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Daily versus alternate day dosing of oral iron supplementation in treatment of iron deficiency anaemia: A systematic review and meta-analysis

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ADMINISTRATIVE INFORMATION

Support - None.

Review Stage at time of this submission - Data analysis.

Conflicts of interest - None declared.

INPLASY registration number: INPLASY2023100046

Amendments - This protocol was registered with the International Platform of Registered Systematic Review and Meta-Analysis Protocols (INPLASY) on 12 October 2023 and was last updated on 12 October 2023.

INTRODUCTION

Review question / Objective Participants: Adult patients with iron deficiency anaemia. Intervention: Alternate day oral iron supplementation. Control: Daily oral iron supplementation. Outcome: Primary outcome measures the change in haemoglobin and ferritin with secondary outcome of incidence of gastrointestinal side effects.

Rationale Iron deficiency anaemia remains the most common cause of anaemia worldwide. Daily supplementation is currently the most common regimen; however, this can be associated with gastrointestinal side effects.

We performed a systematic review and metaanalysis aiming to assess whether the efficacy of alternate daily dosing of iron is comparable to daily dosing in the treatment of iron deficiency anaemia.

Condition being studied Iron deficiency anaemia.

METHODS

Search strategy Electronic search was performed using Medline, EMBASE, and Cochrane Central Register of controlled trials database from inception to 1st June 2023 using the following MeSH terms or free text: "oral iron", "oral ferrous", "dose", "hepcidin".

Participant or population Adult patients age 18 or more, non-pregnant.

Intervention Alternate daily oral iron supplementation.

Comparator Daily oral iron supplementation.

Study designs to be included Randomised control trials.

Eligibility criteria The exclusion criteria were as follows: Studies on pregnant women, on-adult studies, animal studies, non-english studies.

Information sources Electronic databases.

Main outcome(s) Mean change in haemoglobin, mean change in ferritin, risk ratio of gastrointestinal side effects.

Quality assessment / Risk of bias analysis Cochrane risk of bias analysis.

Strategy of data synthesis A random effects model, as described by DerSimonian and Laird, was used to calculate pooled mean difference and risk ratios. Forest plots were used to display the outcomes.l2, Tau2 statistics and p-value were used to assess the heterogeneity.

Subgroup analysis Subgroup analysis looking at follow up time greater than 8 weeks compared to follow up time of less than 8 weeks was performed.

Sensitivity analysis Test of chi with 95% confidence interval.

Country(ies) involved Australia.

Keywords Iron deficiency anaemia; alternate daily; oral iron.

Contributions of each author

Author 1 - Ashley Gaw - Author 1 did data collection, study quality assessment, and drafted manuscript.

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Author 2 - Jin Tan - This author assisted in literature search and article screening, editing manuscript.

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