Physical activity is associated with increased resting-state functional connectivity in networks predictive of cognitive decline in clinically unimpaired older adults

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Abstract

Introduction: Physical activity (PA) promotes resilience with respect to cognitive decline, although the underlying mechanisms are not well understood. We examined the associations between objectively measured PA and resting-state functional connectivity magnetic resonance imaging (rs-fcMRI) across seven anatomically distributed neural networks.

Methods: rs-fcMRI, amyloid beta (Aβ) positron emission tomography (PET), PA (steps/day × 1 week), and longitudinal cognitive (Preclinical Alzheimer’s Cognitive Composite) data from 167 cognitively unimpaired adults (ages 63 to 90) were used. We used linear and linear mixed-effects regression models to examine the associations...
between baseline PA and baseline network connectivity and between PA, network connectivity, and longitudinal cognitive performance.

**Results:** Higher PA was associated selectively with greater connectivity in three networks previously associated with cognitive decline (default, salience, left control). This association with network connectivity accounted for a modest portion of PA’s effects on A\(\beta\)-related cognitive decline.

**Discussion:** Although other mechanisms are likely present, PA may promote resilience with respect to A\(\beta\)-related cognitive decline, partly by increasing connectivity in a subset of cognitive networks.

**KEYWORDS**
Alzheimer’s disease, amyloid, cognition, functional connectivity, physical activity

**INTRODUCTION**

Alzheimer’s disease (AD) pathology develops over decades,\(^1\) and recent studies suggest that the eventual emergence of cognitive impairment in people with AD pathology is strongly influenced by individual risk and lifestyle factors.\(^2\)\(^-\)\(^5\) The long preclinical phase of AD has become a major focus of clinical research, as intervention at this stage of disease may prevent the widespread synaptic loss and neurodegeneration that is characteristic of symptomatic phases of AD. Intervening on modifiable risk and lifestyle factors during the preclinical phase of disease may have a significant impact on the emergence of clinical cognitive impairment and dementia at low cost and with minimal risk. Physical activity (PA) level is a modifiable risk factor known to promote resilience with respect to developing dementia,\(^2\)\(^,\)\(^6\)\(^,\)\(^7\) although the mechanisms underlying this effect are not well understood. Prior work suggests that greater levels of PA may be particularly protective in people with elevated brain amyloid \(\beta\) (A\(\beta\)) levels, both with respect to cognitive decline and to the progression of cortical atrophy, effects that appear to be independent of vascular risk factors.\(^6\)

Resting-state functional connectivity magnetic resonance imaging (rs-fcMRI) is a non-invasive tool that can be used to quantify the integrity of distributed neural networks that are somewhat specialized for particular cognitive functions. Building on the observation that decreased functional connectivity in the default network is seen in AD as compared to normal aging,\(^8\) a broad array of rs-fcMRI–based measures have been examined as potential biomarkers and predictors of disease progression. Our group and others have demonstrated that lower levels of connectivity in the default, salience, and control networks presage cognitive decline in cognitively unimpaired older adults.\(^9\)\(^,\)\(^10\) Our prior work also demonstrated that lower functional connectivity in these networks was most predictive of cognitive decline in cognitively unimpaired individuals with elevated A\(\beta\), suggesting a synergistic effect of low connectivity and greater A\(\beta\) with respect to cognitive decline.\(^7\) Furthermore, these networks, especially the default network, share significant anatomical overlap with structures involved in early AD pathophysiology, including the posterior cingulate and precuneus. Progressive decrements in connectivity in the default network correspond to worsening clinical impairment, including diagnoses of mild cognitive impairment and AD dementia.\(^11\) The extent to which default and salience network connectivity is disrupted correlates with episodic memory performance, both in cognitively unimpaired adults and in adults challenged with administration of scopolamine, an anticholinergic drug that impairs memory.\(^12\) Although more work is needed, these data suggest that rs-fcMRI may be a useful non-invasive tool in both predicting who may be at risk for future cognitive decline as well as evaluating interventions aimed at delaying or preventing the development of dementia.

In this study, we investigated whether levels of PA in older adults were associated with connectivity across a broad set of networks, including three networks (default, salience, and left control) shown previously to be related to cognitive performance on tasks typically affected in AD\(^13\)\(^,\)\(^14\) and in which low connectivity predicts future cognitive decline.\(^9\) In addition, we analyzed the primary visual, visual association, dorsal attention, and motor networks for comparison. Given the association of greater PA with less cognitive decline over time, we also investigated the extent to which increased network connectivity may account for the protective effects of PA against cognitive decline. To do this, we leveraged data from the Harvard Aging Brain Study (HABS), a longitudinal study of cognitive aging and preclinical AD. As HABS participants undergo A\(\beta\) positron emission tomography (PET) imaging, we are also able to examine the extent to which the effects of connectivity and PA may be different in older adults with elevated A\(\beta\) burden.

**METHODS**

**2.1 Participants**

A total of 167 cognitively unimpaired participants from HABS were included in the study. Inclusion criteria for HABS required a Clinical Dementia Rating (CDR) Scale of 0, a Mini-Mental State Examination
Amyloid PET 3O f8 Imaging

2.3.1 (fil.ion.ucl.ac.uk/spm/) using methods described previously. Briefly, rs-fcMRI data processing utilized statistical parametric mapping 12 2.3.2 degrees, echo time 30 ms, matrix 72 × 72, field of view 216 mm, × 3 mm, slice thickness 3 mm, giving an in-plane resolution of 3 × 3 mm. Average was obtained using repetition time 3 seconds, flip angle 85 degrees, echo time 30 ms, matrix 72 × 72, field of view 216 × 216 mm, and 47 × 3 mm axial slices, which resulted in isotropic voxels of 3 mm. A total of 124 volumes were acquired in each run. During the scan, participants were asked to remain awake, lie flat with their eyes open, and with a visible crosshair projected on a screen.

2.3.3 A Siemens 3T Trio Tim MRI scanner with a 12-channel phased-array coil was used to collect structural and functional MRI data. Functional MRI acquisition employed gradient-echo echoplanar imaging sequence sensitive to blood oxygen level–dependent contrast. Whole-brain coverage was obtained using repetition time 3 seconds, flip angle 85 degrees, echo time 30 ms, matrix 72 × 72, field of view 216 × 216 mm, and 47 × 3 mm axial slices, which resulted in isotropic voxels of 3 mm. A total of 124 volumes were acquired in each run. During the scan, participants were asked to remain awake, lie flat with their eyes open, and with a visible crosshair projected on a screen.

rs-fcMRI processing and analysis

rs-fcMRI data processing utilized statistical parametric mapping 12 (fil.ion.ucl.ac.uk/spm/) using methods described previously. Briefly, each run was normalized to the Montreal Neurological Institute (MNI) 152 EPI template using SPM-defined normalization parameters. We employed a 6-mm Gaussian kernel for smoothing. Additional processing involved movement correction using co-registration parameters and first derivatives to reduce movement artifacts, as well as removing frequencies outside of the 0.01 to 0.08 Hz band with temporal band-pass filtering. Additional details regarding rs-fcMRI data processing have been published previously.

2.3.3 Amyloid PET

11C PiB-PET was used to assess Aβ burden, using previously published procedures. Briefly, PiB-PET images were acquired using a dynamic
acquisition protocol that generated 39 volumes during a 1-hour period following an 8.5 to 15 mCi bolus injection of $^{11}$C-PiB. Reconstruction, attenuation correction, and evaluation for head motion were performed prior to co-registering each image to the corresponding T1 image for each participant using a rigid-body registration employing six degrees of freedom. The FreeSurfer-defined cerebellar gray region-of-interest was used as the reference region. As in previous studies from HABS, partial volume correction was applied using geometric transformation and a distribution volume ratio (DVR) of the frontal, lateral, and retrosplenial regions (FLR) was used as the primary measure of Aβ burden. Based on prior work in HABS using a Gaussian mixture modeling approach, a PiB FLR DVR of greater than 1.3 was considered Aβ positive.

### 2.5.1 Baseline physical activity and network connectivity

Linear regression models implemented in R (R Foundation for Statistical Computing, version 3.5.3) were used for primary analyses. PA (mean steps per day) was log-transformed prior to entry into models. To assess the cross-sectional relationship of PA to network connectivity, PA was used as a predictor of connectivity in the default, salience, and left control networks (model: network connectivity ∼ PA + covariates). We included age, sex, and apolipoprotein E ε4 (APOE4) carrier status as covariates. Mean movement during MRI and the number of usable volumes were included as additional covariates to control for the quality of rs-fcMRI measurement. False discovery rate (FDR) was employed to adjust for the number of comparisons performed. To determine the network specificity of PA associations with default, salience, and left control network connectivity, we repeated these models in four networks (dorsal attention, primary visual, visual association networks, and motor) in which connectivity strength at study baseline was not associated with cognitive decline as well as with global connectivity.

Because PA and vascular risk are often correlated, we performed additional sensitivity analyses to examine the degree to which PA is independently associated with connectivity measures. We first calculated a Pearson correlation between log mean steps with the well-described and validated body mass index (BMI)-based Framingham cardiovascular risk score (FHS-CVD). We then included the FHS-CVD score as an additional covariate in the same models described above (connectivity ∼ PA + FHS-CVD + covariates).

### 2.5.2 Physical activity, network connectivity, and longitudinal cognition

Next, we used linear mixed-effects models (R version 3.5.3) to examine the extent to which the protective effects of PA on cognitive decline in those with high Aβ burden may be accounted for by the effects of PA on connectivity (Figure 2). To reduce the number of comparisons being made in the longitudinal analyses of cognitive data, we derived a composite measure of cognitive network connectivity by averaging the z-scored connectivity values in the default, salience, and left control networks. Interactions with Aβ level at baseline (assessed using PiB-PET) were included in these models based on prior work. Model 1 (below) was used to assess the interaction of PA with baseline Aβ relative to longitudinal cognitive performance. Models 2 and 3 followed on results from Model 1 and assessed whether the effects of PA and baseline Aβ on longitudinal cognitive performance may be accounted for by the effects of PA on network connectivity. In addition to MR quality measures, years of education, age, sex, and APOE4 carrier status were included as covariates, as in prior work from the HABS cohort.

1. Cognition ∼ $PA_Aβ^\beta$*time + covariates*time.
2. Cognition ∼ composite connectivity$PA_Aβ^\beta$*time + covariates*time.
3. Cognition ∼ $PA_Aβ^\beta$*time + composite connectivity$PA_Aβ^\beta$*time + covariates*time.

### 3 RESULTS

#### 3.1 Effect of physical activity on functional network connectivity

Baseline demographic and clinical characteristics are summarized in Table 1. We observed an association between greater PA and higher connectivity in the default, salience, and left control networks after correcting for multiple comparisons (default: $t = 2.81(160), P = .009$, salience: $t = 2.60(160), P = .017$, left control network: $t = 3.91(160), P < .001$) (Figure 1). As expected, the same association was not observed between PA levels and connectivity in the dorsal attention ($P = .313$), motor ($P = .419$), primary visual ($P = .074$), and visual association ($P = .859$) networks, or with global connectivity (Figure S1), indicating that PA relationships to connectivity were limited to the three networks shown previously to be associated with longitudinal cognitive changes in HABS.

In analyses performed to account for vascular risk factors, we found PA and FHS-CVD to be weakly correlated in the anticipated direction ($r = -0.16, P = .03$). Associations of PA with connectivity in the default ($P = .02$), salience ($P = .02$), and control networks ($P < .01$) remained significant when including FHS-CVD as a covariate.
FIGURE 1  Associations between level of physical activity and network connectivity. Log-transformed measures of mean steps per day were compared with connectivity within the default (left), salience (middle), left control (right), motor, primary visual, and visual association networks. Greater levels of physical activity were selectively associated with higher connectivity strengths in each of these networks.

TABLE 1  Baseline demographic and clinical characteristics

| Characteristic | All participants (n = 167) |
|----------------|--------------------------|
| Age in years, mean (SD) | 74.1 (6.1) |
| Women, n (%) | 90, 53.9% |
| Education in years, mean (SD) | 15.6 (3) |
| Mean steps per day, mean (SD) | 5367 (2868) |
| Aβ PET FLR DVR (PVC) | 1.4 (0.4) |
| Aβ PET FLR DVR (PVC) > 1.3, n (%) | 56, 33.5% |
| Aβ PET FLR DVR (PVC) < 1.3, n (%) | 111, 66.5% |
| APOE ε4 carriers, n (%) | 52, 31% |

Abbreviations: APOE ε4, apolipoprotein E ε4, Aβ, amyloid beta; DVR, distribution volume ratio; FLR, frontal, lateral temporal and parietal, and retrosplenial regions; PET, positron emission tomography.

suggesting PA has an effect independent of systemic vascular risk on network connectivity.

3.2  Effect of physical activity, Aβ amyloid, and network connectivity on longitudinal cognition

As both higher PA levels and higher connectivity in the default, salience, and control networks have been associated previously with less cognitive decline,6 we next examined whether the potential protective effects of PA may be accounted for by increased mean network connectivity (Figure 2). As the effects of PA on longitudinal cognitive performance are seen largely in older adults with higher Aβ, interactions with Aβ levels were included for both PA and connectivity. As shown previously,6 greater PA was associated with slower cognitive decline in individuals with higher Aβ (PA*Aβ, F[1,917] = 12.74, P < .001). Similarly, we observed a significant interaction of connectivity with Aβ in predicting longitudinal cognitive performance (P = .035). A third model in which both PA and connectivity were included suggests that PA effects on connectivity may account for a modest degree of the association of PA and Aβ with longitudinal cognitive performance (Figure 2, Supplementary table S2). Together, these results suggest that connectivity moderates the association of PA with cognition, although this result should be interpreted in the context of the Aβ and PA interaction included in these models.

4  DISCUSSION

We examined the associations between PA, network connectivity, Aβ, and longitudinal cognitive performance in a deeply phenotyped, longitudinal cohort of older adults. We demonstrate that objectively measured levels of PA are positively correlated with the integrity of resting state connectivity networks. Notably, the association of PA and connectivity was limited to a set of three networks—the default, salience, and left control networks—in which connectivity strength has been shown previously to predict longitudinal cognitive performance, particularly in individuals with elevated Aβ.9 Next we observed suggestive evidence that the association between PA and network connectivity may account for a modest portion of the protective effects of PA on longitudinal cognitive change in preclinical AD, although this conclusion requires further confirmation. This study suggests that connectivity is one link between two previous findings in preclinical AD; first that increased PA is associated with resilience to cognitive decline,6,7 and second that higher connectivity within a subset of cognitive networks may be associated with less cognitive decline over time, particularly in persons with higher Aβ levels.9 Together, our findings suggest that one mechanism by which PA promotes resilience to cognitive decline may be through promoting the integrity of neural networks underlying key aspects of memory and attention.

Previous studies suggest that neurodegenerative diseases may involve the selective vulnerability and deterioration of large-scale functional networks within the brain.32 In AD, changes in default, salience, and control networks have been well described.18–20,33–35 dovetailing with the observation here that increased connectivity is associated with PA in these same networks. A recent study demonstrated that interactive aberrant connectivity in the overlapping nodes of the default, salience, and control networks is seen in AD dementia and associated with worse scores on the MMSE.36 These findings and those of the present study are consistent with the hypothesis that greater connectivity is one possible contributor to the protective
FIGURE 2  Inter-relationships between physical activity, connectivity, and longitudinal cognitive performance. Greater physical activity (PA) in participants with elevated amyloid beta (Aβ) was associated both with lesser cognitive decline over time and with greater network connectivity. PA-associated connectivity changes may account for a modest portion of the observed association of PA and Aβ with longitudinal cognitive performance.

effects of PA against cognitive decline and AD dementia. The effects of both PA and rs-fcMRI on cognitive decline are most clear in individuals with higher levels of Aβ burden, perhaps because this population is at the highest risk of progressive neurodegeneration and loss of cognitive performance.

PA was associated only with increased connectivity in networks related to memory and attention, although the cause for this selectivity is unknown. Going forward, future studies that elucidate the mechanisms underlying this selectivity will be needed to further inform the relationship between PA and cognition. One possibility is through network or cell-targeted molecular mechanisms via release of activity-induced neurotrophic factors that act on proteins expressed in specific neurons, structures, or networks in the brain. For example, Fibronec tin type III domain-containing protein 5 (FNDC5)/irisin is an exercise-associated myokinin expressed in the hippocampus that when knocked down impairs long-term potentiation, and conversely, when overexpressed, rescues memory impairment in AD mouse models.37 Although more work is needed, our study suggests that rs-fcMRI may be useful in evaluating non-pharmacologic interventions in preclinical AD and mild cognitive impairment, particularly those with a PA-related intervention. Boraxbekk et al. demonstrated a PA score to be positively associated with connectivity in the posterior default network in a study of healthy older individuals, consistent with our results.38 Notably, the results here are consistent with a small prospective study of MCI participants engaged in a 12-week walking exercise program, where significant increases in connectivity were observed in 10 brain regions (including the posterior cingulate and precuneus) over the course of the intervention.39 Connectivity levels and changes in response to interventions in AD-relevant brain regions could potentially be used to help determine or predict the intensity, frequency, and type of PA or other non-pharmacologic interventions that have the greatest potential to benefit cognition and delay the onset of dementia.

There are important limitations in this study to consider, and this work should be interpreted in the context of the participants studied. HABS participants tend to be more highly educated than the general population, which may affect the generalizability of the results here. In addition, people with severe or unstable cerebrovascular disease are excluded from the study. Furthermore, this study makes use of baseline Aβ PET, PA, and rs-fcMRI measures, and future work using longitudinal data may strengthen the causal inferences suggested by this study. Similarly, PA may need to be measured in greater detail (e.g., for intensity) or for longer periods to understand the PA more completely and rs-fcMRI relationships. Future work should aim to understand these important variables to inform the design of non-pharmacologic lifestyle interventions intended to prolong healthy cognition.

5 | CONCLUSION

Although greater levels of PA are widely thought to confer resilience for late-life cognitive decline, the linkage between PA and brain health have not been fully elucidated. By utilizing data in which objective measures of PA, functional connectivity, AD pathology, and longitudinal cognitive performance were available, we examined how variations...
in PA are reflected in the integrity of anatomically distributed neural networks known to be important for cognition. We observed an intriguing relationship between greater levels of PA and higher connectivity in a subset of neural networks in older adults at an elevated risk of developing AD. These relationships between PA and connectivity were limited to three networks (default, salience, and left control) that have been shown previously to be degraded in early AD and are linked to performance in memory, attention, and executive function intensive tasks, suggesting that PA may confer resilience to cognitive decline, partly through increased integrity within a subset of AD-sensitive neural networks. Together with prior work, the results here also suggest that rs-fcMRI may be a useful tool in optimizing and evaluating the effectiveness of PA-modifying interventions in cognitive aging and pre-clinical AD.

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CONFLICT OF INTEREST
The authors declare no conflicts of interest.

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**SUPPORTING INFORMATION**

Additional supporting information can be found online in the Supporting Information section at the end of this article.