



ORIGINAL ARTICLE

PEEP titration by EIT strategies for patients with ARDS: A systematic review and meta-analysis



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KEYWORDS

Acute Respiratory Distress Syndrome; Electrical Impedance Tomography; Meta-analysis; Positive End-Expiratory Pressure

Abstract

Objective: To determine which method of Positive End-expiratory Pressure (PEEP) titration is more useful, and to establish an evidence base for the clinical impact of Electrical Impedance Tomography (EIT) based individual PEEP setting which appears to be a promising method to optimize PEEP in Acute Respiratory Distress Syndrome (ARDS) patients.

Design: A systematic review and meta-analysis.

Setting: 4 databases (PUBMED, EMBASE, Web Of Science, and the Cochrane Library) from 1980 to December 2020 were performed.

Participants: Randomized clinical trials patients with ARDS.

Main variables: PaO₂/FiO₂-ratio and respiratory system compliance.

Intervention: The quality of the studies was assessed with the Cochrane risk and bias tool.

Results: 8 trials, including a total of 222 participants, were eligible for analysis. Meta-analysis demonstrates a significantly EIT-based individual PEEP setting for patients receiving higher PaO₂/FiO₂ ratio as compared to other PEEP titration strategies [5 trials, 202 patients, SMD 0.636, (95% CI 0.364–0.908)]. EIT-driven PEEP titration strategy did not significantly increase respiratory system compliance when compared to other PEEP titration strategies, [7 trials, 202 patients, SMD –0.085, (95% CI –0.342 to 0.172)].

Conclusions: The benefits of PEEP titration with EIT on clinical outcomes of ARDS in placebo-controlled trials probably result from the visible regional ventilation of EIT. These findings offer clinicians and stakeholders a comprehensive assessment and high-quality evidence for the safety and efficacy of the EIT-based individual PEEP setting as a superior option for patients who undergo ARDS.

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Abbreviations: ARDS, Acute Respiratory Distress Syndrome; PEEP, Positive End-Expiratory Pressure; EIT, Electrical Impedance Tomography.

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PALABRAS CLAVE

Síndrome de distrés respiratorio agudo; Tomografía de impedancia eléctrica; Metaanálisis; Presión espiratoria final positiva

Evaluación sistemática y metaanálisis de la estrategia eit para la titulación Peep en pacientes con sdra**Resumen**

Objetivo: Para determinar qué método de valoración de la presión espiratoria final positiva (PEEP) es más útil, y para establecer una base de evidencia para el impacto clínico de la tomografía de impedancia eléctrica (EIT) basada en el ajuste individual de PEEP que parece ser un método prometedor para optimizar la PEEP en pacientes con síndrome de dificultad respiratoria aguda (ARDS).

Diseño: Una revisión sistemática y metaanálisis.

Ámbito: Se realizaron 4 bases de datos (PUBMED, EMBASE, Web Of Science y Cochrane Library) de 1980 a diciembre de 2020.

Participantes: Ensayos clínicos aleatorizados de pacientes con SDRA.

Variables principales: PaO₂/FiO₂ ratio y compatibilidad respiratoria.

Intervención: La calidad de los estudios se evaluó con la Cochrane risk and bias tool.

Resultados: Ocho ensayos, incluyendo un total de 222 participantes, fueron elegibles para el análisis. El análisis de \geq eta demuestra una configuración individual significativamente basada en MEIT para pacientes que reciben una mayor proporción EE₂/P PiO₂ en comparación con otras estrategias de titulación FOPEEP SMD CI. La estrategia de titulación de PEEP derivada del tie no aumentó significativamente el cumplimiento del sistema respiren comparación con otras estrategias de titulación de PEEP, [7 ensayos, 202 pacientes, DME -0,085, (IC del 95%: -0,342-0,172)].

Conclusiones: Los beneficios de la valoración de la PEEP con EIT en los resultados clínicos de SDRA en ensayos controlados con placebo probablemente sean el resultado de la ventilación regional visible del EIT. Estos hallazgos ofrecen a los médicos y a las partes interesadas una evaluación integral y evidencia de alta calidad para la seguridad y eficacia de la configuración individual de PEEP basada en EIT como una opción superior para los pacientes que se someten a SDRA.

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Introduction

Acute Respiratory Distress Syndrome (ARDS) was first described in 1967 and is characterized by the abrupt onset of clinically significant hypoxemia with the presence of diffuse pulmonary infiltrates.¹ The criteria were updated in 2012 in the so-called Berlin definition of ARDS in adults based on the severity of hypoxemia represented by the ratio of the partial pressure of oxygen in arterial blood to inspired oxygen.² ARDS is a life-threatening condition of seriously ill patients and is a prevalent cause of acute respiratory failure, carrying high mortality of 30–40%.³ The pathophysiological features were the ratio of ventilation to perfusion mismatch, decreased thoracic compliance, increased dead space, and elevated pulmonary arterial pressure. Therefore, early recognition of ARDS modifiable risk factors and the avoidance of aggravating factors during the patient's hospital stay can help decrease its development.⁴ Of all adjunctive therapies, lung-protective ventilation is still the key to a better outcome in ARDS. Recent guidelines on mechanical ventilation in ARDS provide evidence-based recommendations related to 6 interventions, including low tidal volume and inspiratory pressure ventilation, prone positioning, high-frequency oscillatory ventilation, higher vs. lower Positive End-Expiratory Pressure (PEEP), lung recruitment maneuvers, and extracorporeal membrane oxygenation.⁵

Setting PEEP appropriately is now recognized as an important aspect of a lung-protective ventilation strategy and not just a strategy to improve oxygenation. Setting PEEP levels 5 cm H₂O may be harmful in the acute phase of ARDS. Setting PEEP appropriately is a balance between maintaining alveolar recruitment and avoiding alveolar over distention.⁶ Several methods have been proposed for PEEP titration in an individual patient with ARDS, including gas exchange, compliance, pressure-volume curve, stress index, esophageal manometry, lung volume, imaging.⁷ Of the imaging method, Electrical Impedance Tomography (EIT) plays an important role.

EIT is a bedside monitoring tool that non-invasively visualizes local ventilation and arguably lung perfusion distribution. EIT images possess a high temporal and functional resolution allowing the visualization of dynamic physiological and pathological changes on a breath-by-breath basis.⁸ Current studies focus mainly on its clinical applications to quantify lung collapse, tidal recruitment, and lung over distention to titrate PEEP, which is new in the PEEP setting.^{9–12}

Previous studies compared the methods of conventional PEEP titration strategies but not EIT-based individual PEEP setting. Thus, to assess the effect of EIT-based individual PEEP setting, we undertook a systematic review and meta-analysis investigating the difference between two methods (EIT and other method) in the final P/F ratio or the final compliance when PEEP is titrated using those methods.

Methods

Four electronic databases (PUBMED, EMBASE, Web Of Science, and the Cochrane Library) were searched from inception to December 2020, for trials investigating any other PEEP titration strategies and EIT based individual PEEP setting of ARDS patients, with MeSH headings and text words (discussed in detail in search strategies, web [appendix](#)). We searched for any additional studies in the references of all identified publications, including previous relevant meta-analyses and narrative reviews. This study is registered with the INPLASY website, number INPLASY202160094.

Selection criteria

For inclusion, any studies except reviews and case report in adults, and examine the effect of any other PEEP titration strategies and EIT based individual PEEP setting on respiratory markers and clinical impacts of ARDS. The outcome was assessed by use of respiratory markers (PaO₂/FiO₂-ratio and respiratory system compliance). Only studies published as full-length articles or letters in peer-reviewed English-language journals were included.

Data extraction

The following information was extracted and entered into databases by three investigators (MNY, TTZ, CW): study design, type of intervention, patients' characteristics, and outcomes (web [appendix](#)). If relevant information regarding the design or ARDS outcomes was unavailable, or doubt existed about duplicate publications, authors were contacted to obtain the necessary information (web [appendix](#)). Uncertainties were resolved by consensus.

This study is reported by the Preferred Reporting Items for Systematic Reviews and Meta-Analysis (PRISMA) statement^{12–14} and the Cochrane Handbook for Systematic Reviews of Interventions.¹⁵ Our detailed study protocol is available online and has been previously published¹⁶. No institutional review board approval was required for this meta-analysis because the study included data that had been published previously.

Statistical analysis

Dichotomous outcomes were reported using relative risks (RR) and 95% confidence intervals (CIs). For continuous outcomes (e.g., PaO₂/FiO₂-ratio and respiratory system compliance), we evaluated the standardized difference in means (SMD). Studies were weighted using inverse variance and data were pooled using random-effects models. We examined funnel plots of treatment effect versus study precision to assess for publication bias. Baseline characteristics of trial participants were summarized using weighted averages. Subgroup analyses were performed by stratifying studies by characteristics of interest (co-interventions, recruitment maneuvers, methods of setting PEEP) and assessing for quantitative interaction using Chi-square tests for heterogeneity between subgroups. We used Stata 15.0 to conduct statistical analyses.

Role of the fundingsource

Funding source of Health special talents program of Suzhou high tech Zone had a role in study design, data collection, data analysis, data interpretation, and writing of the report. All authors had full access to all the data in the study and all took full responsibility for the decision to submit for publication.

Results

Included studies and study quality

Eight trials met the inclusion criteria for this review, including 222 adults with ARDS ([Fig. 1](#)). The basic information and methodological quality of the included studies are reported in [Table 1](#). [Table 2](#) summarizes baseline patient characteristics and the varied PEEP protocols used between studies.

Respiratory system compliance

Seven from eight included studies reported respiratory system compliance and were included in the merger. After the data were merged, the heterogeneity of the studies was large ($I^2 = 95.1\%$), so the random-effects model was used and the subgroup analysis was carried out according to the different control PEEP strategies. Two trials compared the effects of PEEP strategy of PEEP/FiO₂ table and EIT-driven PEEP titration on respiratory system compliance, of which the final meta-analysis result showed a significant difference between two groups that respiratory system compliance increased in EIT group [SMD = 0.582, 95% CI (0.044, 1.121), $p < 0.05$] ($n = 30$); Three trials compared the effects of the strategy of a low PEEP and EIT-driven PEEP titration on respiratory system compliance and the meta analysis either showed no significant difference [SMD = -1.216, 95% CI (-1.585, 0.847), $p > 0.05$] ($n = 102$); One trial comparing the effects of PEEP strategy of pressure-volume loop and EIT-driven PEEP titration on respiratory system compliance revealed a significant rise in respiratory system compliance with the EIT-driven PEEP titration [SMD = 0.974, 95% CI (0.410, 1.538), $p = 0.001$] ($n = 55$); One trial which compared the effects of PEEP strategy of pressure-volume loop and EIT-driven PEEP titration on respiratory system compliance uncovered a significant difference that the respiratory system compliance with EIT-driven PEEP titration increased [SMD = 2.106, 95%CI (1.203, 3.010), $p = 0.000$] ($n = 15$) ([Fig. 2](#)).

PaO₂/FiO₂-ratio

Five from eight included studies reported the PaO₂/FiO₂-ratio and were included in the merger. After the data were merged, the heterogeneity of the studies was large ($I^2 = 82.5\%$), so the random-effects model was used and the subgroup analysis was carried out according to the different control PEEP strategies. One trial compared the effects of PEEP strategy of PEEP/ FiO₂ table and EIT-driven PEEP titration on PaO₂/FiO₂-ratio, of which the final meta-analysis result showed a significant difference between two groups

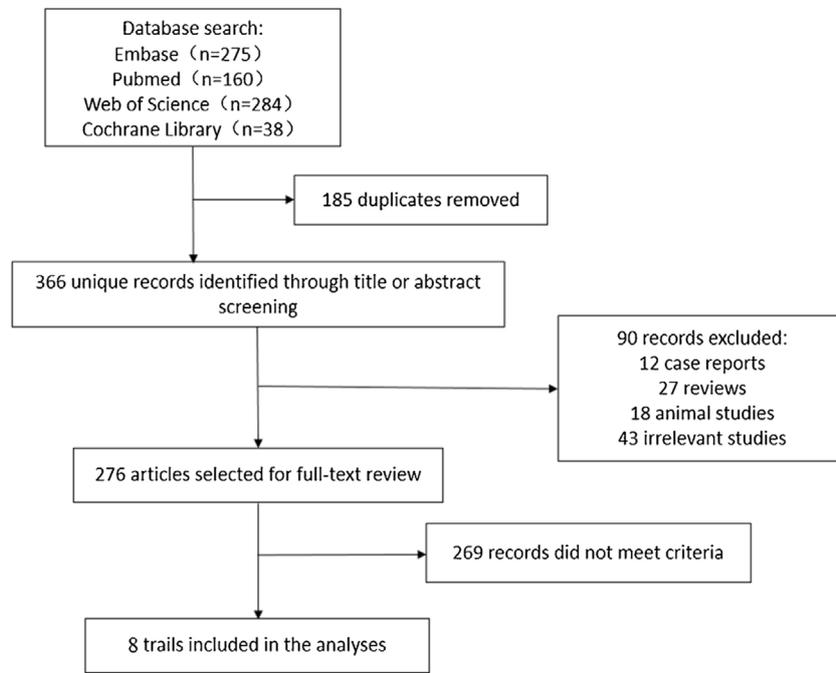


Figure 1 Flow chart of study selection.

Table 1 Methodological quality of included studies.

Author	Zhao	Eronia, N	Weber	Kastern	Heines	Becher	Jacopo	Scaramuzzo
Year	2019	2017	2020	2018	2018	2016	2019	2020
Country	China	Italy	Germany	Germany	The Netherlands	Germany	USA	Italy
Study design	RCT	NRCT	RCT	NRCT	NRCT	NRCT	NRCT	NRCT
Scale	Jadad 6	MINORS 18	Jadad 6	MINORS 18	MINORS 18	MINORS 18	MINORS 18	MINORS 18

Table 2 Baseline patient characteristics and the varied PEEP protocols.

Study	Zhao	Eronia, N	Weber	Kastern	Heines	Becher	Jacopo	Scaramuzzo
Sample (C/T)	31/24	14/14	23/25	15/15	39/39	15/15	14/14	20/20
Gender (M/F)	37/18	14/2	30/18	9/6	25/14	/	6/8	13/7
Interpretation:								
Control group	Pressure–volume loop	PEEP/FiO ₂ table	PEEP 5 cmH ₂ O	Best Compliance approach	PEEP 4 cmH ₂ O	The ARDSnet protocol	PEEP/FiO ₂ table	End expiratory PL/FiO ₂ sliding table
EIT group	ODCL	EELI	CRS	ODCL	ODCL	EELI	ODCL	SStot
Outcomes	①②	①②	①	①	①②	①②	①②	②

ODCL, percentage of over distension/collapse; PL, transpulmonary pressure; SStot, total Silent Spaces; ①, lung compliance; ②, PaO₂/FiO₂-ratio.

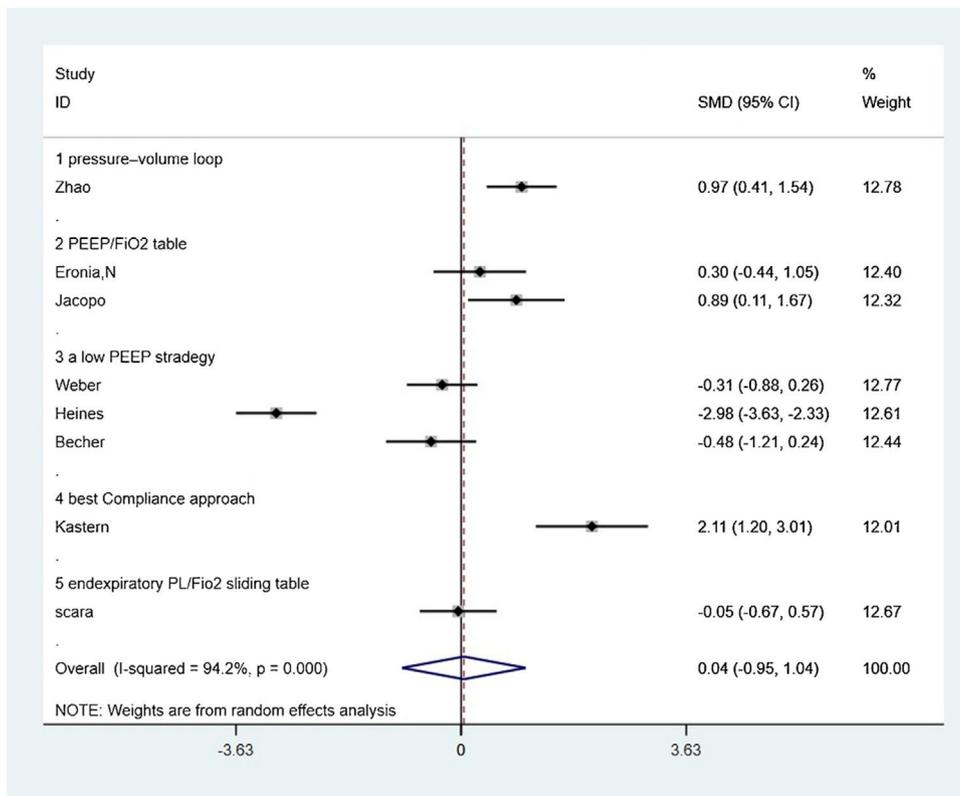


Figure 2 The subgroup analysis of respiratory system compliance.

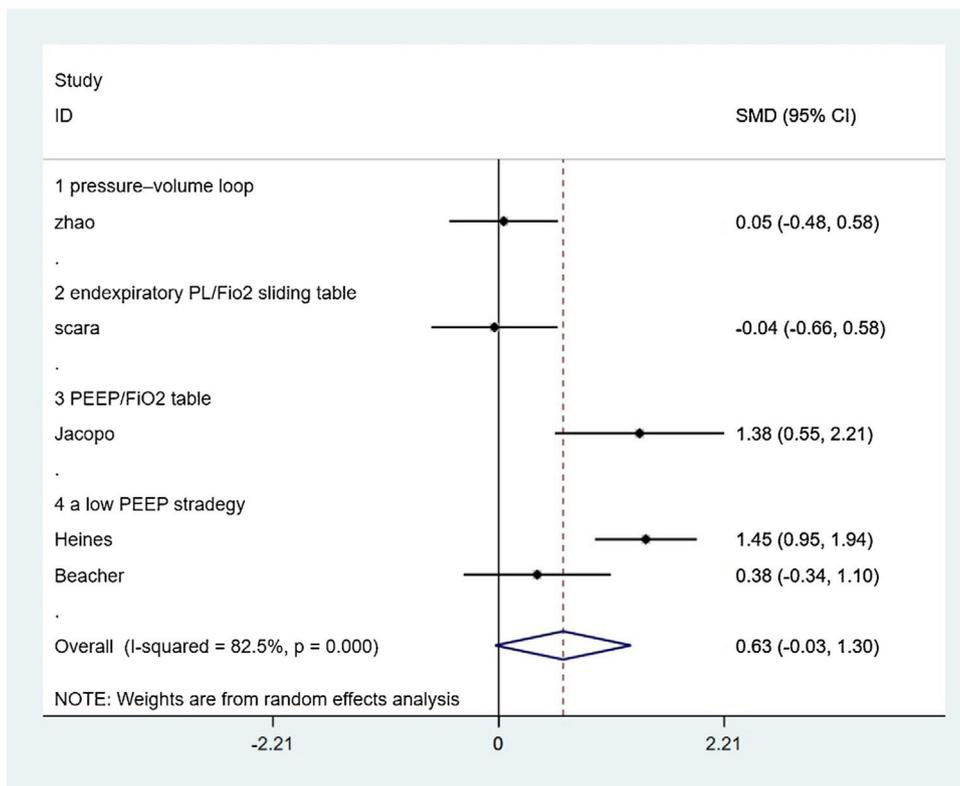


Figure 3 The subgroup analysis of PaO₂/FiO₂-ratio.

that PaO₂/FiO₂-ratio increased in EIT group [SMD = 1.383, 95% CI (0.533, 2.214), *p* < 0.05] (*n* = 14); Two trials compared the effects of the strategy of a low PEEP and EIT-driven PEEP titration on PaO₂/FiO₂-ratio and the meta-analysis either showed no significant difference [SMD = 0.945, 95% CI (-0.097, 1.988), *p* > 0.05] (*n* = 54); one trial comparing the effects of PEEP strategy of the pressure-volume loop and EIT-driven PEEP titration on PaO₂/FiO₂-ratio revealed no significant difference between the two groups [SMD = 0.050, 95% CI (-0.483, 0.583), *p* > 0.05] (*n* = 55). One trial which compared the effects of PEEP strategy of end-expiratory transpulmonary pressure (PL)/FiO₂ sliding table and EIT-driven PEEP titration uncovered no significant difference on PaO₂/FiO₂-ratio [SMD = -0.041, 95% CI (-0.661, 0.579), *p* > 0.05] (*n* = 20) (Fig. 3).

Publication bias

Considering not enough studies enrolled, it is not available for the funnel plot method to publication bias. There is a potential risk of publication bias.

Discussion

Our study indicates that ARDS patients who have optimized PEEP with Electrical Impedance Tomography have a significantly higher PaO₂/FiO₂-ratio than those titrated with the conventional type. However, the respiratory system compliance was similar in the two groups. PEEP was not significantly decreased in the EIT group compared with the conventional group indicating that EIT-driven PEEP has similar efficacy to conventional strategies.

Raised compliance compared to subgroups and raised PaO₂/FiO₂-ratio in the EIT group was maintained across important subgroups associated with patient demographics, the severity of hypoxemia, patient position, imaging feature, tidal volume, other clinician therapies, and use of prophylactic measures. This observation suggests that our findings reflect a true effect of Electrical Impedance Tomography, rather than an artifact of statistical heterogeneity or specific patient or procedural characteristics. Notably, non-invasive and visible are the specific feature of EIT. Despite this difference, the conventional strategies are maintained, suggesting that the effect of dynamic imaging design does not take place of that of conventional strategies in treatment guidance.

In the recent decade, Electrical Impedance Tomography (EIT) is becoming emerging as an easily accessible imaging radiation-free techniques that allow continuous and functional respiratory monitoring at the bedside, based on the repeated measurement of the surface voltages resulting from a rotating injection of high frequency and low-intensity alternating current that circulates between the electrodes located around the chest. The electrodes collect the information on impedance by forming a relative image concerning a reference. The tissues allow the passage of current with little resistance, however, with gas the opposite occurs. In this way, it is possible to perform plethysmography of the volume entering upon each inspiration and in each region of interest.¹³ Previous studies mainly focus on randomized controlled trials of ventilation strategies and

monitoring, animal models on cardio-pulmonary perfusion, and reviews of clinical practice,^{8,9,13-21} but without a meta-analysis of EIT-driven PEEP titration for ARDS patients yet. No meta-analysis of EIT presents up to December 2020, probably because of EIT coming out not long and being not widely used. Thus we performed this meta-analysis investigating the use of EIT-driven PEEP titration for the management of patients with ARDS and establish evidence. The previous meta-analysis reported a lot of conventional PEEP settings (including Gas exchange, compliance, pressure-volume curve, stress index, esophageal manometry, lung volume, Imaging) for patients with ARDS, of which mostly compared the effect of high PEEP level and low PEEP level on outcomes.²²⁻³¹ Our findings stand in clear distinction from those of past studies that did not reach a consensus on this topic. A previous meta-analysis²²⁻²⁶ had limitations and focused on the conventional and indisputable PEEP setting, whereas our study is broad in its scope with robust analyses done by a multidisciplinary team.

However, our study is not without limitations. First, only eight studies were included, all of which are small-sample trials, resulting in poor test efficiency. At the same time, all trials did not report the allocation concealment scheme, which may have selection bias. Only two of the studies were RCT. Six studies did not report their blinded implementation schemes, and the measurement biases might exist. These defects will affect the internal and external authenticity of the results.

In conclusion, we found that EIT-guided PEEP titration was associated with a raise respiratory system compliance and significant raise in PaO₂/FiO₂-ratio, and had similar efficacy to their conventional counterparts. Our findings suggest that the EIT group retains a favorable balance between safety and efficacy when compared with the conventional type. These results provide clinicians and health-care policymakers with a comprehensive assessment and high-quality evidence on the safety and efficacy of atraumatic needles as a superior option for patients with ARDS.

Disclosure statement

This research received grant from health special talents program of Suzhou high tech Zone. The authors report no conflicts of interest.

CRedit authorship contribution statement

Mengnan Yu: Conceptualization, Formal analysis, Investigation, Project administration, Validation, Visualization, Writing – original draft, Writing – review & editing. **Yanjun Deng:** Conceptualization, Investigation, Project administration, Validation, Writing – original draft, Writing – review & editing. **Jun Cha:** Data curation. **Lingyan Jiang:** Data curation, Formal analysis, Validation, Visualization. **Ming-deng Wang:** Data curation, Supervision. **Shigang Qiao:** Data curation, Formal analysis, Investigation, Supervision. **Chen Wang:** Conceptualization, Investigation, Project administration, Supervision, Visualization, Writing – review & editing.

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Appendix A Supplementary data

Supplementary material related to this article can be found, in the online version, at <https://doi.org/10.1016/j.medin.2022.06.009>.

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