

# Systematic review and meta-analysis of the efficacy and safety of amfepramone and mazindol as a monotherapy for the treatment of obese or overweight patients

Rosa Camila Lucchetta,<sup>1,\*</sup> Bruno Salgado Riveros,<sup>1</sup> Roberto Pontarolo,<sup>1</sup> Rosana Bento Radominski,<sup>II</sup> Michel Fleith Otuki,<sup>III</sup> Fernando Fernandez-Llimos,<sup>IV</sup> Cassiano Januário Correr<sup>I</sup>

<sup>I</sup>Laboratório de Serviços Clínicos e Evidências em Saúde, Departamento de Farmácia, Universidade Federal do Paraná (UFPR), Curitiba, PR, BR. <sup>II</sup>Serviço de Endocrinologia e Metabolismo, Hospital de Clínicas, Universidade Federal do Paraná (UFPR), Curitiba, PR, BR. <sup>III</sup>Departamento de Ciências Farmacêuticas, Universidade Estadual de Ponta Grossa, Ponta Grossa, PR, BR. <sup>IV</sup>Research Institute for Medicines (iMed.Ulisboa), Department of Social Pharmacy, Faculty of Pharmacy, University of Lisbon, Lisbon, Portugal.

The aim of this study was to evaluate efficacy and safety of amfepramone, fenproporex and mazindol as a monotherapy for the treatment of obese or overweight patients. A systematic review of primary studies was conducted, followed by a direct meta-analysis (random effect) and mixed treatment comparison. Medline and other databases were searched. Heterogeneity was explored through  $I^2$  associated with a  $p$ -value. Of 739 identified publications, 25 were included in the meta-analysis. The global evaluation of Cochrane resulted in 19 studies with a high level of bias and six with unclear risk. Due to the lack of information in primary studies, direct meta-analyses were conducted only for amfepramone and mazindol. Compared to placebo, amfepramone resulted in higher weight loss in the short-term ( $< 180$  days; mean difference (MD)  $-1.281$  kg;  $p < 0.05$ ;  $I^2: 0.0\%$ ;  $p=0.379$ ) and long-term ( $\geq 180$  days; MD  $-6.518$  kg;  $p < 0.05$ ;  $I^2: 0.0\%$ ;  $p=0.719$ ). Only studies with long-term follow up reported efficacy in terms of abdominal circumference and 5-10% weight reduction. These results corroborated the finding that the efficacy of amfepramone is greater than that of placebo. Treatment with mazindol showed greater short-term weight loss than that with placebo (MD  $-1.721$  kg;  $p < 0.05$ ;  $I^2: 0.9\%$ ;  $p=0.388$ ). However, metabolic outcomes were poorly described, preventing a meta-analysis. A mixed treatment comparison corroborated the direct meta-analysis. Considering the high level of risk of bias and the absence of important published outcomes for anti-obesity therapy assessments, this study found that the evaluated drugs showed poor evidence of efficacy in the treatment of overweight and obese patients. Robust safety data were not identified to suggest changes in their regulatory status.

**KEYWORDS:** Obesity; Weight loss; Treatment Outcome; Evidence-Based Practice.

Lucchetta RC, Riveros BS, Pontarolo R, Radominski RB, Otuki MF, Fernandez-Llimos F, et al. Systematic review and meta-analysis of the efficacy and safety of amfepramone and mazindol as a monotherapy for the treatment of obese or overweight patients. *Clinics*. 2017;72(5):317-324

Received for publication on November 11, 2016; First review completed on December 21, 2016; Accepted for publication on February 14, 2017

\*Corresponding author. E-mail: rc.lucch@yahoo.com.br

## INTRODUCTION

According to the World Health Organization, approximately 39% of adults were obese or overweight in 2014 (1). This population is subject to associated co-morbidities, such as diabetes mellitus type 2, hypertension, coronary heart disease and high risk of death (1–3). Thus, how obesity and its complications burden the health system is readily apparent. The Finkelstein et al. study conducted in the United States found

that medical costs in patients with obesity in 2006 were 41.5% higher than that for patients of normal weight. In Brazil, the estimated annual costs for all diseases related to overweight and obese patients are \$2.1 billion; approximately 10% of these costs can be attributed to being overweight or obese (4).

Treatment of obese and overweight patients is based on behavioral changes, diet and exercise, with or without pharmacotherapy or bariatric surgery, aiming to lose weight and decrease risk factors (5). Pharmacological options are being studied and have been available in the market since the 1930s. Despite its long history, pharmacotherapy remains the main tenet of scientific and political debate. There is great concern about effectiveness and safety, as many drugs previously available in the market were withdrawn due to the increased incidence of psychiatric and cardiac disorders. Moreover, the addictive effects of drugs, such as rimonabant,

**Copyright** © 2017 CLINICS – This is an Open Access article distributed under the terms of the Creative Commons License (<http://creativecommons.org/licenses/by/4.0/>) which permits unrestricted use, distribution, and reproduction in any medium or format, provided the original work is properly cited.

No potential conflict of interest was reported.

**DOI:** 10.6061/clinics/2017(05)10



fenfluramine and sibutramine, are a major concern (6). Currently, divergences among regulatory agencies are observed. The US Food and Drug Administration (FDA) approved the use of amfepramone (diethylpropion), benzphetamine, bupropion + naltrexone, phendimetrazine, phentermine, phentermine + topiramate, liraglutide, lorcaserin and orlistat. The European Medicines Agency (EMA) only recorded bupropion + naltrexone, liraglutide and orlistat. In Brazil, the Agência Nacional de Vigilância Sanitária (Anvisa) approved orlistat, sibutramine, liraglutide and lorcaserin. Moreover, amfepramone is marketed in Mexico, Chile and other countries in Latin America, and mazindol is used in Canada, Japan and several countries in Latin America. For many years, Brazil was mentioned as one of the largest consumers of appetite suppressants in the world, with evidence of irrational use of this drug class (7–10). Therefore, the country was the site of a debate that divided the Anvisa and medical societies over the maintenance record of amfepramone, mazindol and fenproporex. The Chamber of Deputies of Brazil is still discussing an attempt to halt the cancellation of amfepramone, mazindol and fenproporex use, with the justification that the population has few therapeutic options (11). A feasible explanation lies in the low quality of primary and secondary studies that have already been published. Although differences between Anvisa, FDA, EMA and others countries are likely to occur, all organizations are strongly dependent on the quality of randomized clinical trials (RCT) (12,13). Treatment discontinuation, small sample size, low methodological quality and high levels of heterogeneity in meta-analyses are the most common limitations (14–16).

It is therefore important to conduct primary and secondary studies with low risks of bias to serve as cornerstones for political decisions regarding the appropriate regulation of drugs and their use in clinical practice.

Therefore, our study aimed to assess the efficacy and safety of amfepramone, fenproporex and mazindol as a monotherapy for the treatment of obese or overweight patients using a systematic review followed by a meta-analysis.

## METHODS

Standard systematic review and meta-analytic methods were used to conduct and report this analysis (17,18).

### Sources of information and search

A search was conducted in Medline (via PubMed), SCOPUS, Scielo and the Directory of Open Access Journals until March 2016. Manual searches in the references of the included studies were also performed. The terms used were *diethylpropion*, *amfepramone*, *diethylpropione*, *anfeptramone*, *tenuate*, *femproporex*, *fenproporex*, *perphoxene*, *mazindol*, *mazindole* and *sanorex* (the search strategy is available in the Supplemental Materials).

The assessed population comprised obese or overweight individuals, taking into account criteria defined by clinical studies, with or without co-morbidities, and without restrictions of age or gender. It excluded studies undertaken in pregnant women; nursing mothers; or patients with hyperthyroidism, pheochromocytoma, glaucoma, prostate adenoma, kidney failure and liver failure. Publications related to congress abstracts, letters, editorials, news and studies that did not exactly report treatment were also excluded.

Primary studies (RCT, cohort, case-control, case report) were included in our systematic review if they assessed the efficacy or safety of amfepramone, fenproporex or mazindol regardless of treatment duration; controlled by placebo, diet, physical activity or another active drug; reporting one or more of weight change, abdominal circumference or frequency of patients who reached 5% or 10% weight loss or metabolic biomarkers; and withdrawal due to adverse reactions (qualitative or quantitative).

### Study selection and risk of bias in each study

Two independent reviewers (BSR and RCL) performed the search and study selection. Data extraction was performed by RCL and fully reviewed by BSR. Disagreements were settled by an opinion of a third researcher (CJC). Only studies reported in Portuguese, English and Spanish were assessed.

The Cochrane tool was adopted to evaluate the quality of the included studies (19).

### Direct meta-analysis and network meta-analysis

Extracted data were organized and analyzed in the Comprehensive Meta-Analysis software for direct meta-analysis, which applied the random-effects model to predict a high level of heterogeneity among studies. Odds ratios (OR) were applied as effect measurements for dichotomous outcomes, whereas mean differences (MD) were used for continuous outcomes. The statistical method for both outcomes was the inverse of variance. The results are presented with 95% confidence intervals (95% CI). Following the Cochrane recommendation, we did not assess publication bias, as none of the meta-analyses included 10 or more studies. Therefore, Egger regression or funnel plot statistical tests were not developed (19). Sensitivity analysis was performed to identify studies likely to increase heterogeneity.

Network meta-analysis using consistency or inconsistency models were built using Addis version 1.16.6 software, (Drugis, Groningen; The Netherlands). A potential scale reduction factor (PSRF) of convergence assessment close to 1.00 ( $1 < \text{PSRF} \leq 1.05$ ) points indicated that convergence was matched among simulations. Bayesian random effect models were used to assess strategies. The results were expressed by means of credibility intervals of 95% (CrI 95%). Effect measurements for continuous outcomes were MD, whereas OR was used for dichotomous outcomes. The placebo was set as the control.

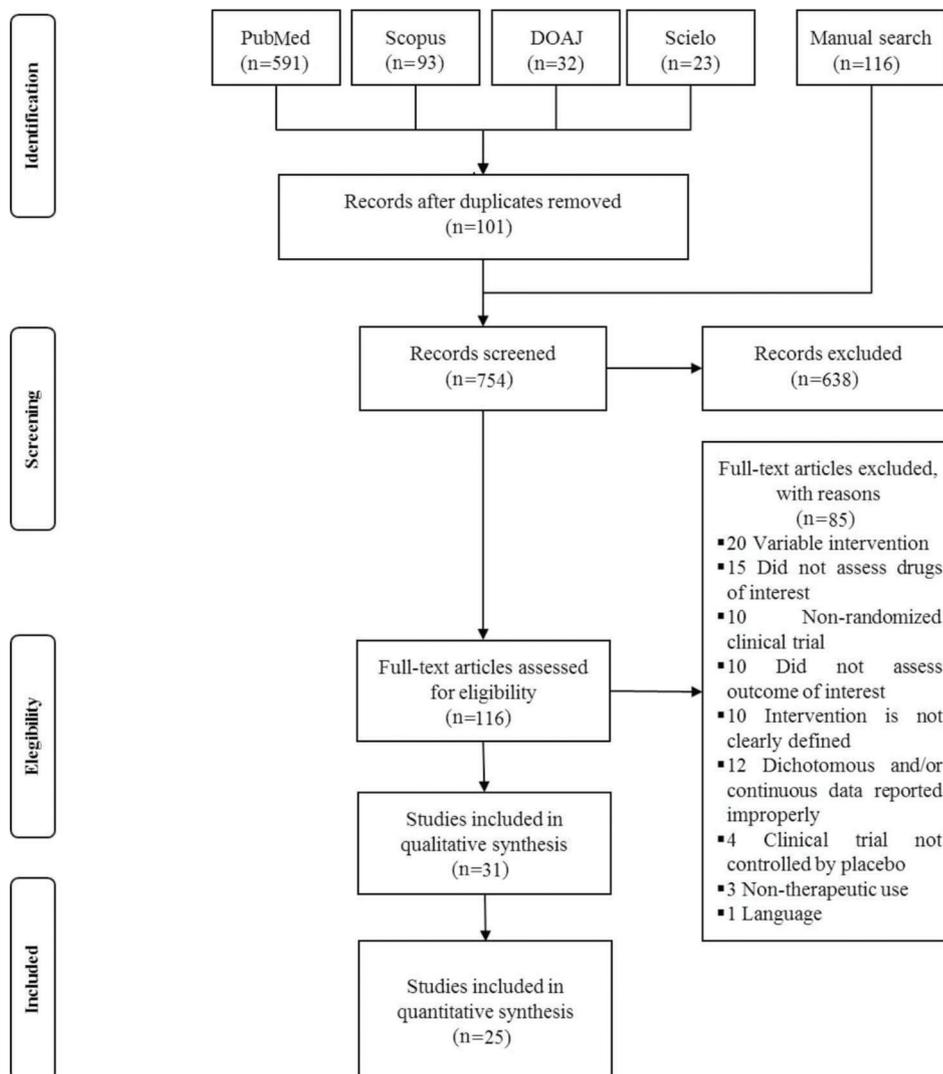
## RESULTS

Our systematic review identified 739 studies published in the assessed databases. Of these, 85 were excluded after full assessment (See Excluded studies in the Supplemental Materials); 31 and 25 were included in our systematic review and the quantitative analysis, respectively (Figure 1).

### Study characteristics

The included studies comprised RCT (n=25) (20–44) and case reports (n=6) (45–50). Cohort and case-control studies that complied with our inclusion criteria were not identified.

Considering only RCTs, the studies were published from 1967 to 2014, mainly in the United States (n=9). Data from 1,965 patients were assessed (median: 50; IIQ 25-75%: 28-80), and 19 studies took into account male and female subjects. However, data from women were more commonly observed



**Figure 1** - PRISMA flow chart. DOAJ: Directory open access journal.

(n=965). Twenty-three studies assessed adults, 12 assessed elderly people, 7 assessed adolescents and 2 assessed children. One study did not report age groups. Only 6 studies reported the presence or absence of co-morbidities. Efficacy, safety and metabolic biomarker findings were collected for amfepramone (n=13), mazindol (n=13) and fenproporex (n=1). The most common comparator was placebo (n=25). There were two head-to-head studies comparing d-amphetamine (n=1), mazindol (n=1), fenproporex (n=1) and sibutramine (n=1).

The majority of studies had a follow up of up to 12 weeks (84 days) (median: 84, I<sup>Q</sup> 25-75%: 56-84). Two studies applied the methodology of intention-to-treat for assessing results (28,41) (see characteristics of the included studies in our systematic review in the Supplemental Materials).

### Risk of bias in RCT

Considering the Cochrane tool for risk of bias, the global evaluation resulted in 19 studies with high risk of bias and 6 with uncertain risk.

The domains presenting more risk of bias were related to selective reporting (n=9) and other sources of bias (n=11). In

the first case, despite no studies reporting any *a priori* project protocol, most studies reported parameters and outcomes of interest in the Methods section, regardless of the lack of results. For the second source of bias, pharmaceutical companies responsible for marketing the drug under assessment were the funding source. Eleven studies mentioned the funding source, whereas 14 did not disclose this information in the conflict of interest statement (see chart of risk of bias in RCT described in the Supplemental Materials).

### Synthesis of Results

#### Direct meta-analysis

**Efficacy.** Only RCTs assessing amfepramone and mazindol compared to placebo were included in the direct meta-analysis (see the analysis in the Supplemental Materials).

Treatment with amfepramone led to greater loss of body weight than treatment with placebo in all studies (28,30,37, 38,41,44) regardless of daily dosage and treatment duration (MD = -1.291 kg (95% CI: -1.548; -1.035), heterogeneity test  $p=0.000$ ;  $I^2$ : 85.5%). After subgroup analysis of treatment duration (short-term: <180 days and long-term: >180 days)



and sensitivity analysis (hypothetic exclusion of Ramos et al. 1964), the result increased to MD=-1.375 kg (95% CI: -1.630; -1.121) and heterogeneity  $I^2$ : 85.5% ( $p=0.000$ ). Nevertheless, homogeneity was identified among studies assessing short-term MD=1.281 kg (95% CI: -1.538; -1.024;  $p=0.379$ ;  $I^2$ : 0.0%). For studies assessing long-term results, MD=-6.518 kg (95% CI: -8.419; -4.617;  $p=0.719$ ;  $I^2$ : 0.0%) (see Summary of findings: efficacy, safety and metabolic outcomes in the Supplemental Materials). Ramos et al. (1964) was identified as the study responsible for a high level of heterogeneity in global analysis. This may have been due to the dosage assessed (25 mg/tid (ter in die) instead of 75 mg/qd (quaque die)) and the methodological quality (37).

The two studies assessing long-term treatment also reported efficacy according to abdominal circumference and 5% and 10% weight reductions, confirming the efficacy of amfepramone over placebo (see summary of findings: efficacy, safety and metabolic outcomes in the Supplemental Materials).

From five studies included in the mazindol meta-analysis, four presented body weight reductions greater than the placebo (MD=-2.396 kg; 95% CI: -3.469; -1.323;  $I^2$ : 77.6%;  $p=0.001$ ) (21,29,32,33,42). In the sensitivity analysis, the result dropped to MD=-1.721 kg after removing Heber et al. (32) (95% CI: -2.164; -1.278;  $I^2$ : 0.9%;  $p=0.388$ ). Heber et al. identified MD strongly favoring mazindol compared to the other studies, without a reasonable explanation for such a difference (see summary of findings: efficacy, safety and metabolic outcomes in the Supplemental Materials).

**Safety.** The treatment included in the safety meta-analysis corresponded to short-term studies only and assessed the following outcomes: individuals presenting one or more adverse reactions, withdrawal due to adverse reactions (see summary of findings: efficacy, safety and metabolic outcomes in the Supplemental Materials) and probability of specific adverse reactions (see summary of findings: adverse reactions in the Supplemental Materials).

Considering the six included studies in the probability of adverse reactions meta-analysis (20,23,24,27,34,35), only one presented results favoring the placebo (52.0% vs. 28.8%) (27), with an OR=1.847 (95% CI: 1.057; 3.229);  $I^2$ : 0.0%;  $p=0.619$ ) (see summary of findings: efficacy, safety and metabolic outcomes in the Supplemental Materials). Studies showing greater withdrawal due to adverse reactions in the amfepramone group were not identified. Studies assessing amfepramone reported nausea, dry mouth, constipation, stomach discomfort, dizziness, insomnia, headache, tremor, somnolence, tension and irritation as the most common adverse reactions. Nevertheless, only dry mouth was associated with amfepramone treatment (OR=2.430; 95% CI: 1.248; 4.729;  $I^2$ : 0.0%;  $p=0.651$ ) (see summary of findings: adverse reactions in the Supplemental Materials).

Out of five studies included in the probability of adverse reactions meta-analysis, only one pointed to a greater risk for such outcomes in the mazindol group (31), generating an OR=4.086 (95% CI: 1.780; 9.376;  $I^2$ : 0.0%;  $p=0.53$ ) (see summary of findings: efficacy, safety and metabolic outcomes in the Supplemental Materials). The meta-analysis results showed a higher risk of discontinuation in the mazindol group (OR=2.760; 95% CI: 1.472; 5.175;  $I^2$ : 0.0%;  $p=0.949$ ) (see summary of findings: efficacy, safety and metabolic outcomes in the Supplemental Materials). The most common adverse

reactions reported were palpitations, rash, nausea, dry mouth, constipation, bad taste sensation, vomiting, dizziness, insomnia, headache, tension, irritation, dysuria and chills. Only insomnia and constipation were associated with mazindol treatment (OR=8.083; 95% CI: 1.782; 36.674;  $I^2$ : 0.0%;  $p=0.910$ ) and OR=3.906 (95% CI: 1.156; 13.197;  $I^2$ : 0.0%;  $p=0.910$ ), respectively (see summary of findings: adverse reactions in the Supplemental Materials).

Adverse reactions described in the identified case reports for amfepramone were paranoid ideation, restlessness, aphasia, transient ischemic attacks due to vasospasm, schizophrenia, psychotic symptoms, dry mouth, constipation, dizziness and chronic bronchitis (45-48,50), and reactions to fenproporex were withdrawal syndrome, aggression, anxiety, irritability, nightmares and insomnia, followed by severe depression and attempted suicide (49).

**Metabolic outcomes.** It was only possible to carry out meta-analyses for total cholesterol change after mazindol treatment. Despite two studies identifying a decrease in total cholesterol ( $p=0.000$ ) (22,39), our meta-analysis showed high heterogeneity (OR=-21.750 mg/dL; 95% CI: -39.514; -3.987;  $I^2$ : 95.3%;  $p=0.000$ ), not justified by dosage difference, treatment duration, sample size or study quality (see summary of findings: efficacy, safety and metabolic outcomes in the Supplemental Materials).

In addition to total cholesterol, the following outcomes were also reported: changes in fasting glucose, serum triglycerides, systolic blood pressure, diastolic blood pressure and heart rate. However, meta-analysis was prevented since primary data were reported incorrectly or only qualitatively.

**Network meta-analysis.** All evaluations were split into short-term (Table 1) and long-term treatments (Table 2). Outcomes that presented statistical significance were change in body weight, favoring amfepramone and mazindol compared to placebo (Table 1), and discontinuation due to adverse reactions, favoring placebo compared to mazindol (Table 1). Regarding adverse reactions, somnolence and insomnia demonstrated statistical significance. The first favored placebo over amfepramone, and the second favored placebo over mazindol (Table 1). All efficacy outcomes assessing long-term treatment favored amfepramone over placebo (Table 2). Due to a closed circuit of evidence, it was not possible to conduct inconsistency analyses. More information on the network meta-analysis is available in the Supplemental Materials.

## ■ DISCUSSION

### Relevance for decision making

Canada, USA, Japan, Brazil, Mexico, Chile and several countries in Latin America have discussed the regulatory status of anorectics as well as updated their regulations for the registration and trade of these drugs. In an official report in 2007 (51), the US FDA required the following criteria for a drug to be registered as an anti-obesity drug: promote statistically significant weight loss compared to placebo in >5% of subjects within one year of treatment or >35% of subjects reaching >5% weight loss. The FDA also requested evidence that new drugs are capable of improving metabolic biomarkers, including blood pressure, lipids and glucose (51).



**Table 1 - Network meta-analysis: short-term treatments.**

Efficacy – Mean difference (CrI 95%) – Cut-off =0				
Change in body weight	AMFEPRAMONE	-	-	
	0.61 (-1.26, 2.27)	MAZINDOL	-	
	<b>-1.51 (-3.16, -0.32)</b>	<b>-2.12 (-3.28, -1.14)</b>	PLACEBO	
Safety and tolerability – Odds ratio (CrI 95%) – Cut-off =1				
Discontinuation due to adverse reactions	AMFEPRAMONE	0.61 (0.12, 3.46)	1.20 (0.44, 2.89)	Participants at least on adverse reaction
	0.51 (0.07, 5.52)	MAZINDOL	1.97 (0.43, 7.38)	
	<b>2.43 (0.36, 19.37)</b>	<b>4.38 (1.89, 16.38)</b>	PLACEBO	
Adverse reactions – Odds ratio (CrI 95%) – Cut-off =1				
Constipation	AMFEPRAMONE	0.20 (0.01, 2.15)	0.93 (0.24, 2.92)	Nausea
	0.38 (0.02, 5.06)	MAZINDOL	4.48 (0.62, 52.81)	
	1.58 (0.16, 13.41)	4.10 (0.81, 38.94)	PLACEBO	
Dizziness	AMFEPRAMONE	0.00 (0.00, 0.03)	1.20 (0.21, 11.90)	Insomnia
	1.39 (0.23, 12.64)	MAZINDOL	<b>6.17x10<sup>8</sup> (48.29, 5.08x10<sup>19</sup>)</b>	
	2.21 (0.64, 9.22)	1.56 (0.34, 6.18)	PLACEBO	
Headache	AMFEPRAMONE	0.12 (0.00, 1.85)	0.94 (0.28, 3.45)	Irritation
	0.99 (0.14, 5.28)	MAZINDOL	7.57 (0.73, 264.30)	
	0.98 (0.31, 2.29)	1.00 (0.21, 4.57)	PLACEBO	
Dry mouth	AMFEPRAMONE	0.00 (0.00, 0.26)	0.00 (0.00, 0.28)	Vomiting
	0.92 (0.09, 7.69)	MAZINDOL	1.63 (0.06, 77.49)	
	2.77 (0.89, 11.87)	3.04 (0.60, 23.89)	PLACEBO	
Somnolence	AMFEPRAMONE	0.48 (0.04, 4.03)	1.21 (0.37, 3.22)	Tension
	2.09 (0.00, 1132452.48)	MAZINDOL	2.40 (0.37, 21.32)	
	<b>1240.03 (1.17, 5.63x10<sup>7</sup>)</b>	402.06 (0.46, 7.63x10 <sup>7</sup> )	PLACEBO	

Lower triangle, left: treatment in the column compared to treatment in the row. Upper triangle, right: treatment in the row compared to treatment in the column. Bolded values indicate statistically significant differences.

**Table 2 - Network meta-analysis: long-term treatments.**

Efficacy – Mean difference (CrI 95%) – Cut-off=0					
Change in body weight	AMFEPRAMONE	-2.09 (-9.21, 4.97)	-3.30 (-9.99, 3.39)	<b>-6.51 (-11.64, -1.51)</b>	Change in waist circumference
	-1.99 (-9.13, 4.99)	FENPROPOREX	-1.23 (-8.82, 6.46)	-4.38 (-11.84, 2.63)	
	-2.36 (-9.05, 4.52)	-0.31 (-7.66, 7.24)	MAZINDOL	-3.20 (-10.10, 3.49)	
	<b>-6.57 (-11.83, -1.36)</b>	-4.53 (-11.72, 2.46)	-4.16 (-11.06, 2.46)	PLACEBO	
Efficacy and safety – Odds ratio (CrI 95%) – Cut-off =1					
10% weight loss	AMFEPRAMONE	1.20 (0.18, 8.15)	0.99 (0.13, 7.00)	<b>5.56 (1.30, 25.11)</b>	5% weight loss
	3.64 (0.13, 84.38)	FENPROPOREX	0.84 (0.10, 6.86)	4.56 (0.66, 34.33)	
	3.06 (0.13, 77.72)	0.86 (0.03, 24.94)	MAZINDOL	5.51 (0.79, 40.82)	
	<b>33.26 (2.73, 577.98)</b>	9.38 (0.38, 291.18)	10.97 (0.42, 324.51)	PLACEBO	
	AMFEPRAMONE	1.64 (0.20, 19.53)	3.89 (0.31, 150.48)	1.24 (0.24, 6.39)	
	FENPROPOREX	2.45 (0.14, 117.99)	0.78 (0.06, 6.97)	Discontinuation due to adverse reactions	
	MAZINDOL		0.33 (0.01, 3.76)		
			PLACEBO		

Lower triangle, left: treatment in the column compared to treatment in the row. Upper triangle, right: treatment in the row compared to treatment in the column. Bolded values indicate statistically significant differences.

Our review identified only two studies, of 6 and 12 months of follow up, that reported the outcome of >5% weight loss (28,41). The first identified 66.7% of patients reaching such an outcome with amfepramone treatment compared to 25% in the placebo group. The second aforementioned study, assessing amfepramone, fenproporex, mazindol, fluoxetine and sibutramine treatment, identified 71.4%, 69.0%, 72.4%, 34.5% and 73.3%, respectively, of patients reaching body weight loss of >5% compared to 34.4% in the placebo group. These studies also reported metabolic biomarkers in a way that prevented their use in a meta-analysis framework. Among the identified studies assessing short-term treatment, none

reported losses of 5% or 10% weight loss, and only two reported data capable of use in meta-analysis, presenting a high level of heterogeneity (22,39). Therefore, according to our systematic review, amfepramone, mazindol and fenproporex do not have enough evidence to comply with the efficacy criteria of the FDA.

Safety parameters, especially related to major adverse reactions, were not identified in clinical trials for the assessed drugs, qualitatively or quantitatively (52). Contributing factors may include the lack of separation of major and minor adverse reactions in primary studies and the small sample size. To corroborate such data, observational studies were searched, identi-



fying only six case reports that included cases of addiction and psychotic disorders. Case report studies often describe unpublished or unknown outcomes of a given treatment (53).

### Importance for research

The systematic review followed by meta-analysis summarized several efficacy and safety results comparing weight loss promoted by pharmacotherapeutic treatment and dieting versus weight reduction only from dieting (54). The main limitations of this analysis reside in the short-term monitoring of clinical trials and the lack of analysis by sub-populations, which limit the use of the results in the clinical population given the heterogeneity of obese and overweight patients. To date, systematic review followed by meta-analysis covering the three drugs evaluated here has not been performed.

The most common efficacy outcome reported in the included studies was mean difference in weight loss between groups. As already discussed, 5% or 10% weight loss was rarely described in RCTs. Jensen et al. recommend an initial target reduction of 5-10% of baseline weight within six months of treatment (5). Metabolic biomarkers were poorly reported, whereas for safety, the risk of withdrawal due to adverse reactions and the description of any adverse event experienced were the most common outcomes. The National Obesity Observatory Standard Evaluation Framework (NOO SEF) (55) recommends that at minimum, weight and height measurements must be reported for the assessment of interventions at three different time points within one year of follow up (55). Another concern is the report of different results during treatment. In most RCTs, the results were not divided across the follow up but instead pooled into one single mean, counteracting NOO SEF recommendations.

The majority of our meta-analysis presented low heterogeneity, which can be explained by the identification bias of heterogeneity, since our meta-analysis comprised few studies and small populations. Hippel (2015) identified that true heterogeneity is more likely to be seen when more studies are included. However, there is not a consensus about the minimum number of studies to carry out a meta-analysis (56,57). Therefore, despite having a small number of meta-analyses with heterogeneity, we cannot exclude the chance of being unable to identify true heterogeneity.

Some RCTs have important limitations that are not sensitive to the Cochrane tool, such as the lack of a thorough description of the population under assessment. Most primary studies assessed obese or overweight adults with or without co-morbidities, reporting pooled results. This practice leads to high heterogeneity and results that are difficult to use in clinical practice. Such data will hardly clarify decisions for decision makers and clinicians, as results such as these are not representative of specific patient profiles (5).

The risk of bias identified through the Cochrane tool is high, which required us to cautiously interpret the results. According to the Cochrane tool, the most important risk of bias lies in the conflict of interest of researchers, as pharmaceutical companies funded the majority of studies. An observational study of 370 RCTs revealed that funding source, treatment effect and double-blinding are the most relevant predictors of statistically significant results (58).

Other observed limitations were small sample size, low number of head-to-head studies, poor description of therapy adherence, frequency of major adverse reactions and changes in metabolic biomarker outcomes. Discontinuation of

treatment, small sample size, low methodological quality and high level of heterogeneity in the meta-analyses were the most common limitations (14–16).

The limitations of this review include the following: a) The review of gray literature (dissertations, theses, abstracts) was absent. Since the three evaluated drugs have over 50 years of marketing and research, and two of them are no longer marketed in major agencies, there was a low probability of identifying studies of interest. Furthermore, a manual search was performed both in the reference lists of included studies and in official materials of companies and organizations, which resulted in 116 identified publications. b) There was also a language limitation; however, its effect was limited to the exclusion of a study published in Polish. c) With respect to inadequate reporting of dichotomous or continuous data in the primary study, we chose to not contact the authors of the studies to obtain raw data, considering the publication dates extended into past decades (23 of 25 studies were published from 1961 to 1987), and we thus had a low probability of success in contact. d) The effects of the combination of drugs with lifestyle changes, including diet and exercise, were not considered in the analysis; although most studies combined medications with lifestyle changes, they did not report the proposed interventions in detail (e.g., calorie deficit, intensity of physical activity, duration). e) The network meta-analysis was performed containing only placebo as the common comparator, which stems from the lack of clinical trials with head-to-head comparisons.

More studies are needed on the safety of the evaluated drugs, principally based in real practice, to have a stronger basis for recommendations to regulatory bodies or health institutions regarding the anorectic drugs evaluated. Future RCTs should be controlled by active drugs, and the results of these trials should be reported in terms of abdominal circumference, change in body weight, the number of participants who achieved a loss of 5-10% of body weight and any impact on metabolic biomarkers over the course of 3, 6, 9 and 12 months. Some population subgroups have plenty of data in the literature. Thus, they have the potential to be further explored in future assessments as obese or overweight adults without co-morbidities, obese adults with diabetes type II, overweight adults with diabetes type II, obese or overweight adults with hypertension and adolescents and children without co-morbidities.

The majority of identified RCTs present a high risk of bias, mostly related to conflict of interest and selective outcome reporting. Regarding short-term treatment (<180 days), amfepramone and mazindol are more effective than placebo for losing weight. Nevertheless, short-term studies assessing a decrease of 5-10% of body weight were not identified. Mazindol was associated with discontinuation due to adverse reactions. The most common adverse reactions caused by this drug were constipation and insomnia. For amfepramone, dry mouth and somnolence were the most common adverse reactions. Long-term treatments (>180 days) only had results favoring amfepramone (compared to placebo). Regarding safety, associations with adverse reactions or discontinuation due to adverse reactions were not identified. Therefore, considering the high risk of bias and the lack of reporting of important outcomes, such as metabolic biomarkers and loss of at least 5% body weight, this study concludes that amfepramone, mazindol and fenproporex have weak evidence of their effectiveness in the treatment of overweight and obese patients. In addition, robust safety data were not



identified, which would allow for suggestions regarding short- and long-term changes in their regulatory status.

## ACKNOWLEDGMENTS

Funding Sources: Ministério da Ciência, Tecnologia e Inovação/ Conselho Nacional de Desenvolvimento Científico e Tecnológico (CNPq), no. 401098/2013-3.

## AUTHOR CONTRIBUTIONS

Lucchetta RC, Riveros BS, Pontarolo R, Radominski RB, Otuki MF, Fernandez-Llimos F and Correr CJ contributed substantially to the study conception, development, analysis and/or interpretation of the data and the drafting of the manuscript.

## REFERENCES

- WHO – World Health Organization. Global Health Observatory. Overweight and obesity [Internet]. 2014 [cited 2015 Nov 9]. Available from: [http://www.who.int/gho/ncd/risk\\_factors/overweight/en/](http://www.who.int/gho/ncd/risk_factors/overweight/en/).
- Field AE, Coakley EH, Must A, Spadano JL, Laird N, Dietz WH, et al. Impact of overweight on the risk of developing common chronic diseases during a 10-year period. *Arch Intern Med.* 2001;161(13):1581-6, <http://dx.doi.org/10.1001/archinte.161.13.1581>.
- Sowers JR. Obesity as a cardiovascular risk factor. *Am J Med.* 2003;115 Suppl 8A:37S-41S, <http://dx.doi.org/10.1016/j.amjmed.2003.08.012>.
- Bahia L, Coutinho ES, Barufaldi LA, Abreu Gde A, Malhão TA, de Souza CP, et al. The costs of overweight and obesity-related diseases in the Brazilian public health system: cross-sectional study. *BMC Public Health.* 2012;12:440, <http://dx.doi.org/10.1186/1471-2458-12-440>.
- Jensen MD, Ryan DH, Apovian CM, Ard JD, Comuzzie AG, Donato KA, et al. 2013 AHA/ACC/TOS Guideline for the Management of Overweight and Obesity in Adults: A Report of the American College of Cardiology/American Heart Association Task Force on Practice Guidelines and The Obesity Society. *Circulation.* 2014;129(25 Suppl 2):S102-38, <http://dx.doi.org/10.1161/01.cir.0000437739.71477.ee>.
- Li MF, Cheung BM. Rise and fall of anti-obesity drugs. *World J Diabetes.* 2011;2(2):19-23, <http://dx.doi.org/10.4239/wjdv.v2.i2.19>.
- Carneiro Mde F, Guerra AA Jr, Acurcio Fde A. [Prescription, dispensing, and regulation of psychoactive anorexigenic drugs in Belo Horizonte, Minas Gerais, Brazil]. *Cad Saude Publica.* 2008;24(8):1763-72, <http://dx.doi.org/10.1590/S0102-311X2008000800005>.
- Nappo SA, Tabach R, Noto AR, Galduróz JC, Carlini EA. Use of anorectic amphetamine-like drugs by Brazilian women. *Eat Behav.* 2002;3(2):153-65, [http://dx.doi.org/10.1016/S1471-0153\(01\)00054-X](http://dx.doi.org/10.1016/S1471-0153(01)00054-X).
- Nappo SA, de Oliveira EM, Morosini S. Inappropriate prescribing of compounded antiobesity formulas in Brazil. *Pharmacoepidemiol Drug Saf.* 1998;7(3):207-12, [http://dx.doi.org/10.1002/\(SICI\)1099-1557\(199805/06\)7:3<207::AID-PDS335>3.0.CO;2-W](http://dx.doi.org/10.1002/(SICI)1099-1557(199805/06)7:3<207::AID-PDS335>3.0.CO;2-W).
- de Lima MS, Béria JU, Tomasi E, Mari JJ. Use of amphetamine-like appetite suppressants: a cross-sectional survey in Southern Brazil. *Subst Use Misuse.* 1998;33(8):1711-9, <http://dx.doi.org/10.3109/10826089809058951>.
- Câmara dos Deputados (Brasil). Projeto de Lei da Câmara no 61, de 2015 [Internet]. 2015. Available from: <https://www25.senado.leg.br/web/atividade/materias/-/materia/122114>.
- Eriksson R, Aagaard L, Jensen LJ, Borisova L, Hørlück D, Brunak S, et al. Discrepancies in listed adverse drug reactions in pharmaceutical product information supplied by the regulatory authorities in Denmark and the USA. *Pharmacol Res Perspect.* 2014;2(3):e00038, <http://dx.doi.org/10.1002/prp2.38>.
- Tafuri G, Stolk P, Trotta F, Putzeist M, Leufkens HG, Laing RO, et al. How do the EMA and FDA decide which anticancer drugs make it to the market? A comparative qualitative study on decision makers' views. *Ann Oncol.* 2014;25(1):265-9, <http://dx.doi.org/10.1093/annonc/mdt512>.
- Arterburn DE, Crane PK, Veenstra DL. The efficacy and safety of sibutramine for weight loss: a systematic review. *Arch Intern Med.* 2004;164(9):994-1003, <http://dx.doi.org/10.1001/archinte.164.9.994>.
- Fabricatore AN, Wadden TA, Moore RH, Butryn ML, Gravallese EA, Erondy NE, et al. Attrition from randomized controlled trials of pharmacological weight loss agents: a systematic review and analysis. *Obes Rev.* 2009;10(3):333-41, <http://dx.doi.org/10.1111/j.1467-789X.2009.00567.x>.
- Chilton M, Dunkley A, Carter P, Davies MJ, Khunti K, Gray LJ. The effect of antiobesity drugs on waist circumference: A mixed treatment comparison. *Diabetes Obes Metab.* 2014;16(3):237-47, <http://dx.doi.org/10.1111/dom.12198>.
- Moher D, Liberati A, Tetzlaff J, Altman DG, PRISMA Group. Preferred reporting items for systematic reviews and meta-analyses: the PRISMA statement. *PLoS Med.* 2009;6(7):e1000097, <http://dx.doi.org/10.1371/journal.pmed.1000097>.
- Hutton B, Salanti G, Caldwell DM, Chaimani A, Schmid CH, Cameron C, et al. The PRISMA extension statement for reporting of systematic reviews incorporating network meta-analyses of health care interventions: checklist and explanations. *Ann Intern Med.* 2015;162(11):777-84, <http://dx.doi.org/10.7326/M14-2385>.
- Higgins J, Green S. *Cochrane Handbook for Systematic Reviews of Interventions.* Version 5.1.0 [updated March 2011]. The Cochrane Collaboration; 2011.
- Andelman MB, Jones C, Nathan S. Treatment of obesity in underprivileged adolescents. Comparison of diethylpropion hydrochloride with placebo in a double-blind study. *Clin Pediatr.* 1967;6(6):327-30, <http://dx.doi.org/10.1177/00092286700600607>.
- Baird IM, Howard AN. A double-blind trial of mazindol using a very low calorie formula diet. *Int J Obes.* 1977;1(3):271-8.
- Bandisod MS, Boshell BR. Double-blind clinical evaluation of mazindol (42-548) in obese diabetics. *Curr Ther Res Clin Exp.* 1975;18(6):816-24.
- Bolding OT. A double-blind evaluation of Tenuate dospan in overweight patients from a private gynecologic practice. *J Med Assoc State Ala.* 1968;38(3):209-12.
- Bolding OT. Diethylpropion hydrochloride: an effective appetite suppressant. *Curr Ther Res Clin Exp.* 1974;16(1):40-8.
- Bradley H, Blum N, Scheib R. Mazindol in obesity with known cardiac disease. *J Int Med Res.* 1974;2(5):347-9, <http://dx.doi.org/10.1177/030006057400200505>.
- Campagnoli M. [Clinical trial with a nonamphetaminic anorectic]. *Rev Iber Endocrinol.* 1976;23(137):463-83.
- Carney DE, Tweddell ED. Double blind evaluation of long acting diethylpropion hydrochloride in obese patients from a general practice. *Med J Aust.* 1975;1(1):13-5.
- Cercato C, Roizenblatt VA, Leança CC, Segal A, Lopes Filho AP, Mancini MC, et al. A randomized double-blind placebo-controlled study of the long-term efficacy and safety of diethylpropion in the treatment of obese subjects. *Int J Obes.* 2009;33(8):857-65, <http://dx.doi.org/10.1038/ijo.2009.124>.
- Defelice EA, Chaykin LB, Cohen A. Double-blind clinical evaluation of mazindol, dextroamphetamine, and placebo in treatment of exogenous obesity. *Curr Ther Res Clin Exp.* 1973;15(7):358-66.
- Elliott BJ. A double-blind controlled study of the use of diethylpropion hydrochloride (Tenuate) in obese patients in a rural practice. *N Z Med J.* 1978;88(622):321-2.
- Hadler AJ. Mazindol, a new non-amphetamine anorexigenic agent. *J Clin Pharmacol New Drugs.* 1972;12(11):453-8, <http://dx.doi.org/10.1002/j.1552-4604.1972.tb00246.x>.
- Heber KR. Double-blind trial of mazindol in overweight patients. *Med J Aust.* 1975;2(14):566-7.
- Maclay WP, Wallace MG. A multi-centre general practice trial of mazindol in the treatment of obesity. *Practitioner.* 1977;218(1305):431-4.
- McQuarrie HG. Clinical assessment of the use of an anorectic drug in a total weight reduction program. *Curr Ther Res Clin Exp.* 1975;17(5):437-43.
- Nolan GR. Use of an anorectic drug in a total weight reduction program in private practice. *Curr Ther Res Clin Exp.* 1975;18(2):332-7.
- Parsons WB Jr. Controlled-release diethylpropion hydrochloride used in a program for weight reduction. *Clin Ther.* 1981;3(5):329-35.
- DERAMOS EC. The use of diethylpropion in the treatment of obesity. *Br J Clin Pract.* 1964;18:210-1.
- SEATON DA, DUNCAN LJ, ROSE K, SCOTT AM. Diethyl-propion in the treatment of "refractory" obesity. *Br Med J.* 1961;1(5231):1009-11, <http://dx.doi.org/10.1136/bmj.1.5231.1009>.
- Slama G, Selmi A, Hautecouverture M, Tchobroutsky G. Double blind clinical trial of mazindol on weight loss blood glucose, plasma insulin and serum lipids in overweight diabetic patients. *Diabete Metab.* 1978;4(3):193-9.
- Smith RG, Innes JA, Munro JF. Double-blind evaluation of mazindol in refractory obesity. *Br Med J.* 1975;3(5978):284, <http://dx.doi.org/10.1136/bmj.3.5978.284>.
- Suplicy H, Boguszewski CL, dos Santos CM, do Desterro de Figueiredo M, Cunha DR, Radominski R. A comparative study of five centrally acting drugs on the pharmacological treatment of obesity. *Int J Obes.* 2014;38(8):1097-103, <http://dx.doi.org/10.1038/ijo.2013.225>.
- Thorpe PC, Isaac PF, Rodgers J. A controlled trial of mazindol (Sanjorex, Teronac) in the management of the obese rheumatic patients. *Curr Ther Res Clin Exp.* 1975;17(2):149-55.
- Wallace AG. AN 448 Sandoz (Mazindol) in the treatment of obesity. *Med J Aust.* 1976;1(11):343-5.
- Williams J. Trial of a long-acting preparation of diethylpropion in obese diabetics. *Practitioner.* 1968;200(197):411-4.
- Crols R, Dierckx R, Saerens J, Deyn PP De. Transient ischemic attacks associated with amfepramone therapy: a case report. *Funct Neurol.* 1993;8(5):351-4.
- Fookes BH. Schizophrenia-like reaction to diethylpropion. *Lancet.* 1976;2(7996):1206, [http://dx.doi.org/10.1016/S0140-6736\(76\)91728-1](http://dx.doi.org/10.1016/S0140-6736(76)91728-1).



47. Hoffman BF. Diet pill psychosis. *Can Med Assoc J.* 1977;116(4):351-5.
48. Kuenssberg E. Diethylpropion. *Br Med J.* 1962;2(5306):729-30, <http://dx.doi.org/10.1136/bmj.2.5306.729-d>.
49. Pelissier-Alicot AL, Piercecchi-Marti MD, Bartoli C, Kuhlmann E, Coiffait PE, Sanvoisin A, et al. Abusive prescription of psychostimulants: a study of two cases. *J Forensic Sci.* 2006;51(2):407-10, <http://dx.doi.org/10.1111/j.1556-4029.2006.00078.x>.
50. Wilson R, Long C. A clinical evaluation of tenuate--a new anti-appetite compound. *J Ir Med Assoc.* 1960;46:86-8.
51. U.S. Department of Health and Human Services, Food and Drug Administration (FDA), Center for Drug Evaluation and Research (CDER). *Guidance for Industry Developing Products for Weight Management (Draft Guidance).* 2007, <http://dx.doi.org/10.1016/j.addr.2009.03.006>
52. Brasil, Agência Nacional de Vigilância Sanitária (Anvisa). *Nota técnica sobre eficácia e segurança dos medicamentos inibidores de apetite.* 2011.
53. Gagnier JJ, Kienle G, Altman DG, Moher D, Sox H, Riley D. The CARE Guidelines: Consensus-based Clinical Case Reporting Guideline Development. *Glob Adv Health Med.* 2013;2(5):38-43, <http://dx.doi.org/10.7453/gahmj.2013.008>.
54. Li Z, Maglione M, Tu W, Mojica W, Arterburn D, Shugarman LR, et al. Meta-analysis: pharmacologic treatment of obesity. *Ann Intern Med.* 2005;142(7):532-46, <http://dx.doi.org/10.7326/0003-4819-142-7-200504050-00012>.
55. Roberts K, Ells L, Adamson A, Flaherty SJ, Hamilton-shield J, Moffett G, et al. Standard Evaluation Framework for weight management interventions [Internet]. 2009; 6. Available from: [http://www.noo.org.uk/uploads/doc721\\_2\\_noo\\_SEF\\_FINAL300309.pdf](http://www.noo.org.uk/uploads/doc721_2_noo_SEF_FINAL300309.pdf).
56. Davey J, Turner RM, Clarke MJ, Higgins JP. Characteristics of meta-analyses and their component studies in the Cochrane Database of Systematic Reviews: a cross-sectional, descriptive analysis. *BMC Med Res Methodol.* 2011;11:160, <http://dx.doi.org/10.1186/1471-2288-11-160>.
57. von Hippel PT. The heterogeneity statistic I(2) can be biased in small meta-analyses. *BMC Med Res Methodol.* 2015;15:35, <http://dx.doi.org/10.1186/s12874-015-0024-z>.
58. Als-Nielsen B, Chen W, Gluud C, Kjaergard LL. Association of funding and conclusions in randomized drug trials: a reflection of treatment effect or adverse events? *JAMA.* 2003;290(7):921-8, <http://dx.doi.org/10.1001/jama.290.7.921>.



**Erratum**

In the article **Systematic review and meta-analysis of the efficacy and safety of amfepramone and mazindol as a monotherapy for the treatment of obese or overweight patients**, on page 319, remove the following phrase from the **RESULTS** section / **Synthesis of Results** / **Direct meta-analysis** / **Efficacy**:

“(see the analysis in the Supplemental Materials)”