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4 **Autonomy support, light physical activity and psychological well-being in Rheumatoid**

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Arthritis: a cross-sectional study

6 **Running title:** Correlates of light-intensity PA in RA

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32

1 **Abstract**

2 **Background:** Participation in physical activity may improve psychological well-
3 being among people with Rheumatoid Arthritis (RA). This study examined the implications
4 of autonomy support for physical activity, on objectively assessed light physical activity
5 (*LPA*) engagement, and in turn, psychological well-being in RA. In addition, the role of
6 lower-limb functional disability in these associations was investigated. **Methods:** RA patients
7 (N = 50) completed questionnaires assessing 1) autonomy support for physical activity [from
8 a patient-specified important other], 2) functional disability to ‘rise’ and ‘walk’ (functional
9 disabilityRW), 3) depressive symptoms, and 4) subjective vitality. Levels of LPA [100-2019
10 counts/minute], were calculated from 7-days of accelerometry. **Results:** Path analysis
11 supported a model ($\chi^2(2) = 2.44, p = .304, CFI = .99, SRMR = .05, RMSEA = .07$) in which
12 important other autonomy support for physical activity significantly and positively predicted
13 LPA engagement. In turn, LPA was significantly and positively associated with subjective
14 vitality, and significantly and negatively linked to depressive symptoms. These associations
15 were observed independently of adverse direct relationships between Functional
16 disabilityRW with depressive symptoms and subjective vitality. **Conclusions:** Important
17 other autonomy support for physical activity may hold positive consequences for LPA
18 engagement and related mental health states in RA, independent of the negative effects of
19 lower-limb functional disability.

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21 **Key words:** Functional disability, Autonomy support, Light physical activity, Accelerometer,
22 Psychological well-being, Rheumatoid Arthritis.

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1 **Introduction**

2 Research underlines the benefits of regular participation in physical activity for promoting
3 more optimal psychological health among both healthy adults and patient cohorts (Bauman,
4 Merom, Bull, Buchner, & Fiatarone Singh, 2016; Cairns & McVeigh, 2009; Penedo & Dahn,
5 2005; Windle, Hughes, Linck, Russell, & Woods, 2010). People living with Rheumatoid
6 Arthritis (RA) frequently report compromised psychological well-being (Gettings, 2010;
7 Murphy, Sacks, Brady, Hootman, & Chapman, 2012). Thus, participation in physical activity
8 may prove beneficial for enhancing psychological health in this patient group.

9 To date, the focus of RA studies has been on the psychological health benefits
10 resulting from participation in physical activity above moderate intensity (i.e., ≥ 3 metabolic
11 equivalents, METS) (Kelley, Kelley, & Hootman, 2015; Verhoeven et al., 2016; Windle et
12 al., 2010). However, the reduced functional ability associated with RA, may restrict
13 individuals' perceived ability to engage and subsequently, overtly participate in moderate
14 intensity physical activity (Hernandez-Hernandez, Ferraz-Amaro, & Diaz-Gonzalez, 2014;
15 Sokka et al., 2008; Veldhuijzen van Zanten et al., 2015). Conversely, participation in lower-
16 intensity physical activities (i.e., light physical activity, 1.6 - 2.9 METS) may be perceived as
17 relatively more feasible and achievable by people living with RA (Manns, Dunstan, Owen, &
18 Healy, 2012), and is being increasingly advocated to improve overall health in several other
19 clinical and ageing populations (Buman et al., 2010; Ekwall, Lindberg, & Magnusson, 2009;
20 Larsen et al., 2014; Manns et al., 2012; Trinity, 2017). However, studies to date are yet to
21 investigate the psychological health implications of engagement in light physical activity
22 (LPA) for people living with RA, as well as factors that may influence engagement in this
23 behaviour (i.e., determinants).

24 The social environment operating within physical activity settings has been proposed
25 as a key determinant of physical activity behaviour. For example, Self-determination theory

1 (SDT), suggests where the social environment supports an individual's sense of autonomy
2 with regards to their physical activity engagement (i.e., it promotes choice and
3 understanding), this is more likely to encourage the adoption and maintenance of physical
4 activity behaviour (Chan, Lonsdale, Ho, Yung, & Chan, 2009; Fortier, Duda, Guerin, &
5 Teixeira, 2012; Milne, Wallman, Guilfoyle, Gordon, & Corneya, 2008). The social
6 environment is largely created by the interpersonal behaviours of 'significant' or 'important'
7 others acting within that setting. When considering physical activity in RA, this 'important
8 other' could be the health care professional (e.g., rheumatology consultant, nurse, or GP) or
9 other individuals the patient considers relevant to their attempts to be physically active (e.g.,
10 a spouse, offspring or friend) (Edmunds, 2007; Hardcastle, Blake, & Hagger, 2012; Williams,
11 2002).

12 Recent research revealed autonomy support for physical activity provided by
13 'important others', was linked to higher levels of self-reported total physical activity
14 (comprising light, moderate and vigorous) among people living with RA (Yu et al., 2015).
15 However, this study did not examine the role of autonomy support for LPA participation
16 specifically, and a reliance on self-report somewhat limits the validity of these findings. Thus,
17 research is required to investigate the implications of autonomy support for objectively
18 assessed LPA engagement in RA, to determine whether the social environment represents a
19 salient and modifiable determinant of LPA in these patients. In turn, investigating the extent
20 to which variability in LPA (predicted by autonomy support) is associated with psychological
21 well-being among people living with RA, will help to establish the potential value of
22 interventions focused on creating autonomy supportive physical activity environments for
23 improving psychological well-being among this patient group.

24 Upon investigating these associations, we must still consider the possibility that the
25 compromised physical function symptomatic of RA may represent a barrier to even low-

1 intensity physical activity engagement for these patients. Of particular relevance is functional
2 disability related to standing and walking – two common light intensity activities. Indeed,
3 walking is reported as the most common behaviour undertaken by people living with RA, and
4 light intensity walking (including standing incidental and sporadic movement) comprises
5 approximately 90% of ambulatory behaviour (Paul et al., 2014). Accordingly, an individual's
6 disability related to 'standing' and 'walking' (i.e., lower-limb functional disability) should be
7 taken into account when seeking to identify modifiable determinants of LPA participation in
8 RA (e.g., the social environment).

9 The primary aim of this research was therefore to examine the implications of
10 autonomy support for physical activity *and* lower-limb functional disability, for levels of
11 objectively assessed LPA engagement, and associated positive and negative indicators of
12 well-being in RA. Specifically, this study sought to examine the sequential associations
13 between perceived autonomy support from a participant specified 'important other', lower-
14 limb functional disability to 'rise' and to 'walk', accelerometer assessed LPA, and in turn,
15 depressive symptoms and subjective vitality among people living with RA (Figure 1). These
16 two outcomes are particularly pertinent to psychological functioning in RA. Specifically,
17 depression represents a highly prevalent co-morbidity in RA (Ang, Choi, Kroenke, & Wolfe,
18 2005; Margaretten, Julian, Katz, & Yelin, 2011; Treharne et al., 2005), and subjective vitality
19 provides an indication of an individuals overall optimal psychological functioning (Rouse et
20 al., 2015; Ryan & Deci, 2001).

21 It was hypothesised that higher lower-limb functional disability (poorer function),
22 would be negatively associated with LPA engagement. It was also expected that perceived
23 'important other' autonomy support would be *independently* and positively associated with
24 LPA, and that LPA would be subsequently positively related to subjective vitality, and
25 negatively associated with the prevalence of depressive symptoms (Figure 1). That is, we

1 propose that autonomy support for physical activity predicts variability in LPA, to the degree
2 it will hold positive implications for psychological well-being among people living with RA,
3 after taking into account lower-limb functional disability.

4 **Methods**

5 **Participants**

6 Patients with RA were recruited as part of the xxxxx study (Trial Number:xxxxx).
7 The xxxxx study was a randomised controlled trial, with the aim of promoting self-
8 determined motivation for exercise engagement and improving cardiorespiratory fitness (xxxx
9 *study reference*). Baseline data were used to answer the current research questions. The study
10 was granted ethical approved by the local National Health Service Research Ethics
11 Committee (*reference: xxxxx*).

12 **Recruitment and protocol**

13 Information sheets were distributed to interested participants attending Rheumatology
14 outpatient clinics at xxxxx Hospital (xxxxx NHS Foundation Trust). In total, 115 participants
15 ($\text{Mage} = 53.98 \pm 12.47$ years) were recruited to the xxxxx study and provided informed
16 consent. Questionnaire data were collected from participants during appointments at xxxxx
17 Hospital. Following this, accelerometer data were collected over 7 days among a sub-
18 subsample of willing participants ($N = 97$). The full xxxxx study protocol is detailed
19 elsewhere (*xxxx study reference*).

20 **Measures**

21 ***Important other Autonomy Support for Physical Activity***

22 Important other support for physical activity (here-on referred to as autonomy
23 support) was assessed using an adapted version of the Important Other Climate Questionnaire
24 (IOCQ) (Williams et al., 2006). Participants were first asked to indicate who they consider to
25 be the ‘most important person in their effort to engage in physical activity’ (e.g., a spouse,

1 sibling, offspring, friend). Following this, participants responded to 6 statements regarding
2 the degree of perceived autonomy for physical activity provided by their important other, as
3 follows; 1) I feel that my important other has provided me with choices and options in
4 regards to my physical activity, 2) I feel my important other understands how I see things
5 with respect to my physical activity participation, 3) my important other conveys confidence
6 in my ability to make changes regarding my physical activity participation, 4) my important
7 other listens to how I would like to do things regarding physical activity, 5) my important
8 other encourages me to ask questions about physical activity, 6) my important other tries to
9 understand how I see my physical activity participation before suggesting any changes.
10 Responses were given on a 7-point Likert scale ranging from 1 (strongly disagree) to 7
11 (strongly agree). The IOCQ demonstrated high internal reliability in this sample ($\alpha = .92$).

12 ***Functional disability to 'rise' and 'walk'***

13 Participants' functional ability to 'rise' and to 'walk' (functional disabilityRW) was
14 determined using the 'rising' and 'walking' subscales of the Stanford Health Assessment
15 Questionnaire (HAQ) (Kirwan & Reeback, 1986) Following the stem, "Are you able to....",
16 respondents were asked to rate on a scale from 0 (without any difficulty) to 3 (unable to do),
17 the extent to which they are able to undertake functions related to *rising* (functions; 1) stand
18 up from an armless straight chair and 2) get in and out of bed) and *walking* (functions; 1)
19 walk outside on flat ground and 2) climb up five stairs). The score given to each subscale is
20 the highest score reported across the two questions. Higher scores represent higher functional
21 disability (i.e., poorer ability to 'rise' and 'walk'). A mean functional disabilityRW score was
22 derived (to represent lower-limb functional disability), as the average score from the two
23 subscales. Overall functional disability was also determined from the HAQ and is reported
24 herein for descriptive purposes.

1 ***Objectively assessed physical activity behaviours***

2 LPA was assessed using GT3X accelerometers (Actigraph). Participants wore the
3 accelerometer on the right hip for 7 consecutive days, removing only for water-based
4 activities (e.g., swimming and bathing) (Semanik et al., 2010; Trost, McIver, & Pate, 2005).
5 The GT3X detected movements over sixty-second epochs in this study. Movement counts
6 within each minute-epoch were summed and converted to activity counts that were
7 interpreted to determine LPA engagement [i.e., ≥ 100 and < 2020 counts per minute, (cpm)]
8 (Troiano et al., 2008).

9 *Accelerometer data reduction*

10 Actilife software (version 6.2) was used to analyse the data. Data pertaining to waking
11 hours [i.e., 7:00am–10:30pm - identified from visual inspection of graphical data (Tudor-
12 Locke et al., 2015)], were downloaded and cleaned to check for spurious values and periods
13 of non-wear. Non-wear time was determined by identifying strings of uninterrupted zero
14 counts recorded by the accelerometer, for periods of > 60 minutes, allowing for 2 minutes of
15 counts < 100 (Troiano et al., 2008). Data were retained for subsequent statistical analyses
16 where participants accumulated ≥ 10 waking hours wear, on ≥ 4 days, including a weekend
17 day (Troiano et al., 2008). On this basis, $N = 36$ participants were excluded from analyses
18 due to invalid accelerometer data. The outcome variable derived was minutes per day spent in
19 LPA. To adjust for variability in accelerometer wear time, LPA min/day was converted to
20 represent a % of daily accelerometer wear spent engaged in LPA (i.e., %LPA per day; [LPA
21 (min/day) \div accelerometer wear time (min/day)] $\times 100$).

22 ***Psychological well-being***

23 *Depressive symptoms*

24 Depression is an independent risk factor for mortality among people living with RA
25 (Ang et al., 2005; Treharne et al., 2005), and is estimated to affect up to 42% of this patient

1 group (Margaretten et al., 2011). Prevalence of depressive symptoms was assessed using the
2 depressive symptom subscale of the Hospital Anxiety and Depression Scale (HADS)
3 (Zigmond & Snaith, 1983). The HADS requires patients to rate the extent to which they agree
4 with 7 statements representing depressive symptoms (e.g., “I feel cheerful”) via a 4-point
5 scoring system (ranging from 0 to 3). The HADS has been validated previously in RA
6 (Treharne, Lyons, Booth, & Kitas, 2007) and internal reliability of the HADS depressive
7 symptom subscale in this study was acceptable ($\alpha = .81$).

8 *Subjective Vitality*

9 Subjective vitality (e.g., feeling alive, full of energy and spirit) provides an indication
10 of the extent to which an individual is experiencing optimal psychological functioning –
11 referred to as *eudaimonic* well-being (Rouse et al., 2015; Ryan & Deci, 2001). Subjective
12 vitality is considered to have an internal locus of causality, which is influenced by both
13 physical (e.g., rheumatic pain) and psychological factors. It is an individual’s own perceived
14 meaning behind these factors that determine the degree of energy, vitality and spirit felt. For
15 people with RA, an individual’s subjective vitality will therefore provide important
16 information regarding their overall psychological functioning, within the context of their
17 rheumatic disease.

18 Participants’ feelings of personal energy were determined using the Subjective
19 Vitality Scale (SVS) (Ryan & Frederick, 1997). Following the stem... “During the past 3-4
20 weeks, in my everyday life...”, participants are asked to respond to 5 statements (e.g., “I feel
21 alive and full of spirit”) on a 5-point Likert scale, ranging from 1 (strongly disagree) to 5
22 (strongly agree). The SVS demonstrated high internal reliability in this study ($\alpha = .93$) and
23 has recently been validated for use in RA (Rouse et al., 2015).

24

25

1 **Statistical analyses**

2 Kolmogorov-Smirnov tests of normality were conducted and non-normally distributed
3 data were log-transformed for use in subsequent analyses. Where transformations did not
4 reduce data skewness (Kolmogorov-Smirnov, $p < .05$, Table 1), non-parametric statistical
5 tests were used in analyses as appropriate.

6 ***Preliminary analysis***

7 All preliminary analyses were conducted on participants providing valid
8 accelerometer data ($N = 61$), using SPSS (version 22). Independent samples t-tests, Mann-
9 Whitney U Tests and chi-squared tests confirmed that participants excluded on the basis of
10 missing accelerometer data ($N = 36$) did not differ from those included in terms of age,
11 gender, self-reported functional disabilityRW, perceptions of autonomy support, subjective
12 vitality and depressive mood (all p 's $> .05$).

13 Descriptive statistics were calculated and independent samples t-tests and correlation
14 analyses conducted to examine whether participant sex and age, were associated with light
15 physical activity and wellbeing variables. Where significant associations were observed,
16 variables were adjusted for in path models.

17 ***Correlation analysis***

18 Bivariate correlations between autonomy support for physical activity, functional
19 disabilityRW, light physical activity and positive/negative well-being outcomes were
20 computed. In order to adjust for inter-participant variability in daily accelerometer wear-time,
21 LPA was modelled as %LPA per day in both correlation and subsequent path analysis.

22 ***Path analyses***

23 Path analysis was employed to examine the associations between autonomy support,
24 functional disabilityRW, LPA, depressive symptoms and subjective vitality. In brief, this
25 approach involves stipulating hypothesised associations or 'paths' between variables of

1 interest, in order to specify a causal model (e.g., Figure 1). The relationships specified within
2 the model are then analysed simultaneously, to investigate the extent to which the current
3 multivariate set of non-experimental data ‘fits’ with the hypothesised causal model.
4 Analytically, this approach is an advance over correlation and traditional regression analysis
5 as it enables exploration of how a set of variables relate to each other, including analysis of
6 multiple dependent variables. For example, it allows us to examine if a hypothesised
7 dependent variable (e.g., LPA), is also an independent variable for other dependent variables
8 (e.g., vitality and depression). In addition, path analysis affords the ability to examine both
9 direct *and* indirect effects. This means the possible indirect contribution of an independent
10 variable on a dependent variable (e.g., via LPA) is not discounted where a direct association
11 is not evident.

12 Path analysis with maximum likelihood estimation was employed in conjunction with
13 the bootstrapping procedure to test the hypothesised model, as depicted in Figure 1. Previous
14 research has shown this approach to be superior to alternative tests with respect to Type 1
15 error rates and power (Preacher & Hayes, 2008; Shrout & Bolger, 2002). Thus, it was
16 deemed appropriate given the study sample size. Model fit was evaluated using the chi-square
17 statistic (χ^2), comparative fit index (CFI), root square mean error of approximation (RMSEA,
18 90% CI and PCLOSE], and standardised root mean square residual (SRMR). A non-
19 significant χ^2 ($p = < .05$), a CFI $> .90$, and an SRMR and RMSEA of $< .10$ specify reasonable
20 fit of the model to the data (Hu, 1999). For the RMSEA, a p of close fit [PCLOSE] statistic
21 $> .05$ also indicates a well-fitting model. In the instance where CFI is $> .95$, the model is
22 considered to demonstrate excellent fit to the data. The strength and direction of path
23 coefficients were also considered in assessing the validity of the models. Standardised path
24 coefficients corresponding to ($\beta =$) 0.1, 0.3 and 0.5 were interpreted as small, medium and
25 large effect sizes, respectively. Indirect effects were determined via examination of the

1 bootstrap bias-corrected 95% confidence intervals. Specifically, the indirect effects of
2 autonomy support and functional disabilityRW, on depressive symptoms and subjective
3 vitality (via LPA) were examined.

4 All path analysis was conducted using AMOS (version 22). As required for AMOS
5 path models, only data representing participants who provided complete valid data points for
6 all targeted variables were retained for inclusion in path analyses (N = 50) (Arbuckle, 1999).
7 Participants were excluded on the basis of invalid accelerometer data as previously described
8 (N = 36), and a further N = 11 participants were excluded due to missing questionnaire data
9 (SVS, N = 1, IOCQ, N = 10). Analyses established that participants excluded from path
10 models on the basis of missing data (N = 47) did not differ from those included in terms of
11 age, gender, self-reported functional disabilityRW, perceptions of autonomy support and
12 depressive mood (all p's > .05). Mann-Whitney U Tests indicated levels of subjective vitality
13 were significantly higher among included compared to excluded participants ($U = -2.06$, p
14 $= .041$, effect size (r) = $-.20$).

15 **Results**

16 *Descriptive statistics*

17 Descriptive statistics for the targeted variables are reported in Table 1. Data are
18 presented for the full sample recruited to the xxxxx study, and separately for those who
19 provided valid accelerometer data (N = 61). Participants' providing valid data were largely
20 female (67.2%) and white Caucasian (85.2%). Of these participants, 73.8% reported being
21 married and/or living with a partner (9.8% single, 1.6% not living with partner, 6.6%
22 divorced, 4.9% widowed, missing data = 3.3%), and 49.2% reported being in current
23 employment (34.4% retired, 4.9% unable to work due to arthritis, 3.3% homemaker, 3.3%
24 unemployed, missing data = 4.9%).

1 Results revealed a degree of functional disabilityRW of between 0 (without any
2 difficulty) and 1 (with some difficulty) [*NB*: overall functional disability from eight HAQ
3 dimensions, $M \pm SD = .67 \pm .58$]. On average, participants engaged in 4.5 hours of LPA per
4 day and reported moderate to high levels of autonomy support for physical activity from their
5 identified important other. Average prevalence of depressive mood was below the proposed
6 clinical cut-off of ≥ 8 for probable depression, and subjective vitality was moderate to high
7 for this sample of RA patients. Independent samples-tests and correlation analysis revealed
8 participants' sex and age were not associated with LPA or wellbeing outcomes (all p 's $>.05$,
9 i.e., no adjustments were made for these variables in path models).

10 ***Correlation analyses***

11

12 Results of bivariate correlations are displayed in Table 2. Analysis revealed
13 perceptions of autonomy support were significantly positively related to %LPA engagement
14 and subjective vitality, but were not significantly associated with depressive symptoms.
15 Functional disabilityRW was not significantly related to %LPA engagement, but was
16 significantly negatively related to subjective vitality, and significantly positively linked to
17 depressive symptoms. Finally, a significant positive association was observed between
18 %LPA and subjective vitality, and a significant negative relationship revealed between LPA
19 and depressive symptoms.

20 ***Path analysis***

21 *Hypothesised model*: The hypothesised model demonstrated a poor fit to the data (χ^2
22 (5) = 22.29 $p = .000$, CFI = .73, SRMR = .19, RMSEA = .27 (90% CI .00 to .26, PCLOSE =
23 .16). Modification indices provided by AMOS (Arbuckle, 1999) were consulted in order to
24 determine if there were problems with the hypothesised model that could be remedied in the
25 context of the current data. Specifically, modification indices were used to identify
26 associations between variables within the data set that were not currently specified within the
27 hypothesised model. Aligned with recommendations regarding model re-specification,

1 modifications to the hypothesised model were made *only* where relationships identified were
2 conceptually justifiable based on previous research and theoretical assumptions (i.e., SDT)
3 (MacCallum, 1995). Evaluation of modification indices demonstrated that re-specification of
4 the model to stipulate direct paths from; 1) functional disabilityRW to depressive symptoms,
5 2) functional disabilityRW to subjective vitality, and 3) autonomy support to subjective
6 vitality, would improve the fit between the model and the data. This is in agreement with
7 results revealed in bivariate correlation analyses and consequently, the hypothesised model
8 was revised and re-tested in accordance with these specifications (Figure 2).

9 *Re-specified model:* The revised model demonstrated an excellent fit to the data
10 (Figure 2, $\chi^2(2) = 2.44$ $p = .304$, CFI = .99, SRMR = .05, RMSEA = .07 (90% CI .00 to .30,
11 PCLOSE = .34). Results revealed autonomy support for physical activity significantly
12 positively predicted %LPA engagement, which in turn, was significantly positively related to
13 subjective vitality, and significantly negatively associated with depressive symptoms.
14 Functional disability RW was not associated with %LPA engagement. All significant
15 associations were of a small to moderate effect size ($\beta = \geq .2$ and $< .5$). Examination of R^2
16 values indicated autonomy support for physical activity accounted for 15% of the variance in
17 %LPA ($R^2 = .15$). This subsequently predicted 4% of the variance in both subjective vitality
18 and depressive symptoms ($R^2 = .04$).

19 *Indirect effects:* Perceptions of autonomy support demonstrated a significant negative
20 indirect effect on depressive symptoms, ($\beta = -.12$, 95% CI: $-.26$ to $-.02$), and a significant
21 positive indirect effect on subjective vitality ($\beta = .10$, 95% CI: $.01$ to $.28$) via LPA. No
22 significant indirect effect of functional disabilityRW on depressive symptoms or subjective
23 vitality via LPA was observed (depressive symptoms, $\beta = -.02$, 95% CI: $-.13$ to $.06$,
24 subjective vitality, $\beta = .02$, 95% CI: $-.05$ to $.13$).

1 *Direct effects:* Model re-specification enabled investigation of direct effects;
2 functional disabilityRW was significantly negatively associated with subjective vitality, and
3 significantly positively associated with depressive symptoms, accounting for 18% and 23%
4 of the variability in these outcomes, respectively (subjective vitality, $R^2 = .18$; depressive
5 symptoms, $R^2 = .23$). Perceptions of autonomy support for physical activity were
6 significantly positively associated with subjective vitality, predicting 16% of the variability in
7 this outcome ($R^2 = .16$).

8 **Discussion**

9 This cross-sectional study is the first to examine the relationships between autonomy
10 support for physical activity, lower-limb functional disability, LPA engagement and
11 indicators of positive and negative psychological well-being in RA. Results revealed that
12 ‘important other’ autonomy support is beneficially linked to LPA engagement, and in turn,
13 lower prevalence of depressive symptoms and higher subjective vitality in RA. These
14 relationships were observed to be independent of the adverse role of self-reported functional
15 disability to ‘rise’ and to ‘walk’ on psychological well-being states in these patients.

16 Past work has revealed autonomy support for physical activity to be positively
17 associated with self-reported physical activity engagement among patient groups and the
18 general population (Duda et al., 2014; Fortier et al., 2012; Milne et al., 2008). Previous
19 research among older adults, has also demonstrated an association between objectively
20 assessed LPA with indices of psychological well-being. (Buman et al., 2010; Rennemark,
21 Lindwall, Halling, & Berglund, 2009). This study extends these findings in three ways. First,
22 by providing new evidence of an association between autonomy support and objectively
23 assessed LPA in RA. Second, by highlighting the potential role of LPA for fostering more
24 optimal psychological well-being in this patient group. Finally, the analytical approach
25 adopted permitted exploration of a hypothesised causal model, by which autonomy support

1 may influence mental health states among people living with RA, via LPA engagement. That
2 is, results suggest autonomy support from an ‘important other’ may encourage daily LPA
3 participation to the extent it may impact positively on psychological health among people
4 living with RA.

5 Our findings also revealed functional disability to ‘rise’ and ‘walk’, was not
6 significantly associated with LPA engagement among this group of RA patients. This
7 supports the contention that LPA (relative to moderate-intensity physical activity) may be
8 more achievable for people with RA, despite the physical dysfunction symptomatic of this
9 condition. However, whilst not related to LPA, lower-limb functional disability was observed
10 to demonstrate direct adverse relationships with both subjective vitality (negatively) and
11 depressive symptoms (positively). Results therefore substantiate findings from existing
12 research, which demonstrate the deleterious consequences of functional disability for mental
13 health in people living with RA (Benka et al., 2014; Wan et al., 2016) (van der Heide et al.,
14 1994). Still, this study demonstrated autonomy support to be related to both subjective
15 vitality (directly and via LPA) and depressive symptoms (via LPA), independently of the
16 potential negative effects of lower-limb physical dysfunction on psychological functioning.

17 Establishing the independence of these associations not only improves our
18 understanding of these relationships, but also serves to advance the management of RA
19 outcomes, providing a framework for the development of effective interventions that aim to
20 facilitate LPA and optimise psychological functioning in Rheumatic disease. Accordingly,
21 when considering potential targets for interventions, strategies which ensure ‘important
22 others’ are equipped with the skills to; support an individual’s choices with regards to
23 physical activity engagement, provide a meaningful rationale (e.g., improved mental health)
24 to encourage physical activity participation, and demonstrate understanding of an individual’s
25 feelings/perspectives towards physical activity (Williams et al., 2006), may exhibit enhanced

1 efficacy for encouraging LPA, and in turn, and improving psychological well-being in this
2 patient group (Fortier et al., 2012; Ng et al., 2012).

3 Nevertheless, it is still important to consider the implications of current findings
4 within the broader context of the xxxxx study. Participants recruited to this RCT were ready
5 to engage in physical activity behavioural change – i.e., they were consenting to be
6 prescribed (and undertake) an exercise programme to improve their cardiovascular health.
7 Study participants therefore likely represent a cohort of RA patients at the ‘preparation’ stage
8 of change in regards to their physical activity (Daley & Duda, 2006; Prochaska &
9 DiClemente, 1983). It is possible that for individuals with RA who are not ready and
10 preparing to initiate behavioural change (e.g., at the preceding pre-contemplation/
11 contemplation stages of change), autonomy support for physical activity may represent a less
12 prominent determinant of LPA behaviour. Exploration of the extent to which an individuals
13 ‘readiness to change’ may interact with social environmental factors and psychological well-
14 being states in regards to their physical activity, represents an interesting avenue for future
15 research.

16 Similarly, xxxxx study participants reported low-to-moderate functional disability,
17 limiting the generalisability of our findings to RA patients with more severe physical function.
18 Moreover, we did not undertake clinical assessment of disease activity (i.e., Disease
19 Assessment Score-28, DAS-28) to characterise the study sample. Studies employing the
20 DAS-28 are required to confirm the extent to which autonomy support may contribute to
21 more optimal mental health (via promoting LPA) among RA patients with more ‘active’ vs.
22 ‘controlled’ disease.

23 Finally, the cross-sectional design of this study and small sample size should also be
24 considered when interpreting current results. Specifically, compliance with the accelerometer
25 protocol (63%) restricted the number of participants available for analyses, and the cross-

1 sectional design limits the extent to which inferences can be made regarding causal direction
2 of the associations examined. For example, it is possible that a patients' mood state (e.g.,
3 depressive symptoms) could influence their perceptions of autonomy support. However,
4 results from experimental studies framed by SDT strongly support the directionality of the
5 associations as investigated herein (Duda et al., 2014; Fortier et al., 2012; Teixeira, Carraca,
6 Markland, Silva, & Ryan, 2012). In addition, the sample size is comparable with past
7 research employing accelerometers coupled with questionnaires to investigate links between
8 physical activity and self-reported health in RA (Khoja, Almeida, Chester Wasko, Terhorst,
9 & Piva, 2016).

10 **Conclusion**

11 Findings suggest that autonomy support for physical activity provided by an
12 'important other', is positively related to levels of LPA engagement among people living with
13 RA. In turn, higher engagement in LPA is beneficially linked to lower prevalence of
14 depressive symptoms and higher vitality in this patient group. These beneficial associations
15 are observed independently of the adverse consequences of lower-limb functional disability
16 for psychological well-being in RA. Results underline the importance of determining avenues
17 through which 'important others' can be encouraged to provide autonomy support for
18 physical activity among people with RA, in order to enhance mental health in this patient
19 group.

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