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## Moderate-to-high intensity exercise with person-centered guidance influences fatigue in older adults with rheumatoid arthritis

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### Abstract

Fatigue is described as a dominant and disturbing symptom of rheumatoid arthritis (RA) regardless of the advances in pharmacological treatment. Fatigue is also found to correlate with depression. The objective was to evaluate the impact of moderate-to-high intensity, aerobic and resistance exercise with person-centered guidance on fatigue, anxiety and depression, in older adults with RA. Comparisons were made between older adults (> 65 years) with RA taking part in a 20-week moderate-to-high intensity exercise at a gym ( $n=36$ ) or in home-based exercise of light intensity ( $n=38$ ). Assessments were performed at baseline, at 20 weeks, and at 52 weeks. Outcomes were differences in Multidimensional Fatigue Inventory (MFI-20), Visual Analog Scale Fatigue (VAS fatigue), and Hospital Anxiety and Depression Scale (HADS). Analysis of metabolomics was also performed. The subscales “physical fatigue” and “mental fatigue” in MFI-20 and symptoms of depression using HADS depression scale improved significantly at week 20 in the exercise group compared with the control group. Exercise did not influence global fatigue rated by VAS or subscales “reduced motivation”, “reduced activity” and “general fatigue” in MFI-20. No significant change was found on the anxiety index of HADS. The improvements in physical fatigue were associated with changes in the metabolism of lipids, bile acids, the urea cycle and several sugars. Moderate-to-high intensity exercise with person-centered guidance decreased fatigue and improved symptoms of depression and were accompanied by metabolic changes in older adults with RA.

**Keywords** Arthritis · Rheumatoid · Patient-centered · Exercise · Fatigue · Depression · Anxiety · Metabolomics

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## Introduction

Rheumatoid arthritis (RA) is a chronic inflammatory disease characterized by joint pain, stiffness and inflammation of joint synovial tissue. In recent years, pharmacological treatments of RA have undergone revolutionary developments that have led to better control of inflammation and reduced joint destruction. The majority of the patients with RA, however, still suffer from fatigue [1], which they find to be a dominant and disturbing symptom [2–4]. There is currently no consensus definition of fatigue. Fatigue is acknowledged as a complex and subjective experience [5] and a commonly used description of fatigue is “an overwhelming sustained sense of exhaustion and decreased capacity for physical and mental work” [6]. Fatigue is multi-factorial; in patients with RA it is partially dependent on inflammation and production of pro-inflammatory proteins such as cytokines [7] and as in other conditions, can be related to inflammation-dependent anemia [8]. Fatigue correlates strongly with pain, depression [9–11] and sleep disturbances. Disability and psychosocial factors can also contribute [12]. The many effects of fatigue on the individual include disability, anxiety, reduced self-efficacy, feelings of helplessness and limitations in social functioning [13]. Unsurprisingly, a third of RA patients also suffer from depression.

Mechanisms explaining the interaction between inflammatory conditions and fatigue have remained largely elusive. Tools such as metabolomics can be used to explore the disease pathogenesis and generate new hypotheses for mechanisms based on the profile of small molecules in a sample. In blood plasma, the metabolites include organic acids, carbohydrates, amino acids and lipids, which together reflect many different metabolic pathways, and crucially, reflect the interaction between patient genetics, their environment, and the intervention. Metabolomics can be used to identify which pathways have been perturbed which is particularly relevant to patients with RA, for whom it has been shown that metabolites resulting from increased oxidative stress were most consistently associated with increased fatigue [14].

There is some evidence that increased physical activity can reduce fatigue and depression in younger patients with RA [15]. However, little is known about fatigue in relation to physical activity in older adults (> 65 years) with RA, an age group where the risk of increased pain and fatigue in the implementation of an exercise program is a concern [16]. To overcome this hurdle we used a person-centered approach [17], with exercise developed together with the participant taking into account their limitations, resources and goals.

Here, we report the result of a secondary analysis of a randomized controlled multi-center trial, with the

objective to study the effect of moderate-to-high intensity aerobic and resistance exercise during 20 weeks, on fatigue and symptoms of anxiety and depression in older adults with RA.

## Methods

This is a sub-study of a randomized controlled trial that evaluated the effect of 20 weeks moderate-to-high intensity exercise on older adults with RA [18]. The study participants were recruited from the Rheumatology clinics at the Sahlgrenska University Hospital, Gothenburg and Skaraborg Hospital, Skövde, Sweden, using the Swedish Rheumatology Quality Register (SRQ).

The participants were randomized, in blocks of 6 subjects, by a person not involved in the examinations or interventions, using a computer-generated sequence concealed in sequentially numbered, sealed, opaque envelopes.

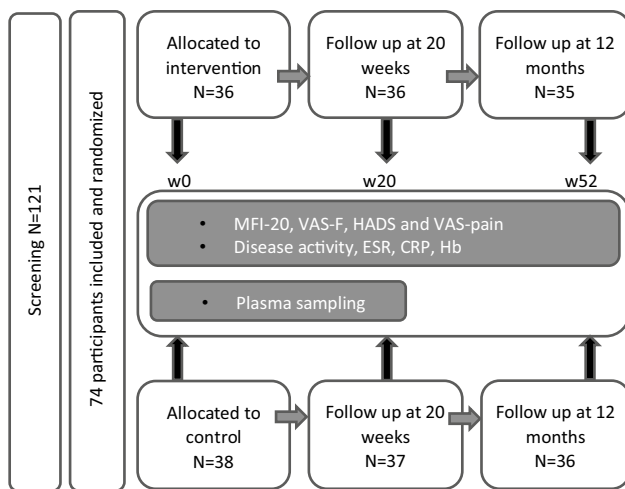
## Study groups

The exercise group ( $n=36$ ) followed a gym-based exercise program with person-centered guidance from a physiotherapist three times a week together with light home-based exercise for 20 weeks. The exercise consisted of moderate-to-high intensity aerobic and resistance exercise with a total of 27 min of aerobic exercise and 5 resistance exercises for large muscle groups. The active control group ( $n=38$ ) performed light home-based exercise for mobility, lower body strength and balance, but no gym-based exercise for 20 weeks. The exercise of both groups is described in detail elsewhere [18].

## Patients

The participants were screened for eligibility by examination of their medical records, a telephone interview and a medical assessment. The medical assessment included physical examination, disease activity and blood samples. The flow-chart of participant enrolment and follow-up, along with the details of tests of physical fitness, is presented elsewhere [18].

Inclusion criteria: RA according to the ACR 1987/EULAR 2010 criteria; age  $\geq 65$  years; disease duration > 2 years; disease activity score  $28 < 5.1$ . Exclusion criteria: co-morbidities such as unstable ischemic heart disease or arrhythmia that might preclude moderate intensity exercise; joint surgery within the 6 months prior to inclusion; ongoing exercise of moderate-to-high intensity  $\geq 2$  times/week; inability to understand or speak Swedish; inability to participate in physical testing that involved walking or bicycling [18].



**Fig. 1** Assessment points of the study with assessed variables and participant flow. *MFI* Multidimensional Fatigue Inventory, *VAS-F* Visual Analog Scale Fatigue, *HADS* Hospital Anxiety Depression Scale, *VAS-Pain* Visual Analog Scale Pain, *ESR* erythrocyte sedimentation rate, *CRP* C-reactive protein, *Hb* hemoglobin

There were 74 older adults with RA included in the study (Fig. 1). The exercise group ( $n = 36$ ) and the active control group ( $n = 38$ ) were considered equal in sociodemographic data [18]. All participants were invited to a post-treatment examination after 20 weeks and 100% ( $n = 36$ ) in the exercise group and 97% ( $n = 37$ ) in the active control group completed the test. At the 12 months follow-up the corresponding figures of attendance for assessment were 97% ( $n = 35$ ) in the exercise group and 95% ( $n = 36$ ) in the control group.

## Main outcome variable

### Assessment of fatigue

Subjective fatigue was quantified using the Multidimensional Fatigue Inventory (Swedish version) (MFI-20) [19]. Twenty statements were rated on a five-point Likert scale and divided into five subscales of fatigue: general fatigue, physical fatigue, mental fatigue, reduced activity, and reduced motivation. Scores range from 4 to 20 for each subscale, with higher scores indicating a higher degree of fatigue [19]. Participants' subjective experience of global fatigue was quantified using a Visual Analog Scale (0–100 mm)-Fatigue (VAS-F). The participants rated global fatigue during the previous week considered due to RA on a scale of 0 (no fatigue) to 100 (the most severe fatigue possible) [20, 21].

### Assessment of anxiety and depression

Anxiety and depression was assessed using the Hospital Anxiety and Depression Scale-Swedish version (HADS). The scale is built up of the subscales anxiety (0–21) and depression (0–21). HADS contains 14 statements scored from 0 to 3, higher scores being indicative of higher degrees of distress [22]. A score of eight or above on any of the subscales has been suggested as a valid cut-off to identify anxiety or depression disorders, respectively [22, 23].

### Measurement of plasma metabolomics

Fasting baseline and 20-week plasma samples ( $n = 48$  subjects) were used to understand potential metabolite associations with the intervention outcomes. Plasma metabolomics was measured using a previously validated gas chromatography–mass spectrometry (GC–MS) method [24]. In brief, 100  $\mu$ L of plasma from fasting blood samples taken at baseline and at 20 weeks was extracted with 900  $\mu$ L methanol:water (9:1 v/v), including a mix of internal standards, and a 200- $\mu$ L aliquot evaporated and derivatized using methoxymation and silylation. The metabolic profile was measured using GC–MS (Shimadzu TQ8030, Shimadzu Europa GmbH, Duisberg, Germany), and metabolites identified based on retention index and mass spectral matching.

### Other variables

Disease activity was assessed using disease activity score 28 (DAS28) [25]. Erythrocyte sedimentation rate (ESR), C-reactive protein (CRP) and hemoglobin (Hb) concentrations were measured by the accredited routine laboratories for clinical chemistry at Sahlgrenska University, Gothenburg, and at Unilabs, Skövde. Participants' objective experience of pain during the previous week considered due to RA was quantified using a Visual Analog Scale (0–100 mm) [21, 26]. Demographic data were collected via a standardized interview. For the analysis of metabolomics, correlations with the variable of maximal oxygen consumption ( $VO_2$ max) presented elsewhere are made [18].

### Procedures

The participants were assessed at baseline, after 20 weeks and at follow-up after 12 months. All assessors were blinded to the treatment groups.

The study complied with the Declaration of Helsinki and was approved by the Regional Ethical Review Board in Gothenburg (2014-11-24/790-14). Informed, written consent was obtained from the participants before the baseline examinations. The study was registered in ClinicalTrials.gov in March 19, 2015 (NCT02397798).

## Statistical analysis

Comparison between groups was performed with the Mann–Whitney *U* test for non-parametric variables and Student's *t* test for parametric variables using GraphPad Prism (GraphPad, San Diego, USA).  $p < 0.05$  was considered as statistically significant. For comparisons within groups the Wilcoxon signed rank test was used. Baseline correlations between DAS28 and outcome variables were performed with Spearman's correlation and a subgroup analysis was performed with DAS28 dichotomized as below 3.2 or 3.2 and above. The subgroup analysis was performed in the SPSS Statistics for Windows, Version 24.0 (IBM corp., Armonk, NY, USA).

Metabolomics data were log-transformed and corrected for baseline (post-intervention values minus baseline values) to correct for inter-individual differences. Metabolic feature reduction was done using orthogonal projection to latent structures discriminant analysis (OPLS-DA) and partial least squares (PLS) in Simca (Umetrics AB, Umeå, Sweden). Differences in metabolic response between groups were identified using the General Linear Model, including change in  $VO_2\max$  (as a proxy for compliance) and analytical batch as covariates.

Analysis were performed on the association between metabolic profile changes (week 20 minus baseline values), with change in physical fatigue and change in maximum oxygen consumption ( $VO_2\max$ ) reported previously [18]. Associations were determined using Pearson's or Spearman's correlation coefficient depending on whether or not the data were normally distributed, and relationships reported if  $p < 0.05$ .

## Results

Demographics and clinical characteristics are presented in Lange et al. 2019 [18]. In brief, in the intervention group 75% were women with a mean age of 69.14 (SD 2.6) and in the control group, 76.3% were women, aged 70.11 (SD 3.10) at baseline. At baseline the variables MFI-20 general fatigue ( $r = 0.419$ ,  $p < 0.001$ ), MFI-20 physical fatigue ( $r = 0.421$ ,  $p < 0.001$ ) and the VAS-F variable ( $r = 0.454$ ,  $p < 0.001$ ) correlated with DAS28.

Two participants in the exercise group and four in the control group used sedatives and the use of sedatives did not change during the study. At baseline two participants in the exercise group and one in the control group used antidepressant medication. After 20 weeks one of the participants in the exercises group had dimidiated the dosage and another two participants in the controls group had started on antidepressants (data not shown).

## 20-week exercise reduced fatigue in older adults with RA

The subscales of MFI-20 physical fatigue ( $p = 0.002$ ) and mental fatigue ( $p = 0.048$ ) was significantly reduced at 20 weeks in the exercise group when compared to the control group (Table 1). The remaining subscales of MFI-20, general fatigue, reduced activity and motivation, remained unaffected by exercise. Global fatigue reported using VAS-F did not significantly differ between exercise and control groups. The subgroup analysis based on dichotomization of DAS28 found no significant differences between the exercise and control group. Within the exercise group there was a significant difference in MFI-20 subscale physical fatigue both in the group with a low ( $n = 26$ , median  $\Delta = -2$ , ICR:  $-3.25$ ;  $-1$ ,  $p = 0.001$ ) and a moderate disease activity ( $n = 27$ , median  $\Delta = 0$ , ICR:  $-1$ ;  $3$ ,  $p = 0.041$ ).

## 20-week exercise alleviated symptoms of depression in older adults with RA

At 20 weeks, there was a significant ( $p = 0.039$ ) reduction in symptoms of depression in the exercise group compared to the control group (Table 1). At baseline one participant in the exercise group and two participants in the control group scored HADS depression above the cut-off of 8 points. After 20 weeks these three individuals still scored above the cut-off and further one of the control group participants (data not shown). The subgroup analysis showed that within the exercise group with a low disease activity, symptoms of depressions were significantly improved ( $n = 27$ , median  $\Delta = -1$ , ICR:  $-1$ ;  $0$ ,  $p = 0.045$ ) when compared to baseline. At baseline six participants of the exercise group and seven participants of the control group scored above the cut-off of 8 points on HADS anxiety. After 20 weeks three of these participants in the exercise group and one in the control group had lowered their score below the cut-off (data not shown). However, there was no significant difference between groups in anxiety (Table 1).

## 20-week exercise did not influence pain, disease activity or inflammatory markers in older adults with RA

The participants were considered to be equivalent in disease activity (DAS28) at baseline; 2.33 (SD 1.10) in the exercise group and 2.41 (SD 0.91) in the control group. After 20 weeks the disease activity was not significantly changed within or between groups with DAS28 in the exercise group at 2.27 (SD 1.05) and in the control group 2.27 (SD 0.92). Hb was also equivalent at baseline 131.4 g/L (SD 11.6) in the exercise group and 131.1 g/L (SD 13.6) in the control group and did not significantly change after the intervention. The participants

**Table 1** Differences between groups in assessment measures at weeks 20 and 12 months compared to baseline

	Intervention group (N=36)			Control group (N=38)			Between-group differences	
	Baseline	Post-treatment—baseline	Follow-up—baseline	Baseline	Post-treatment—baseline	Follow-up—baseline	Post-treatment—baseline	Follow-up—baseline
		Δ 20-0 w	Δ 52-0 w		Δ 20-0 w	Δ 52-0 w	p value	p value
<b>MFI-20</b>								
General fatigue	8 (6.3 to 13.8)	−1 (−3 to 0)	0 (−3 to 1)	12 (9.3 to 13.8)	0 (−3 to 2)	0 (−2 to 2)	0.345	0.952
Physical fatigue	9 (6 to 13)	−2*** (−3 to −0.8)	−2* (−3.3 to 1)	10 (7.5 to 13.5)	0 (−1.8 to 2)	0 (−2 to 1)	0.002**	0.084
Mental fatigue	8 (5 to 11)	0 (−1.3 to 4)	0 (−3 to 2)	9 (4.5 to 11)	0 (0 to 1)	0 (−1 to 2)	0.048*	0.466
Reduced motivation	6 (4 to 9)	0 (−1 to 0.3)	0 (−1 to 1)	7 (4 to 9.5)	0 (−1 to 2)	1 (−1 to 2)*	0.106	0.156
Reduced activity	9 (5.3 to 13)	−1* (−2 to 1)	−1 (−3 to 0)	11 (7 to 13)	0 (−1.3 to 2)	0 (−2 to 2)	0.074	0.095
VAS-F, 0–100 mm	26 (7.3 to 50.5)	−2 (−11.5 to 4.8)	−1 (−16 to 8)	28.5 (9.8 to 42)	0 (−11 to 11.8)	−3 (−10 to 17)	0.355	0.707
<b>HADS (0–21)</b>								
Anxiety	2.5 (0.3 to 6.8)	−1 (−2 to 0)	0 (−1 to 1)	2 (1 to 6)	0 (−1 to 1)	0 (−1 to 1)	0.064	0.370
Depression	2 (1 to 4)	0 (−1 to 0)*	0 (−1 to 0)	2 (1 to 4)	0 (−0.5 to 2)	0 (−1 to 1)	0.039*	0.328

Variables are presented as medians (IQR)

*MFI* Multidimensional Fatigue Inventory, *VAS-F* Visual Analog Scale Fatigue, *HADS* Hospital Anxiety Depression Scale

Delta values were compared using the Mann–Whitney *U* test. Within-group comparisons were made with Wilcoxon

\*Significant difference,  $p < 0.05$

\*\*Significant difference,  $p < 0.01$

\*\*\*Significant difference,  $p < 0.001$

were considered to be equivalent in levels of inflammatory markers (CRP and ESR) and rated VAS-Pain at baseline [18] and there were no significant between- or within-group differences in these parameters at 20 weeks compared to baseline (data not shown).

### Changes to physical fatigue in MFI-20 and $VO_2$ max were associated with several metabolic changes

Based on targeted analysis of GC–MS data, 146 unique metabolites were identified in plasma samples from subjects in this study. Seventeen of these metabolites were significantly associated with changes in the MFI-20 physical fatigue scale, and aerobic capacity ( $VO_2$ max) (Table 2). These metabolites represent diverse compound classes and metabolic pathways monosaccharides, bile acids and metabolites related to lipid metabolism and the urea cycle.

### 20-week exercise reduced circulating concentrations of some lipids in older adults with RA

Twenty weeks of exercise led to decreases in plasma levels of 1-myristoyl-glycerol ( $p = 0.001$ ), 24-methylcholesterol ( $p < 0.001$ ), cholesterol ( $p = 0.046$ ), campesterol ( $p < 0.001$ ) and  $\alpha$ -tocopherol ( $p = 0.003$ ). Circulating levels of agmatine ( $p = 0.027$ ) and phenylpyruvic acid ( $p = 0.020$ ) were also increased in the intervention group (Table 3).

### 52-week follow-up

Withdrawal of the supervision of the exercise intervention program at 20 weeks resulted in the loss of the significant improvements in fatigue and depression recorded at that point in the study (Table 1). These findings coincide with diminished intensity and duration of exercise at 52 weeks [18]. Additionally, none of the outcomes from the assessments for HADS anxiety, VAS-P, or the remaining MFI-20



**Table 2** Correlations between change in physical fatigue or aerobic capacity, with change in metabolite concentrations between baseline and 20 weeks

Metabolite	Change in aerobic capacity	<i>p</i> value	Change in physical fatigue	<i>p</i> value
Arabinose	−0.333	0.034*		
Lyxose	−0.348	0.026*		
Pyruvic acid	−0.327	0.037*		
Agmatine	−0.317	0.043*	0.302	0.042*
Arginine	−0.410	0.008**		
Cysteine	−0.324	0.039*		
Indole-3-acetic acid	−0.307	0.051*		
<i>N</i> -Acetylglutamic acid	−0.320	0.041*		
Ribose	−0.353	0.024*		
Gluconic acid 1,5-lactone	0.331	0.025*		
24-Methylcholesterol			0.382	0.009*
Alpha-tocopherol			0.361	0.014*
Campesterol			0.403	0.006**
Deoxycholic acid			−0.304	0.040*
Glyceric acid			0.304	0.040*
Quinic acid			−0.314	0.034*
Sorbitol-6-phosphate			0.314	0.034*

Only statistically significant correlations are shown

Data on aerobic capacity from Lange et al. [18]

\*Significant difference,  $p < 0.05$

\*\*Significant difference,  $p < 0.01$

**Table 3** Metabolites significantly affected by exercise intervention at 20 weeks compared to baseline

All participants	Intervention group ( $N=24$ )		Control group ( $N=24$ )		<i>p</i> value
	$\Delta$ 20-0 w		$\Delta$ 20-0 w		
	Mean	SE	Mean	SE	
1-Myristoyl-glycerol	−3782.99	1528.857	6807.943	1528.857	0.001**
24-Methyl-cholesterol	−1790.35	725.679	4112.523	725.679	<0.001***
Agmatine	−83,912.7	25,804.98	33,310.1	25,804.98	0.027*
Alpha-tocopherol	−12,470.4	5794.299	23,300.33	5794.299	0.003**
Campesterol	−1385.58	833.4642	4547.694	833.4642	<0.001***
Cholesterol	−42,907.5	23,636.28	53,526.94	23,636.28	0.046*
Phenylpyruvic acid	−19,582.9	7740.779	17,587.55	7740.779	0.020*

SE standard error

\*Significant difference,  $p < 0.05$

\*\*Significant difference,  $p < 0.01$

\*\*\*Significant difference,  $p < 0.001$

subscale categories (Table 1), was different at 52 weeks from the baseline readings. In the subgroup analysis the MFI-20 subscale physical fatigue (median  $\Delta = -2$ , ICR:  $-3.5; 0.5$ ,  $p = 0.019$ ) and reduced activity (median  $\Delta = -1.5$ , ICR:  $-4; 0$ ,  $p = 0.013$ ) were significantly improved within the exercise group with a low disease activity ( $n = 25$ ) but not in the other groups.

## Discussion

To our knowledge, this is the first study to show that moderate-to-high intensity exercise reduces physical and mental dimensions of fatigue and has positive effects on symptoms of depression in older adults with RA, at the same time recording significant associations between these changes and differences in plasma concentrations of

metabolites from a wide range of compound classes and pathways, reflecting the wide-ranging effects of effective exercise on metabolic status. The exercise intervention led to significant improvement in aerobic capacity, endurance and physical strength of participants [18], which have separately been found to reduce fatigue in people with RA or fibromyalgia less than 65 years of age [27, 28]. The observed effect of exercise on fatigue is to be contrasted with treatments that reduce disease activity profoundly, such as methotrexate and biologic/targeted synthetic disease modifying anti-rheumatic drugs (DMARDs), alone or in combination, as well as preventing joint destruction, but that only partly reduce [29] and seldom eliminate fatigue [1]. The participants in the present study were in remission or had low disease activity, but still scored clinically relevant fatigue (VAS > 20) [12]. Their exercise affected neither disease activity nor inflammatory markers [18, 30], yet contributed to a clinically relevant [31] reduction of physical fatigue, suggesting that it is relevant to evaluate non-pharmacological treatment options against fatigue in older adults with RA.

The subgroup analysis indicates that the participants with a lower disease activity at baseline got the best effects of moderate-to-high intensity exercise on fatigue and depression. Disease activity has been found to correlate with fatigue [32], but has not been documented as a stronger predictor for fatigue in RA [12, 33]. However, patients have described struggling with the unpredictability of the disease activity to be physically active [34] and reduced physical health is a barrier for exercise among older persons with RA [35].

Fatigue ratings by participants in this study were low relative to those reported by others [3, 36], one explanation for which could be selection bias; persons volunteering to participate in a study involving exercise 3 times a week are likely to be at the lower end of the fatigue spectrum for people with RA. Nonetheless there was a significant decrease in physical and mental fatigue directly after the intervention period (at 20 weeks) in the intervention group compared to the control group. Physical and mental fatigue are defined here by their respective MFI-20 subscales; “physical fatigue” records the person’s feeling of physical wellness combined with their ability to perform different tasks, and is supposed to be related to significantly improved physical fitness in the participants in the intervention group [18]; “mental fatigue” on the other hand focuses on concentration and coping with thoughts. It is likely that the lessening of mental fatigue seen here is part of the alleviation of symptoms of depression rated with the HADS scale in the same participants. This provides at least a partial explanation of the positive effects of physical activity on depression found elsewhere [37–39], and reconfirms the link between fatigue and depression that has already been suggested [9]. A small minority of

participants (3% of the intervention group and 5% in the control group at baseline) had symptoms of depression defined as HADS depression of more than 8 points. Nevertheless, the significant reduction of the HADS depression score in the intervention group at week 20 indicate that the mood of the participants was improved.

When the degree of exercise was reduced after 20 weeks the improvement in fatigue and symptoms of depression disappeared, along with enhanced physical fitness; only endurance was sustained to week 52 [18]. This outcome identifies a link between the supported maintenance of moderate-to-high intensity exercise and its success in improving fatigue and depression, important determinants of quality of life of persons with RA [40], and should be taken into account in the design of therapies for the future.

We found no significant effects of exercise on either of the VAS-F or MFI-subscale “general fatigue” parameters, probably due to construction of the scales, as they rate global fatigue that may include several other feelings, such as pain and sleep. Neither were there significant differences between the two groups after the intervention in the subscales “reduced activity” and “reduced motivation”, maybe reflecting a high degree of motivation in the selected cohort; in support of this possibility, the compliance with the study protocol was high and 98.6% of the participants completed the 20 weeks examination [18]. Both the selection of participants and the person-centered approach are likely to have contributed to this success [18]. Additionally, study participants were retired, and having fewer mandatory duties (working hours) and less demanding expectations from others might help older adults to cope with fatigue more easily [41].

The metabolomic profiling included in this study revealed changes attributable to the exercise intervention in eight of the identified metabolites, with only one of these metabolites also being significantly associated with the decrease in physical fatigue shown by the MFI-20 analysis. The metabolic changes found in the current study were largely related to lipid and specifically sterol metabolism, where circulating concentrations of some lipids were lower in the intervention group than in control group. Some of these are known to be diet-related (24-methylcholesterol, campesterol, and  $\alpha$ -tocopherol); the exercise intervention may have led to modifications in participants’ food intake, or metabolism of these compounds may have increased. Exercise has elsewhere been found to raise concentrations of campesterol and sitosterol concentrations, in association with improved  $VO_2$ max [42].  $\alpha$ -Tocopherol has been found to increase in high-intensity exercise, though rapidly returns to normal concentrations on resting [43, 44]. As diet was not measured in this study, we can only speculate that physiological effects of exercise rather than dietary changed altered these lipids. Other studies using metabolomics to probe the effects



of exercise interventions report effects on lipid metabolism, though often in addition to purine and central energy metabolism [45]. The association of pyruvic acid and ribose with  $VO_2$ max suggests that both purine and central energy metabolism pathways were impacted by the intervention, in addition to the urea cycle, as demonstrated by the relationship between both arginine and agmatine with change in  $VO_2$ max. These data suggest that the exercise intervention led to general metabolic changes that are consistent with the previous literature on exercise interventions, suggesting that RA does not alter metabolic response to exercise.

Especially for the metabolomics analysis, the low overall number of subjects and free-living nature of the current study are likely to be the cause of a considerable degree of inter-individual variation and potential confounding from lifestyle factors that have not been measured here. We also noted that there were significant gender effects which should be accounted for (data not shown). To control for this, future studies would benefit from including more subjects and using lipidomics analysis (methodology that focuses specifically on the lipid fraction of blood), since lipid metabolism appeared to be widely affected by exercise here, and many lipids play a role in inflammation pathways. This exploratory component of this study suggests that there are several metabolic changes related to improvement in physical fatigue in RA, and the use of such biomarkers will become increasingly useful in the tailoring of therapies including supervised exercise to individuals who are likely to see the greatest benefit.

Fatigue and depression are subjective symptoms with multi-factorial causes; for instance it has been observed that persons with RA suffer more with fatigue during the winter [46]. In documenting the lessening of fatigue and depression resulting from exercise, independent of disease status, this report emphasizes the importance of multi-modal approaches to both pharmacological and non-pharmacological aspects of treatment; understanding mechanisms may help to guide multi-pronged approaches to therapy for patients with RA.

## Limitations

The studied population was highly selected [18] and therefore not representative for all older adults with RA. Most of the participants' had low disease activity and were positive about participating in a study regarding physical activity. Alongside this it would have been of interest to compare moderate-to-high intensity exercise not only with low-intensity exercise but with a control group performing no exercise. The study lacks logs of the participants' sleep and self-reporting sleeping questionnaires concerning the participants' sleeping habits that could have contributed to the understanding of changes in fatigue.

With the present study being a secondary analysis, power was calculated based on primary outcome of the primary analysis previously published [18]. The number of participants was considered to be sufficient for the aim of this analysis and the significant results indicate that they were.

## Conclusion

Person-centered moderate-to-high intensity exercise decreased fatigue and lessened symptoms of depression in older adults with RA, improvements that were accompanied by metabolic changes. Our study strongly suggests that moderate-to-high intensity exercise should be implemented in standard care in older adults with RA. Further, to maintain the beneficial effects, the exercise needs also to be maintained.

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## Compliance with ethical standards

**Ethical standards** The study complied with the Declaration of Helsinki and was approved by the Regional Ethical Review Board in Gothenburg (2014-11-24/790-14).

**Informed consent** Informed, written consent was obtained from the participants before the baseline examinations.

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