

## Review Article

# The Clinical Efficacy of Imiglucerase versus Eliglustat in Patients with Gaucher's Disease Type 1: A Systematic Review

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## INTRODUCTION

Gaucher's disease (GD) is one of the most common lysosomal diseases in humans. It results from  $\beta$ -glucosidase deficiency and leads to necrosis, especially in macrophages with the accumulation of glucosylceramidase in cells.<sup>[1,2]</sup> The risk of developing GD increases with consanguinity in the family. Its frequency varies with different populations being most prevalent at 1:45 birth incidence individuals of Ashkenazi Jewish descent.<sup>[3]</sup>

The disease has three subtypes: Type 1 or non-neuropathic form making up more than 90% of cases (with the prevalence of 1/20000–1/40000), Type 2

**ABSTRACT** Gaucher's disease (GD) is one of the most common lysosomal diseases in humans. It results from  $\beta$ -glucosidase deficiency and leads to necrosis, especially in macrophages with the accumulation of glucosylceramidase in cells. Most of the deleterious effects of the disease are seen in the liver, spleen, and bone marrow. The aim of this study was to compare the efficacy of Imiglucerase with Eliglustat in treating patients with GD. PubMed/Medline, Cochrane Library, Scopus, Web of Science, Embase, and Google Scholar were searched from inception to August, 2018. Predefined inclusion criteria for included studies were based on search methodology and are as follows: All randomized, quasi-randomized controlled, and cohort studies about patients with GD Type 1 that Imiglucerase was compared with Eliglustat were included. Two authors independently choose the papers based on the inclusion criteria. From 2979 recognized studies, three studies including two randomized clinical trials and one cohort study were recognized to meet the inclusion criteria. The primary outcomes were hemoglobin level, platelets count, liver, and spleen size, and the secondary outcomes were the immunological side effects of the medicines and bone complications. The results showed that there is no meaningful difference between the two medicines in terms of increasing blood hemoglobin, platelets count, and reducing the liver and spleen size. The findings of this review showed that both medicines are effective in the treatment of GD Type 1 and there is no statistically significant difference between their efficacies.

**KEYWORDS:** Eliglustat, Gaucher disease, Imiglucerase

or the acute neuropathic form (with the prevalence of 1/100000), and Type 3 or the chronic neuropathic form (with the prevalence of 1/100000). All types of the disease are inherited in an autosomal recessive pattern.<sup>[4-6]</sup> Mutation in alleles including the N370s substitution is affiliated with the non-neuropathic Type 1 GD. It is generally found in Ashkenazi Jewish and non-Jewish Europeans.<sup>[7,8]</sup> The symptoms of the disease are anemia, thrombocytopenia, bone involvement, hepatomegaly, splenomegaly, lung, heart, kidney involvements, and

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growth disorders.<sup>[9-15]</sup> The available treatments for GD include Enzyme Replacement Therapy (ERT) and Substrate Reduction Therapy (SRT).<sup>[6]</sup> ERT medications are including Imiglucerase, Velaglucerase, and Taliglucerase,<sup>[6,10,16]</sup> while SRT medications consisted of Eliglustat and Miglustat.<sup>[6,17,18]</sup>

Imiglucerase (200 units/5 ml vials and 400 units/5 ml vials) is a recombinant DNA-produced analogue of human  $\beta$ -glucocerebrosidase (GBA) and is only indicated for treatment of Type 1 GD (the non-neuropathic form) and Type 3 GD (the chronic neuropathic form); it is not effective in the treatment of Type 2 GD (the acute neuropathic form). Eliglustat (hard gelatin capsule, 100 mg Eliglustat tartrate equivalent of 84 mg Eliglustat) minimizes the accumulation of excess material by inhibiting material synthesis. The most important advantages of SRT are its oral administration, easier crossing the blood-brain barrier, and reaching other organs.<sup>[19,20]</sup> Lifelong intravenous administration, high cost, and not entering to the nervous system are among the disadvantages of ERT.<sup>[17,21]</sup>

Evidences for efficacy of ERT and SRT in GD are unusually sparse. The recent development of new products such as Eliglustat, as an alternative for Imiglucerase, raise curiosity about biological benefits, and cost savings of it. The effectiveness of all ERTs and SRTs has been evaluated in the previous systematic review. As a result, assessing the role of Eliglustat as an alternative to Imiglucerase in treating patients with GD Type 1 by conducting a systematic review of published relevant studies seems rational. Hence, the aim of this study was to compare the efficacy of Imiglucerase with Eliglustat in treating patients with GD Type 1.

## METHODS

This systematic review was conducted in accordance with the Preferred Reporting Items for Systematic Reviews and Meta-Analyses guideline<sup>[22]</sup> and was designed methodologically according to the “Standards for Systematic Reviews.”<sup>[23]</sup>

The study protocol was submitted in PROSPERO website (<http://www.crd.york.ac.uk/PROSPERO>) with the registration number CRD42018093219.

### Data source and search strategy

A systematic review of related texts was performed in Medline/PubMed, Cochrane Library, Scopus, Web of Science, Embase, and Google Scholar from inception to August, 2018 using selected MeSH terms related to the studied topic, including “Gaucher's Disease,” “Glucosylceramide lipidosis,” “Imiglucerase,” “Glucosylceramidase,” and “Cerezyme.” In addition, the list of references of the key studies and the review

papers, which could have been overlooked in web search were scanned for more citations. Grey literature search was also conducted for unpublished sources.

### Study selection and data extraction

The studies were selected regarding predefined participants/intervention/comparison/outcome (PICO) for this review. All randomized, quasi-randomized controlled, clinical trials and retrospective cohort studies evaluating the efficacy of Imiglucerase against Eliglustat in patients (male/female) of any age with GD Type 1 were included in the review. The review was performed by completing the “defining a question and eligibility criteria” checklist. Imiglucerase (of any dose) was compared with Eliglustat (of any dose). The primary outcomes were including hemoglobin concentration, platelet count, liver size, and spleen size. The secondary outcomes were the immunological side effects of medicines and bone complications. The discovered studies were excluded from qualitative analysis if: (a) study population was GD Types 2 and 3; (b) study evaluate the efficacy of Imiglucerase and Eliglustat as combined therapy with other regimes; (c) study outcomes of the measure were not similar to ours; (d) it is conference abstracts, case reports, letters, reviews, or comments; and (g) study language was other than English.

After dropping the repeated cases, two authors (A.N. and B.A.) reviewed the search results independently. First, the results were screened by their titles and irrelevant results were excluded. Second, the abstracts of the selected results were reviewed to eliminate conference abstracts, case reports, letters, reviews, or comments. Third, the full text of the chosen studies were reviewed separately. Then, the two authors held a face-to-face meeting to compare their results with each other. The differences between both authors were resolved through the discussion and mediation of the third researcher (M.D.). Finally, the requested data were extracted and summarized in the data sheet. The relevant data including the name of first author, publication date, place of the study, intervention, comparator, study design, sample size, follow-up period, and outcomes (hemoglobin level, platelet count, the liver, and spleen size) were presented in the tables. After primary analysis, it was clear that because of different metrics of the same outcome, dissimilar study designs, non-normality of data, we were not able to conduct a meta-analysis or a quantitative analysis. Therefore, a qualitative analysis of the data was conducted.

### Quality and risk of bias assessment

Methodological quality was assessed independently by two authors according to the Cochrane Collaboration

Handbook for randomized clinical trial (RCT) studies and Critical Appraisal Skills Program for cohort study.<sup>[24]</sup> Likewise, the risk of bias within each included study was assessed based on the random sequence generation, allocation concealment, blinding, incomplete outcome data, and selective outcome reporting domains and reported with ratings of “low risk of bias,” “high risk of bias,” and “unclear” (uncertain risk of bias).

## RESULTS

### Characteristics of included studies

Out of 2979 studies (1788 from PubMed, 84 from Cochrane Library, 462 from Web of Science [Institute for Scientific Information], 561 from Embase, and 67 from Scopus), 23 were qualified for the review in the first step. However, in the second step, twenty more studies were excluded based on the exclusion criteria. Finally, three studies were recognized to meet the inclusion criteria; two RCTs and one retrospective cohort study. The features of these studies have been shown in Tables 1 and 2. The flow diagram of the selection process has been shown in Figure 1. And electronic search strategy in PubMed database are shown in Figure 2.

### Study quality and risk of bias

The risk of the incidence of possible errors in RCTs is summarized in Table 3. The details of randomization method had been described in the selected two RCTs and show that these studies had low risk of bias. Since the participants were taking an oral drug or intravenous Imiglucerase infusions, it was impossible to mask to the treatment allocation or blinding the participants. Therefore, these studies had a high risk of blinding bias in this stage. Nonetheless, since all components of the composite efficacy end-points were examined by central readers were masked, this risk of bias was handled appropriately. The bias of incomplete outcome data was unknown in both studies.

### Efficacy outcome

The results showed that there is no meaningful difference between the two medicines in terms of increasing the hemoglobin level, platelets count, as well as in reducing the liver and spleen size. In the study by Cox *et al.*, reducing the liver and spleen size were the same in patients treated by Eliglustat compared to those treated by Imiglucerase. The mean bone mineral density was in the normal range and maintained; mean bone marrow burden scores showed moderate infiltration of hemopoietin marrow and were also maintained.<sup>[25]</sup> In a 12-month ENCORE trial, Pleat *et al.* found that Eliglustat was non-inferior to Imiglucerase in maintaining stability in adult patients who had reached

Table 1: The characteristics of all included eligible studies

Study	Year	Country	Design	n	Follow-up, Intervention/Control	Outcome measures		Safety				References
						eliglustat	Imiglucerase	Diarrhea	Arthralgia	Fatigue	Headache	
Cox <i>et al.</i>	2015	UK	RCT	160	12 month	Imiglucerase/30-130 U/kg every 2 weeks/50, 100, 150 mg twice daily*	Eliglustat	12%	15%	14%	14%	[25]
						Imiglucerase/Mean: 35, range: 15-60/50, 100, 150 mg twice daily*	Imiglucerase	4%	17%	2%	2%	
Ibrahim <i>et al.</i>	2016	USA	Cohort	121	12	Imiglucerase/30-130 U/kg every 2 weeks/50, 100, 150 mg twice daily*	Eliglustat	NR	NR	NR	NR	[26]
Pleat <i>et al.</i>	2016	USA	RCT	30	12	Imiglucerase/30-130 U/kg every 2 weeks/50, 100, 150 mg twice daily*	Eliglustat	NR	NR	NR	NR	[27]
						Imiglucerase/Mean: 35, range: 15-60/50, 100, 150 mg twice daily*	Imiglucerase	NR	NR	NR	NR	

\*Depending on the plasma concentration of eliglustat ( $\geq 5$  ng/mL ( $\geq 12$  nmol/L) or  $< 5$  ng/mL ( $< 12$  nmol/L)). NR=Not recorded, RCT=Randomized clinical trial

**Table 2: The outcomes of all included eligible studies**

Outcome	Intervention	Cox study, mean change (%)	Ibrahim study, mean baseline (SD)	Pleat study
Hemoglobin concentration (g/L)	Imiglucerase	0.4 (0.4)	12.2±1.66	100% patient met stability criterion
	Eliglustat	-2.1 (-1.1)	11.1±1.67	95% patient met stability criterion
Platelet (×10 <sup>9</sup> /L)	Imiglucerase	6.0 (2.9)	74.7±19.97	100% patient met stability criterion
	Eliglustat	9.5 (3.8)	66.4±20.14	95% patient met stability criterion
Liver volume (MN)	Imiglucerase	0.03 (3.6)	1.5±0.48	83% patient met stability criterion
	Eliglustat	0.02 (1.8)	1.8±0.63	100% patient met stability criterion
Spleen volume (MN)	Imiglucerase	-0.1 (-3.0)	14.4±9.8	100% patient met stability criterion
	Eliglustat	-0.2 (-6.2)	20±12.8	100% patient met stability criterion
Total BMB1 score	Imiglucerase	8.12 (2.63)	NA <sup>3</sup>	In moderate infiltration range
	Eliglustat	8.22 (2.66)	NA	In moderate infiltration range
Femur BMD2 T score	Imiglucerase	-0.54 (1.38)	NA	In reference (normal) range
	Eliglustat	-0.15 (1.09)	NA	In reference (normal) range

BMB=Bone marrow burden, BMD=Bone mineral density, NA=Not available, SD: Standard deviation, MN=Multiple of normal

**Table 3: Risk of bias summary: Author's judgment about each risk of bias item for each included study**

Study	Random sequence generation (selection bias)	Allocation concealment (selection bias)	Blinding of participants and personnel (performance bias)	Blinding of outcome assessment (detection bias)	Incomplete outcome data (attrition bias)	Selective reporting (reporting bias)	Other bias
Cox 2015	Low	High	High	Low	Unclear	Low	Unclear
Pleat 2016	Low	High	High	Unclear	Unclear	Low	Unclear

stability by the administration of Imiglucerase or Velaglucerase. This *post hoc* analysis was studied safety and efficacy of Velaglucerase in 30 ENCORE patients. The patients were randomly divided into two groups; Eliglustat ( $n = 22$ ) or Imiglucerase ( $n = 8$ ). In this study, 90% of the patients who switched to Eliglustat and 88% of individuals who switched to Imiglucerase showed a stable hemoglobin level, platelet count, as well as liver and spleen size. The results showed that the mean baseline bone mineral density scores for both lumbar spine and femur were in the normal range. Likewise, the mean bone marrow burden scores were in the moderate infiltration range. The patients who transitioned from Velaglucerase alfa to Eliglustat had stable bone measures after 12 months.<sup>[26]</sup>

In the study by Ibrahim *et al.*, parameters improved from baseline in both treatment groups, with a time course and degree of improvement in Eliglustat-treated patients were similar to Imiglucerase-treated patients.<sup>[27]</sup> The outcomes of all included studies are summarized in Table 2. Among the evaluated Registry cohort of the patients, adverse event data were not recorded and sufficient data on bone were not available.

### Safety outcome

The most common adverse events were diarrhea, arthralgia, fatigue, and headache. ECG analysis showed no significant effect of Eliglustat on heart rate or cardiac repolarization. Eliglustat had a 2–3 ms (upper bound of 90% confidence interval 4.8 ms) effect on cardiac depolarization (QRS duration), which was not

time-dependent or dose-dependent, and only slightly greater than that of Imiglucerase.<sup>[25,26]</sup>

Four serious adverse events including appendicitis, syncope, ischemic colitis, and uterine leiomyoma were also reported. All adverse events were considered unrelated to treatment and none resulted in study withdrawal.<sup>[26]</sup>

### DISCUSSION

The aim of this study was to compare the efficacy of Imiglucerase with Eliglustat in patients with GD Type 1. There are few randomized controlled studies, which compare Imiglucerase with Eliglustat. Moreover, they have insufficient sample size and short follow-up period, and they report inadequate therapeutic outcomes in people with GD1. However, all selected studies have concluded that Imiglucerase and Eliglustat are the same in terms of their effect on hemoglobin level, platelets count, as well as the liver and spleen size. The narrative review by Belmatoug *et al.* in 2016 showed that Eliglustat is effective in bone synthesis while Imiglucerase has little role in bone synthesis. Also, it was found that there is no difference between Imiglucerase and Eliglustat in terms of their effect on hemoglobin level, platelets count, as well as the liver and spleen size, which is the same as the findings of the present study.<sup>[28]</sup>

In 2014, a systematic review by Smid *et al.* demonstrated that Imiglucerase and Eliglustat are the same in terms of their effect on hemoglobin level, platelets count, as well as the liver and spleen size. In addition, it was found

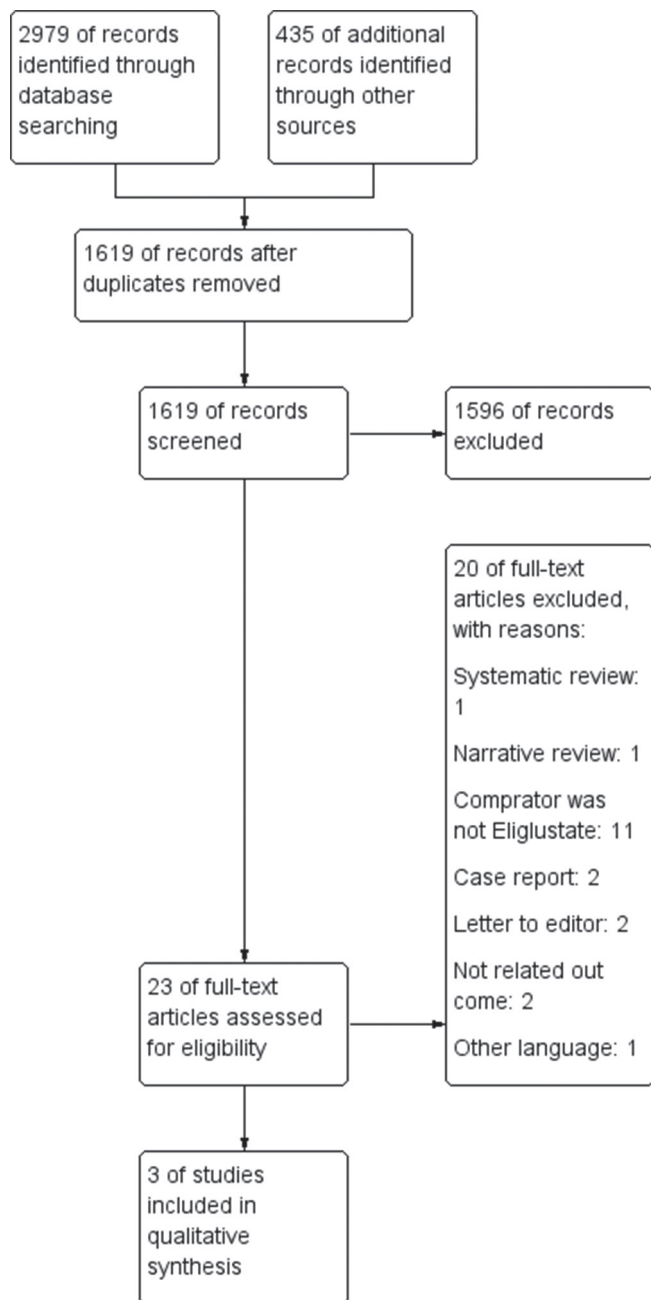


Figure 1: Process of the screening of studies

that Eliglustat is more effective in bone synthesis than Imiglucerase, which is supporting by our findings.<sup>[29]</sup>

The most common adverse events were diarrhea, arthralgia, fatigue, and headache. However, frequency of side effects was 10% more with Eliglustat than that with Imiglucerase.<sup>[25,26]</sup> The main side effects of Imiglucerase are immune response (15%) and hypersensitivity reactions (50%).<sup>[28,30,31]</sup> Gastrointestinal disorders were not recorded in Eliglustat.<sup>[29,32]</sup>

The latest study published by Cox *et al.* in 2017 revealed that patients received Eliglustat for 4 years showed no serious side effects and the drug were well tolerated.<sup>[33]</sup>

Search	Add to builder	Query	Items found	Time
#13	Add	Search (((Gaulcher[Title/Abstract]) OR "Gaulcher Disease"[Mesh]) OR Glucosylceramide Lipidosis[Title]) AND (((Imiglucerase) OR Cerezyme) OR Glucosylceramidase[MeSH Terms]) OR Glucosylceramidase[Title/Abstract]	1788	01:38:02
#14	Add	Search (((Gaulcher[Title/Abstract]) OR "Gaulcher Disease"[Mesh]) OR Glucosylceramide Lipidosis[Title]) AND (((Imiglucerase) OR Cerezyme) OR Glucosylceramidase[MeSH Terms]) OR Glucosylceramidase[Title/Abstract] Filters: Humans	1661	01:34:58
#12	Add	Search (((Imiglucerase) OR Cerezyme) OR Glucosylceramidase[MeSH Terms]) OR Glucosylceramidase[Title/Abstract]	2256	01:33:46
#11	Add	Search ((Gaulcher[Title/Abstract]) OR "Gaulcher Disease"[Mesh]) OR Glucosylceramide Lipidosis[Title]	5618	01:33:00
#10	Add	Search Glucosylceramidase[Title/Abstract]	124	01:32:07
#9	Add	Search Glucosylceramidase[MeSH Terms]	2120	01:31:21
#7	Add	Search Cerezyme	389	01:25:09
#6	Add	Search Imiglucerase	371	01:24:49
#5	Add	Search Glucosylceramide Lipidosis[Title]	2	01:24:22
#4	Add	Search "Gaulcher Disease"[Mesh]	4409	01:22:48
#2	Add	Search Gaulcher[Title/Abstract]	5029	01:21:02

Figure 2: Electronic search strategy in PubMed database

Before early 1990s, when ERT was recognized as the exclusive treatment for GD “symptomatic treatment” (that is any medical therapy of a disease that only affects its symptom, not its cause) had been used with the support of many observational studies. Enzyme infusion improved the blood and visceral complications of the patients,<sup>[34]</sup> and this improvement meaningfully enhanced their quality of life, in comparison with chronic patients. However, intravenous injection for people with hard-to-find veins and the long-term use of catheter for children could be troublesome. Generally, lifelong intravenous injection, high cost, and lack of entrance to the nervous system are the main limitations of treatment with ERT.<sup>[34]</sup>

SRT products have some advantages, which may make them superior to ERT. SRT minimizes the accumulation of excess material by inhibiting material synthesis. The biggest advantage of SRT is its oral administration, which makes it easier to cross the blood–brain barrier and reach other organs. The effect of Eliglustat on bone metabolism is another advantage of this agent over Imiglucerase.<sup>[17]</sup>

Although extensive knowledge about efficacy of Eliglustat are currently sparse, it is undeniable that the hematological and visceral responses of GD1 patients to Eliglustat are clinically related. However, since there was no direct and head-to-head comparison with sufficient sample size and lengthy follow-up period with treatment-naïve study population that consist of sever patients with bone disease and splenectomies, it is not reasonable to state sharply that these results are exactly equivalent with Imiglucerase. Moreover, the potential effects of Eliglustat in long-term complications and its associated conditions need more investigation. For example in ENGAGE trial, only mild to moderate patients were included in the study and there was no results for sever patients.<sup>[29]</sup>

It is claimed that heterozygous mutations in GBA gene is a common risk factor for the development of

Parkinson's disease in GD patients and in heterozygous GBA mutation positive carriers. There is no evidence for supporting the effectiveness of Imiglucerase and Eliglustat in the treatment of Parkinson's disease and peripheral neuropathy, which are the main and most dangerous outcomes in middle-aged and adult patients with GD.<sup>[35-38]</sup>

Furthermore, to predict CYP2D6 metabolizer status and to find appropriate dosing of Eliglustat, determination of the CYP2D6 genotype by testing a blood sample at a nationally accredited laboratory is essential. Eliglustat is approved in the European Union and Food and Drug Administration for adult patients who are predicted to be extensive, intermediate, or poor metabolizers. Eliglustat is not approved for patients in whom genotyping indicates CYP2D6 ultra-rapid and indeterminate metabolize since these patients may not reach adequate Eliglustat concentrations to achieve appropriate therapeutic effect. Nevertheless, Eliglustat interacts with drugs metabolized by CYP2D6 and it cannot be used in patients with heart, liver disease, kidney diseases, breastfeeding, pregnant women, and people above 65 years of age.<sup>[28]</sup> This is important to not that there are no such restrictions for prescription of Imiglucerase.

Although we performed an extensive literature search and consulted with the study authors to ensure that all the relative data were included and accurately interpreted, this review is nonetheless significantly constrained. There are few randomized controlled studies, which compare Imiglucerase with Eliglustat. Moreover, they have insufficient sample size and short follow-up period, and they report inadequate therapeutic outcomes in people with GD1. Although we did a quality assessment of studies, because of the limited number of studies, we did not meet our included criteria according to the quality of studies. In addition, as a result of small number of participants included in the study, the low quality of the methodology and inadequacy of information reported in the studies, as well as the difference in methodologies of the studies, it was impossible to conduct a meta-analysis or a quantitative analysis.

## CONCLUSION

Eliglustat is a very promising alternative for ERT with regard to its effects on hematological and visceral abnormalities. Currently, further investigations are needed to determine whether it is as effective in patients with severe disease, especially with symptomatic bone disease or whether it is safe in patients with polyneuropathy. Its superiority to ERT with respect to prevention of long-term complications and associated conditions needs further study as well. Frequently prescribed concomitant

medications, which are metabolized by CYP2D6 and cardiovascular disease will probably restrict the prescription of Eliglustat.

## AUTHORS' CONTRIBUTION

Azita Nabizadeh, Bahman Amani, and Majid Davari contributed in searching databases and extracted data from selected articles. Maliheh Kadivar, Akbar Abdollahi Asl, Mehdi Toreski, and Yahya Baiazidi contributed in the study concept and quality analysis of selected articles. Azita Nabizadeh, Bahman Amani, and Majid Davari contributed in quality analysis of selected articles. All authors contributed in manuscript preparation and final editing.

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## Conflicts of interest

There are no conflicts of interest.

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