomarkers, e.g., HbA1C, C-reactive protein, brain derived neurotrophic factor (BDNF), interleukin-6.
- Cognitive functioning, e.g. performance in neuropsychological tests
- Brain structure and connectivity, e.g., determined through magnetic resonance imaging or diffusor tension imaging changes.
- Quality of life and functioning.
- Dropout rates and predictors from physical activity interventions.
- Adverse events (e.g., injuries sustained from PA)
- Economic evaluations.

Data extraction

Data extraction was conducted by two authors and reviewed by a third author. We extracted data from systematic reviews and meta-analyses of RCTs/ CCTs investigating exercise interventions, including:

Number of studies included, number of participants in each arm, participant demographics, length of follow-up, details of the exercise intervention, statistical analyses conducted, effect size information, heterogeneity, publication bias and details of any meta regression and subgroup analyses.

Where a systematic review was not available for an outcome, we narratively summarised information from newer RCTs/ CCTs in the manuscript text, noting the study design, sample size, description of the intervention and control group, intervention effect and any harms and dropout rates.

Quality assessment and grading of evidence

Two independent authors assessed the quality of systematic reviews and meta-analyses with the AMSTAR tool ^{64,65}. In addition, we assessed the quality of evidence using the SIGN (2011) recommendations as indicated by the EPA guidelines ⁶⁰ and in accordance with a recent published EPA guidelines on psychotherapy for depression ²⁴. Specifically, each included study was graded from 1++ (highest quality) to 4 (lowest quality) (see table 1). Based on the quality of evidence, we graded recommendations according to best practice (table 2).

Table 1 Grading of evidence in accordance with SIGN (2011 (and adapted from ²⁴)

Grade	Description
1++	High-quality meta-analyses, systematic reviews of RCTs or RCTs with a very low risk of bias
1+	Well-conducted meta-analyses, systematic reviews or RCTs with a low risk of bias
1-a	Meta-analyses, systematic reviews or RCTs with a high risk of bias
2 ++	High-quality systematic reviews of case control or cohort studies. High-quality case control or cohort studies with a very low risk of confounding or bias and a high probability that the relationship is causal
2+	Well-conducted case control or cohort studies with a low risk of confounding or bias and a moderate probability that the relationship is causal
2-a	Case control or cohort studies with a high risk of confounding or bias and a significant risk that the relationship is not causal
3	Nonanalytic studies, e.g. case reports, case series
4	Expert opinion

Table 2 Grading of recommendations, modified from the SIGN (2011) and adopted from 66

Grade	Description
A	At least one meta-analysis, systematic review, or other study rated at low risk of bias
В	A body of evidence including studies rated as 2, directly applicable to the target population, and
	demonstrating overall consistency of results; or extrapolated
	evidence from studies rated as I or II
C	A body of evidence including studies rated as II–III, directly applicable to the target population and
	demonstrating overall consistency of results.
D	Good practice point recommended good practice based on the clinical experience of the Guidance
	development group and arrived at through consensus

Results

Search results and included studies

The initial database searches acquired 2,722 hits and 2,081 were reviewed at the abstract level. Subsequently, we reviewed 145 full texts, and 125 were excluded with reasons (see figure 1). Overall, 20 systematic reviews and meta-analyses were included in this guidelines document, which included studies that provided effect sizes for the benefit of exercise for SMI ⁵¹ (N=1), schizophrenia-spectrum disorders ⁶⁷⁻⁷³ (N=7), BD ^{74,75}(N=2) and MDD ⁷⁶⁻⁸⁵ (N=10).

Figure 1 here

Details of the included systematic reviews representing participants, interventions, outcomes, main results, considerations, AMSTAR scores and level of evidence scores are summarized in table 3 for schizophrenia-spectrum disorders and mixed SMI participants, and in table 4 for MDD and BD participants. Details of the results are described below for each population group.

Insert table 3 and table 4 here

Pooled SMI

One meta-analysis provided non-overlapping data on exercise for people with pooled SMI 51,86 . It was demonstrated that aerobic exercise improved fitness in pre and post-test studies among people with schizophrenia-spectrum disorders and MDD (Hedges' g = 0.33, 95 % CI 0.21–0.45, p = 0.001, 13 studies) 51 (AMSTAR 7, evidence level -1). Improvements in fitness were not accompanied by improvements in BMI. Meta regression showed that higher (vs, low or moderate) intensity interventions and those delivered by qualified professional predicted better outcomes. In 5 RCTs, exercise improved fitness versus control (Hedges' g = 0.43, 95 % CI 0.10–0.76, p = 0.01; n = 109).

Dropout, adverse events and economic evaluations of exercise interventions in pooled SMI

No information was reported on dropout rates, adverse events or economic evaluations in people with SMI.

Schizophrenia-spectrum disorders

In total, 7 systematic reviews and meta-analyses reported the benefits of exercise for schizophrenia-spectrum disorders ⁶⁷⁻⁷³. There was broad variation in the frequency, intensity, type and time of the interventions across the included systematic reviews and meta-analyses (Table 3). The mean AMSTAR score for the included reviews was 6 (range 4-10).

A summary of the main results for exercise interventions in schizophrenia are presented in table 5. In a meta-analysis of 8 RCTs ⁶⁸ (AMSTAR 5, evidence level -1) and 659 patients with schizophrenia-spectrum disorders, exercise reduced total psychiatric symptoms when interventions used at least 90 min of moderate-to-vigorous physical activity (MVPA) per week (SMD: 0.72, 95% confidence interval -1.14 to -0.29). Ninety minutes of MVPA was also associated with reduced positive (SMDs -0.54, 95% CI -0.95 to -0.13) and negative symptoms (SMD-0.44, 95% CI -0.78 to -0.09). Pooled data from four RCTs found that exercise had no significant effect on reducing BMI (MD: -0.98 kg/m2; 95% CI -3.17 to 1.22 kg/m2, N=4). Narrative findings regarding the influence of exercise on additional cardiometabolic parameters, such as waist circumference, high density lipoprotein levels and triglycerides, were inconsistent. Data from a narrative synthesis across 3 RCTs found that exercise also improved quality of life in people with schizophrenia-spectrum disorders. The key consideration from this systematic review was that optimal effects of exercise interventions are derived when people achieve over 90 minutes of MVPA per week.

A meta-analysis from 10 studies ⁶⁷ (AMSTAR 5, evidence level -1) including a sample of 186 in the intervention and 199 in the control group, found that exercise improved global cognition (g = 0.33, 95%CI = 0.13-0.53, P = .001). Meta-regression analyses indicated that greater improvements in global cognition were moderated by greater amounts of weekly exercise and when interventions were delivered by exercise professionals (e.g., physiotherapists, BSc level qualified exercise scientists). Subgroup analyses found that exercise improved cognitive domains of working memory (g = 0.39, P = .024, N = 7, n = 282), social cognition (g = 0.71, P = .002, N = 3, n = 81), and attention/vigilance (g = 0.66, P = .005, N = 3, n = 104). A systematic review of 14 trials ⁶⁹ investigated the pro-cognitive mechanisms of exercise in schizophrenia-spectrum disorders (AMSTAR 4, evidence level -1). The review suggested that exercise may increase brain volume, although there was a lack of consistency regarding which regions. One region that has been implicated is the hippocampus 87 although a meta-analysis of four studies did not support this hypothesis 44. This review 69 also found inconsistent evidence for an effect of exercise on BDNF levels, which is consistent with a meta-analysis 72 (AMSTAR 10, evidence level -1) also finding that exercise did not influence BDNF levels. Thus, whilst there seems reasonable evidence that exercise does improve cognition in schizophrenia-spectrum disorders ⁶⁷, the underlying mechanisms are not yet clear.

A recent systematic narrative review ⁸⁸ investigated the influence of strength training on outcomes in schizophrenia-spectrum disorders (AMSTAR 4, evidence level -1). Across 6 studies (3 RCTs) the authors found tentative evidence that strength training over time can improve muscle strength, walking performance and reduce psychopathology symptoms. This systematic review was complimented by another systematic review ⁷¹ (AMSTAR 7, evidence level -1) that found evidence that combined aerobic and strength training (7 RCTs) may improve strength, symptoms of schizophrenia and mental health, but not cardiovascular fitness.

A narrative systematic review ⁷⁰ across 10 intervention studies (5 RCTs) (AMSTAR 5, evidence level -1) found that in those with schizophrenia-spectrum disorders, walking may reduce weight, BMI and body fat in the short-term. The authors noted that walking appeared to be safe, although the evidence was noted to be of medium-high risk of bias in the included studies.

Finally, a meta-analysis of seven studies (3 RCTs) 73 (AMSTAR 8, evidence level -1) found that over a 12 week exercise intervention, aerobic exercise improved cardiorespiratory fitness versus control conditions (g = 0.41, 95% = CI 0.05 to 0.82, P = 0.028).

Insert table 5

Dropout, adverse events and economic evaluations of exercise interventions in schizophreniaspectrum disorders

A meta-analysis of 19 RCTs and 594 schizophrenia-spectrum disorder patients ⁸⁹ demonstrated that 26.7% [95% confidence interval (CI)=19.7%-35.0%] of patients dropped out of interventions, more than double reported in non-active control interventions (odds ratio=2.15, 95% CI=1.29-3.58, P=.003). Factors influencing lower dropout in meta-regression analyses included supervision from a recognized exercise professional and including a motivational component.

Surprisingly, very little information is available on the potential harms or adverse events from exercise in schizophrenia-spectrum disorder patients. Specifically, the following papers did not report harms ^{67-69,71,72,88}. Only one study ⁷⁰ reported potential harms and found that walking was not associated with adverse events in schizophrenia-spectrum disorders.

To the best of our knowledge, there have been no studies with an economic evaluation of the cost and value of delivering exercise interventions in schizophrenia-spectrum disorders.

Major depression

In total 10 systematic reviews and meta-analyses reported the benefits of exercise for MDD ⁷⁶⁻⁸⁵. The mean AMSTAR score for the included reviews was 9.1 (range 3-10). Full details of the included systematic reviews are summarized in table 4.

A summary of the effects of exercise in MDD are reported in table 5. In a comprehensive Cochrane review 83 including exercise trials, the authors found that exercise was effective in reducing depressive symptoms versus control conditions (Hedge's g=-0.68, 95% CI -0.92 to -0.44, p=0.001, I²=67.99, k=23) (AMSTAR 11, evidence level 1+). Exercise was effective in studies with (Hedge's g=-0.40, 95% CI = -0.70 to -0.11, P = 0.01, k=10) or without (Hedge's g=-0.91, 95% CI = -1.22 to -0.61, P < 0.001, k =

13) blinded outcome, with (Hedge's g=-0.56, 95% CI -0.87 to -0.25, p<0.001, k=12) or without (Hedge's g=-0.80, 95% CI = -1.15 to -0.46, p<0.001, k=11) intention-to-treat, and in studies comparing exercise with no intervention (Hedge's g = -1.24, 95% CI = -1.83 to -0.65, k = 4), and with usual care (Hedge's g = -0.48, 95% CI = -0.8 to -0.16, k = 4). There was no difference in exercise versus psychological interventions (Hedge's g = -0.22, 95% CI -0.65 to 0.21, k=4), or antidepressant treatment (Hedge's g = -0.08, 95% CI = -0.33 to 0.18, k = 3), indicating exercise demonstrated comparable efficacy versus these established interventions. However, information on symptom severity was not available. Of note, beneficial effects were confirmed when considering only studies with blinded outcomes (Hedge's g = -0.40, 95% CI -0.69 to -0.11, P = 0.01, k = 10), but not when considering blinded outcomes with both adequate allocation concealment and intention-to-treat together (Hedge's g = -0.26, 95% CI = -0.61 to 0.08, P = 0.14, k = 6). Also, at longer follow-up (mean 10.75 month) no significant effect was reported (Hedge's g = -0.22, 95% CI -0.53 to 0.09, P = 0.16, k = 7).

More recently, a meta-analysis 81 (AMSTAR 10, evidence level 1+) attempted to address some of the criticisms of the Cochrane review 83, such as the inclusion of a number of in-eligible trials, comparing exercise versus other exercise interventions, and the absence of adjusting for publication bias. The authors 81 found across 25 RCTs (intervention 757, control 730) that exercise versus non-active interventions over a mean of 14 weeks resulted in a large improvements of depressive symptoms (SMD = 1.11, 95% CI % = 0.79-1.43, k = 25) with a failsafe number of 1057. Exercise was effective both when the diagnosis was made according to DSM⁶¹ (SMD = 1.14, 95% CI = 0.46-1.81, P < 0.001, $I^2 =$ 88.5, k = 9) or depressive symptoms (SMD = 0.801, 95% CI = 0.49-1.11, P < 0.001, $I^2 = 68.5$, k = 14), high (SMD = 0.882, 95% CI = 0.221-1.544, P = 0.009, I² = 90.1, k = 4) and low (SMD = 1.033, 95% CI = 0.66-1.41, P < 0.001, $I^2 = 79.2$, k = 21) quality studies, outpatients (SMD = 1.123, 95% CI = 0.77-1.001.47, p<0.001, I2=84.6, k=21) and inpatients (SMD = 0.553, 95% CI = 0.167-0.938, P = 0.005, $I^2 = 0$, k = 3), but not in a nursing home (k=1). Exercise was effective at moderate (SMD=1.33, 95% CI 0.46-2.197, P = 0.003, $I^2 = 83.4$, k = 6) and vigorous (SMD = 1.342, 95% CI = 0.437-2.246, P = 0.004, $I^2 = 1.342$, $I^2 = 1.342$ 91.1, k = 7), but not at light intensity (k = 3), and was effective as aerobic exercise (SMD = 1.045, 95% CI = 0.653-1.437, P < 0.001, $I^2 = 80.97$). The same author group also reported that exercise was effective in improving depressive symptoms in older adults (SMD = -0.90, 95%CI = 0.28-1.51, P = 0.004, $I^2 = 81.63$, P = 0.001, k = 8). In addition, a recent meta-analysis ⁸² (AMSTAR 10, evidence level 1+) confirmed that exercise was effective in reducing depressive symptoms, including those with and without blinded outcomes and with or without intention-to-treat analysis (see table 4). The authors 82 also confirmed results from the Cochrane review that exercise was as effective as psychological intervention and antidepressant treatment for depressive symptoms (see table 4). In another review regarding mental health symptoms 84 (AMSTAR 7, evidence level -1), exercise increase response but not remission of MDD (see table 4). Finally, a trial sequential meta-analysis 85 (AMSTAR 10, evidence level -1) confirmed earlier results that exercise was effective for MDD (SMD -0.66, 95% CI -0.86 to -0.46; p<0.001). Moreover, the authors tentatively suggested that when pooling four RCTs of "high quality" with a trial sequential meta-analysis (two of which were their own studies and concerns have been raised about the inappropriate use of active control groups ⁹⁰) that exercise may no longer be effective.

A narrative systematic review ⁷⁶ (AMSTAR 4, evidence level -1) suggested that exercise did not influence anxiety symptoms, oxygen uptake, blood glucose, body fat or functioning versus control conditions, although all of these outcomes were limited to one study only. A recent meta-analysis of 8 RCTs ⁷⁷ (AMSTAR 10, evidence level -1) found that aerobic exercise was not effective for global cognition (Hedges' g = 0.07, 95% CI = -0.08 to 0.24, I² =0%) nor for any cognitive subdomains. A meta-analysis of 7 RCTs ⁷⁸ demonstrated that aerobic exercise over approximately 12 weeks improved cardiorespiratory fitness (Hedge's g = 0.64, 95%CI = 0.32–0.96, P = 0.001, I² = 67%, k = 8) (AMSTAR 10, evidence level 1+). Exercise also improved quality of life (QOL) overall (SMD = 0.39, 95%CI = 0.47–0.74, P = 0.002, I² = 0.0, k = 5) in addition to physical (SMD = 0.53, 95%CI = 0.22–0.84, P = 0.001, I² = 00.00, k = 5) and psychological (SMD = 0.53, 95%CI = 0.22–0.85, P = 0.001, I² = 4.89, k = 5) QoL domains. However, no effect was seen in social (SMD=0.28, 95%CI = -0.13 to 0.71,p=0.18, I² = 13.04, k = 5) or environmental QoL domains (SMD = 0.36, 95%CI = -0.12 to 0.85,P = 0.14, I² = 30.75, k = 5).

Dropout, adverse events and economic evaluations of exercise interventions in major depression

A meta-analysis of 31 RCTs 91 reported that the dropout rate adjusted for publication bias was 17.2% (95%CI = 13.5-21.7) in MDD only. In MDD participants, higher baseline depressive symptoms (β = 0.0409, 95%CI = 0.0809-0.0009, P = 0.04) predicted greater dropout, whilst supervised interventions delivered by physiotherapists (β = -1.2029, 95%CI = -2.0967 to -0.3091, P = 0.008) and exercise physiologists (β = -1.3396, 95%CI = -2.4478 to -0.2313, P= 0.01) were associated with lower dropout. A comparative meta-analysis (N = 29) established dropout was lower in exercise than control conditions (OR = 0.642, 95%CI = 0.43-0.95, P = 0.02).

Information on adverse events was limited in the reported systematic reviews. One review ⁸³ reported that 6 RCTs reported adverse events. These included muscular pain/symptoms, fatigue and increased depressive symptoms, but there was no difference versus those reported in control conditions. Another review confirmed there were no differences in serious or non-serious adverse events versus control conditions ⁸⁵.

To the best of our knowledge, there have been no economic evaluations of the cost effectiveness of delivering exercise interventions for people with MDD.

Bipolar disorder

In total, 2 systematic reviews reported the benefits of exercise for BD ^{74,75}. Both systematic reviews were rated with an AMSTAR score of 4. The evidence was very limited because of the number of eligible studies in BD patients. A summary of the effects of exercise in BD are reported in table 5. One review identified only two studies that included people with BD, ⁷⁴ finding some evidence that exercise may improve depressive symptoms and reduce stress, but observing no influence on function, general symptoms or clinical global impression. In the other systematic review, only one study was identified in BD patients, ⁷⁵ which suggested that exercise decreased body weight more than in the control group.

Dropout, adverse events and economic evaluations of exercise interventions in bipolar disorder

We found no reports in the included systematic reviews of dropout rates, adverse events or economic evaluations.

Recommendations

Based on our extensive systematic review of the literature, we have made ten recommendations for future research (where gaps are evident) and clinical practice based on the degree of evidence and grades in line with the SIGN methodology checklist. In line with the SIGN and previous EPA guidelines, ⁶⁶ recommendations were graded as A (meta-analysis, systematic review or other level 1 evidence), B (meta-analysis, systematic review or RCT with moderate or high risk of bias), C (large body of evidence from intervention studies, observational data), or D (expert opinion).

Research Recommendations

1. For all SMI groups, more research is needed into the effects of PA interventions during the prodromal and early phase of illness onset (Based on expert opinion, D)

The majority of research on the effect of PA has been conducted in patients with established SMI. However, in the case of physical health outcomes, there is evidence from the general population that using lifestyle changes to prevent deterioration in metabolic health is a more feasible and effective approach than attempting to reverse chronic cardio-metabolic diseases ⁹². Research is needed to determine whether exercise interventions in the early stages of SMI likewise have a positive effect on physical and mental health outcomes. Furthermore, research is needed to assess whether using exercise to improve mental health and psychosocial functioning shortly after illness onset may reduce the likelihood of long-term functional disability, or increase the likelihood of achieving a full and sustained recovery. Given this situation, we recommend a greater research emphasis on evaluating PA interventions during the prodromal phase and first/early presentation of SMI.

2. To optimize treatment 'reach', research should focus on establishing pragmatic, scalable methods for delivering PA as a treatment for SMI (Based on expert opinion, D).

Research priorities should shift from efficacy studies to pragmatic effectiveness trials. Specifically, there is a need to develop replicable and scalable methods for delivering PA interventions to people with SMI, in a format which is accessible, engaging and effective for large numbers of patients. There is sufficient evidence to indicate that the most effective and engaging interventions will be those that are (i) delivered by qualified exercise professionals (rather than mental health staff), and (ii) delivered at sufficient levels of intensity to significantly improve physical capacities, such as cardiorespiratory fitness. Whilst the evidence is clear that better treatment outcomes and less dropout are achieved across SMI for PA interventions delivered by qualified exercise professionals, the potential benefits of training mental health staff (e.g., psychologists, psychiatric nurses - of whom there are many more) on principles of PA in delivering exercise interventions should be evaluated.

3. Controlled trials of exercise for SMI should aim to determine the optimal frequency, mode and dose-response of PA needed to treat SMI and should focus on implementation and culture in clinical practice and prevent adverse events (Based on expert opinion, D).

Whilst there is some evidence from exercise trials in MDD that moderate/high intensity PA is associated with larger effect sizes than low-intensity PA for mental health and cardiorespiratory fitness ⁸¹ and for better cognitive outcomes in established schizophrenia ⁶⁷, more research is specifically required to ascertain the optimum frequency, intensity, time and type of interventions in each SMI, although it is important to note this may vary for each individual based on their preferences and/or illness characteristics. In addition, very few systematic reviews reported adverse events. Future studies should carefully assess any adverse events, particularly in light of any cardiovascular risk and events. This may include screening for cardiovascular risk and appropriately individualizing the optimal frequency and intensity of PA, which has been covered extensively elsewhere ^{93,94}. Of most importance, future translational research focussing on implementation into clinical practice is urgently required. This should involve determining optimal motivational frameworks and understanding cultures in practice that enable the optimal uptake and maintenance of PA.

4. Research examining the effects of interventions to reduce sedentary behaviors and increase PA is needed (Based on expert opinion, D).

Recent meta-analyses have demonstrated that people with SMI are highly sedentary ⁴⁹. With increasing evidence that sedentary behaviours have independent adverse effects on somatic health³⁷, research is needed to investigate if sedentary behaviors can be reduced in this population and tease out the relative importance of reducing sedentary behaviours in the context of structured exercise interventions.

5. The underlying neurobiological mechanisms of PA in SMI are inadequately understood and require further investigation (Based on expert opinion, D).

Several tenable theories have been proposed to explain the positive effects of PA in SMI, but the results are unclear. In particular, it is uncertain what 'dose' of exercise is needed to elicit the purported biological mechanisms, and how these interact with the psycho-social benefits of exercise. Future research should explore the underlying mechanisms of PA efficacy in all SMIs.

6. Undertake long term and cost-effectiveness analyses of PA interventions (Based on expert opinion, D).

Whilst there is an increasing evidence base on the efficacy of PA for SMI, there is an absence of cost effectiveness studies. Undertaking cost-effectiveness research should be a priority. There are also relatively few long term (\geq 12 months) trials, although a recent large RCT lasting 12 months

demonstrated that PA was as effective as internet-delivered CBT, and both were better than the control group ⁹⁵.

Clinical practice recommendations

7. Physical activity should be used as a treatment for mild-moderate depression to improve symptoms and physical fitness (good evidence, A).

There is now an abundance of data demonstrating that as an adjunctive treatment, PA is effective in reducing depressive symptoms in people with MDD. Specifically, based on current evidence, we recommend that intervention consists of 2-3 sessions of supervised aerobic and/or aerobic and resistance training exercise a week of 45-60 minutes duration of moderate intensity. The literature regarding resistance training as a standalone intervention for MDD is limited, but growing. To date, there have been limited direct head-to-head comparisons of different types of exercise interventions (in terms of frequency, intensity, time and type), which precludes definitive conclusions being made about the superiority or better acceptability (or lack thereof) of specific forms of PA, and our current knowledge is derived primarily from interventions comparing "PA" versus non-active interventions. In order to achieve optimal outcomes and less dropouts, the evidence base indicates that physiotherapists/qualified exercise professionals should lead and supervise PA interventions. However, since qualified exercise professionals are not available in each setting, the efficacy and effectiveness of training other personnel in the principles and delivery of PA requires further study. Finally, there is also good evidence that PA can improve cardiorespiratory fitness in people with depression⁷⁸.

8. Physical activity should be utilised as an adjunctive treatment for schizophreniaspectrum disorders, to improve symptoms, cognition and QoL (good evidence, B)

Our review indicates that PA is an effective adjunct to improve symptoms, global cognition and various subdomains, and quality of life in schizophrenia-spectrum disorders. In fact, at least in indirect comparison, the benefits of PA for cognition seem to be comparable to other psychological interventions (e.g., cognitive remediation) and should be a core part of multidisciplinary treatment. The most consistent evidence to date is available for aerobic exercise that meets 150 minutes of moderate to vigorous PA per week. There are limited data available specifically on resistance training for people with schizophrenia-spectrum disorders and data are inadequate comparing the potential superiority of various forms of PA to each other. Nonetheless, interventions supervised by qualified professionals, with a motivational component result in less dropout.

9. Physical activity should be used to improve physical health in people with SMI (Some evidence, C)

Whilst there is an abundance of evidence for PA to prevent the onset of a number of physical diseases (e.g., cardiovascular disease) and to have the potential to improve these conditions in the general population, the evidence base in individual SMIs is mixed, possibly due to poor adherence and the paucity of studies that have targeted this important topic. However, there is some indication that PA can improve cardiovascular disease risk markers in people with SMI. Given the weight of evidence in the general population and some initial data in people with SMI, we advocate that PA forms a core part of preventing and managing poor physical health in people with SMI. Nevertheless, more research should be performed in this area to further strengthen the support for this recommendation.

10. People with SMI should be screened for PA habits in primary and secondary care (Based on expert opinion, D).

In light of the evidence of the health benefits of PA, the use of brief PA screening in primary/secondary care in the general population as a health indicator and to manage chronic conditions is common ^{96,97}. A previous systematic review ⁹⁸ identified, no specific self-report PA tool has robust psychometric properties and can be advocated for routine use in clinical practice. Nonetheless, given that people with all SMIs are known to engage in low levels of PA and high SB ⁴⁹ a routine measure of PA to monitor PA habits could prove useful. To the best of our knowledge, only one PA tool, the SIMPAQ ⁹⁹ has been developed specifically for use to measure PA and SB in clinical practice. Given the above literature, we advocate that people with SMI are screened for PA and advised on increasing activity levels in primary and secondary services. Further research is needed to identify a practical and valid measure that can be used in clinical care.

Conclusion

There is now considerable empirical evidence supporting the use of PA interventions in the treatment of MDD and schizophrenia-spectrum disorders. Specifically, our meta-review supports the use of aerobic exercise of moderate-vigorous intensity at a frequency of 2-3 times a week, ideally supervised by qualified professionals and achieving 150 minutes of MVPA per week in order to improve outcomes in people with MDD and schizophrenia-spectrum disorders. There is also some evidence that a combination of aerobic and resistance training meeting the above frequency, intensity and time criteria can improve outcomes in people with MDD and schizophrenia-spectrum disorders. There is growing interest in the use of resistance training as a standalone PA intervention. However, the literature is equivocal in both MDD and schizophrenia-spectrum disorders. Currently, the evidence for use of PA in patients with BD is promising, but limited by the paucity of studies, and it is therefore not possible to reach any firm conclusions regarding population-specific recommendations above and beyond those aimed at the general population. Across all disorders, there have been limited interventions that have specifically compared various forms of PA versus one another, which again precludes reaching any firm conclusions based on the relative superiority of the different forms of PA, and one should be aware that our knowledge to date is largely drawn from studies comparing PA versus non-active control conditions. The underlying psychobiological mechanisms of the benefits derived from PA are only poorly understood, and further research is needed to elucidate causal relationships between exercise and improvements in the mental and physical domains of people with SMI. Finally, due to absence of data, it is currently unclear whether PA is a cost-effective treatment option, and more work is needed to establish the financial aspects of this mode of treatment compared with conventional approaches, taking into account the various domains of lives of people with SMI that PA can affect.

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Conflict of Interest

BS, DV, JF, MH, MS, MG, NV, FK have no conflicts of interest to declare.

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References

- 1. Whiteford HA, Ferrari AJ, Degenhardt L, Feigin V, Vos T. The global burden of mental, neurological and substance use disorders: an analysis from the Global Burden of Disease Study 2010. *PLoS One*. 2015;10(2):e0116820.
- 2. Walker ER, McGee RE, Druss BG. Mortality in Mental Disorders and Global Disease Burden Implications: A Systematic Review and Meta-analysis. *JAMA Psychiatry*. 2015.
- 3. Nielsen RE, Uggerby AS, Jensen SOW, McGrath JJ. Increasing mortality gap for patients diagnosed with schizophrenia over the last three decades--a Danish nationwide study from 1980 to 2010. *Schizophrenia Research*. 2013;146(1-3):22-27.
- 4. McGrath J, Saha S, Chant D, Welham J. Schizophrenia: a concise overview of incidence, prevalence, and mortality. *Epidemiol Rev.* 2008;30:67-76.
- 5. Hayes JF, Miles J, Walters K, King M, Osborn DPJ. A systematic review and meta-analysis of premature mortality in bipolar affective disorder. *Acta Psychiatrica Scandinavica*. 2015.
- 6. Whiteford HA, Degenhardt L, Rehm J, et al. Global burden of disease attributable to mental and substance use disorders: findings from the Global Burden of Disease Study 2010. *Lancet*. 2013;382(9904):1575-1586.
- 7. Chesney E, Goodwin GM, Fazel S. Risks of all-cause and suicide mortality in mental disorders: a meta-review. *World Psychiatry*. 2014;13(2):153-160.
- 8. Correll CU, Detraux J, De Lepeleire J, De Hert M. Effects of antipsychotics, antidepressants and mood stabilizers on risk for physical diseases in people with schizophrenia, depression and bipolar disorder. *World Psychiatry: Official Journal Of The World Psychiatric Association (WPA).* 2015;14(2):119-136.
- 9. Correll CU, Solmi M, Veronese N, et al. Prevalence, incidence and mortality from cardiovascular disease in patients with pooled and specific severe mental illness: a large-scale meta-analysis of 3,211,768 patients and 113,383,368 controls. *World Psychiatry*. 2017;16(2):163-180.
- 10. Vancampfort D, Correll CU, Galling B, et al. Diabetes mellitus in people with schizophrenia, bipolar disorder and major depressive disorder: a systematic review and large scale meta-analysis. *World Psychiatry*. 2016;15(2):166-174.
- 11. Vancampfort D, Stubbs B, Mitchell AJ, et al. Risk of metabolic syndrome and its components in people with schizophrenia and related psychotic disorders, bipolar disorder and major depressive disorder: a systematic review and meta-analysis. *World Psychiatry: Official Journal Of The World Psychiatric Association (WPA)*. 2015;14(3):339-347.
- 12. Partti K, Vasankari T, Kanervisto M, et al. Lung function and respiratory diseases in people with psychosis: population-based study. *Br J Psychiatry*. 2015;207(1):37-45.
- 13. Olfson M, Gerhard T, Huang C, Crystal S, Stroup TS. Premature Mortality Among Adults With Schizophrenia in the United States. *JAMA Psychiatry*. 2015;72(12):1172-1181.
- 14. Stubbs B, De Hert M, Sepehry AA, et al. A meta-analysis of prevalence estimates and moderators of low bone mass in people with schizophrenia. *Acta Psychiatrica Scandinavica*. 2014.
- 15. Stubbs B, Koyanagi A, Veronese N, et al. Physical multimorbidity and psychosis: comprehensive cross sectional analysis including 242,952 people across 48 low- and middle-income countries. *BMC Med.* 2016;14(1):189.
- 16. Metzler S, Dvorsky D, Wyss C, et al. Changes in neurocognitive functioning during transition to manifest disease: Comparison of individuals at risk for schizophrenic and bipolar affective psychoses. *Psychological Medicine*. 2015;45(10):2123-2134.
- 17. Amann B, Gomar JJ, Ortiz-Gil J, et al. Executive dysfunction and memory impairment in schizoaffective disorder: a comparison with bipolar disorder, schizophrenia and healthy controls...[corrected][published erratum appears in PSYCHOL MED 2012 Oct; 42: 2136]. *Psychological Medicine*. 2012;42(10):2127-2135 2129p.

- 18. Elias LR, Miskowiak KW, Vale AM, et al. Cognitive Impairment in Euthymic Pediatric Bipolar Disorder: A Systematic Review and Meta-Analysis. *J Am Acad Child Adolesc Psychiatry*. 2017;56(4):286-296.
- 19. Koster LS, Carbon M, Correll CU. Emerging drugs for schizophrenia: an update. *Expert Opin Emerg Drugs*. 2014;19(4):511-531.
- 20. Carbon M, Correll CU. Thinking and acting beyond the positive: the role of the cognitive and negative symptoms in schizophrenia. *CNS Spectr.* 2014;19 Suppl 1:38-52; quiz 35-37, 53.
- 21. Leucht S, Cipriani A, Spineli L, et al. Comparative efficacy and tolerability of 15 antipsychotic drugs in schizophrenia: a multiple-treatments meta-analysis. *Lancet.* 2013;382(9896):951-962.
- 22. Correll CU, Rubio JM, Inczedy-Farkas G, Birnbaum ML, Kane JM, Leucht S. Efficacy of 42 Pharmacologic Cotreatment Strategies Added to Antipsychotic Monotherapy in Schizophrenia: Systematic Overview and Quality Appraisal of the Meta-analytic Evidence. *JAMA Psychiatry.* 2017.
- 23. Turner DT, van der Gaag M, Karyotaki E, Cuijpers P. Psychological interventions for psychosis: a meta-analysis of comparative outcome studies. *Am J Psychiatry*. 2014;171(5):523-538.
- 24. Jobst A, Brakemeier EL, Buchheim A, et al. European Psychiatric Association Guidance on psychotherapy in chronic depression across Europe. *Eur Psychiatry*. 2016;33:18-36.
- 25. Cuijpers P, Cristea IA, Karyotaki E, Reijnders M, Huibers MJ. How effective are cognitive behavior therapies for major depression and anxiety disorders? A meta-analytic update of the evidence. *World Psychiatry*. 2016;15(3):245-258.
- 26. Huhn M, Tardy M, Spineli LM, et al. Efficacy of pharmacotherapy and psychotherapy for adult psychiatric disorders: a systematic overview of meta-analyses. *JAMA Psychiatry*. 2014;71(6):706-715.
- 27. Fusar-Poli P, Smieskova R, Kempton MJ, Ho BC, Andreasen NC, Borgwardt S. Progressive brain changes in schizophrenia related to antipsychotic treatment? A meta-analysis of longitudinal MRI studies. *Neurosci Biobehav Rev.* 2013;37(8):1680-1691.
- 28. Cella M, Preti A, Edwards C, Dow T, Wykes T. Cognitive remediation for negative symptoms of schizophrenia: A network meta-analysis. *Clin Psychol Rev.* 2017;52:43-51.
- 29. Naci H, Ioannidis JPA. Comparative effectiveness of exercise and drug interventions on mortality outcomes: metaepidemiological study. *BMJ (Clinical Research Ed)*. 2013;347:f5577-f5577.
- 30. Nelson ME, Rejeski WJ, Blair SN, et al. Physical Activity and Public Health in Older Adults: Recommendation from the American College of Sports Medicine and the American Heart Association. *Medicine & Science in Sports & Exercise*. 2007;39(8):1435-1445.
- 31. Sattelmair J, Pertman J, Ding EL, Kohl HW, 3rd, Haskell W, Lee IM. Dose response between physical activity and risk of coronary heart disease: a meta-analysis. *Circulation*. 2011;124(7):789-795.
- 32. Haskell WL, Lee IM, Pate RR, et al. Physical activity and public health: updated recommendation for adults from the American College of Sports Medicine and the American Heart Association. *Med Sci Sports Exerc.* 2007;39(8):1423-1434.
- 33. Hayashino Y, Jackson JL, Hirata T, et al. Effects of exercise on C-reactive protein, inflammatory cytokine and adipokine in patients with type 2 diabetes: a meta-analysis of randomized controlled trials. *Metabolism: Clinical And Experimental.* 2014;63(3):431-440.
- 34. Bergström G, Behre CJ, Schmidt C. Moderate Intensities of Leisure-Time Physical Activity Are Associated With Lower Levels of High-Sensitivity C-Reactive Protein in Healthy Middle-Aged Men. *Angiology.* 2012;63(6):412-415.
- 35. Kohler CA, Freitas TH, Maes M, et al. Peripheral cytokine and chemokine alterations in depression: a meta-analysis of 82 studies. *Acta Psychiatr Scand.* 2017;135(5):373-387.

- 36. Sedentary Behaviour Research N. Letter to the Editor: Standardized use of the terms 'sedentary' and 'sedentary behaviours'. *Applied Physiology, Nutrition & Metabolism.* 2012;37(3):540-542.
- 37. Biswas A, Oh PI, Faulkner GE, et al. Sedentary Time and Its Association With Risk for Disease Incidence, Mortality, and Hospitalization in Adults: A Systematic Review and Meta-analysis. *Annals of Internal Medicine*. 2015;162(2):123-132.
- 38. Kodama S, Saito K, Tanaka S, et al. Cardiorespiratory fitness as a quantitative predictor of all-cause mortality and cardiovascular events in healthy men and women: a meta-analysis. *JAMA: Journal of the American Medical Association*. 2009;301(19):2024-2035.
- 39. Voelcker-Rehage C, Niemann C. Structural and functional brain changes related to different types of physical activity across the life span. *Neuroscience and Biobehavioral Reviews*. 2013;37(9, Part B):2268-2295.
- 40. Stubbs B, Chen LJ, Chang CY, Sun WJ, Ku PW. Accelerometer-assessed light physical activity is protective of future cognitive ability: A longitudinal study among community dwelling older adults. *Exp Gerontol*. 2017;91:104-109.
- 41. Snowden M, Steinman L, Mochan K, et al. Effect of exercise on cognitive performance in community-dwelling older adults: Review of intervention trials and recommendations for public health practice and research. *Journal of the American Geriatrics Society*. 2011;59(4):704-716.
- 42. Prakash RS, Voss MW, Erickson KI, Kramer AF. Physical Activity and Cognitive Vitality. *Annual Review of Psychology*. 2015;66(1):769-797.
- 43. Stanmore E, Stubbs B, Vancampfort D, de Bruin ED, Firth J. The effect of active video games on cognitive functioning in clinical and non-clinical populations: A meta-analysis of randomized controlled trials. *Neurosci Biobehav Rev.* 2017;78:34-43.
- 44. Firth J, Stubbs B, Vancampfort D, et al. Effect of aerobic exercise on hippocampal volume in humans: A systematic review and meta-analysis. *Neuroimage*. 2018;166:230-238.
- 45. Schuch F, Vancampfort, D., Firth., J., Rosenbaum., Ward, PB., Silva, E., Hallgren, Dunn, AL., Deslandes., A., Fleck, MC., Carvalho, AF., Stubbs, B. Physical activity and incident depression: A meta-analysis of prospective cohort studies. In. American Journal of Psychiatry In press.
- 46. Stubbs B, Firth J, Berry A, et al. How much physical activity do people with schizophrenia engage in? A systematic review, comparative meta-analysis and meta-regression. *Schizophr Res.* 2016.
- 47. Vancampfort D, Firth J, Schuch F, et al. Physical activity and sedentary behavior in people with bipolar disorder: A systematic review and meta-analysis. *J Affect Disord*. 2016;201:145-152.
- 48. Schuch F, Vancampfort D, Firth J, et al. Physical activity and sedentary behavior in people with major depressive disorder: A systematic review and meta-analysis. *J Affect Disord*. 2016;210:139-150.
- 49. Vancampfort D, Firth J, Schuch FB, et al. Sedentary behavior and physical activity levels in people with schizophrenia, bipolar disorder and major depressive disorder: a global systematic review and meta-analysis. *World Psychiatry*. 2017;16(3):308-315.
- 50. (WHO) WHO. 10 facts on physical activity. In:2014.
- 51. Vancampfort D, Rosenbaum S, Schuch F, et al. Cardiorespiratory Fitness in Severe Mental Illness: A Systematic Review and Meta-analysis. *Sports Med.* 2016.
- 52. Vancampfort D, Correll CU, Probst M, et al. A review of physical activity correlates in patients with bipolar disorder. *J Affect Disord*. 2013;145(3):285-291.
- 53. Vancampfort D, Knapen J, Probst M, Scheewe T, Remans S, De Hert M. A systematic review of correlates of physical activity in patients with schizophrenia. *Acta Psychiatrica Scandinavica*. 2012;125(5):352-362.
- 54. Vancampfort D, Stubbs B, Sienaert P, et al. What are the factors that influence physical activity participation in individuals with depression? A review of physical activity correlates from 59 studies. *Psychiatria Danubina*. 2015;27(3):210-224.

- 55. Firth J, Rosenbaum S, Stubbs B, Gorczynski P, Yung AR, Vancampfort D. Motivating factors and barriers towards exercise in severe mental illness: a systematic review and meta-analysis. *Psychol Med.* 2016;46(14):2869-2881.
- 56. Vancampfort D, Stubbs B, Ward PB, Teasdale S, Rosenbaum S. Integrating physical activity as medicine in the care of people with severe mental illness. *The Australian And New Zealand Journal Of Psychiatry*. 2015;49(8):681-682.
- 57. Vancampfort D, Rosenbaum S, Probst M, et al. What are the top 10 physical activity research questions in schizophrenia? *Disabil Rehabil.* 2016;38(22):2235-2243.
- 58. Vancampfort D, Rosenbaum S, Probst M, et al. Top 10 research questions to promote physical activity in bipolar disorders: A consensus statement from the International Organization of Physical Therapists in Mental Health. *J Affect Disord*. 2016;195:82-87.
- 59. Moher D, Liberati A, Tetzlaff J, Altman DG. Preferred Reporting Items for Systematic Reviews and Meta-Analyses: The PRISMA Statement. *PLoS Clinical Trials.* 2009;6(7):1-6.
- 60. Heun R, Gaebel W. The relevance of EPA guidance papers in the framework of the European Psychiatric Association. *Eur Psychiatry*. 2015;30(3):357-359.
- 61. Association AP. Diagnostic and Statistical Manual of Mental Disorders DSM-IV-TR. 4th edition. In: American Psychiatric Association; 2000.
- 62. organisation Wh. The ICD-10 Classification of Mental and Behavioural Disorders Diagnostic Criteria for Research. In:1993.
- 63. Caspersen CJ, Powell KE, Christenson GM. Physical activity, exercise, and physical fitness: definitions and distinctions for health-related research. *Public Health Reports (Washington, DC: 1974).* 1985;100(2):126-131.
- 64. Shea BJ, Hamel C, Wells GA, et al. AMSTAR is a reliable and valid measurement tool to assess the methodological quality of systematic reviews. *Journal of Clinical Epidemiology*. 2009;62(10):1013-1020.
- 65. Shea BJ, Grimshaw JM, Wells GA, et al. Development of AMSTAR: a measurement tool to assess the methodological quality of systematic reviews. *BMC Medical Research Methodology*. 2007;7:10-17.
- 66. Schmidt SJ, Schultze-Lutter F, Schimmelmann BG, et al. EPA guidance on the early intervention in clinical high risk states of psychoses. *Eur Psychiatry*. 2015;30(3):388-404.
- 67. Firth J, Stubbs B, Rosenbaum S, et al. Aerobic Exercise Improves Cognitive Functioning in People With Schizophrenia: A Systematic Review and Meta-Analysis. *Schizophr Bull.* 2016.
- 68. Firth J, Cotter J, Elliott R, French P, Yung AR. A systematic review and meta-analysis of exercise interventions in schizophrenia patients. *Psychological Medicine*. 2015;45(7):1343-1361.
- 69. Firth J, Cotter J, Carney R, Yung AR. The pro-cognitive mechanisms of physical exercise in people with schizophrenia. *Br J Pharmacol.* 2017;174(19):3161-3172.
- 70. Soundy A, Muhamed, A., Stubbs, B., Probst, M., Vancampfort, D. The benefits of walking for individuals with schizophrenia spectrum disorders: A systematic review. In. International Journal of Therapy and Rehabilitation 21 (9), 410-4202014.
- 71. Martin H, Beard, S., Clissold, N., Androas, K., Currey, L. Combined aerobic and resistance exercise interventions for

individuals with schizophrenia: A systematic review. In. Mental Health and Physical Activity, March, pg 147-1552017.

- 72. Sanada K, Zorrilla I, Iwata Y, et al. The Efficacy of Non-Pharmacological Interventions on Brain-Derived Neurotrophic Factor in Schizophrenia: A Systematic Review and Meta-Analysis. *Int J Mol Sci.* 2016;17(10).
- 73. Vancampfort D, Rosenbaum S, Ward PB, Stubbs B. Exercise improves cardiorespiratory fitness in people with schizophrenia: A systematic review and meta-analysis. *Schizophrenia Research*. 2015.
- 74. de Sa Filho AS, de Souza Moura AM, Lamego MK, et al. Potential Therapeutic Effects of Physical Exercise for Bipolar Disorder. *CNS Neurol Disord Drug Targets*. 2015;14(10):1255-1259.

- 75. Thomson D, Turner A, Lauder S, et al. A brief review of exercise, bipolar disorder, and mechanistic pathways. *Front Psychol.* 2015;6:147.
- 76. de Souza Moura AM, Lamego MK, Paes F, et al. Comparison Among Aerobic Exercise and Other Types of Interventions to Treat Depression: A Systematic Review. *CNS Neurol Disord Drug Targets*. 2015;14(9):1171-1183.
- 77. Brondino N, Rocchetti M, Fusar-Poli L, et al. A systematic review of cognitive effects of exercise in depression. *Acta Psychiatr Scand.* 2017;135(4):285-295.
- 78. Stubbs B, Rosenbaum S, Vancampfort D, Ward PB, Schuch FB. Exercise improves cardiorespiratory fitness in people with depression: A meta-analysis of randomized control trials. *J Affect Disord*. 2016;190:249-253.
- 79. Schuch FB, Vancampfort D, Rosenbaum S, Richards J, Ward PB, Stubbs B. Exercise improves physical and psychological quality of life in people with depression: A meta-analysis including the evaluation of control group response. *Psychiatry Res.* 2016;241:47-54.
- 80. Schuch FB, Vancampfort D, Rosenbaum S, et al. Exercise for depression in older adults: a metaanalysis of randomized controlled trials adjusting for publication bias. *Rev Bras Psiquiatr*. 2016;38(3):247-254.
- 81. Schuch FB, Vancampfort D, Richards J, Rosenbaum S, Ward PB, Stubbs B. Exercise as a treatment for depression: A meta-analysis adjusting for publication bias. *J Psychiatr Res.* 2016;77:42-51.
- 82. Kvam S, Kleppe CL, Nordhus IH, Hovland A. Exercise as a treatment for depression: A metaanalysis. *J Affect Disord*. 2016;202:67-86.
- 83. Cooney GM, Dwan K, Greig CA, et al. Exercise for depression. *The Cochrane Database Of Systematic Reviews*. 2013;9:CD004366.
- 84. Silveira H, Moraes H, Oliveira N, Coutinho ES, Laks J, Deslandes A. Physical exercise and clinically depressed patients: a systematic review and meta-analysis. *Neuropsychobiology*. 2013;67(2):61-68.
- 85. Krogh J, Hjorthoj C, Speyer H, Gluud C, Nordentoft M. Exercise for patients with major depression: a systematic review with meta-analysis and trial sequential analysis. *BMJ Open.* 2017;7(9):e014820.
- 86. Rosenbaum S, Tiedemann A, Sherrington C, Curtis J, Ward PB. Physical activity interventions for people with mental illness: a systematic review and meta-analysis. *The Journal Of Clinical Psychiatry*. 2014.
- 87. Pajonk FG, Wobrock T, Gruber O, et al. Hippocampal plasticity in response to exercise in schizophrenia. *Arch Gen Psychiatry.* 2010;67(2):133-143.
- 88. Keller-Varady K, Varady PA, Roh A, et al. A systematic review of trials investigating strength training in schizophrenia spectrum disorders. *Schizophr Res.* 2017.
- 89. Vancampfort D, Rosenbaum S, Schuch FB, Ward PB, Probst M, Stubbs B. Prevalence and predictors of treatment dropout from physical activity interventions in schizophrenia: a meta-analysis. *Gen Hosp Psychiatry*. 2016;39:15-23.
- 90. Ekkekakis P. Honey, I shrunk the pooled SMD! Guide to critical appraisal of systematic reviews and meta-analyses using the Cochrane review on exercise for depression as example. *Mental Health and Physical Activity.* 2015;8:21-36.
- 91. Stubbs B, Vancampfort D, Rosenbaum S, et al. Dropout from exercise randomized controlled trials among people with depression: A meta-analysis and meta regression. *J Affect Disord*. 2016;190:457-466.
- 92. Schellenberg ES, Dryden DM, Vandermeer B, Ha C, Korownyk C. Lifestyle interventions for patients with and at risk for type 2 diabetes: a systematic review and meta-analysis. *Ann Intern Med.* 2013;159(8):543-551.
- 93. Vancampfort D, Rosenbaum S, Probst M, et al. Promotion of cardiorespiratory fitness in schizophrenia: a clinical overview and meta-analysis. *Acta Psychiatr Scand.* 2015;132(2):131-143.

- 94. De Hert M, Dekker JM, Wood D, Kahl KG, Holt RIG, Möller HJ. Cardiovascular disease and diabetes in people with severe mental illness position statement from the European Psychiatric Association (EPA), supported by the European Association for the Study of Diabetes (EASD) and the European Society of Cardiology (ESC). *European Psychiatry*. 2009;24(6):412-424.
- 95. Hallgren M, Kraepelien M, Öjehagen A, et al. Physical exercise and internet-based cognitive behavioural therapy in the treatment of depression: randomised controlled trial. *The British Journal of Psychiatry*. 2015:bjp. bp. 114.160101.
- 96. Golightly YM, Allen KD, Ambrose KR, et al. Physical Activity as a Vital Sign: A Systematic Review. *Prev Chronic Dis.* 2017;14:E123.
- 97. Carlson SA, Maynard LM, Fulton JE, Hootman JM, Yoon PW. Physical activity advice to manage chronic conditions for adults with arthritis or hypertension, 2007. *Preventive Medicine*. 2009;49(2-3):209-212.
- 98. Soundy A, Roskell C, Stubbs B, Vancampfort D. Selection, use and psychometric properties of physical activity measures to assess individuals with severe mental illness: a narrative synthesis. *Archives Of Psychiatric Nursing*. 2014;28(2):135-151.
- 99. Rosenbaum S, Ward PB. The Simple Physical Activity Questionnaire. *Lancet Psychiatry*. 2016;3(1):e1.