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# More evidence is needed. Iron, incident cognitive decline and dementia: a systematic review

Diane E. Hosking, Scott Ayton, Nigel Beckett, Andrew Booth and Ruth Peters

## Abstract

**Background:** Our aim was to systematically review the relationship between iron and incident cognitive decline or dementia from midlife onwards.

**Methods:** Systematic review of eligible studies using Medline, Embase and PsycINFO® for the period from 1 January 1986 to 2 December 2016 (CRD42016023800), where study populations had a mean age of over 50 years and were free of cognitive impairment or dementia at baseline. Two authors independently extracted data according to eligibility criteria and assessed study characteristics, quality and outcomes. Disagreement was resolved by discussion.

**Results:** A total of 1185 relevant records were identified with 12 full-text articles eligible for review. Six studies were excluded, leaving six texts to be included. Sample size ranged from 90 to 7173, with an average follow up of approximately 11.5 years. Baseline iron measures included brain iron ( $n = 2$ ), iron-related biomarkers in blood and plasma ( $n = 2$ ), and iron intake estimates from dietary records ( $n = 2$ ). Outcomes were dementia incidence ( $n = 2$ ) and longitudinal outcomes on neuropsychological tests ( $n = 4$ ). Bias was evident across studies in one or more of the following: recruitment, iron exposure, outcome assessments, potential confounders, missing data or attrition.

**Conclusions:** Diversity across the small number of identified studies precludes conclusions regarding the role of iron in cognitive decline or dementia. Our review highlights substantial gaps in the evidence base and the need for more comprehensive, higher quality studies in this area.

**Keywords:** cognitive ageing, dementia, incidence, iron, review

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Correspondence to:

**Ruth Peters**

Lifecourse Ageing Research Centre, Neuroscience Research Australia, University of New South Wales, Sydney NSW 2031, Australia

[r.peters@neura.edu.au](mailto:r.peters@neura.edu.au)

**Diane E. Hosking**

Centre for Research into Ageing Health and Wellbeing, Australian National University, Canberra, Australia

**Scott Ayton**

The Florey Institute of Neuroscience and Mental Health, University of Melbourne, Melbourne, Australia

**Nigel Beckett**

Care of the Elderly, Imperial College London, London, UK

**Andrew Booth**

School of Health and Related Research, University of Sheffield, Sheffield, UK

## Introduction

Older age is associated with increased risk of iron deficiency (ID), elevated body iron stores and increased brain iron levels.<sup>1,2</sup> Complex regulatory mechanisms fine-tune iron metabolism and homeostasis for both heme and nonheme iron.

Iron has a key role in multiple pathways, for example, as a constituent part of proteins needed for oxygen transport, oxidative phosphorylation, myelin production, and production and breakdown of neurotransmitters.<sup>3–5</sup> Consequently there are multiple plausible mechanisms by which iron deficiency or overload may elevate the risk of age-related cognitive decline and dementia. A detailed mechanistic review is beyond the scope of this article, however for recent reviews in this area, see Hare and colleagues<sup>3</sup> and Waldvogel- Abramowski and colleagues<sup>4</sup>

## Iron deficiency

In older adults, absolute ID is caused by insufficient dietary iron intake, gastrointestinal malabsorption, or increased blood losses attributable to gastrointestinal pathologies.<sup>2</sup> Potential cognitive impacts related to inadequate iron might stem from cerebral hypoxia,<sup>6</sup> insufficient neurotransmitter synthesis<sup>5</sup> or poor myelin integrity.<sup>7</sup>

There is also a growing body of evidence that anaemia in older age (defined by haemoglobin level <13 g/dl in men and <12 g/dl in women) is associated with poor cognition, cognitive decline and dementia.<sup>8</sup> ID anaemia accounts for around 16% of total anaemia cases.<sup>9</sup>

## Iron overload

Body iron levels may be elevated in middle-aged and older adults due to consumption of highly bioavailable forms of iron (supplemental iron and red meat) and of fruit, an enhancer of nonheme iron absorption (vitamin C).<sup>10</sup> In two meta-analyses of prospective cohort studies, high body iron stores and consumption of heme iron were associated with increased risk of coronary heart disease (21 studies)<sup>11</sup> and type 2 (T2) diabetes (11 studies),<sup>12</sup> which may highlight a further pathway between iron overload and cognitive impairment and dementia.<sup>13–15</sup>

## Increased brain iron

The accumulation of iron in the brain is an established hallmark of ageing.<sup>16</sup> Excess iron stores may increase pro-oxidant reactions and generation of free radicals,<sup>1,3</sup> which could contribute to neurodegeneration. In cross-sectional studies, magnetic resonance imaging (MRI) measures of iron are associated with amyloid load,<sup>17</sup> smaller hippocampal volume<sup>18</sup> and poorer cognitive performance.<sup>15,19</sup> Hippocampal iron content has also been correlated with cognitive performance and disease duration in Alzheimer's disease<sup>20,21</sup> and MRI-measured iron in the basal ganglia in cognitively normal adults predicted greater rates of cognitive decline and cortical atrophy.<sup>22,23</sup> Brain iron deposits have also been positively associated with white matter hyperintensities, suggesting a potential link between brain iron and vascular pathology.<sup>15,24</sup>

The proportion of the population at risk for cognitive decline and dementia is rapidly increasing, and there is no effective pharmacological cure or preventive intervention.

The promotion of clinical understanding through collation of evidence for modifiable risk factors is an important preventive strategy. Iron has been raised as a potentially modifiable risk factor for cognitive decline or dementia but the area is complex. There are multiple ways of measuring both iron and cognition, and to date there is no critical appraisal of the evidence depth, breadth or quality. Our aim was to systematically review and evaluate the current evidence base that relates to the relationship between iron and incident cognitive decline or dementia.

## Methods

### Search strategy and selection criteria

The databases MEDLINE, Embase and PsycINFO were searched over a 30-year period from 1986 to 2 December 2016. Because this is a first review of the literature in this area, search terms were chosen to be as broad as possible to prevent inadvertent exclusion of particular cognitive assessment or particular iron measures.

Search terms included dementia, alzheimers disease, vascular dementia, multi-infarct dementia, cognit\* and iron, heme, haem, ferritin and serum transferrin. See supplementary text for details. Reference lists were screened and experts in the field were consulted. In addition, searches of the Cochrane Library, the ISRCTN Register and the ClinicalTrials.gov website were carried out to look for ongoing or completed relevant trials.

All identified abstracts, or titles, where abstracts were unavailable, were independently read by two reviewers (DH, RP) and a list of potential evidence for inclusion was compiled by the two analysts (DH, RP). These lists were then compared and any differences resolved by discussion. Full text copies of the selected texts were independently read and assessed for relevance by both analysts and in accordance with the following inclusion and exclusion criteria.

### Inclusion criteria.

Prospective longitudinal studies of primary research (cohorts or clinical trials) reporting on the following:

1. An assessment of exposure to iron (assessed *via* biomarkers, e.g. blood, cerebral spinal fluid or neuroimaging) or external sources, for example, diet or iron supplementation.
2. Evidence, or clear implication, that participants were free of cognitive decline or dementia at baseline assessment.
3. Use of formal assessment of cognitive function.
4. Reporting outcomes of cognitive decline or incident dementia.
5. Adult populations with a mean age over 50 years

### Exclusion criteria:

1. Publications relating only to populations selected on the basis of clinical comorbidity when that comorbidity may also affect cognitive function, for example, requirement for renal dialysis.
2. Non English publications (in the absence of resources available for translation).
3. Inclusion of nonadults.

Papers identified as relevant were then independently assessed for quality and data relating to study characteristics, assessment of cognitive function and iron exposure were extracted by each analyst. A formal quality scoring scheme was not used as these lack discriminatory power, however each paper was assessed against the key factors given in Critical Appraisal Skills Programme checklists for evaluating trials and longitudinal studies (Critical Appraisal Skills Programme: <http://www.casp-uk.net/> accessed 15th February 2017).

The review was carried out to best practice and in accordance with PRISMA guidelines.<sup>25</sup> Protocol registration no. CRD42016023800.

## Results

A total of 2001 records were identified from the searches and a further two from reference screening/expert recommendation; 1185 records remained once duplicates were removed. These were screened and 12 full text articles representing 12 separate studies were reviewed. Six studies were included with the remaining six excluded because they lacked results related to iron exposure ( $n = 2$ ),<sup>26,27</sup> included prevalent cases of cognitive impairment ( $n = 3$ ),<sup>15,28,29</sup> or had a cross-sectional design ( $n = 1$ ).<sup>30</sup> See Figure 1 for the PRISMA study flow chart.

### Study characteristics

Of the studies that were included, four reported results from North American populations (United States of America); two using data from the National Health and Nutrition Examination Survey (NHANES),<sup>31,32</sup> one from a Detroit-based cohort study,<sup>22</sup> and one using participant data from the Alzheimer's Disease Neuroimaging Initiative (ADNI) cohort.<sup>33</sup> The remaining studies were from Australia, a longitudinal cohort study based in Canberra and Queanbeyan,<sup>34</sup> and the Netherlands, a cohort analysis of clinical trial data from the Folic Acid and Carotid Intima-Media Thickness (FACIT) trial.<sup>35</sup> For further study characteristics, see Table 1.

### Measures of iron

Studies used widely differing measures of baseline iron (Table 2). Of the six identified studies, two used measures of brain iron<sup>22,33</sup> and four used measures of peripheral iron.<sup>31,32,34,35</sup> The NHANES 40-plus cohort reported on serum transferrin,<sup>32</sup> the NHANES 60-plus cohort and the Australian cohort study used dietary iron,<sup>34</sup> the FACIT trial collected total iron-binding capacity, transferrin saturation, ferritin and nontransferrin bound iron.<sup>35</sup> Of those that collected data on brain iron, the ADNI cohort used ferritin in cerebrospinal fluid (CSF)<sup>33</sup> and the Detroit study used imaging (MRI).<sup>22</sup> The Detroit cohort and the FACIT trial cohort were the only studies to report assessment of iron at follow up. The FACIT trial collected serum iron, ferritin and nontransferrin bound iron and the Detroit study repeated their baseline MRI scan.<sup>22,35</sup>

### Cognitive outcome measures

The NHANES cohorts reported on dementia outcomes (Table 3). Data on dementia and Alzheimer's disease outcomes were collected from death certificates, hospital and nursing home records for the 40-plus cohort.<sup>32</sup> Diagnoses of Alzheimer's disease for the 60-plus cohort were collected from death certificates only;<sup>31</sup> neither

carried out cognitive testing. The Detroit cohort,<sup>22</sup> FACIT trial cohort,<sup>35</sup> ADNI cohort<sup>33</sup> and Australian cohort<sup>34</sup> used standard cognitive assessment tools at baseline and follow up.

### *Association between iron measures and cognitive outcomes as reported by the included studies*

Overall the results were variable across studies (Table 4):

1. In the NHANES 40-plus cohort ( $n = 6427$ ), an association was found between co-occurring high levels of transferrin (TS) and cholesterol, and increased risk of Alzheimer's disease [both TS and cholesterol at the 75th percentile, hazard ratio (HR) 3.19, 95% confidence interval (CI) 1.31–7.75]; but no association between transferrin alone and Alzheimer's disease<sup>32</sup> (transferrin >75% compared with < 75% Alzheimer's disease: HR 1.47, 95% CI 0.82–2.63).
2. In the FACIT trial ( $n = 800$ ) the potential of folic acid supplementation to modify the effects of iron measures on cognitive change was tested prior to examining associations between serum iron parameters and longitudinal cognitive functioning. No associations were found. When stratifying for status as a blood donor, higher iron status was associated with less decline in sensorimotor (parameter estimate = 0.005,  $p = 0.010$ ) and word fluency tasks (parameter estimate = 0.006,  $p = 0.012$ ).<sup>35</sup>
3. The Australian PATH cohort study ( $n = 1354$ ) found an increased risk of cognitive impairment with higher levels of iron intake (HR 1.54, 95% CI 1.03–2.29 for transition from normal cognition to mild cognitive impairment (MCI), per 1 mg iron) and the potential for an interaction with sex such that higher intake may be associated with lower risk in women but higher risk in men.<sup>3</sup>
4. The PATH study reported a mean iron intake of 18 mg per day, more than double the daily intake of 8 mg recommended by the Australian National Health and Medical Research Council (NHMRC) for those in their sixties. 4. The Detroit cohort ( $n = 78$ ) (examining the hippocampus, caudate and putamen) reported no association between baseline iron or change in iron and episodic memory or nonverbal working memory, with the exception of baseline iron in the caudate nucleus which was associated with less improvement in verbal working memory,  $\beta -0.18$  ( $p = 0.01$ ).<sup>22</sup>
5. The NHANES 60-plus cohort ( $n = 4639$ ) found no association between Alzheimer's disease related death and dietary iron intake ( $p = 0.1022$ ) but an increased risk of Alzheimer's disease mortality found for those with low haemoglobin (HR 8.4, 95% CI 1.4–50.8).<sup>31</sup>
6. The study using the ADNI cohort ( $n = 43$ ) reported an increased risk of decline associated with higher levels of ferritin, but suggested that the result may be confined to those with the apolipoprotein (APOE)  $\epsilon 4$  allele. For those who were APOE  $\epsilon 4$  positive, Alzheimer's disease Assessment Scale (ADAS Cog)  $\beta 0.09$  [standard error (SE) 0.04],  $p = 0.02$ , Rey Auditory Verbal Learning Test (RAVLT)  $\beta -1.49$  (SE 0.4),  $p < 0.001$  compared with APOE  $\epsilon 4$  negative, ADAS Cog  $\beta -0.04$  (SE 0.016),  $p = 0.02$ , RAVLT nonsignificant.<sup>33</sup>

**Figure 1.** Study selection flowchart.  
Source: <http://prisma-statement.org/>

### Study quality

All studies reported a clear research question (Table 5). However, studies utilized diverse designs with the potential for bias in recruitment, measures of iron exposure, outcome assessment, potential confounders, missing data or attrition.

The applicability of three studies was limited by recruitment from specific population groups; one from an ongoing longitudinal study of ageing,<sup>22</sup> one from the ongoing ADNI study<sup>33</sup> and one from a clinical trial population.<sup>35</sup> The other three studies used representative population samples from Australia<sup>34</sup> and the USA.<sup>31,32</sup> However, Mainous and colleagues<sup>32</sup> ( $n = 6427$ ) selected a cohort aged from 40 years upwards, thus potentially minimizing the potential to observe meaningful cognitive change. Four of the studies reported on populations with a mean age over 60 years, that is those more likely to manifest cognitive change,<sup>31, 33–35</sup> and recruited large numbers (NHANES 60  $n = 4639$ , PATH  $N = 1354$ , FACIT  $n = 800$ ), with the exception of the ADNI study ( $n = 43$ ). None focused solely on older adults and one, the Daugherty imaging study, included a wide age range from 19 to 77 years ( $n = 78$ ), which would have included a varied population; in particular, pre- and postmenopausal women.<sup>22</sup> Furthermore, full details of statistical methods and adjustment for potential confounders was limited and inconsistent, and three studies lacked adequate adjustment for the confounding variables likely to influence cognitive function.<sup>22,31,32</sup>

The measurement of iron was similarly varied. None of the studies attempted to take account of prior iron exposure and iron status at follow up was assessed only in the FACIT trial<sup>35</sup> and the Detroit Aging study.<sup>22</sup> Exposure measures were selective, with no overlap and no combined dietary and biomarker assessments. Only two studies measured brain iron levels.<sup>22,33</sup>

Measurement of cognitive function was also limited, both in breadth and sensitivity. Outcome assessments were selected from standard test batteries for four of the studies. <sup>33–36</sup> The two studies using the NHANES database collected outcome data from death certificates<sup>31,32</sup> and healthcare records,<sup>32</sup> thus raising the possibility of misclassification and under reporting due to missed cases in the community, lack of high-quality diagnoses in nursing home environments and less than rigorous death certificate coding.

The interpretation of study results is also limited by the varying research questions and statistical focus in the different studies. For example, the publication by Mainous and colleagues aimed to look at cholesterol in combination with transferrin rather than using iron measures alone,<sup>32</sup> and the publication by Min and colleagues focused on haemoglobin, folate and vitamin B12, reporting on iron only in passing as a covariate and as part of the baseline

**Table 1.** Study characteristics.

Author and study	Design	Participants and setting	Baseline n	Follow up n	Follow-up years	Reason losses to follow up	Baseline age or mean (SD)	Baseline% female
Mainous <i>et al.</i> <sup>32</sup> NHANES 40	Longitudinal cohort study	The National Health and Nutritional Examination Survey (NHANES survey): USA						
Schiepers <i>et al.</i> <sup>35</sup> FACIT	Longitudinal cohort study	The Folic Acid and Carotid Intima-Media Thickness (FACIT) Trial: The Netherlands						
Cherbuin <i>et al.</i> <sup>34</sup> PATH	Longitudinal cohort study	The PATH through Life study: Canberra, Australia						
Daugherty <i>et al.</i> <sup>22</sup> DETROIT	Longitudinal cohort study	Detroit cohort study: Detroit, USA						
Min <i>et al.</i> <sup>31</sup> NHANES 60	Longitudinal cohort study	NHANES survey linked to mortality data (1999–2006): USA						
Ayton <i>et al.</i> <sup>33</sup> ADNI	Longitudinal cohort study	The Alzheimer's Disease Neuroimaging Initiative (ADNI) cohort: USA						

6558 6427 18 No 40+ ~ 45–63%  
across  
measures

818 800 3 Yes Mean  
60.3 (5.6)  
28.4%

2096 1354 8 Yes 60–64  
Mean  
62.54  
(1.53)  
51.7%

125 78 2 Yes 19–77  
 Mean  
 52.53  
 (14.91)  
 70%

4688 4639 5–12 N/A: loss  
 due to  
 mortality  
 60–89 51%

90 43 7 No Mean  
 75.7 (5.5)  
 51%  
 N/A, not applicable.

**Table 2.** Measurements of iron.

Author	Peripheral or brain iron measures	Measure of iron at baseline	Measure of iron at follow up
Mainous <i>et al.</i> <sup>32</sup> NHANES 40	Peripheral	Serum transferrin	N/A
Schiepers <i>et al.</i> <sup>35</sup> FACIT	Peripheral	Total serum iron, serum concentration of transferrin*, Transferrin saturation\$, serum ferritin‡, nontransferrin-bound iron§	Same as baseline with the exception of nontransferrin-bound iron

Cherbuin *et al.*<sup>34</sup>

PATH

Peripheral Dietary iron calculated from the Commonwealth Scientific Industrial Research Organisation (CSIRO FFQ)

N/A

Daugherty *et al.*<sup>22</sup>

DETROIT

Brain *In vivo* estimation with MRI adds regions of interest hippocampus, caudate and putamen

*In vivo* estimation with MRI

Min *et al.*<sup>31</sup>

NHANES 60

Peripheral Dietary iron (calculated from 24 h recall) N/A

Ayton *et al.*<sup>33</sup>

ADNI

Brain Ferritin in cerebrospinal fluid N/A

\*Transferrin: iron transport protein.

‡Transferrin saturation: the ratio of serum iron to serum concentration of transferrin.

‡Ferritin: indicator of total body iron stores.

§Nontransferrin-bound iron: free iron in the circulation not bound to transferrin.

MRI, magnetic resonance imaging; N/A, not applicable; FFQ, food frequency questionnaire.

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characteristics.<sup>31</sup> None of the studies focused on a potential U-shaped relationship with iron and cognitive outcome. Participant numbers, and subsequently the numbers per cell in analyses were also particularly small in some studies; in the study by Min and colleagues only 49 participants from 4688 had died from Alzheimer's disease at follow up.<sup>31</sup> Daugherty included only 125 participants<sup>22</sup> and in the study by Ayton and colleagues there were only 90 participants

in the cognitively normal group.<sup>33</sup> Finally, participant dropout and missing data<sup>31,34</sup> further limit the interpretation of the results.

### Discussion

To our knowledge, this is the first systematic review of the evidence relating to iron, cognitive decline and dementia. Overall, the available studies were inconsistent in terms of length of follow up, composition of study population, percentage female or male, age of study sample, iron assessment, outcome measure and results. Two studies assessed the influence of dietary iron in relatively similar age groups, one (the USA NHANES 60-plus cohort<sup>31</sup>) found no association with dementia. The other (the Australian PATH study<sup>34</sup>) found an increased risk of cognitive

**Table 3.** Measurement of cognitive function.

#### Author Baseline assessment cognitive function

#### Follow-up assessment cognitive function

Mainous *et al.*<sup>32</sup>

NHANES 40

N/A N/A

Schiepers *et al.*<sup>35</sup>

FACIT

Five cognitive domains assessed. Domains constructed by using either the average of individual test z scores or for two constructs, single tests only

Memory (visual word learning test; immediate recall, maximal immediate recall, delayed recall)

Sensorimotor speed (concept shifting tests 'empty', 'letters', 'numbers', colour Stroop, 'reading words')

Complex speed (concept shifting, 'letters & numbers', colour Stroop, 'naming ink')

Information processing speed (letter digit substitution test)

Verbal fluency (verbal fluency test)

Five cognitive domains assessed. Domains constructed by using either the average of individual test z scores or for two constructs, single tests only

Memory (visual word learning test; immediate recall, maximal immediate recall, delayed recall)

Sensorimotor speed (concept shifting tests, 'empty', 'letters', 'numbers', colour Stroop, 'reading words')

Complex speed (concept shifting, 'letters & numbers', colour Stroop, 'naming ink')

Information processing speed (letter digit substitution test)

Verbal fluency (verbal fluency test)

Cherbuin *et al.*<sup>34</sup>

PATH

Participants were selected for clinical assessment if they had any of the following: a MMSE cutoff  $\leq 25$ , a score below 5th percentile on the CVLT, on SDMT or the Purdue pegboard with both hands or RT

Clinical assessment included neuropsychological testing using the following: trials A&B; clock drawing; Boston naming task; constructional praxis (CERAD); the RAVLT, recall of constructional praxis for nonverbal memory

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Daugherty *et al.*<sup>22</sup>

DETROIT

Latent cognitive constructs were tested for: working memory (listening span, verbal N-back task), nonverbal working memory (spatial recall, visual N-back tasks), episodic memory (logical memory subset of the Wechsler memory scales revised)

Latent cognitive constructs were tested for: working memory (listening span, verbal N-back task), nonverbal working memory (spatial recall, visual N-back tasks), episodic memory (logical memory subset of the Wechsler memory scales revised)

Min *et al.*<sup>31</sup>

NHANES 60

N/A N/A

Ayton *et al.*<sup>33</sup>

ADNI

The ADAS Cog, RAVLT ADAS Cog, RAVLT

ADAS Cog, Alzheimer's disease Assessment Scale; CERAD, Consortium to Establish a Registry for Alzheimer's disease; CVLT, California Verbal

Learning Test; N/A, not applicable; RAVLT, Rey Auditory Verbal Learning Test; RT, reaction time; SDMT, Digit Symbol Modalities Test;

MMSE, mini-mental state exam.

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**Table 4.** Results reported by included studies.

**Author** **N** **Results** **Additional comment** **Confounders**

Mainous *et al.*<sup>32</sup>

NHANES 40

6427 Unadjusted (transferrin alone) transferrin

>75% compared with <75%

Dementia HR 1.11 (95% CI 0.79–1.56)

AD HR 1.47 (95% CI 0.82–2.63)

High cholesterol and high transferrin saturation levels consistently had the highest risk of developing AD compared with the reference group (low cholesterol and low transferrin). Having only high cholesterol or only high transferrin did not increase risk for AD. Unadjusted relative risk of developing AD when both TS and cholesterol were at the 75th percentile was 3.19 (95% CI 1.31–7.75). In adjusted models when only one marker was elevated, there was no significant increased risk for AD. The risk of AD increased as both markers increased

Age, sex, race, education, BMI, exercise, diabetes, hypertension, vitamin

C level

Schiepers *et al.*<sup>35</sup>

FACIT

800 When analyses were stratified by blood donor status, higher serum iron predicted less decline in nondonors in sensorimotor speed (parameter estimate = 0.005,  $p = 0.010$ ) in word fluency (parameter estimate = 0.006,  $p = 0.012$ )

Cognitive performance declined over 3 years for sensorimotor speed, complex speed and information processing. Memory improved and word fluency did not

change. Ferritin concentrations increased significantly but serum iron decreased, although not significantly. No significant longitudinal associations were found between any of the iron parameters and cognitive functioning. The *HFEC28Y* mutation (produces elevated iron levels) was not associated with cross-sectional or longitudinal cognitive performance, indicating lifelong exposure to elevated iron does not affect cognitive function in later life. The potential effect modification by treatment with folic acid in the trial was assessed prior to longitudinal analyses and no interaction between folic acid supplementation and iron status was found. All subsequent analyses were adjusted for covariates

Age, sex, education (formal schooling), alcohol consumption, current smoking, BMI, physical activity, apolipoprotein E, genotype C-reactive protein, haemoglobin, blood donor status, group status (for folic acid treatment)

Cherbuin *et al.*<sup>34</sup>

PATH

1354 For each 1 mg increase in iron at baseline, risk of MCI but not MCD increased over 8 years of follow up

For transition from normal cognition to MCI, HR 1.54 (95% CI 1.03–2.29),  $p = 0.03$

For transition from normal cognition to MCD, HR 1.10 (95% CI 0.87–1.37),  $p = 0.418$ .

Unadjusted

For each 1 mg increase in iron at baseline the risk of MCI but not MCD increased over 8 years

For transition from normal cognition to MCI, HR 1.50 (95% CI 1.07–2.10),  $p = 0.019$

For transition from normal cognition to MCD, HR 1.05 (95% CI 0.95–1.16),  $p = 0.309$

Trend towards sex interaction such that higher iron intake associated with decreased risk of MCI incidence in women, HR 0.81 (95% CI 0.61–1.07),  $p = 0.144$ ) and increased risk of MCI incidence in men, HR 1.08 (95% CI 0.94–1.18),  $p = 0.287$

age, sex education, alcohol intake, smoking, BMI, stroke, diabetes, hypertension (including medication use) *APOE*

£ 4status depressive symptoms physical activity

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**Author N/Results Additional comment Confounders**

Daugherty *et al.*<sup>22</sup>

DETROIT

78 Higher baseline iron in the caudate nucleus was associated with less improvement in working memory,  $\beta -0.18$  ( $p = 0.01$ )

No association between putamen or hippocampal baseline iron or change in iron with working memory. No association

between brain iron at baseline or change in brain iron and episodic memory

Age and sex retained in final models

Min *et al.*<sup>31</sup>

NHANES 60

4639 Increased risk of AD mortality found for those with low haemoglobin (HR 8.4, 95% CI 1.4–50.8)

There were 49 AD-related deaths at follow up. No relationship between AD death and dietary iron intake, but those with AD mortality had significantly lower haemoglobin levels. Those who had AD had lower haemoglobin (14.0g/dl compared with 14.1g/dl,  $p = 0.0089$  *t* test).

Significant reduction in AD mortality risk with increases in haemoglobin for those with low levels of both folate and vitamin B12

Age, sex, ethnicity, education, smoking history, BMI, diabetes or hypertension, dietary intake of iron

Ayton *et al.*<sup>33</sup>

ADNI

43 Baseline ferritin was associated with cognitive decline in a three-way interaction with time and presence of APOE  $\epsilon$ 4

ADAS Cog  $\beta$  0.11 (SE 0.04),  $p = 0.01$

RAVLT  $\beta$  -1.58 (SE 0.54),  $p = 0.004$

Examined by presence of APOE  $\epsilon$ 4

APOE  $\epsilon$ 4 positive:

ADAS Cog  $\beta$  0.09 (SE 0.04),  $p = 0.02$

RAVLT  $\beta$  -1.49 (SE 0.4),  $p < 0.001$

APOE  $\epsilon$ 4 negative:

ADAS Cog  $\beta$  -0.04 (SE 0.016),  $p = 0.02$

RAVLT NS

All models included sex and years of education. The covariates CSF APOE, factor H and haemoglobin were not predictive in any models. No unadjusted results. Models initially included all covariates before minimal models were obtained empirically *via* empirical testing of model fit

Age, sex, BMI, education, CSF tau/amyloid  $\beta$ , CSF APOE, CSF ferritin, CSF factor H (a measure of inflammation)

AD, Alzheimer's disease; ADAS Cog, Alzheimer's disease Assessment Scale; APOE, apolipoprotein E; BMI, body mass index; CI, confidence interval; CSF, cerebrospinal fluid; HR, hazard ratio; NS, nonsignificant; RAVLT, RAVLT, Rey Auditory Verbal Learning Test; SE, standard error; MCD, mild cognitive disorders; MCI, mild cognitive impairment.

**Table 4.** (Continued)

**Table 5.** Assessment of study quality.

**Author**

**Recruitment bias**

**Exposure bias**

**Bias in outcome measurement**

**Bias in assessment of confounders**

**Bias in follow up, patient loss**

**Bias in follow up, length**

**Overall quality assessment**

Mainous

*et al.*<sup>32</sup>

NHANES

40

Moderate. NHANES was designed to be representative of the US population with additional oversampling of key population subgroups including those aged 65 and over. However, baseline included those in midlife (40 years+) which limits opportunity to detect association between exposure and dementia incidence over 20 years

Moderate. Baseline measures only and longer term exposure not considered

Moderate. Dementia/AD measured through health, hospital and nursing home records plus death certificates.

Misclassification possible and cases living in the community may have been overlooked

Moderate. Multiple key confounders are included but assessment methods were not noted except for vitamin C usual intake not necessarily captured by 24 h dietary history

Low. 98% follow-up rate for NHANES.

Follow-up data collected by interview or proxy if participant unable to respond or deceased

Moderate. Potential for some participants to have contributed fairly short



					<b>loss</b>		
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Cherbuin

*et al.*<sup>34</sup>

PATH

Low. Recruitment of a random selection from the electoral role membership of which is mandatory in Australia

Moderate. Baseline measures only and longer term exposure not considered and those with missing data were excluded. Iron status assessed through self-report FFQ which does not have the reliability of serum markers for iron status

Moderate.

Neuropsychological battery and clinical assessment of dementia used to assess cognitive function but those with missing data were excluded

Moderate. Key confounders are included but those with missing data were excluded. BMI was calculated using self-report height and weight

Moderate.

There were 2551 recruited and 1275 included after exclusion of missing data etc.

Moderate.

Follow up is sufficient to detect incident change,

however exclusion of those with missing data may have caused bias

Moderate

Daugherty

*et al.*<sup>22</sup>

DETROIT

Moderate.

Recruitment was part of an ongoing longitudinal study of ageing  
 Moderate. Baseline and follow-up measures are used but longer term exposure not considered  
 Moderate. Working and episodic memory measures were taken from a larger neuropsychological battery assessment.  
 No theoretical justification was given as to why these measures were chosen  
 Moderate A selection of key confounding variables were included. In particular, time-varying confounders were measured by objective blood markers and included in the model as a latent metabolic factor, thereby excluding variable-specific error  
 High; ~37% of sample lost to follow up  
 Moderate.  
 Reverse causality possible in 2-year follow up, however reverse effects models were empirically tested  
 Moderate

**Table 5.** (Continued)  
 (Continued)

Author	Recruitment bias	Exposure bias	Bias in outcome measurement	Bias in assessment of confounders	Bias in follow up, patient loss	Bias in follow up, length	Overall quality assessment

Min *et al.*<sup>31</sup>  
 NHANES  
 60  
 Low. NHANES was designed to be representative of

the US population with additional oversampling of key population subgroups including those aged 65 and over

Moderate. Baseline measures only and longer term exposure not considered.

Measurement methods for dietary iron not clear

Moderate. Death certificate data may not be accurate or complete. At follow up those without AD mortality may nonetheless have had AD

High. Insufficient details of analysis methods for iron to allow evaluation. Small cell sizes for each iron quartiles in subjects with AD mortality

Moderate. Those without measures of haemoglobin, or red-cell folate/B12 and those with any other missing data were excluded

Moderate. Potential for some participants to have contributed fairly short follow up, meaning reverse causality cannot be excluded

Moderate

Ayton *et al.*<sup>33</sup>

ADNI

Moderate. Selection was from the mild cognitive impairment and control groups of the Alzheimer's Disease Neuroimaging Initiative (ADNI) data

sets. ADNI recruited from multiple sites across North America. Moderate. Baseline measures only and longer term exposure not considered. Moderate. A standard neuropsychological test and cognitive screening test were used and data collected annually. However, the ADAS Cog (screening test) may not be a sensitive measure of cognitive change in those who are cognitively normal. Some bias may also have been possible as participant data were included until death or loss to follow up.

Low. Key confounders including a measure of inflammation. Moderate. However, all participants were included until loss.

Moderate. Some participants left the study prior to the end of the 7-year follow up, however overall follow up was long enough to measure change in cognitive function. Moderate.

AD, Alzheimer's disease; ADAS Cog, Alzheimer's disease Assessment Scale; BMI, body mass index; NHANES, National Health and Nutrition Examination Survey; FFQ, food frequency questionnaire.

**Table 5.** (Continued)

decline associated with greater iron intake, possibly more relevant in men.

Several of the studies did report a stronger relationship in a particular subgroup, although the subgroups also varied. The older NHANES cohort found a relationship between higher transferrin and increased risk of Alzheimer's disease in

those with higher cholesterol.<sup>32</sup> In the ADNI cohort the relationship between higher cerebrospinal ferritin and cognitive decline was strongest in those who were also *APOE*  $\epsilon$ 4 positive,<sup>33</sup> and in the FACIT trial cohort, higher serum iron levels were associated with less decline in sensorimotor speed and word fluency in those who did not donate blood.<sup>35</sup>

A recent comprehensive review of iron deficiency and risk of cardiovascular disease included 13 longitudinal studies of serum iron, 15 studies of ferritin and 14 of transferrin saturation and total iron binding capacity, but also observed heterogeneity and inconclusive findings across studies. Although they concluded that 'extreme conditions of iron deficiency as well as iron overload were associated with increased cardiovascular disease risk'.<sup>37</sup>

Limitations associated with our review and its conclusions stem mostly from the paucity of studies rather than the methods used for the review process. It is possible that in restricting the language to English and in excluding study populations with serious clinical comorbidities, potentially relevant evidence has been overlooked. However, including populations with specific comorbidities where the comorbidity itself may influence iron metabolism, risk of mortality and risk of cognitive decline would have been to address a different research question. The lack of evidence, and the varying study designs and populations used within the available evidence base, present a more serious limitation.

#### **Limitations in the assessment of iron**

None of the studies used a fully comprehensive assessment of iron status. This is important because cerebrospinal fluid and blood reporters of iron only have a modest association,<sup>29</sup> blood iron markers do not predict MRI measured brain iron content<sup>38</sup> and, in older adults in particular, levels of the peripheral iron reporters, ferritin and haemoglobin, may also signify peripheral inflammation such as anaemia of chronic inflammation,<sup>39</sup> or indeed noniron-deficient anaemia. In the case of the latter, tissue deposition of iron may occur even when presenting with low iron markers in the blood.

For example, the iron-trafficking disease, aceruloplasminemia, results in tissue iron deposition and lower blood iron markers because iron cannot efficiently be exported out of the cell.<sup>40</sup> If cellular iron trafficking, rather than iron exposure, is the lesion that results in brain iron elevation in Alzheimer's disease, it is possible that depletion of iron in the blood accompanies iron elevation in tissues. Furthermore, only two of the studies assessed dietary iron and no studies evaluated the impact of iron supplementation. No studies directly followed a potential link between iron intake and iron blood or neuroimaging biomarkers and subsequent cognitive function, and none attempted to evaluate iron impact by baseline iron level or by clinically relevant categories such as iron deficiency anaemia or iron deficiency no anaemia. We were unable to identify any trials of dietary iron exposure, or of iron supplementation, in normal or iron-deficient adults in mid or late life and risk of cognitive decline.

### **Limitations in the assessment of outcome**

Cognition is complex and multifaceted and it is possible that iron status differentially affects change or decline across the spectrum of ability domains. Clearly, in the cognitively healthy a more comprehensive and psychometrically driven approach to assessing cognitive change is necessary if the impact of iron status on cognitive decline is to be more fully understood. The use of medical and death certificate records for dementia diagnoses in the NHANES analyses presents a further limitation since their accuracy depends upon robust and comprehensive assessment and record keeping.

### **Gaps in the evidence base and future work**

Our systematic review of the current evidence base relating to iron and cognitive function and dementia found a sparse and limited field of evidence.

Much more work is needed to evaluate the links between iron intake, brain deposition, ageing and cognitive function over time. In particular, a detailed focus is needed on regional brain iron and related cognitive abilities with multiple follow ups, MRI measures or historical health measures taking closer account of potential causal and associative relationships, including key confounding measures. Obvious examples include ensuring that analyses with ferritin (also an acute phase protein) are, at minimum, adjusted for other inflammatory markers, and taking account of obesity and its association with iron dysregulation.

Multimodal approaches should also be used, such as using CSF iron biomarkers in combination with MRI measures of brain iron. Larger, longer and more thorough epidemiological studies are also needed to fully explore the potential pathways as is a greater emphasis on collection of biomarkers to allow more accurate evaluation of the homeostatic iron picture in its entirety, taking account of the varied iron types and storage mechanisms.

This would also be an important part of the safety considerations prior to conducting any randomized clinical trials of iron supplementation or iron chelation. In particular, we recommend that future studies in this area focus on reducing risk of bias derived from varied participant populations; that is, select those of a particular age range and stratify by sex. Studies should collect at least two measures of cognitive function over a minimum follow up of 12 months to allow assessment of change; when possible, two matching iron measures should be taken at the same interval, ideally using imaging and linking region of interest to the cognitive domain under assessment. This should be supplemented by peripheral measures, including inflammatory markers; when possible, additional data on usual diet, and clinical and cardiovascular history should be obtained.

### **Conclusion**

The current evidence base is sparse and limited. The diversity of the existing evidence limits interpretation and applicability by either healthcare providers or researchers and precludes any conclusion relating to the relationship between iron and cognition. Much more work is needed before we can develop our understanding of the role of iron in cognitive function.

## Author contributions

RP: study concept; RP, DH, AB: study design; RP, DH, SA: data analysis; RP, DH, SA, NB: data interpretation. All authors contributed to manuscript preparation and revision, and approved the final version. No sponsor had any role in the design, methods, subject recruitment, data collection, analysis or preparation of the paper.

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## Conflict of interest statement

The authors declare that there is no conflict of interest.

## Supplemental Material

Supplemental material for this article is available online.

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