SYSTEMATIC REVIEW

Application of genus Cassia in the treatment of Constipation: A systematic review [version 1; peer review: 2 approved with reservations]

Muhammad Shahzad Aslam
Department of Chemistry, Xiamen University Malaysia, Sepang, Selangor, 43900, Malaysia

Abstract
Purpose: Role of genus cassia in the treatment of Constipation
Methods: Methodological analysis, systematic review, and meta-analysis of identified studies using RevMan
Result and Discussion: Cassia fistula was partially effected in treating constipation however there is a need for improvement in the protocol of studies to reduce biases. These results were only limited to one species so it cannot be generalized among all species of Cassia.
Conclusion: Cassia fistula is partially effective in reducing the pain and consistency of stool during constipation among children.

Keywords
Cassia, Cassia fistula, Constipation, Pediatric gastroenterology

Open Peer Review

Reviewer Status ️
Invited Reviewers
version 1
1 Massimo Bellini, University of Pisa, Pisa, Italy
2 Farhad Shokraneh, University of Nottingham, Nottingham, UK

Any reports and responses or comments on the article can be found at the end of the article.

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Author roles: Aslam MS: Conceptualization, Data Curation, Formal Analysis, Funding Acquisition, Investigation, Methodology, Project Administration, Resources, Software, Supervision, Validation, Visualization, Writing – Original Draft Preparation, Writing – Review & Editing

Competing interests: No competing interests were disclosed.

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**Introduction**

Constipation is a clinical disorder attributed to ineffectual colonic function and/or increased resistance to the proliferation of colonic markers. Approximately 20% of the world population suffers from chronic constipation. It is one of the most common pediatric problems. It was found to be the second most stated disorder in the field of pediatric gastroenterology. Treatment costs for children with constipation will be around three times higher than children without constipation in the United States. African American children, particularly girls, are greatly affected by constipation, which has been associated with poor hygiene conditions.

Commonly, constipation is treated by Cisapride in children, other treatments include polyethylene glycol 3350 and lactulose, however polyethylene glycol 3350 has been found to be more effective. Supplemented and non-supplemented yoghurt helps in reducing abdominal pain and to enhance defecation frequency. It has been observed that different species of *Cassia* act as a laxative such as *Cassia fistula*, *Cassia alata*, and *Cassia augustifolia*. The genus *Cassia* is well known in alternative medicine as hepatoprotective, laxative, and in the treatment of ringworm infection, skin diseases and leprosy. It has many pharmacological properties including acting as a hypolipidemic agent, anti-microbial, anti-fungal, and anti-cancer agent. *Genus Cassia* contains a number of bioactive compounds such as anthraquinone, tannin, coumarins, triterpene, volatile oil, phenolic glycoside, flavonoids from different parts of the plant. Different species of *Cassia* possess laxative properties due to various anthraquinone derivative such as aloe-emodin, rhein, chrysophanol and chrysobutin. In this review, we systematically assessed the laxative potential of different species of *Cassia*.

**Methods**

**Literature search strategy**

A systematic literature search was conducted in accordance with the Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA) guidelines. Using the keywords (Senna AND Laxatives AND Clinical trial) (Cassia AND Laxatives AND Clinical trial) (Senna AND Clinical trial) AND publication range from 01 January 1960 till 31 December 2018 for identification of the records. Table 1 shows the search strategy for PubMed Central. During screening of the records only full-length open access articles were considered. Abstract only or closed access articles were excluded. Only articles involving children aged between 2–15 were included. All review articles, *in-vivo* studies and those >10 years from the search data were excluded. A preliminary search of the PubMed, CNKI, Scopus, Web of Science, Google Scholar and PsydINFO databases and digital archive such as PubMed Central yielded 2207 papers published in English from the last ten years. Duplicate and irrelevant articles were removed (n=2203). One article was further removed during screening due to closed access (n=3). One publication was removed because the article did not meet the eligibility criteria (n=2). A PRISMA Flow Diagram is given in Figure 1.

**Literature screening**

Identification of articles were performed at level 1 using the search strategy as mentioned in Table 1. Duplicate articles, irrelevant articles such as polyherbal formulation, review articles, or any article other than Cassia or Senna were removed at level 2. Only four articles were identified as being relevant. One record was excluded due to not being a full text article. Abstracts were being reviewed for the following inclusion and exclusion criteria at level 4 and one article was removed for not meeting the eligibility criteria i.e. Randomized, clinical trial on Constipation, full-length open access articles, Pediatric Functional Constipation (age range: 2–15 years).

**Eligibility criteria**

**Types of studies.** The author has selected studies of randomized open label, prospective, controlled, parallel-group clinical trial for meta-analysis. Baseline characteristics of randomized trials of studies included on pediatric functional constipation are presented in Table 4. Characteristics of the studies included are mentioned in Table 5.

**Types of participants.** The author included studies involving patients (aged 2–15 years) with Functional constipation. The diagnosis of Functional constipation was according to according to the Rome III criteria. Inclusion and exclusion criteria were based on Study design, participants, intervention, outcome (SPIO) criteria and indicated in Table 2.

**Table 1. Search strategy.**

<table>
<thead>
<tr>
<th>PubMed Central</th>
<th>CNKI</th>
</tr>
</thead>
</table>
Types of interventions. Included studies were focused on the role Cassia in the treatment of Functional constipation. Unfortunately, there were only two studies identified.

Types of outcomes. Eligible studies included consisted of the following outcomes: improvement in the episodes of fecal incontinence per week, improvement in the episodes of retentive posturing per week, improvement in the average of severity of pain of defecation (by VAS), improvement in defecation frequency per week, patient’s drug compliance and improvement in the average of consistency of stool defecated (by VAS).

Methodological quality assessment (MQ)
Methodological quality assessment was made on the basis of following criteria. 1) Aims and Hypothesis clearly defined, adequate sample representation, patient care quality, ethical approval protocol, outcomes assessment, validity and reliability of outcome measure, attempt to blind researcher, follow-up, appropriate statistical analysis and missing data reported. Ten item defined evaluation of methodological quality (MQ) is presented in Figure 2. Risk of Bias were assessed using Cochrane collaboration’s tool on the basis of the following criteria such as selection bias, performance bias, attrition bias, reporting bias and miscellaneous. Cochrane Collaboration’s tool for assessing the risk of bias was used and the results are presented in Table 6.

Data extraction
The following data were extracted according to study characteristics (e.g., first author, year of publication, search dates, and number of included studies), patient characteristics (functional constipated children, aged between 2–15), sample size, study type (e.g., Randomized open label, prospective, controlled,
parallel-group clinical trial study), randomization methods (e.g., "systematic randomization and simple randomization") and outcome measures/variables (e.g., improvement in defecation frequency per week, improvement in the episodes of fecal incontinence per week, improvement in the episodes of retentive posturing per week, improvement in the average of severity of pain of defecation (by VAS), improvement in the average of consistency of stool defecated (by VAS) and patient’s drug compliance). Data extraction was performed by Muhammad Shazad Aslam. Transcripts were analysed, coded and data was extracted using the demo version of qualitative data analysis software Atlas.ti 8.0. Table 3 represent all the data that was extracted. All the meta-data are available as Dataset 1.

**Statistical analysis**

Meta-analysis was conducted using the Review Manager (RevMan) 5.3 software. The summary measures were reported as odds ratios (ORs) or as a standard mean difference (SMD) with 95% confidence intervals (CI). The presence of heterogeneity among trials was assessed using the Chi-square test.

### Table 2. Study design; participants; intervention; outcome (SPIO) criteria.

<table>
<thead>
<tr>
<th>Inclusion Criteria</th>
<th>Exclusion Criteria</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Study design</strong></td>
<td>Randomized, clinical trial on Constipation, full-length open access articles, Pediatric Functional Constipation</td>
</tr>
<tr>
<td><strong>Participants</strong></td>
<td>Children (age range: 2–15 years)</td>
</tr>
<tr>
<td><strong>Intervention</strong></td>
<td>Cassia fistula was delivered to Pediatric with Functional Constipation</td>
</tr>
<tr>
<td><strong>Outcomes</strong></td>
<td>Role of the Cassia fistula emulsion in Pediatric Functional Constipation</td>
</tr>
</tbody>
</table>

**Figure 2.** Methodological quality assessment of the 2 studies included in the meta-analysis (0=No/not reported, 1=Yes).
test, and the extent of the inconsistency was measured by I2 statistics. Output file from RevMan is available as Underlying data.

### Results and discussion

Both of the selected studies were not blinded during intervention and outcome assessment that will result in performance bias and detection bias respectively. These biases occur where the investigators know about the participant’s treatment group. Performance bias can also refer to the fact that participants can change their responses or behaviour if they know which group they are allocated in. Blinding of outcome assessment may decrease the risk of the investigator or participant being aware of the treatment that a patient is receiving. If the participants and the caregivers are aware of the intervention and outcome that may affect the behavior of the participants, these behavioral changes may affect the performance of the treatment. Clinical trials on adults were also excluded, such as a randomized clinical trial of a phytotherapeutic compound containing *Pimpinella anisum, Foeniculum vulgare, Sambucus nigra*, and *Cassia augustifolia* for chronic constipation. Results of both included studies were non-significant when comparing their baseline characteristics of pediatric functional constipation as presented in Table 5. During analysis of study characteristics, it was found that both of studies demonstrated *Cassia fistula* is helping to treat constipation among the children as shown in Table 3, but there is a risk of bias according to Cochrane Collaboration’s tool (Table 6).

Moreover, both studies found were from one country (Iran). During a methodological assessment, many flaws were identified such as inadequate patient care, attempt to blind the researcher and missing data (Figure 2). During meta-analysis, the comparison was made before and after treatment among different variables such as defaecation, fecal incontinence, retentive posturing, the severity of pain, and consistency of stool. All the variable (before and after treatment) were found to be symmetrical when plotted on a funnel plot as shown in Figure 4, Figure 6, Figure 8, Figure 10, Figure 12, and Figure 14 respectively. The overall effect for some variables is statistically insignificant (P=0.11, P=0.49, P=0.24) such as fecal incontinence, retentive posturing, and acceptance, tolerance respectively. High heterogeneity was found in two variables i.e severity of pain (90%) and consistency of stool (77%). All the forest plot of defaecation,

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**Table 3. Data extraction.**

<table>
<thead>
<tr>
<th>Characteristics</th>
<th>Seyyed Ali Mozaffarpur</th>
<th>Mohammad Reza Esmaeilidooki</th>
</tr>
</thead>
<tbody>
<tr>
<td>Publication Year</td>
<td>2012</td>
<td>2016</td>
</tr>
<tr>
<td>Type of Study</td>
<td>Randomized, clinical trial</td>
<td>This randomized open label, prospective, controlled, parallel-group clinical trial study</td>
</tr>
<tr>
<td>Age</td>
<td>Age between 4–13 years</td>
<td>Aged between 2 – 15 years</td>
</tr>
<tr>
<td>Randomization</td>
<td>systematic randomization</td>
<td>Simple randomization</td>
</tr>
<tr>
<td>Total Sample Size</td>
<td>81</td>
<td>51</td>
</tr>
<tr>
<td>Variables</td>
<td>Frequency of defecation, consistency of stools, and severity of pain during defecation, retentive posturing and fecal incontinence per week</td>
<td>Improvement in defecation frequency per week, improvement in the episodes of fecal incontinence per week, improvement in the episodes of retentive posturing per week, improvement in the average of severity of pain of defecation (by VAS), improvement in the average of consistency of stool defecated (by VAS) and patient’s drug compliance</td>
</tr>
<tr>
<td>Length of each contact with the participant/caregivers</td>
<td>Clinical efficacy and tolerance were assessed using weekly sheets, parents completed every night. They were given three sheets (included seven questions in seven columns) to complete them daily for 3 weeks.</td>
<td>During the study, we had regular phone calls with the parents to check the probable complications, treatment (taking the prescribed drugs) and data filling process. If there were any serious questions or problems, we visited the child. At the end of 4 weeks of treatment, the children were visited and the filled out forms were taken and evaluated.</td>
</tr>
<tr>
<td>Blinding of Experiment</td>
<td>The investigators, the children and their parents were aware of the study group assignment.</td>
<td>Fortunately, due to the developing socioeconomic conditions of the people in these regions in recent years, we were able to keep in touch with all the patients during the study period by phone call.</td>
</tr>
<tr>
<td>Diagnosis</td>
<td>Rome III criteria of functional constipation</td>
<td>Rome III criteria of functional constipation</td>
</tr>
<tr>
<td>Diagnostic test</td>
<td>Paraclinics like anorectal manometry, thyroid function tests, anti-tTG, and etc. If it remained any doubt, barium study and anorectal manometry would be performed</td>
<td>thyroid function tests, anti-tTG, and etc.</td>
</tr>
</tbody>
</table>
Table 4. Baseline characteristics of randomized trials of studies included in pediatric functional constipation.

<table>
<thead>
<tr>
<th>Variable in Treatment (T)</th>
<th>Mozaffarpur, 2012 (T)</th>
<th>Esmaeilidooki, 2016 (T)</th>
<th>Mozaffarpur, 2012 (C)</th>
<th>Esmaeilidooki, 2016 (C)</th>
<th>Variable in Control (C)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sample size in Treatment (N)</td>
<td>41</td>
<td>52</td>
<td>40</td>
<td>57</td>
<td>Sample size in Control (N)</td>
</tr>
<tr>
<td>Age, months, Mean±SD</td>
<td>69.4(±24.3)</td>
<td>64.6(±25.2)</td>
<td>65.9(±19.1)</td>
<td>55.2(±31.2)</td>
<td>Age, months, Mean±SD</td>
</tr>
<tr>
<td>Sex, Male, (n) (%)</td>
<td>29(70.7%)</td>
<td>33</td>
<td>23(57.5%)</td>
<td>30</td>
<td>Sex, Male, (n) (%)</td>
</tr>
<tr>
<td>Weight, Kg Mean±SD</td>
<td>21.7 (±7.2)</td>
<td>20.5(±7.2)</td>
<td>20.7(±7.8)</td>
<td>18.5(±8.9)</td>
<td>Weight, Kg(±SD)</td>
</tr>
<tr>
<td>Duration of constipation, months, Mean±SD</td>
<td>34.2(±25.9)</td>
<td>31.1(±24.6)</td>
<td>30.8(±22.8)</td>
<td>23.5(±21.8)</td>
<td>Duration of constipation, months, Mean±SD</td>
</tr>
<tr>
<td>Defecation ≤ 2 per week, n (%)</td>
<td>32(78%)</td>
<td>41</td>
<td>30(75%)</td>
<td>52</td>
<td>Defecation ≤ 2 per week, n (%)</td>
</tr>
<tr>
<td>Incontinence, n (%)</td>
<td>31(75.6%)</td>
<td>34</td>
<td>27(67.5%)</td>
<td>37</td>
<td>Incontinence, n (%)</td>
</tr>
<tr>
<td>History of previous treatment, n (%)</td>
<td>32 (78%)</td>
<td>43</td>
<td>28(70%)</td>
<td>51</td>
<td>History of previous treatment, n (%)</td>
</tr>
<tr>
<td>Retentive posturing, n (%)</td>
<td>32(78%)</td>
<td>40</td>
<td>29(72.5%)</td>
<td>37</td>
<td>Retentive posturing, n (%)</td>
</tr>
</tbody>
</table>

Table 5. Study characteristics.

<table>
<thead>
<tr>
<th>Author, year</th>
<th>Study Design</th>
<th>Hypothesis</th>
<th>Statistical analysis</th>
<th>Software</th>
<th>Outcomes</th>
</tr>
</thead>
<tbody>
<tr>
<td>Mozaffarpur, 2012</td>
<td>Randomized, clinical trial</td>
<td>The author hypothesized that <em>Cassia fistula</em> emulsion (CFE) would be as effective or better than Mineral oil (MO) in treating FC.</td>
<td>The statistical analyses included the determination of means and Standard deviation (SDs), t test, ( \chi^2 ) test, ANOVA repeated measures and Fisher's exact test, with significance accepted at the 5% level.</td>
<td>SPSS (version 17),</td>
<td>CFE was most effective than MO in the 3-week treatment of children with FC.</td>
</tr>
<tr>
<td>Esmaeilidooki, 2016</td>
<td>Randomized, clinical trial</td>
<td>N/A</td>
<td>The statistical analyses included the determination of means and SDs, t test, ( \chi^2 ) test, ANOVA repeated measures and Fisher's exact test, with significance accepted at the 5% level.</td>
<td>SPSS IBM20 and STATA 11.2</td>
<td>Significant improvement when compared with the control group however unable to find substantial evidence of the role of identified bioactive compounds due to limitation as it requires further investigation</td>
</tr>
</tbody>
</table>

Table 6. Cochrane Collaboration’s tool for assessing risk of bias.

<table>
<thead>
<tr>
<th>Study (author, year)</th>
<th>Selection bias</th>
<th>Performance bias</th>
<th>Detection bias</th>
<th>Attrition bias</th>
<th>Reporting bias</th>
<th>Other bias</th>
</tr>
</thead>
<tbody>
<tr>
<td>Mozaffarpur, 2012</td>
<td>Unclear</td>
<td>Yes</td>
<td>Yes</td>
<td>No</td>
<td>No</td>
<td>Unclear</td>
</tr>
<tr>
<td>Esmaeilidooki, 2016</td>
<td>Unclear</td>
<td>Yes</td>
<td>Yes</td>
<td>No</td>
<td>No</td>
<td>Unclear</td>
</tr>
</tbody>
</table>

Fecal incontinence, retentive posturing, severity of pain, consistency of stool and acceptance and tolerance are represented in Figure 3, Figure 5, Figure 7, Figure 9, Figure 11 and Figure 13 respectively.

Conclusion

After evaluation of results, it was found that *Cassia fistula* was not completely effective. It was partly effective in reducing the pain and consistency of stool during constipation. However, these results cannot be generalized among all population. A well designed, expert validated protocol is required in the future. There is a need to develop an instrument that will be free from bias. Moreover, the results cannot be generalized among all species of Cassia as the studies available are only for one species. There is a need to isolate identified bioactive compounds from different species of Cassia and evaluate the effect of different factors such as duration of constipation, defecation, incontinence or retentive posturing under clinical trial.
Figure 3. Forest plot in defaecation before and after treatment.

Figure 4. Funnel plot showing overall standardized mean difference in defaecation before and after treatment.

Figure 5. Forest plot in fecal incontinence before and after treatment.
Figure 6. Funnel plot showing overall standardized mean difference in fecal incontinence before and after treatment.

Figure 7. Forest plot in retentive posturing before and after treatment.

Figure 8. Funnel plot showing overall standardized mean difference in retentive posturing before and after treatment.
Figure 9. Forest plot in severity of pain before and after treatment.

Figure 10. Funnel plot showing overall standardized mean difference in severity of pain before and after treatment.

Figure 11. Forest plot in consistency of stool before and after treatment.
Declarations

Data availability


This project contains the following underlying data:
- Cassia Senna for Constipation.rm5 (study RevMan file)
- Quotation Manager.xlsx (study characteristics of citations included in this study)

Reporting guidelines

Open Science Framework: PRISMA diagram and flowchart for the study “Application of genus Cassia in the treatment of constipation: A systematic review”. https://doi.org/10.17605/OSF.IO/PKR4N
References


Grant information
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The funders had no role in study design, data collection and analysis, decision to publish, or preparation of the manuscript.
Open Peer Review

Current Peer Review Status: 

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Farhad Shokraneh
University of Nottingham, Nottingham, UK

The author investigates the effect of *Cassia fistula* on three outcomes based on two included studies. It seems a valuable research that could potentially direct the pharmaceutical companies toward a new agent to treat the constipation.

**Title**
The title should be “Cassia fistula for treatment of children and adolescents with constipation: A systematic review of randomized controlled trials”.

**Authorship**
The systematic review requires a process that involves at least two people for screening and data extraction. I suggest the author to find another systematic reviewer to be involved in this work.

**Abstract**
The abstract is very brief and sentences are incomplete. Abstract should be re-written following the example from similar systematic reviews and following PRISMA for Abstract.

**Minor edits**
1. “act an effective as a laxative” should be “act as effective as laxatives”.
2. “search data” should be “search date”.

**Methods**
1. The standard search filter for finding RCTs as reported in Cochrane Handbook should be used.
2. It is advised in Cochrane Handbook searching Embase, MEDLINE/PubMed, and CENTRAL for systematic reviews of RCTs and this review reports searching only MEDLINE/PubMed.
3. Excluding closed access papers without trying to order them from library or contacting the authors introduces full-text or open access bias.
4. The structure of the method should change and start with 1: Eligibility Criteria 2: Databases and search strategy 3: Screening process 4: Data extraction process 5: Assessment of risk of bias process 6: Data analysis plan. Currently, the first section of the methods reports info on search and eligibility criteria while there are specific subheadings for these sections.

5. The rationale behind excluding studies older than 10 years from search date is not clear.

6. The reported number of search results 2207 is not reasonable. A proper search should not find more than 100-200 results for this topic.

7. The search date has not been reported.

8. Following a published Cochrane review of RCTs in terms of structure and all the items in PRISMA Checklist for full systematic reviews is recommended.

9. Constipation is missing from search strategy and MeSH terms in the table.

10. The numbers in PRISMA flow diagram does not match the numbers reported in the text and it does not seem to be correct. Following one of the existing published systematic reviews of RCTs from Systematic Reviews journal or Cochrane Database of Systematic Reviews is recommended.

11. The quality of included studies is being assessed using Cochrane Risk of Bias tool. The role of MQ and using Atlas.ti is not clear in this review. These two are not usually used for SR of RCTs.

12. Studying the meta-analysis section in the Cochrane Handbook which is freely available online could help revising the statistical analysis.

13. Table 2 should report PICOS not SPIO.

14. The time point for the outcome is not clear.

15. The outcome measure has not been described properly.

16. The complications and adverse effects have not been reported in the review.

**Results**

1. The sample size reported in the Data Extraction table does not match the numbers in meta-analysis. For example the total for experimental and control group in Review Manager file is 109 for Esmaeilidooki 2016 and 71 for Mozaffarpur 2012 while in the table is 51 and 81 respectively.

2. The analysis for before treatment is not required. All before treatments can be deleted.

3. Funnel plots only work with about 10-20 studies. All funnel plots can be deleted.

4. Risk of bias based on Cochrane tool is not clear. It should be low risk, unclear risk and high risk. The current description of Yes/No does not provide enough information.
5. In meta-analysis, “one randomized always analysed” while the numbers in Table 4 are not the numbers in meta-analysis in Review Manager.

6. The control groups in two studies are different so I am not sure if combining the data from two studies in a meta-analysis is a right when there is such heterogeneity in PICOS components.

7. SD is greater than mean is some outcomes which is an indicator of skewed data or error in reporting. Contacting the trials authors might make some of the points clear.

Discussion and Conclusion
Current data and status of the review may not be appropriate for discussion or making a conclusion. The current evidence is limited and more rigorous studies will be required to make any conclusion or recommendations.

I understand the author’s passion for conducting a systematic review and so I request the author to work with an expert supervisor in the field before revising the work.

Are the rationale for, and objectives of, the Systematic Review clearly stated?
Partly

Are sufficient details of the methods and analysis provided to allow replication by others?
Partly

Is the statistical analysis and its interpretation appropriate?
No

Are the conclusions drawn adequately supported by the results presented in the review?
No

Competing Interests: No competing interests were disclosed.

Reviewer Expertise: Evidence synthesis, Systematic review, Overview, Scoping review, Rapid review, Randomized controlled trials, Open data, Open access, Open source, Medical journalism, Peer-review, Open source, Outcomes, Interventions, Cochrane

I confirm that I have read this submission and believe that I have an appropriate level of expertise to confirm that it is of an acceptable scientific standard, however I have significant reservations, as outlined above.
Massimo Bellini  
Gastrointestinal Unit, Department of Translational Research and New Technologies in Medicine and Surgery, University of Pisa, Pisa, Italy

I’m really puzzled about the usefulness of a review which takes into consideration only two papers and altogether only 132 patients. Constipation, as the author himself states, is a very frequent condition, so the conclusions are inevitably flawed by this bias. The author should address this issue in the conclusions. Hence, I have significant reservations.

Moreover, [here] enclosed you’ll find a copy of the paper. The authors can find some suggestions, as tracked changes and comments to the original manuscript, which could improve the quality of the manuscript.

Are the rationale for, and objectives of, the Systematic Review clearly stated?  
Yes

Are sufficient details of the methods and analysis provided to allow replication by others?  
Yes

Is the statistical analysis and its interpretation appropriate?  
Yes

Are the conclusions drawn adequately supported by the results presented in the review?  
Yes

**Competing Interests:** No competing interests were disclosed.

**Reviewer Expertise:** Gastroenterology: Functional Digestive Disorders.

I confirm that I have read this submission and believe that I have an appropriate level of expertise to confirm that it is of an acceptable scientific standard, however I have significant reservations, as outlined above.

Reviewer Response 23 May 2019  
Muhammad Shahzad Aslam, Xiamen University Malaysia, sepang, Malaysia

Dear,

I have read the report. The selected studies were according to the criteria given inside the paper. So, there were only two selected studies for the systematic review. The comments you have mentioned are appreciable. Thanks

**Competing Interests:** No
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