Original Research Article

Weekly iron supplementation in 2-year-olds is effective in combating anaemia

ABSTRACT

Introduction: Iron deficiency anaemia is a serious public health problem in developing countries, especially among children, as it is associated with serious developmental problems. Objective: To assess the effects of weekly ferrous sulfate supplementation on haemoglobin (Hb) levels and the prevalence of anaemia in children aged 2 to 3 years. Methods: A cluster_cluster-randomized clinical trial was conducted,—itwo schools were randomly chosen,—i. In the first school, the children received 6mg/kg of elemental iron in the form of iron sulfate once a week (intervention),—i. In the other school, the children received a placebo (control). The intervention group had 44 participants at the end of the study, and the control group had 48 children. Blood samples were taken at baseline and at the end of the study to assess serum Hb levels and anaemia prevalence. The intervention lasted 14 weeks. Results: There was a mean increase in Hb of 0.85g/dL (p=0.0003) in the intervention group and a decrease of 0.74g/dL (p=0.0001) in the control group. The prevalence of anaemia significantly decreased in the weekly supplementation group with p=0.0002. Conclusion: Weekly iron supplementation in preschool-age children promoted a significant increase in Hb levels and a decrease in the prevalence of anaemia.

Keywords: Iron-deficiency anaemia; Hemoglobin Haemoglobins; Ferrous sulfate; Preschool child; Clinical trial.

1. INTRODUCTION

The World Health Organization (WHO) defines the term-angemia as a condition in which the serum concentration of haemoglobin is below the reference values, to the point of not meeting the physiological needs according to age, sex, pregnancy and altitude(1). Within eof the causes of angemia, approximately 50% are attributed to a diet deficient in iron, which is considered the most prevalent nutritional deficiency in the world, affecting mainly children under 5-fiveyears of age (preschoolers), women of childbearing age, pregnant and lactating women, in greater numbers in developing countries (2,3).

Iron is the most found metal in the human body, playing a crucial role in all phases of protein synthesis, cellular respiration, <u>and</u> oxidative and immunological processes (4,5,6). Iron deficiency is associated with bone fragility and distortions, hepatosplenomegaly (possibly from extramedullary hematopoiesis), delayed growth and puberty, neurodevelopmental changes, cardiomegaly, and electrocardiographic abnormalities (7). A recent study associated iron deficiency in childhood and adolescence with the increased prevalence of Attention Deficit Hyperactivity Disorder (ADHD), Anxiety Disorder and Bipolar Mood Disorder, highlighting the long-term importance of iron deprivation (8).

In this sense, one of the strategies recommended by the WHO to control iron deficiency is the use of weekly (intermittent) supplementation for risk groups, infants and preschoolers, which worldwide have a prevalence of angemia close to 40% (9,10). Our

study used weekly iron supplementation in public schools as a strategy to try to improve <a href="children's-c

2. METHODOLOGY

2.1. Study design

To address the research purpose, the authors designed and implemented a cluster randomized clinical trial studyhe authors designed and implemented a cluster randomized clinical trial study to address the research purpose. The study sample was derived from the population of preschoolers aged between 24 and 36 months, from public Infant Education Centers, in the municipality of Sobral - Ceará, a middle-sized city, in the northeast of Brazil, between August and December 2019.

Prior to intervention, three public Infant Education Centers were chosen using a table of randomized numbers; the first formed Group A, the second Group B. Group A received 6 mg/kg of elemental iron once weekly (intervention); and Group B was designated as control.

All preschoolers aged 24 to 36 months from the two Infant Education Centers were invited to participate in our study. Exclusion criteria were <u>parents' parents'</u> refusal to participate and infants already using iron supplementation.

2.2. Intervention

The preschoolers in Group A received 6 mg/kg elemental iron once weekly (Mondays); intervention was administered using an individual plastic medical syringe with a scale, previously prepared according to child's weight, to gently squirt the solution into the side of the child's mouth by graduate medical trainees. <a href="mailto:lntervention-The intervention-The intervention-

2.3. Primary outcomes and other variables

The study included <u>2-two</u>primary outcome variables: 1) change in Hb concentration measured in g/dL; and 2) an<u>a</u>emia prevalence before and after the intervention. Hb concentration <11.0g/dL was used as a cutoff point to define anaemia (10).

According to information provided by parents, a standardized data sheet was filled in containing information on (other study variables): age, gender, exclusive breastfeeding (EBF) up to 6 months, mother's mother's schooling, and family income.

2.4. Sample size

According to previous studies conducted in this region, an<u>a</u>emia prevalence in the study population was estimated at 40-50% (11). To <u>achieve a reduction inreduce</u> global an<u>a</u>emia prevalence from 50 to 25%, with 80% power, 2-sided, type I error of 5 %, accounting for 10% losses to follow-up, each group required a minimum of 43 participants (12).

2.5. Data collection

Two biochemical evaluations were performed, to determine Hb concentrations, before and after the intervention. Hb concentrations were premptly analyzed with a portable HemoCue B-hemoglobin-photometer (Hb 301 - HemoCue AB, Ängelholm, Sweden) by technicianA technician promptly analyzed Hb concentrations with a portable HemoCue B-haemoglobin photometer (Hb 301 - HemoCue AB, Ängelholm, Sweden). Finger prick capillary blood was collected under aseptic conditions using Carelet® Safety Lancets (Facet Technologies, Atlanta, GA, USA). Members of the study team who collected outcome data were blinded to the different interventions.

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2.6. Data analyses

To compare means we used, the paired student's student's t-test to assess the difference in Hb concentration before and after the intervention within the groups, and Fisher's exact test to assess the difference between good and bad outcomes (absence or presence of angemia). Data had a normal distribution. The statistical software package SPSS for Windows, version 17.0, was used for all analyses (SPSS Inc., Chicago, IL). The limit for statistical significance was set at p=0.05. Analyses were by intention to treat

This study was approved by the Ethics Committee for Research at the Universidade Federal do Ceará following the ethical principles established by the National Health Council Resolution #466/2012, with necessary prior written consent from school directors and parents/guardians. Medical support was available upon request. After the_intervention, anaemic children were referred for treatment.

3. RESULTS

At baseline, 14 preschoolers were excluded before blood analysis, nine from group A (3 refused and six already using iron supplementation), and five from group B (3 refused and 2-twoalready using iron supplementation) (Figure 1).

Before the second biochemical evaluation (at the end of the intervention), there were <u>10-tendropouts</u> from Group A (5 left Infant Education Center, <u>2-two</u>absentees, <u>3-three</u>non-compliant); in Group B₁ there were 11 dropouts (7 left Infant Education Center, <u>2-two</u>absentees, <u>2-two</u>non-compliant) (Figure 1).

At baseline, Hb concentration and the other study variables were analyzed. There were no statistically significant differences for age, gender, EBF, mother's mother's schooling, and family income. Mean age (in months) for group A was 29.9 ± 3.51 for group A_7 and 30.4 ± 3.36 for group B, p= .30; in group A, 27 participants were male_ and 27 were female,—i._In group B_ 28 were male and 31were female, p= .96. The p-values between groups for EBF, mother's mother's schooling and family income were .96 $_7$ and .90, respectively. However, there was a significant difference between the groups for mean Hb values; the mean Hb for group A was 11.34 ± 1.31 for group A_7 and 11.88 ± 0.78 g/dL for group B, p= .003 (Table 1).

In Group A, the mean baseline Hb concentration was 11.19 ± 1.42 g/dL, and after intervention mean Hb concentration increased to 12.04 ± 0.96 g/dL, p= .0003; and anaemia prevalence was 20 out of 44, 45.5% at baseline, and 4 out of 44 (9.1%) at the end of the study, p= .0002. In the control group (Group B), the mean baseline Hb concentration was 11.85 ± 0.86 g/dL, and after intervention mean Hb concentration decreased to 11.11 ± 0.87 g/dL, p< .0001; and anaemia prevalence was 8 out of 48, 16.7% at baseline, increasing to 12 out of 48, 25.0% at the end of the study, without a statistical difference, p= .452. (Table 2).

When considering alterations in mean Hb concentrations, there was an increase in mean Hb values for Group A (0.85±1.42); however, Group B registered a reduction in mean Hb concentration (-0.74±0.96), p< .0001 (Table 3).

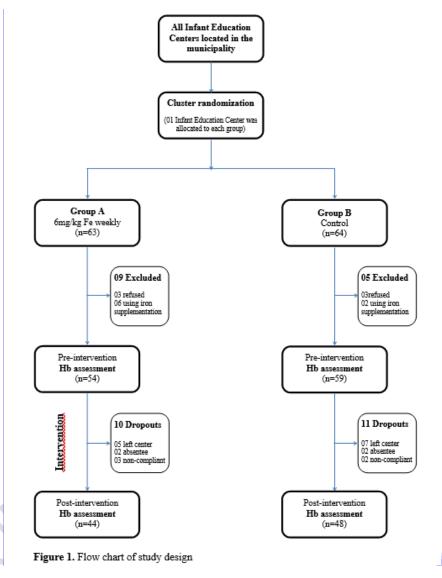
When considering only the anaemic participants, in Group A (n=20), the mean Hb concentration was 9.82±0.60 at baseline and 11.58±0.45 after the intervention, p< .0001; at baseline, 20 participants were anaemic; however, after the intervention, this number reduced to 4, p< .0001. In the control group (Group B), the mean Hb concentration was 10.78±0.14 g/dL at baseline, decreasing to 10.60±1.27 after the intervention, without statistical significance, p= .077. Intervention_The intervention group presented an increase in mean Hb concentration, 1.76±0.85 g/dL, while the control group (Group B) presented a slight decrease inslightly decreased mean Hb concentration, p= .677. (Table 3).

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In this study the following indicators were compared The following indicators were compared in this study: intervention group A versus control, for a favourable or adverse outcome (absence of angemia versus angemia). At the endpoint, adverse outcome was preseThe adverse outcome was present at the endpoint in 100% of control subjects and 20% (group A) of experimental subjects. The difference, the Reduction of Absolute Risk (RAR), Reduction of Absolute Risk (RAR) difference was 80% for group A. The 95% confidence interval for this difference is difference 95% confidence interval ranged from 62.5 to 97.5% (group A). Relative Risk (RR) was 0.36 for the weekly supplementation group. The

Number number Needed to Treat (NNT) was 2 for group A. This means that one in every 2 two preschoolers in the intervention group benefited from the intervention. The 95% confidence interval for the NNT ranged from 1.0 to 1.6 (group A).

Table 1. Baseline characteristics of study participants, by intervention group and control

	Group A (n=54)	Group B (n=59)		
Variables	Weekly iron Control		p-value ^a	
Age (months)	29.9±3.51	30.4±3.36	.30ª	
Mean±SD	29.9±3.51	30.4±3.30	.30	
Hemoglobin Haemoglobin	11.34±1.31	11.88±0.78	.003 ^a	
(g/dL)	11.34±1.31	11.00±0.70	.003	
Gender M:F	27:27	28:31	.96 ^b	
EBF	25	24	.46 ^b	
Mother with	22	23	.96 ^b	
≥9y schooling	22	23	.50	
Family income ≥300USD	23	25	.90 ^b	

All numbers are absolute; SD standard deviation; M:F male:female; EBF exclusively breastfed up to 6 months of age; ^aBased on unpaired <u>Student's-Student's f-tests</u>; ^b Based on <u>Fisher's Fisher's exact test</u> (2-tailed).

Table 2. Effects of weekly iron supplementation and control on $hagemoglobin levels_{\tau}$ and an an an an an an an aremia prevalence before and after the intervention.

	Group A (n=44)			Group B (n=48)		
	Weekly iron			Control		
Variables	Before	After	p	Before	After	p
Hb (g/dL) Mean±SD	11.19±1.42	12.04±0.96	.0003ª	11.85±0.86	11.11±0.87	<.0001 ^a
CI	10.76, 11.62	11.75, 12.33		11.60, 12.10	10.85, 11.36	
Mean increase in Hb Mean±SD		0.85±1.42			-0.74±0.96	<.0001 ^a
CI		0.413, 1.278			-1.020, - 0.463	

Anemia Anaemia 20 (45.5) 4 (9.1) .0002° 8 (16.7) 12 (25.0) .452°

All numbers are absolute except numbers in brackets, which represent percentages; Hb Hemoglobin-Haemoglobin; SD standard deviation; CI 95% Confidence interval; ^a Based on paired Student's Student's t-tests; ^b AnemiaAnaemia defined as Hb concentration <11.0 g/dL; ^c Based on Fisher's Fisher's exact test (2-tailed).

Table 3. Effects of weekly iron supplementation and control on haemoglobin levels, and anomia prevalence for anaemic preschoolers, before and after the intervention.

	Group A (n=20)			Group B (n=8)		
	Weekly iron			Control		
Variables	Before	After	р	Before	After	p
Hb (g/dL)	0.00.0.60	11 50.0 15	<.0001 ^a	10.78±0.14	10 00 1 07	.677 ^a
Mean±SD	9.82±0.60	11.58±0.45	<.0001	10.78±0.14	10.60±1.27	.077
CI	9.58, 10.06	11.34, 11.82		10.09, 11.46	9.91, 11.29	
Mean increase in		4.70.005		\	0.40.4.44	<.0001 ^a
Hb Mean±SD		1.76±0.85			-0.18±1.14	
CI		1.363, 2.157			-1.126, 0.776	
Anemia Anaemia b	20	4	<.0001°	8	4	.077 ^c

All numbers are absolute; Hb HemoglobinHaemoglobin; SDstandard deviation; CI 95% Confidence interval; Based on paired Student's-Student's t-tests; AnemiaAnaemia defined as Hb concentration <11.0 g/dL; Based on Fisher's Fisher's exact test (2-tailed).

4. DISCUSSION

In Brazil, in 2005, the Ministry of Health implemented the National Iron Supplementation Program, which aimed to reduce the prevalence of iron deficiency anemiaanaemia, through preventive iron supplementation in children aged 6-six months to 2 years, and pregnant women, and women in the postpartum period (13). This type of program has been conducted for more than 60 years in developed countries, but only in the last decade has it been implemented on a larger scale in the rest of the world (14,15).

Even with these interventions, we still have anemia_naemia as a public health problem in our country. In our study, the prevalence of anemia_naemia was 30.4%, classified as moderate, unlike the prevalence found in a recent systematic review on the prevalence of anemia in preschool children in Brazil, 40.2%, a level considered to in a recent systematic review on the prevalence of anemiaanaemia in pre-school children in Brazil, 40.2%, a level considered to be considered be a severe public health problem (10). Our study, despite being conducted in a poor region, our study presented a lower prevalence of anemiaanaemia than the more developed regions of the country (38.7%) (11). This perhaps may be explained by effective local public policies, iron rich school lunch menus and extra governmental interventions carried out locallymay be

explained by effective local public policies, iron-rich school lunch menus, and extragovernmental interventions (16-18).

In the present study, the weekly use of iron led to a significant decrease in the prevalence of anemiaanaemia in children, from 45.5 to 9.1%. Whereas, in the control group, there was an increase in the anaemic population from 16.7 to 25.0% there was an increase in the anaemic population from 16.7 to 25.0% in the control group. When analyzing only anaemic participants, a large reduction of anemiaanaemia was observed, 80%, in the group that had weekly iron supplementation. We achieved an NNT of 2,—that is, for every 2 two-children exposed to the intervention, 4one-child was recovered from the condition of anaemic. Such data show that weekly iron supplementation in the anaemic participants was very effective.

Still analyzing these same groups, a significant increase in serum Hb levels was verified in the group that received weekly ferrous sulfate supplementation (0.85g/dL) compared to the control group (-0.74g/dL), which presented a decrease in Hb levels. This result agrees with the systematic review by De-Regil (2011), who found a mean increase in Hb of 0.5g/dL when compared to placebo. In this review, greater adherence to treatment was found with weekly supplementation when compared to the daily use of iron (19). It is understood that intermittent supplementation may be an alternative both to increase patients' adherence to treatment and reduce the costs that daily supplementation demands.

Numerous studies have analyzed the specific benefits of preventive iron supplementation in children. The convergence in the reduction of the prevalence of anemiaanaemia, in the reduction of the morbidity of infectious diseases and of-infant mortality, in addition, to and the contribution to the integral development of the tissues has been observed (7,9,10). However, there are few studies that few studies analyze in a practical manner practically the possible damage from this conduct, such as possible losses in the absorption of some micronutrients, such as zinc, and the possibility of excessive accumulation of iron in the body, which could be maximized with the daily use of iron and perhaps minimized with intermittent supplementation (20).

Most of the studies that relate iron supplementation with the reduction of anemiaanaemia do not specifically analyze the 2-year-old age group that was addressed in the present study, g. Generally, they assess older children (11,14,16-19,21). It is observed in these studies, which includeincluding a systematic review, randomized clinical trials and community trials, that it is frequent to increase Hb levels and reduce anemiaanaemia with weekly supplementation, as verified in the present study, in addition to increasing iron deposits (11,14,16-19,22-24).

5. CONCLUSION

In our study, we have some approaches that make it innovative: low cost due to the weekly use of ferrous sulfate, use of supplementation in a community manner with the school lunch space in schools, reducing the chances of failures that could occur in the family environment, and intervention conducted with the intent to treat, with significant results in the short time spanperiod of just 14 weeks. Thus, it becomes a plausible strategy to be implemented on a large scale in developing countries with a high prevalence of anemiaanaemia.

CONSENT

Informed consent forms (ICF) were distributed to parents or guardians, in accordance withper Resolution No 466, of December 12, 2012. All students who signed the assent term and whose ICF was completed and signed by their parents or guardians were included in the study.

ETHICAL APPROVAL

This study was approved by the ethics committee for research at the Universidade Federal do Ceará following the ethical principles established by the National Health Council resolution #466/2012, with necessary prior written consent from school directors and parents/guardians. Medical support was available upon request. After the intervention, anaemic children were referred for treatment.

All authors hereby declare that all experiments have been examined and approved by the appropriate ethics committee and have therefore been performed in accordance with the ethical standards laid down in the 1964 Declaration of Helsinki.

COMPETING INTERESTS DISCLAIMER:

Authors have declared that no competing interests exist. The products used for this research are commonly and predominantly use products in our area of research and country. There is absolutely no conflict of interest between the authors and producers of the products because we do not intend to use these products as an avenue for any litigation but for the advancement of knowledge. Also, the research was not funded by the producing company rather it was funded by personal efforts of the authors.

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