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Regular exercise during haemodialysis promotes an anti-inflammatory leucocyte profile

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Abstract

Background: Cardiovascular disease is the most common cause of mortality in haemodialysis (HD) patients and is highly predicted by markers of chronic inflammation. Regular exercise may have beneficial anti-inflammatory effects, but this is unclear in HD patients. This study assessed the effect of regular intradialytic exercise on soluble inflammatory factors and inflammatory leucocyte phenotypes.

Methods: Twenty-two HD patients from a centre where intradialytic cycling was offered thrice weekly and 16 HD patients receiving usual care volunteered. Exercising patients aimed to cycle for 30 min at rating of perceived exertion of ‘somewhat hard’. Baseline characteristics were compared with 16 healthy age-matched individuals. Physical function, soluble inflammatory markers and leucocyte phenotypes were assessed again after 6 months of regular exercise.

Results: Patients were less active than their healthy counterparts and had significant elevations in measures of inflammation [interleukin-6 (IL-6), C-reactive protein (CRP), tumour necrosis factor-α (TNF-α)], intermediate and non-classical monocytes; all P < 0.001. Six months of regular intradialytic exercise improved physical function (sit-to-stand 60). After 6 months, the proportion of intermediate monocytes in the exercising patients reduced compared with non-exercisers (7.58 ± 1.68% to 6.38 ± 1.81% versus 6.86 ± 1.45% to 7.88 ± 1.66%; P < 0.01). Numbers (but not proportion) of regulatory T cells decreased in the non-exercising patients only (P < 0.05). Training had no significant effect on circulating IL-6, CRP or TNF-α concentrations.

Conclusions: These findings suggest that regular intradialytic exercise is associated with an anti-inflammatory effect at a circulating cellular level but not in circulating cytokines. This may be protective against the increased risk of cardiovascular disease and mortality that is associated with chronic inflammation and elevated numbers of intermediate monocytes.

Key words: cytokines, exercise, haemodialysis, inflammation, monocytes

Introduction

Haemodialysis (HD) patients have a vastly elevated risk of all-cause, and specifically cardiovascular, mortality [1], which is the most common cause of death in these patients. This increased risk of cardiovascular disease (CVD) morbidity and mortality is multifactorial but includes factors relating directly
to kidney disease (e.g. the malnutrition–inflammation complex syndrome) and to the process of HD itself [2]. HD patients have chronically elevated circulating soluble markers of inflammation [including interleukin-6 (IL-6), C-reactive protein (CRP) and tumour necrosis factor-α (TNF-α)] that strongly associate with mortality and cardiovascular events [3]. Altered inflammatory immune cell populations and phenotypes are also evident in HD patients, including a shift towards the ‘pro-inflammatory’ CD16⁺ monocyte subpopulations [4, 5], which are also strongly associated with increased cardiovascular mortality in chronic kidney disease (CKD) [6].

Regular exercise is associated with reduced mortality and morbidity in various chronic diseases and is associated with a plethora of cardioprotective benefits [7]. The so-called ‘anti-inflammatory’ effect of regular exercise may play a key role in this through a number of different mechanisms. These include altering the balance of inflammatory immune cell phenotypes, reducing visceral adipose tissue mass and inflammatory cytokine release and, at higher intensities, through release of a potent anti-inflammatory cascade stimulated by muscle-derived cytokines [8, 9]. The importance of promoting an active lifestyle in the management of kidney disease is recognized in national and federal guidelines and in the scientific literature, as evidenced by a recent Cochrane Review [10], but the incorporation of exercise into clinical service has been slow and lags national and federal guidelines and in the scientific literature, as evidenced by a recent Cochrane Review [10], but the incorporation of exercise into clinical service has been slow and lags

Exercise programme

The exercise group participated in a progressive intradialytic exercise programme for 6 months; the non-exercising control group continued HD treatment as per their usual care. Exercise was provided in the form of a specially designed recumbent cycle (Lotto series; Motomed, Reck, Germany). Patients aimed to cycle at a ‘somewhat hard’ rating of perceived exertion (RPE 12–14) [16] with supervised intradialytic exercise offered thrice weekly. The exercise programme was designed with a pragmatic approach; more details are provided in the Supplementary Material.

Randomization

An HD unit providing exercise and a unit not providing exercise were selected based on practicality and, therefore, the study was not randomized. All non-exercising patients were eligible to exercise if such a programme were available at their unit. This design reduced the sampling bias of using patients unable to, or not interested in, exercise as controls and ‘exercise-contamination’ of control patients that occurs when control and exercise patients share shifts.

Outcome measures

Blood samples were collected prior to HD at baseline and after 6 months (at least 48 h after previous exercise session); estimated glomerular filtration rate (eGFR) was calculated using the simplified Modification of Diet in Renal Disease equation [17]. Medical details were extracted from the patients’ clinical notes. Plasma concentrations of CRP (IBL International GmbH, Hamburg, Germany), IL-6 and TNF-α (R&D systems, Abingdon, UK) were determined using commercially available high-sensitivity enzyme-linked immunosorbent assays. Assessment of monocyte phenotypes (classical: CD14⁺CD16⁻, intermediate: CD14⁺CD16⁺ and non-classical: CD14⁻CD16⁺) and regulatory T cells (Tregs: CD4⁺CD25⁻CD127low/−) via flow cytometry are described in the Supplementary Material.

Activity levels were determined using a tri-axial accelerometer (SenseWear, BodyMedia Inc., Pittsburgh, PA, USA). The participant wore the accelerometer for a single 7-day period collecting minute-by-minute data. Physical function was assessed using the sit-to-stand 60 (STS 60) test, which is validated against more detailed and lengthy assessment methods [18].

Omitted samples

Patients on immunosuppressive therapy were excluded from analysis of inflammatory factors (Figure 1). Four exercise patients were omitted from analyses of monocyte phenotype due to unclear fluorescence patterns. Some patients declined physical function and activity outcomes; numbers for each outcome are presented in the Results.

Sample size

Sample size for this study was pragmatic: all eligible patients in the two dialysis units were invited to participate (Figure 1).

Statistics

Data are presented as mean ± standard deviation or median (25th–75th percentiles) unless otherwise stated. Baseline

Materials and methods

Participants

The study was approved by the East Midlands Research Ethics Committee (Northampton; 11/EM/0149). Additional ethical approval for the healthy cohort was granted by the Loughborough University Ethical Advisory Committee. All participants gave written informed consent.

Patients were recruited from two satellite HD units within the same renal network: one offered an intradialytic exercise programme as part of clinical care and one did not. An independent Consultant Nephrologist verified the medical suitability of all patients to exercise. Exclusion criteria for the study were: <18 years; established contraindications to exercise [15]; lower limb vascular access; recent clinically overt infection; already taking part in exercise programme; or an insufficient command of English to consent.

Following recruitment of the patient group, age-matched healthy participants were recruited from the local area. Healthy individuals completed a health screen and were ineligible if they had any history of kidney disease, diabetes, recent infection, CVD, smoking, or took any drugs known to affect digestion or metabolism. Healthy individuals donated a one-off serum blood sample and completed the same outcome measures as the HD patients (described below) within a laboratory setting.

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Comparisons between groups were completed using independent t-tests, non-parametric Mann–Whitney tests or chi-square tests where applicable. Two-factor mixed-measures analysis of variance (ANOVA) was used to analyse the effect of exercise: group (exercise versus control) × time (baseline versus 6 months). Where a significant group × time interaction was found, post hoc analysis explored the differences using paired t-tests within groups and independent t-tests for between-group differences in change from baseline at 6 months. Comparisons between treatment and non-treatment day activity levels were assessed using Wilcoxon signed-rank test. Effect sizes (ES) were calculated using Cohen’s D [19]. All statistical analyses were completed on Statistical Package for Social Sciences (SPSS v.21, IBM, New York, NY, USA). Statistical significance was accepted at $P < 0.05$.

**Results**

**Recruitment and participant flow**

A total of 38 HD patients participated in the study (Figure 1). Twenty-two patients were recruited into the exercise group, of whom 16 completed the 6-month study period (27.3% attrition rate). Sixteen non-exercising control HD patients volunteered, of whom 15 completed the study. Sixteen healthy participants volunteered and blood samples were obtained from 15.

**Implementation of intervention**

The exercising patients participated in 38 ± 12 exercise sessions. The mean session duration was 35 (30–42) min with patients exercising at 63 ± 8 r.p.m. and perceiving the exercise to be ‘somewhat hard’ [RPE: 12 (11–13)]. Mean power output was 16 ± 7 W.

**Participant characteristics**

The age and sex of the HD patients and healthy participants were similar (Table 1). The age of the exercising and non-exercising HD groups were different (57.0 ± 10.5 versus 70.2 ± 13.7 years; $P = 0.005$), with more South Asian patients in the exercise group (7/16 versus 1/15; $P = 0.02$), which reflected the demographic of the centres where recruitment took place. However, there were no differences in circulating or cellular inflammatory markers between the exercising and non-exercising groups.

**Outcomes**

Inflammatory markers. HD patients had significantly higher concentrations of circulating IL-6 (ES = 1.32), TNF-α (ES = 1.37) and CRP (ES = 1.35) than the healthy participants (Table 2).
Circulating concentrations of IL-6, CRP and TNF-α did not differ between the exercising and non-exercising patients at baseline (Table 2). ANOVA found no interaction or main effects in response to the 6-month exercise programme (Figure 2a–c).

Monocytes. The proportion and number of intermediate (ES = 1.23 and ES = 1.17, respectively) and non-classical monocytes (ES = 1.56 and ES = 1.31) were significantly greater in HD patients compared with healthy participants (Table 2).

There were no significant baseline differences between the exercising and non-exercising HD patients in the proportion of classical, intermediate or non-classical monocytes (Table 2).

### Table 1. Participant characteristics

<table>
<thead>
<tr>
<th></th>
<th>Healthy group (n = 16)</th>
<th>Combined HD patients (n = 31)</th>
<th>P-value</th>
<th>Exercising HD patients (n = 16)</th>
<th>Non-exercising HD patients (n = 15)</th>
<th>P-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (years)</td>
<td>61.5 ± 10.9</td>
<td>63.4 ± 13.7</td>
<td>0.63</td>
<td>57.0 ± 10.5</td>
<td>70.2 ± 13.7</td>
<td>0.005*</td>
</tr>
<tr>
<td>Gender (male/female)</td>
<td>8/8</td>
<td>18/13</td>
<td></td>
<td>8/8</td>
<td>10/5</td>
<td>0.35</td>
</tr>
<tr>
<td>Weight (kg)</td>
<td>75.3 ± 16.6</td>
<td>75.9 ± 23.1</td>
<td>0.92</td>
<td>71.0 ± 24.9</td>
<td>81.2 ± 20.6</td>
<td>0.23</td>
</tr>
<tr>
<td>BMI (kg/m²)</td>
<td>26.1 ± 5.0</td>
<td>27.1 ± 6.4</td>
<td>0.58</td>
<td>25.9 ± 6.7</td>
<td>28.4 ± 6.1</td>
<td>0.29</td>
</tr>
<tr>
<td>Systolic blood pressure (mmHg)</td>
<td>129 ± 13</td>
<td>136 ± 21</td>
<td>0.16</td>
<td>135 ± 24</td>
<td>138 ± 19</td>
<td>0.68</td>
</tr>
<tr>
<td>eGFR (ml/min/1.73 m²)</td>
<td>102 ± 22</td>
<td>b</td>
<td>–</td>
<td>b</td>
<td>b</td>
<td>–</td>
</tr>
<tr>
<td>Number of comorbidities</td>
<td>–</td>
<td>–</td>
<td>–</td>
<td>4 ± 2</td>
<td>4 ± 2</td>
<td>0.57</td>
</tr>
</tbody>
</table>

*Data are presented as mean ± standard deviation.

**All HD patients had established end-stage renal disease and had received HD treatment for at least 3 months.

†Denotes a significant difference between exercising HD patient and non-exercising HD patient groups.

### Table 2. Comparisons in markers of inflammation and haematology at baseline between a healthy group and HD patients, and between the exercising and non-exercising HD groups

<table>
<thead>
<tr>
<th></th>
<th>Healthy group</th>
<th>HD group</th>
<th>P-value</th>
<th>Exercising HD group</th>
<th>Non-exercising HD group</th>
<th>P-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>IL-6 (pg/mL)</td>
<td>0.81 (0.43–1.76)</td>
<td>4.63 (2.73–7.09)</td>
<td>&lt;0.001*</td>
<td>5.39 ± 3.05</td>
<td>4.92 ± 2.79</td>
<td>0.68</td>
</tr>
<tr>
<td>TNF-α (pg/mL)</td>
<td>0.96 (0.60–1.73)</td>
<td>3.21 (2.66–4.62)</td>
<td>&lt;0.001*</td>
<td>3.22 (2.74–4.24)</td>
<td>3.20 (2.64–4.47)</td>
<td>0.77</td>
</tr>
<tr>
<td>CRP (mg/L)</td>
<td>0.83 (0.21–1.90)</td>
<td>4.61 (2.68–9.78)</td>
<td>&lt;0.001*</td>
<td>3.99 (2.34–6.86)</td>
<td>7.89 (3.39–10.4)</td>
<td>0.37</td>
</tr>
<tr>
<td>Classical monocytes (%)</td>
<td>89.5 (87.0–92.1)</td>
<td>80.3 (72.6–82.2)</td>
<td>&lt;0.001*</td>
<td>80.5 (75.4–82.3)</td>
<td>79.8 (72.6–80.9)</td>
<td>0.53</td>
</tr>
<tr>
<td>Classical monocytes (cells/µL)</td>
<td>506 ± 130</td>
<td>516 ± 112</td>
<td>0.78</td>
<td>435 (410–525)</td>
<td>580 (558–599)</td>
<td>0.09</td>
</tr>
<tr>
<td>Intermediate monocytes (%)</td>
<td>4.34 (3.74–4.95)</td>
<td>6.77 (5.98–8.47)</td>
<td>&lt;0.001*</td>
<td>7.58 ± 1.68</td>
<td>6.86 ± 1.45</td>
<td>0.27</td>
</tr>
<tr>
<td>Intermediate monocytes (cells/µL)</td>
<td>21.8 (19.4–29.1)</td>
<td>45.9 (40.5–52.3)</td>
<td>&lt;0.001*</td>
<td>46.3 ± 14.7</td>
<td>49.4 ± 14.2</td>
<td>0.61</td>
</tr>
<tr>
<td>Non-classical monocytes (%)</td>
<td>6.45 (4.2–8.5)</td>
<td>13.6 (11.2–17.7)</td>
<td>&lt;0.001*</td>
<td>12.4 (10.8–16.4)</td>
<td>13.8 (12.7–17.5)</td>
<td>0.36</td>
</tr>
<tr>
<td>Non-classical monocytes (cells/µL)</td>
<td>30.0 (23–50)</td>
<td>90.9 (72.2–124.5)</td>
<td>&lt;0.001*</td>
<td>75.9 (59.9–105.7)</td>
<td>96.9 (89.0–139.2)</td>
<td>0.11</td>
</tr>
<tr>
<td>Tregs (%)</td>
<td>7.67 ± 0.98</td>
<td>6.75 ± 1.53</td>
<td>0.02*</td>
<td>6.93 ± 1.77</td>
<td>6.55 ± 1.24</td>
<td>0.53</td>
</tr>
<tr>
<td>Tregs (cells/µL)</td>
<td>20.1 (18.6–25.0)</td>
<td>23.7 (17.3–36.0)</td>
<td>0.53</td>
<td>23.6 (16.0–38.0)</td>
<td>23.8 (19.1–32.4)</td>
<td>0.57</td>
</tr>
<tr>
<td>CD4⁺ lymphocytes (%)</td>
<td>20.2 (17.4–23.9)</td>
<td>24.1 (20.1–28.9)</td>
<td>0.18</td>
<td>23.9 ± 8.0</td>
<td>24.7 ± 7.2</td>
<td>0.79</td>
</tr>
<tr>
<td>CD4⁺ lymphocytes (cells/µL)</td>
<td>264 (227–369)</td>
<td>366 (269–493)</td>
<td>0.05*</td>
<td>317 (253–496)</td>
<td>369 (301–483)</td>
<td>0.53</td>
</tr>
</tbody>
</table>

*Data are presented as mean ± standard deviation or median (25th–75th percentiles). P-values are derived from independent t-tests or Mann-Whitney test, where applicable. Classical monocytes refer to CD14⁺CD16⁻; intermediate monocytes to CD14⁺CD16⁺; non-classical monocytes to CD14⁺CD16⁻. Tregs (%) and (cells/µL) refer to the proportion and number of CD4⁺ lymphocytes that are CD25⁺CD127⁻low⁻.

†For monocytes n = 24 for HD group; n = 11 for exercise HD group and n = 13 for non-exercising HD group.

‡For percent Tregs and percent CD4⁺ lymphocytes n = 16 for healthy group.

†Denotes a significant difference between exercising and non-exercising groups.

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Tregs. The number of circulating CD4\(^+\) lymphocytes were higher in the HD patients compared with the healthy participants (ES = 0.71, Table 2). However, the proportion of circulating CD4\(^+\) lymphocytes that were Tregs was lower in the HD patients (ES = 0.72, Table 2). The number of Tregs did not differ between groups.

At baseline, there were no significant differences between the exercising and non-exercising HD patients in the proportion or number of Tregs or CD4\(^+\) lymphocytes (Table 2).

ANOVA revealed group × time interactions in the proportion and number of CD4\(^+\) lymphocytes that were Tregs was lower in the HD patients (ES = 0.72, Table 2). The number of Tregs did not differ between groups.

At baseline, there were no significant differences between the exercising and non-exercising HD patients in the proportion or number of Tregs or CD4\(^+\) lymphocytes (Table 2).

ANOVA revealed group × time interactions in the proportion and number of CD4\(^+\) lymphocytes (F = 11.3, P = 0.002 and F = 12.7, P = 0.001, Figure 2h). In both cases, post hoc paired t-tests revealed a significant decrease in the non-exercising patients from baseline to 6 months (P < 0.001, ES = 0.67 and P = 0.001, ES = 0.61, respectively) with no significant change in the exercise group.

ANOVA also revealed a group × time interaction in the number of Tregs (F = 9.89, P = 0.004). Post hoc paired t-tests revealed a trend for an increase in exercising patients (P = 0.10, ES = 0.23) and a decrease in non-exercising patients (P = 0.003, ES = 0.50; Figure 2g). The magnitude of change in the number of Tregs observed over the 6 months in the exercising patients was significantly different from that seen in the non-exercising patients (P = 0.02). There was no change in the proportion of CD4\(^+\) lymphocytes that were Tregs (interaction: P = 0.74, Figure 2i).

Physical function. Compared with healthy participants, HD patients had significantly lower levels of physical function, as measured by the STS 60 test (ES = 1.28, Table 3).

At baseline, there were no significant differences between the exercising and non-exercising HD groups (P = 0.88). Change in physical function was different between groups (F = 28.1, P < 0.001, Figure 3), with increases in the exercise group (P < 0.001, ES = 1.47), but not the non-exercising group (P = 0.21, ES = 0.18).
6-month intradialytic exercise programme promotes a shift away from the pro-inflammatory and pro-atherogenic intermediate monocyte phenotype towards an anti-inflammatory circulating leucocyte profile, as evidenced by an increase in the number of the anti-inflammatory Treg phenotype, compared with non-exercising patients. We also confirm elevated circulating and cellular markers of systemic inflammation in HD patients compared with an age-matched healthy cohort: CRP, IL-6 and TNF-α were significantly elevated in the HD cohort, the proportion of Tregs was reduced and monocyte populations shifted towards intermediate and non-classical phenotypes. Physical activity levels and physical function were also lower in HD patients.

This is the first study in any clinical population to show that exercise training selectively diminishes the proportion of monocytes within the inflammatory intermediate subset. Elsewhere, in an inactive elderly, but healthy group, 12 weeks of thrice weekly endurance and resistance exercise training significantly reduced the percentage of CD14+CD16+ monocytes and decreased the TNF-α response to lipopolysaccharide stimulation [20]. However, CD16+ monocytes were not analysed separately as intermediate and non-classical subsets, as this is a relatively new classification system [21]. Whether the present findings are applicable to populations other than HD patients is not known, but may explain these previous observed changes in ‘CD16+’ monocytes.

Elevations in circulating intermediate monocytes are associated with an increased cardiovascular risk in non-CKD, CKD and HD patients [4, 6, 22] and have been described as a promising therapeutic target for CVD in CKD [23]. In agreement with the present study, the number and proportion of intermediate monocytes are increased in HD patients compared with healthy cohorts [4, 5]. Intermediate monocytes secrete pro-inflammatory cytokines (TNF-α, IL-1β and IL-6), express toll-like receptor 4 and adhesion molecules (CCR5 and CX3CR1) and exhibit spontaneous reactive oxygen species production [24–26], all considered mechanisms for cardiovascular dysfunction. The reduction in intermediate monocytes that we report here is encouraging as it may be a mechanism by which exercise protects against systemic inflammation and associated cardiovascular disorders in HD patients.

In the present study, the proportion of CD4+ lymphocytes that were Tregs was lower in HD patients than in the healthy group. Reduced Treg capacity has been reported previously in HD patients [27, 28]. Abnormalities in Treg activity are harmful; deficiencies are associated with autoimmunity, inflammation and allergy, whereas over-activation is associated with increased risk of chronic infections and tumour growth [29]. The diminished Treg capacity in HD patients may contribute to chronic immune activation and inflammation typically exhibited in this population.
The favourable change in the number of Tregs in the exercise group after 6 months compared with the non-exercising group may represent enhanced anti-inflammatory capacity. Sedentary but otherwise healthy individuals have demonstrated lower numbers of Tregs than physically active groups and antigen-stimulated IL-10 release correlates with this cell population [30]. However, as we found no change in the proportion of CD4\(^+\) lymphocytes that were Tregs, the increase we observed may simply relate to differences in overall CD4\(^+\) numbers. This is further supported by the observation that the number of CD4\(^+\) lymphocytes decreased significantly in the non-exercising group but not in the exercise group. Therefore, it is not clear whether the enhanced Treg population in the exercise group actually represents a true anti-inflammatory adaptation.

Circulating CRP, IL-6 and TNF-\(\alpha\) were not improved after 6 months of regular intradialytic exercise in this study. Examination of the literature to date reveals that no well-controlled trials have reported significant meaningful reductions in these soluble inflammatory markers after exercise training in HD patients [14]. Well-designed studies have reported no change in CRP after 3–4 months of intradialytic cycling or extradialytic resistance training [31, 32], or with maintained CRP but no improvement in IL-6 or TNF-\(\alpha\) [33, 34]. In predialysis CKD cohorts, resistance exercise that increases muscle mass [35] and regular walking exercise have been shown to favourably alter circulating inflammatory factors [13].

In a large cohort of healthy individuals (\(n = 4289\)), physical activity was associated with reduced CRP and IL-6, and increasing activity levels reduced markers of inflammation over a 10-year follow-up [36]. Therefore, for these soluble inflammatory markers, a large cohort or a long exercise intervention may be required to detect meaningful change. This is likely exacerbated in HD patients by the large intra- and inter-individual variation of IL-6, CRP and TNF-\(\alpha\). The day-to-day variation (i.e. due to fluid overload, recurrent infections, HD itself) in this population may limit the use of CRP and circulating cytokines in isolation to detect long-term change.

Previous studies have demonstrated sedentary behaviour in HD patients [11, 37], but the extent of inactivity is worth highlighting. Significantly, the least active HD patients have a 62% increased mortality risk [12] and those with the lowest physical function have higher hospitalization rates and all-cause mortality [38, 39]. A recent Cochrane Review concludes there are various benefits of increasing physical activity for health and well-being in CKD [10], and the present study adds to this body of evidence. Despite the seemingly low power output the exercise group improved physical function. This improvement was probably augmented by low baseline values giving greater potential for improvement. Increasing physical function has a positive impact on quality of life and is valued very highly by patients; therefore, promotion of this key outcome is likely to be more effective in encouraging continued participation in physical activity.

This exercise programme was limited to one mode and prescription of exercise and was not standardized per patient; different exercise modalities, at different times or intensities, may have different results. Nonetheless, this study makes a major contribution because it was pragmatic in design, thereby ascertaining a realistic response to an achievable intradialytic programme per se in a naturally heterogeneous HD cohort. There was, therefore, expected variation in the absolute exercise intensity achieved in this study (comparable to other studies [31, 40]); however, subjective intensity remained quite consistent, as evidenced by RPE.

For reasons of practicality and to avoid sampling bias and ‘exercise-contamination’ of control patients that occurs when control and exercise patients share shifts, this study was not randomized. This led to different demographics between the exercising and non-exercising groups, including ethnicity and age, but did not confound the key outcome measures. Likewise, haemoglobin was different between groups at baseline, but as this did not change over time, it is unlikely to have influenced the findings. Finally, the small sample size is due to the practicalities of running such a study despite approaching all eligible patients. This pragmatic evaluation of intradialytic exercise provides important novel evidence into anti-inflammatory adaptations that provides a primary point of reference for future investigations into the therapeutic benefits of exercise in this population.

Conclusions

HD patients are sedentary and demonstrate high levels of chronic inflammation, yet we show for the first time that undertaking regular exercise during HD has an anti-inflammatory effect at a circulating cellular level. The reduction in intermediate monocytes may be protective against the substantially increased risk of cardiovascular morbidity and mortality and further supports the therapeutic potential of regular exercise in these patients.

Supplementary data

Supplementary data are available online at http://ckj.oxfordjournals.org.

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Authors’ contributions

Research idea and study design: N.C.B. and A.C.S.; data acquisition: M.D., H.M.L.Y and D.R.C.; data analysis/interpretation: M.D.; and supervision or mentorship: N.C.B., A.C.S. and J.O.B. Each author contributed important intellectual content during manuscript drafting or revision and accepts accountability for the overall work by ensuring that questions pertaining to the accuracy or integrity of any portion of the work are appropriately investigated and resolved.
Conflict of interest statement

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