Should HFE p.C282Y homozygotes with moderately elevated serum ferritin be treated? A randomised controlled trial comparing iron reduction with sham treatment (Mi-iron)

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To cite: Ong SY, Dolling L, Dixon JL, et al. Should HFE p.C282Y homozygotes with moderately elevated serum ferritin be treated? A randomised controlled trial comparing iron reduction with sham treatment (Mi-iron). BMJ Open 2015;5:e008938. doi:10.1136/bmjopen-2015-008938

ABSTRACT

Introduction: HFE p.C282Y homozygosity is the most common cause of hereditary haemochromatosis. There is currently insufficient evidence to assess whether non-specific symptoms or hepatic injury in homozygotes with moderately elevated iron defined as a serum ferritin (SF) of 300–1000 µg/L are related to iron overload. As such the evidence for intervention in this group is lacking. We present here methods for a study that aims to evaluate whether non-specific symptoms and hepatic fibrosis markers improve with short-term normalisation of SF in p.C282Y homozygotes with moderate elevation of SF.

Methods and analysis: Mi-iron is a prospective, multicentre, randomised patient-blinded trial conducted in three centres in Victoria and Queensland, Australia. Participants who are HFE p.C282Y homozygotes with SF levels between 300 and 1000 µg/L are recruited and randomised to either the treatment group or to the sham treatment group. Those in the treatment group have normalisation of SF by 3-weekly erythrocytapheresis while those in the sham treatment group have 3-weekly plasmapheresis and thus do not have normalisation of SF. Patients are blinded to all procedures. All outcome measures are administered prior to and following the course of treatment/sham treatment. Patient reported outcome measures are the Modified Fatigue Impact Scale (MFIS—primary outcome), Hospital Anxiety and Depression Scale (HADS), Medical Outcomes Study 36-item short form (SF36v2) and Arthritis Impact Measurement Scale 2 short form (AIMS2-SF). Liver injury and hepatic fibrosis are assessed with transient elastography (TE), Fibrometer and Hepascore, while oxidative stress is assessed by measurement of urine and serum F2-isoprostanes.

Ethics and dissemination: This study has been approved by the Human Research Ethics Committees of Austin Health, Royal Melbourne Hospital and Royal Brisbane and Women’s Hospital. Study findings will be disseminated through peer-reviewed publications and conference presentations.

Trial registration: Trial identifier: NCT01631708; Registry: ClinicalTrials.gov

INTRODUCTION

Hereditary haemochromatosis (HH), an iron overload disorder, is the most common genetic condition in Northern Europeans among whom approximately 1 in 200 are homozygous for the HFE p.C282Y amino acid substitution, the cause of most HH. Not all p.C282Y homozygotes develop morbidity. About 20% of males and 40% of females with this genotype do not develop iron overload. At least 28% of males and 1% of females develop iron overload-related disease such as liver cirrhosis, diabetes mellitus and cardiomyopathy. These individuals generally have severe iron overload as indicated by a serum ferritin (SF) of greater than 1000 µg/L. Therefore, the question arises as to whether the largest group of p.C282Y homozygotes, those with raised SF but SF less than 1000 µg/L (defined here as moderate iron overload), require venesection treatment to normalise SF. Answering this question is important since if treatment is beneficial, introduction of community screening for HH should be considered whereas if there is no benefit from treatment, p.C282Y homozygotes with moderate iron overload do not need to undertake this somewhat onerous intervention.
Major management guidelines for HH recommend treatment of those with HH and SF above the upper limit of normal and recommend that SF be reduced to 50–100 µg/L. The rationale for normalisation of SF in those with severely elevated SF (>1000 µg/L) is clear as severe morbidity and mortality can be prevented. There is little evidence that treatment is beneficial in those with moderate iron overload however. There have been no randomised controlled trials to objectively assess whether returning iron levels to normal improves symptoms in p.C282Y homozygotes with moderately elevated SF. Reasons why such studies have not been carried out likely include the fact that treatment of HH is relatively safe and therefore commentators have adopted the stance that treatment is unlikely to result in harm while there are theoretical reasons why harm may result from not normalising SF. In addition, blinding is far more complex in a trial of venesection than in a placebo-controlled pharmaceutical trial.

HFE p.C282Y homozygotes with moderately elevated SF do not have an increase in frequency of HH-related symptoms when compared to controls in cross-sectional studies. Such cross-sectional studies are not designed to identify subtle symptoms such as fatigue, however. Anecdotal evidence suggests that fatigue benefits from treatment of HH. Fatigue is a non-specific symptom that is commonly reported in individuals with HH and has a negative impact on the physical and psychological quality of life. There are conflicting data about the relationship between fatigue and levels of excess iron. Adams et al found an association between fatigue and hepatic iron index in 410 p.C282Y homozygotes. In contrast, fatigue has been reported in p.C282Y homozygotes with normal SF and has been found to be worse in some individuals following normalisation of SF by venesection treatment. One population study identified significantly higher Modified Fatigue Impact Scale (MFIS) scores in p.C282Y homozygotes who knew their diagnosis and had their iron levels normalised compared to those who were unaware of the diagnosis and had significantly higher SF. There are three possible explanations for these observations: (1) There is no relationship between fatigue and iron levels in HH; (2) there is a significant psychosomatic effect of diagnosis on how these individuals perceive fatigue; or (3) there is a subgroup of individuals who have fatigue and are more likely to be diagnosed with HH.

A recent study suggested that treated HFE p.C282Y homozygotes with moderate iron overload have decreased cardiovascular and extrahepatic cancer-related mortality rates compared to the general population, while p.C282Y homozygotes with normal SF have the same mortality rates as the general population. The relationship between the mortality findings and treatment of iron overload was questioned, however.

AIMS AND HYPOTHESIS
The aim of this multicentre randomised trial is to compare the prevalence of symptoms and objective markers of disease between those in the treatment group and those in the sham treatment group.

Our hypothesis is that HFE p.C282Y homozygotes with moderately elevated SF will have few symptoms and signs of disease and decreasing SF to normal levels will not result in a greater change in patient reported outcomes or objective markers of liver injury or hepatic fibrosis compared to those whose SF levels are not lowered to the normal range.

METHODS AND ANALYSIS
The Mi-iron (Moderately increased iron levels) study is a multicentre, randomised single-blinded trial being conducted in Victoria (Austin Hospital and the Royal Melbourne Hospital) and Queensland (Royal Brisbane and Women’s Hospital) that started in August 2012 and is due to conclude in December 2015. Figure 1 summarises the methodology of the Mi-iron Study.

Participants
Inclusion criteria
1. Age 18–70 years inclusive
2. HFE p.C282Y homozygous
3. SF between 300 and 1000 µg/L
4. Previously or currently raised TS
Exclusion criteria
1. HH due to other genotypes
2. Venesection in the past 2 years for treatment of HH
3. Other risk factor(s) for liver injury including hepatitis B (HBV) or C (HCV), excess alcohol consumption (>60 g/day in males, 40 g/day in females), body mass index (BMI) of ≥35 kg/m²
4. Pregnant females

Study intervention
Apheresis is being used as the study intervention. All procedures are conducted using the Haemonetics MCS Plus apheresis system.

Randomisation and stratification
Participants are randomised to either the treatment group to have erythrocytapheresis or the sham treatment group to have plasmapheresis. Randomisation is by computer generated random number sequence. Randomisation is stratified by gender, initial SF (300–599 µg/L or 600–1000 µg/L as a binary variable) and site.

Maintenance of blinding
The participant is blinded to which arm of the study she/she has been randomised by being connected to the apheresis machine with the machine and the tubing not visible to the individual. This is achieved by the participant’s arm being passed through an opaque black curtain (figure 2). Thus participants are unaware of whether they are having red blood cells (RBCs) or plasma removed and are unaware of whether or not they
are having their iron levels reduced. Staff performing
the apheresis are trained to not inadvertently reveal the
treatment arm of the participant through strict adher-
ence to study protocol and careful use of language in
describing what is being performed. A member of the
research team is present during the intervention to
ensure blinding is maintained by monitoring the pro-
cEDURE and conversation between apheresis staff with
the participant.

Those undergoing plasmapheresis have the same
volume of plasma removed as the volume of RBCs
removed had they been randomised to the erythrocyta-
pheresis arm. Therefore, the risk of symptoms due to
reduction in circulating blood volume is the same for
individuals in both arms of the study.

Intervention
Treatments are administered every 3 weeks. The volume
of RBCs/plasma removed is individualised based on the
individual’s blood volume and haematocrit. Haematocrit
is measured at the start of each erythrocytapheresis
treatment while in the sham treatment group, mock
blood tests are taken to ensure the participant’s experi-
ence is identical irrespective of the treatment group to
which they have been randomised. The volume removed
is calculated based on the height, weight, pretreatment
haematocrit and the target haematocrit (30–35%; figure
3). Treatments are ceased in the treatment group when
SF is below 300 µg/L. For the sham group, the calcu-
lated number of treatments is equivalent to as if their SF
was normalised in the RBC group.

Participants in the sham arm are offered erythrocyta-
pheresis or phlebotomy to normalise SF at their choice
of venue on completion of the study.

Safety blood test monitoring
All participants have SF measured 2–3 weeks after the
expected last treatment to ensure the SF level has
decreased to the normal range (<300 µg/L) for those in
the treatment group before proceeding to the end of
trial assessment. Participants in both arms of the study
have the same blood test to ensure blinding. Serum B12,
Folate, iron studies and full-blood count are checked approximately mid-treatment in both groups for safety monitoring.

**Target SF**

There have been no definitive studies conducted to demonstrate what the target SF should be at the end of treatment in an individual with HH. Guidelines recommend a target SF of less than 50–100 μg/L. In this study, we have chosen to reduce SF to anywhere in the normal range, that is, a SF of 20–300 μg/L, based on the expectation that an individual with HH whose SF is in the normal range should have similar symptoms to those without HH and who have a normal SF.

**Outcome measures**

The outcome measures being assessed are patient-reported outcome scales to assess symptoms, as well as markers of liver injury, hepatic fibrosis and oxidative stress. These are administered at baseline and the end of erythrocytapheresis/sham apheresis treatment.

**Patient-reported outcome scales**

1. **Modified Fatigue Impact Scale (MFIS)** is the primary outcome measure for this study. The MFIS is a 21-item scale that measures the impact of fatigue on three independent subscales of physical, cognitive and psychosocial functioning. Participants rate their fatigue in the past month on a five-point Likert-type scale. The total score ranges from 0 to 84 with higher scores reflecting greater fatigue. Subscale scores, physical (0–36); cognitive (0–40); and psychosocial (0–8), can also be derived.

2. **Medical Outcomes Study 36-item short form V.2 (SF36v2)** is a 36-item generic health survey to measure health and well-being that has been previously used in various HH studies to measure quality of life. It assesses eight different health components (physical functioning, role-physical, bodily pain, general health, vitality, social functioning, role-emotional, mental health) and provides a summary score for both physical and mental components. It is a norm-based scoring system and thus can be used to compare participant scores to the general population.

3. **Hospital Anxiety and Depression Scale (HADS)** is composed of a 14-item total scale (HADS-T) consisting of two seven-item independent subscales, the Anxiety (HADS-A) and Depression (HADS-D) subscales. Participants rate how they have felt in the past week on a four-point Likert-type scale. Scores on each scale can be interpreted in ranges: normal (0–7); mild (8–10); moderate (11–14); and severe (15–21). Higher scores on each subscale or the entire scale indicate...
greater anxiety, depression or both. This scale has been found to be valid and reliable in various populations.23 24

4. Arthritis Impact Measurement Scales 2 short form (AIMS2-SF) is a 26-item validated scale that assesses the impact of arthritis on five core domains of the participants.25 26 It measures physical functioning, symptoms, affect, role and social interactions of the individuals. A five-point Likert-type scale is used to rate how participants have felt in the past month. The higher the raw score, the greater the impact of arthritis on the participant. Use of arthritis medication at baseline and end of trial will also be compared.

5. To assess the fidelity of the blinding process, the participants are asked which treatment group they believe they were allocated to at the completion of the study, before unblinding.

Liver injury and hepatic fibrosis markers

Transient elastography (TE) and blood tests for components of Hepascore and Fibrometer3G V are collected from individuals at baseline and end of the trial.

Transient elastography

Fibroscan is a method of TE that evaluates liver stiffness using an ultrasound probe to measure the velocity of a mechanical wave that is pulsed through the liver. As the liver becomes progressively more fibrotic, it becomes harder and less elastic. The velocity of the wave correlates directly with tissue stiffness and results are reported in kilopascals (kPa).27 TE has been evaluated in a number of different liver diseases, including HBV and HCV, alcoholic liver disease, non-alcoholic fatty liver disease and HH.28 29 A recent meta-analysis of nine studies involving TE showed excellent results for diagnosing cirrhosis, with a sensitivity of 87% and specificity of 91%.30 Adhoute et al31 have shown that TE measurements correlate with Hepascore measurements in individuals with HH. The results of liver stiffness are acquired from at least 10 successful valid measurements with a success rate of at least 60% within the IQR of ≤30%. A cut-off value of 8.7 kPa was sensitive for the diagnosis of those with significant fibrosis (≥F2), with an area under the curve (AUC) for the receiver operating characteristic (ROC) of 0.7932 and a reading of more than 13 kPa was highly predictive that cirrhosis of the liver was present in a cohort with iron overload due to β-thalassemia.33

Hepascore

Hepascore is derived from an age-specific and gender-specific model that inputs parameters of serum bilirubin, γ glutamyl transferase (GGT), hyaluronic acid and α2-macroglobulin. The test results in a score between 0 and 1 with a higher score being associated with more severe liver disease. In a HCV cohort, Hepascore demonstrated an AUC for the ROC of 0.8 for predicting significant fibrosis (≥F2) and 0.90 for predicting cirrhosis.34 A score >0.5 was found to have a specificity of 70% and sensitivity of 77% to detect significant fibrosis (≥F2) in a large HCV cohort.35 A Hepascore <0.25 can exclude significant fibrosis with a negative predictive value of 0.9.36 In a study that included the Hepascore in HFE related HH, 44 p.C282Y homozygotes had a median score of 0.1.36

Fibrometer3G V

Fibrometer3G V is formulated from the platelet count (PLT), prothrombin index (PI), and the alanine amino transaminase (ALT), aspartate amino transaminase (AST), GGT, α2-macroglobulin and urea levels. This biomarker had an AUROC of 0.85 for predicting significant fibrosis and an AUROC of 0.9 for predicting cirrhosis in a HCV cohort.37 Its robustness has been evaluated in different studies and has been recommended by the French National Authority for Health for the estimation of liver fibrosis in HCV.

The combination of Fibrometer3G V and TE has recently been shown to increase the accuracy of diagnosing significant fibrosis and cirrhosis to 92% compared with Fibrometer3G V (84% accuracy) or TE (88% accuracy) alone. The combination has an AUROC of 0.89, improving the reliability and precision of diagnosis of significant fibrosis in chronic liver disease.38 39 Fibrometer has not been tested in HH.

Oxidative stress marker

Iron is a strong pro-oxidant and there is evidence that markers of oxidative stress are elevated in individuals with elevated iron indices due to HH.40–45 To assess oxidative stress, F2-isoprostanes, a validated marker of cellular lipid oxidative damage,44 are being measured in urine and blood. While elevated markers of oxidative stress are not necessarily related to symptoms of disease, we will be able to assess the relationship between this early marker of tissue injury and the other markers being measured, including iron indices, Hepascore, Fibrometer, TE score and the scores for the various clinical scales being administered. We will also assess whether F2-isoprostanes are positively impacted by normalisation of iron indices in the erythrocytapheresis group.

Sample size calculation

Data from the Melbourne HealthIron study2 were used to calculate SDs of the MFIS score of 14.1 and 17.8 for male and female C282Y homozygotes, respectively. Using a conservative value for the SD of 18, a sample size of 50 in each treatment group ensures an 80% chance (statistical power) that a treatment effect of a mean difference of 10 MFIS units (well above a clinically relevant difference on this scale which runs from 0 to 84) is reflected in a p value less than 0.05. Summary statistics from figure 2 of Adams et al44 show the mean Hepascore changing from 0.20 in patients with META VIR fibrosis grade 0 or 1 (F0 or F1), through 0.45 in those with F2, to close to 1.0 in
those with F3 or F4. The within-fibrosis grade SD of Hepascore is approximately 0.20 in the F2 group and it is much lower in the remaining groups. Using this SD, a sample size of 50 in each treatment group of the trial delivers statistical power of 85% to detect a treatment effect of 0.12 on the Hepascore scale. A change of this magnitude is similar to the observed mean difference in Hepascore between adjacent fibrosis groups presented in Adams et al.34 Accommodating the stratified design, the regression-based statistical analysis will result in minimal loss of power provided that only the average initial measures and not the treatment effects are different between strata.

Statistical analysis
The primary analysis addressing the research hypothesis will be a comparison of the change in scores for all outcome scales, biochemical tests and TE scores, from baseline to end of treatment, between those who have had their iron levels returned to normal and those who were in the sham treatment group, with assignment to the comparison groups based on intention to treat. This analysis will be implemented using a linear regression model of the final measure on each scale including as covariates the value of the initial measure, gender, initial SF (300–599 μg/L or 600–1000 μg/L as a binary variable) and site (Melbourne or Brisbane as a two category variable). In a separate analysis, this model with the Hepascore as the outcome measure will be extended to include the quantity of iron removed (calculated as 1 g iron per litre of RBCs removed by erythrocytapheresis) to determine whether there is an association between the change in Hepascore and the reduction in iron level. Similar analyses will be performed for F2-isoprostanes, TE and patient-reported outcome scales.

CONCLUSION
This is the first randomised controlled trial of treatment for HH. It will demonstrate whether there is any benefit in the short term from normalisation of SF in HFE p. C282Y homozygotes with moderately elevated SF. This has implications for management of this group of individuals and may assist in determining whether introduction of community screening for HH should be considered.

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