### Association between gastroesophageal reflux disease and depression disorder

A protocol for systematic review and meta-analysis

Yu Liu, MD\(^a\), Panpan Zhou, MD\(^b\), Shixiong Zhang, MD\(^a\), Huiqing Wu, MS\(^b\), Zeqi Yang, MD\(^a\), Miaochan Xu, MD\(^a\), Shaowei Liu, MD\(^a\), Yangang Wang, PhD\(^{a,b}\)

**Abstract**

**Background:** This study will systematically synthesize the evidence on the potential association between gastroesophageal reflux disease (GERD) and depression disorder (DD).

**Methods:** We will search the following electronic bibliographic databases: PubMed/MEDLINE, Embase, Cochrane Library, Web of Science, the Chinese Bio Medical Literature Database, China National Knowledge Infrastructure (CNKI), the China Science and Technology Journal database (VIP) and Wanfang Data. In addition, ongoing trials will be retrieved from the WHO ICTRP Search Portal, the Chinese Clinical Trial Register and The Clinical Trials Register. Articles related to gastroesophageal reflux disease and depression will be searched. And language and time will be unlimited.

**Results:** The study will afford additional insight into the investigation the association between GERD and DD.

**Conclusions:** The results of this study will provide helpful evidence to explore the association between GERD and DD.

**Registration number:** INPLASY202090026.

**Abbreviations:** DD = depression disorder, GERD = gastroesophageal reflux disease, RCTS = randomized controlled trials.

**Keywords:** association, depression disorder, gastroesophageal reflux disease, meta-analysis

1. **Introduction**

According to the Montreal definition, gastroesophageal reflux disease (GERD) is defined as a disease which is associated with troublesome symptoms and/or complications on account of reflux of stomach contents into the esophagus. Most patients with GERD presents with heartburn and effortless regurgitation.[1] GERD is one of the most prevalent gastrointestinal disorders globally, showing an increasing prevalence in several developing countries, but there is considerable geographic variation.[2] The result of a direct pairwise comparison and NMA shows that esophageal reflux monitoring or endoscopy is superior to other diagnostic tests for GERD, which is in accordance with the recommendation in current guidelines.[3,4] It has been reported that the pooled prevalence of at least weekly GERD symptoms reported from population-based studies worldwide is approximately 13%, but there is considerable geographic variation.[5] The disease has a chronic course that leads to a significant decline in the quality of life of patients and is associated with a high economic burden worldwide.[6]

Major depressive disorder, also known as depression, is a common neuropsychiatric disorder that affects more than 300 million people of all ages and is one of the leading causes of global disease burden.[7,8] It is a very common medical condition that is associated with a wide range of emotional, cognitive, and physical symptoms. Depressive disorders involve all major bodily functions, mood, and thoughts, affecting the ways in which a depressed individual, eats and feel about themselves, and thinks. Without treatment, depressive symptoms can last for weeks, months, or a life-time. Measured by years that people spend disabled with depression, it is the biggest blight on human society—bar none.[9–11]

Many studies have investigated the relationship between functional gastrointestinal disorder and psychological factors.[12–14] A close relationship has been established between the brain and the gastrointestinal tract. For example, stress and emotions can affect gastrointestinal function, as well as the occurrence of gastrointestinal symptoms and disease.[13] Likewise, the state of the gastrointestinal organs may affect a persons...
emotional status. Psychological factors may influence the severity of a functional gastrointestinal disorder by affecting the perception of pain through an action on the gut-brain axis—a concept that is also applicable to patients with GERD.[16] To date, a few studies describing GERD have shown that psychological factors, particularly anxiety and depression, play an important role in patients with GERD.[17–19] However, a comprehensive explication of this association had not yet been given. Thus, this study aims to systematically investigate the association between GERD and DD.

2. Methods

2.1. Objective

This study will aim to explore the association between GERD and DD systematically and comprehensively.

2.2. Study registration

We have registered this study on INPLASY202090026. It has been reported according to the guideline of preferred reporting items for systematic reviews and meta-analysis protocol statement. And we followed the PRISMA-P guidelines.[20,21]

2.3. Inclusion criteria for study selection

2.3.1. Type of studies. All potential case-controlled studies will be included, which identified the association between GERD and DD, regardless language and publication status limitations.

2.3.2. Type of participants. The patients with clinically diagnosed GERD, regardless of race, gender, and age.

2.3.3. Type of exposures. Exposure reviewed is the diagnosis of DD which is defined by each individual study. And the comparator is no diagnosis of GERD.

2.3.4. Outcomes. Relative risk of DD in people with GERD compared to people without GERD.

2.4. Search methods

We will search the following electronic bibliographic databases: PubMed/MEDLINE, Embase, Cochrane Library, Web of Science, the Chinese Bio Medical Literature Database, China National Knowledge Infrastructure (CNKI), the China Science and Technology Journal database (VIP) and Wanfang Data. In addition, ongoing trials will be retrieved from the WHO ICTRP Search Portal, the Chinese Clinical Trial Register and The Clinical Trials Register. Articles related to gastroesophageal reflux disease and depression will be searched. And language and time will be unlimited.

2.5. Search strategy

Two authors will screen the titles and abstracts of the all records retrieved in above electronic databases independently to find potentially eligible reviews. According to the inclusion and exclusion criteria outlined above, the full texts of them will be retrieved for further identification. Any disagreement will be resolved by discussion or by consultation with a third author. The search strategy for PubMed is presented in Table 1 and the strategy will be modified upon the requirement of other databases.

2.6. Data collection and analysis

2.6.1. Study selection. Two reviewers will perform literature screening, study selection, and data extraction independently. The literature obtained will be imported into EndnoteX9 to screen the title and abstract, the duplications and studies failing to meet the pre-specified inclusion criteria will be excluded. After reading the full text of the remained literature and discussing within the group, the final included studies will be determined.

| Table 1 |
| --- |
| Search strategy used in PubMed database. |
| Number | Search terms |
| --- | --- |
| #1 | Gastroesophageal reflux[MeSH Terms] |
| #2 | Gastric acid reflux[Title/Abstract] OR acid reflux, gastric[Title/Abstract] OR reflux, gastric acid [Title/Abstract] OR gastric acid reflux disease [Title/Abstract] OR gastro-esophageal reflux[Title/Abstract] OR gastro esophageal reflux[Title/Abstract] OR reflux, gastro-esophageal [Title/Abstract] OR gastroesophageal reflux disease [Title/Abstract] OR gerd [Title/Abstract] OR reflux, gastroesophageal [Title/Abstract] OR esophageal reflux [Title/Abstract] OR gastro-esophageal reflux [Title/Abstract] OR reflux, gastro-esophageal [Title/Abstract] |
| #3 | #1 OR #2 |
| #4 | Depression Disorder[MeSH Terms] |
| #5 | Complementary and alternative therapies[Title/Abstract] OR Disorder, Depressive[Title/Abstract] OR Disorders, Depressive[Title/Abstract] OR Neurosis, Depressive [Title/Abstract] OR Depressive Neuroses[Title/Abstract] OR Depressive Neurosis[Title/Abstract] OR Neuroses, Depressive[Title/Abstract] OR Depression, Endogenous [Title/Abstract] OR Depressions, Endogenous[Title/Abstract] OR Endogenous Depression[Title/Abstract] OR Endogenous Depressions[Title/Abstract] OR Depressive Syndrome[Title/Abstract] OR Depressive Syndromes[Title/Abstract] OR Syndrome, Depressive[Title/Abstract] OR Syndromes, Depressive[Title/Abstract] OR Depression, Neurotic[Title/Abstract] OR Neurotic Depression[Title/Abstract] OR Neurotic Depressions[Title/Abstract] OR Neurotic Depressions[Title/Abstract] OR Melancholia[Title/Abstract] OR Unipolar Depression[Title/Abstract] OR Depression, Unipolar[Title/Abstract] OR Depressions, Unipolar[Title/Abstract] OR Unipolar Depression[Title/Abstract] |
| #6 | #4 OR #5 |
| #7 | randomized controlled trial[Publication Type] |
| #8 | controlled clinical trial[Publication Type] |
| #9 | randomized[Title/Abstract] |
| #10 | randomly[Title/Abstract] |
| #11 | #7 OR #8 OR #9 OR #10 |
| #12 | #3 AND #6 AND #11 |
The corresponding author of original RCT will be contacted when the full text is unavailable. Disagreements will be solved by consulting a third-party arbitrator or discussing within a group.

2.6.2. Data extraction and management. Two authors will screen the titles and abstracts of the all records retrieved in above electronic databases independently to find potentially eligible reviews. According to the inclusion and exclusion criteria outlined above, the full texts of them will be retrieved for further identification. Any disagreement will be resolved by discussion or by consultation with a third author.

Data will be extracted by 2 reviewers independently using a pre-designed data extraction form. A third reviewer will validate data. The following data will be extracted: General information, Trial characteristics, Intervention(s) and control(s), Participants, Study methodology, Outcomes, Results, etc.

2.6.3. Risk of bias in included studies. The methodological quality of eligible studies will be assessed by 2 review authors independently according to the the Cochrane Handbook for Systematic Reviews of Interventions. The following characteristics will be assessed: 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. Based on the assessments of the studies against these 7 domains, they will be classified as being of “low risk”, “high risk” or “unclear risk” of bias. Any disagreements will be resolved by discussion or discussed with another reviewer if necessary.

2.6.4. Data analysis. Meta-analysis was conducted using Review Manager software (version 5.3). Odds ratio (OR) with 95% confidence intervals (CI) was reported for the dichotomous data, and mean differences (MD) with 95% CI for the continuous data. Statistical heterogeneity between studies was tested by calculating Higgins’ I² values or using the χ² test. P I² > 50%, and I² > 75% were respectively defined to indicate moderate, substantial, and considerable heterogeneity. When the P value of χ² test was < .1, an I² test was carried out. If the I² test showed a value > 50%, a random effects model was carried out. Otherwise, a fixed effects model was carried out. A P value lower than 0.05 was considered to be statistically significant.

2.6.5. Patient and public involvement. This is a meta-analysis study based on previously published data, so patient and public involvement will not be included in this study.

2.6.6. Grading the quality of evidence. The Grading of Recommendations Assessment, Development and Evaluation (GRADE) guidelines will be utilized to grade the quality of evidence as very low, moderate, or high.

3. Discussion

This will be the first study to investigate the association between GERD and DD. It will systematically and comprehensively search electronic databases, and other literature sources to avoid missing potential studies. This study will summarize the most recent eligible studies of the association between GERD and DD. The findings of this study will provide evidence to judge the association between GERD and DD, which may benefit for the clinical practice and future studies.

Author contributions

Conceptualization: Yu Liu, Yangang Wang.
Data curation: Yu Liu, Panpan Zhou.
Formal analysis: Shixiong Zhang.
Funding acquisition: Yangang Wang.
Methodology: Shaowei Liu, Zeqi Yang, Miaochan Xu.
Resources: Yangang Wang.
Software: Shaowei Liu, Zeqi Yang.
Supervision: Huiqing Wu.
Writing – original draft: Yu Liu.
Writing – review & editing: Yu Liu, Miaochan Xu.

References

[1] Vakil N, van Zanten SV, Kahrilas P, et al. Globale Konsensusgruppe. Die montreal-definition und -klassifkation der gastrooesophagealen refux-rankheit: ein globales evidenzbasiertes Konsensus-Papier [the montreal definition and classification of gastroesophageal reflux disease: a global, evidence-based consensus paper]. Z Gastroenterol 2007;45:1125–40.
[2] Richter JE, Rubenstein JH. Presentation and Epidemiology of Gastroesophageal Reflux Disease. Gastroenterology 2018;154:267–76.
[3] Zhang M, Pastolinoje M, Zhou X, et al. Assessing different diagnostic tests for gastroesophageal reflux disease: a systematic review and network meta-analysis. Therap Adv Gastroenterol 2019;12:1756284819890337.
[4] Aziz Q, Fass R, Goyal CP, et al. Functional esophageal disorders. Gastroenterology 2016;150:1366–79.
[5] Eusebi LH, Ramakumar R, Yuan Y, et al. Global prevalence of, and risk factors for, gastro-oesophageal reflux symptoms: a meta-analysis. Gut 2018;67:430–40.
[6] Gawron AJ, French DD, Pastolinoje M, et al. Economic evaluations of gastroesophageal reflux disease medical management. Pharmacoeconomics 2014;32:745–58.
[7] Mathers CD, Loncar D. Projections of global mortality and burden of disease from 2002 to 2030. PLoS Med 2006;3:e442.
[8] Chong PS, Fung ML, Wong KH, et al. Therapeutic potential of hericium erinaceus for depressive disorder. Int J Mol Sci 2019;21:163.
[9] Blackburn TP. Depressive disorders: treatment failures and poor prognosis over the last 30 years. Pharmacol Res Pract 2019;7:e00472.
[10] Cui R. Editorial: a systematic review of depression. Curr Neuropharmacol 2015;13:480.
[11] Smith K. Mental health: a world of depression. Nature 2014;515:181.
[12] Lee SY, Ryu HS, Choi SC, et al. Psychological factors in psychiatric patients in South Korea. Psychiatry Investig 2020;17:262–7.
[13] Karling P, Maripuu M, Wikgren M, et al. Association between gastrointestinal symptoms and affectivity in patients with bipolar disorder. World J Gastroenterol 2016;22:8340–8.
[14] Xiong N, Duan Y, Wei J, et al. Antidepressants vs. Placebo for the treatment of functional gastrointestinal disorders in adults: a systematic review and meta-analysis. Front Psychiatry 2018;9:659.
[15] Rudzki L, Maes M. The Microbiota-Gut-Immune-Glia (MGIG) axis in major depression. Mol Neurobiol 2020;10:1007.
[16] Simpson CA, Schwartz OH, Simmons JG. The human gut microbiota and depression: widely reviewed, yet poorly understood. J Affect Disord 2020;274:73–5.
[17] Choi JM, Yang JH, Kang SJ, et al. Association between anxiety and depression and gastroesophageal reflux disease: results from a large cross-sectional study. J Neurogastroenterol Motil 2018;24:593–602.
[18] Suganami Y, Oka K, Hanayama Y, et al. Correlations between depressive condition and gastroesophageal reflux symptoms in patients visiting a department of general medicine. Acta Med Okayama 2019;73:479–86.
[19] Wong MW, Bair MJ, Chang WC, et al. Clinical and psychological characteristics in gastroesophageal reflux disease patients overlapping with laryngopharyngeal reflux symptoms. J Gastroenterol Hepatol 2019;34:1720–6.
[20] Hutton B, Salanti G, Caldwell DM, 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 2013;162:777–84.
[21] Zheng W, Gao T, Huang H, et al. Thirteen kinds of Chinese medicine injections for acute exacerbation of chronic obstructive pulmonary disease: protocol for a systematic review and network meta-analysis. Medicine (Baltimore) 2019;98:e16200.