ABSTRACT
Objectives To evaluate whether early intensive care transthoracic echocardiography (TTE) can improve the prognosis of patients with mechanical ventilation (MV).

Design A retrospective cohort study.

Setting Patients undergoing MV for more than 48 hours, based on the Medical Information Mart for Intensive Care III (MIMIC-III) database and the eICU Collaborative Research Database (eICU-CRD), were selected.

Participants 2931 and 6236 patients were recruited from the MIMIC-III database and the eICU database, respectively.

Primary and secondary outcome measures The primary outcome was in-hospital mortality. Secondary outcomes were 30-day mortality from the date of ICU admission, days free of MV and vasopressors 30 days after ICU admission, use of vasoactive drugs, total intravenous fluid and ventilator settings during the first day of MV.

Results We used propensity score matching to analyse the association between early TTE and in-hospital mortality and sensitivity analysis, including the inverse probability weighting model and covariate balancing propensity score model, to ensure the robustness of our findings. The adjusted OR showed a favourable effect between the early TTE group and in-hospital mortality (MIMIC: OR 0.78; 95% CI 0.65 to 0.94, p=0.01; eICU-CRD: OR 0.76; 95% CI 0.67 to 0.86, p<0.01). Early TTE was also associated with 30-day mortality in the MIMIC database (OR 0.71, 95% CI 0.57 to 0.89, p=0.001). Furthermore, those who had early TTE had both more ventilation-free days (only in eICU-CRD: 23.48 vs 24.57, p=0.01) and more vasopressor-free days (MIMIC: 18.22 vs 20.64, p=0.005; eICU-CRD: 27.37 vs 28.59, p<0.001) than the control group (TTE applied outside of the early TTE and no TTE at all).

Conclusions Early application of critical care TTE during MV is beneficial for improving in-hospital mortality. Further investigation with prospectively collected data is required to validate this relationship.

STRENGTHS AND LIMITATIONS OF THIS STUDY
⇒ The large sample size from two ICU databases increased the credibility of the research.
⇒ Inverse probability weighting model and covariate balancing propensity score model were used to analyse the association between early TTE and in-hospital mortality.
⇒ Although a sensitivity analysis was performed to make the results more reliable, some biases are inevitable.
⇒ Only patients from the MICU and SICU were selected to rule out the impact of severe pre-existing heart diseases or cardiac surgery.

INTRODUCTION
Echocardiography, which can be conveniently performed in intensive care units (ICUs), provide more information on cardiac abnormalities, including anatomical abnormalities and functional abnormalities, and quickly and accurately assess haemodynamic changes in the ICU. Compared with other ICU diagnostic tools, the frequency of transthoracic echocardiography (TTE) use in the ICU has increased rapidly owing to the advantages of non-invasiveness, easy availability and safety. In the span of 10 years from 1999 to 2008, the use of TTE has almost doubled, but the resulting economic and guardianship costs are also increasing. The research on TTE mainly focuses on the management changes caused by TTE, but the impact of these changes is unclear. Besides, despite the release of practice guidelines for cardiac ultrasound in 2011, many clinicians still use TTE based on their clinical experience. In the ICU, where information from randomised controlled trials is a relative deficiency, it is significant to understand the clinical value of a commonly used diagnostic tool.

MV is a common and important procedure for patients in the ICU. Although respiratory failure is the greatest problem to be solved in patients undergoing MV, haemodynamic also plays a role that cannot be ignored, and a recent study also showed that haemodynamic
changes caused by the influence of airway pressure affect the prognosis of patients. Several studies have suggested that cardiac assessment should be included in management strategies in patients undergoing mechanical ventilation (MV). TTE can be used to assess the patient’s fluid response under MV and is ideally suited to diagnose the weaning failure of cardiac origin. However, there is no strong evidence to support that TTE should be performed during MV. Thus, understanding the clinical value of early TTE in mechanically ventilated patients is enormously important.

Based on the above questions, our study was implemented to investigate the impact of earlier TTE performance on the outcomes of critically ill adult patients with MV.

MATERIALS AND METHODS

Database

This study was reported in accordance with the STrengthening the Reporting of OBservational studies in Epidemiology statement. We collected data from Medical Information Mart for Intensive Care (MIMIC)-III v1.4 and the eICU Collaborative Research Database (eICU-CRD) v2.0. Both are extensive, free, public databases containing hospitalisation information. MIMIC covers 61,392 ICU admissions for 46,476 patients at the Beth Israel Deaconess Medical Center in Boston, MA, USA. The eICU-CRD covers 200,859 ICU admissions from 139,226 patients at 208 US hospitals. Xueshu Yu completed the required courses for the use of the database and obtained the corresponding certificate (researcher certification number 1605699 and record id 27 752 407).

Study cohort

We conducted a retrospective study of mechanically ventilated adult patients from MIMIC-III; only admissions to medical intensive care units (MICUs) or surgical intensive care units (SICUs) were included, patients in a coronary care unit and cardiac surgery recovery unit were excluded. We used only the first ICU admission data for the first hospitalisation and patients 16 years of age or older who had been continuously ventilated for at least 48 hours. Patients who had incomplete datasets were excluded. Patients who underwent echocardiography less than 24 hours before MV or within 24 hours after MV were classified as the early TTE group, and the remaining patients constituted the control group (patients who had no early TTE, including TTE applied outside of the early TTE or no TTE at all).

Data were extracted from the database using structured query language code established by Serpa Neto et al. The following demographic data (using data from the first 24 hours of admission) were collected: age, sex, weight, race, comorbidities (chronic obstructive pulmonary disorder (COPD), asthma, congestive heart failure (CHF), chronic kidney disease (CKD), hypertension, sepsis, acute respiratory distress syndrome (ARDS)), Sequential Organ Failure Assessment (SOFA) score, Oxford Acute Severity of Illness Score (OASIS), vital signs (mean arterial pressure (MAP), heart rate (HR)) and laboratory values (white blood cell (WBC) count, haemoglobin (Hb), blood urea nitrogen (BUN), pH, pO2, pCO2 and lactate). In addition, we also collected the management data for the first day of MV (tota intravenous fluid, ventilator settings, use of dobutamine and norepinephrine).

Patient and public involvement

No patients and the public were involved in any part of this study.

Outcomes

The primary outcome of the study was in-hospital mortality. Secondary outcomes were 30-day mortality from the date of ICU admission; days free of MV and vasopressors 30 days after ICU admission, use of vasodilating drugs, total intravenous fluid and ventilator settings during the first day of MV.

Statistical analysis

In order to control the confounding factors, propensity score matching (PSM) was performed. The baseline characteristics of the original cohort were stratified by TTE. The propensity score for an individual was determined based on the covariates age, sex, weight, race, HR, COPD, asthma, CHF, CKD, hypertension, ARDS, sepsis, SOFA score, OASIS, WBC, Hb, pH, pO2, pCO2 and lactate using a standard software package (matching package) with a PSM methodology. These variables were selected due to their clinical relevance. This method consisted of ranking the MV patients in the early TTE group and the control group, then selecting the TTE patients who had the highest propensity score and finding the control group patients with the closest propensity score (maximum calliper, 0.2). Both patients were then removed from consideration for matching, and the next highest patient was selected (matched 1:1 using the nearest-neighbour algorithm).

After matching, to assess the balance between the two groups, the standardised mean differences (SMDs) between the early TTE cohort and the control cohort were calculated. SMDs eliminate not only the influence of the absolute values from a study but also the influence of the unit of measurement on the results. Continuous variables are shown as the means and SD, and categorical variables are represented as the total and proportion. For continuous variables, we used a non-parametric test or the Wilcoxon rank-sum test. For the categorical variables, we used a χ2 test or Fisher’s exact test.

Secondary outcomes were observed after matching as well. We used paired t-tests for continuous outcomes and χ2 tests for categorical outcomes.

We used the random forest model to impute missing data (online supplemental eFigures 1 and 2).

Sensitivity analysis

We conducted a series of sensitivity analyses with the cohort with missing data, the cohort after imputation and
the cohort after PSM to assess the outcomes. In addition, we used multiple logistic regression, the inverse probability of treatment weight (IPTW) and the covariate balancing propensity score (CBPS) to further validate the primary outcome. To adjust for these covariates, the doubly robust estimation method was used to deduce the independent associations between TTE and in-hospital mortality and 30-day mortality (details about the IPTW and CBPS can be found in the online supplemental material). In addition, we divided the patients into three groups (Early TTE, late TTE (TTE time ≥ MV time +24 hours) and non-TTE (no TTE after MV)), and further analyses were conducted to identify predictors of early TTE using multiple logistic regression. Finally, we carried out a sensitivity analysis through multivariate logistic regression focussing on patients with ARDS and sepsis.

Statistical significance was assessed to be determined by a two-sided p<0.05. All statistical analyses mentioned above were performed using R version 3.5.3.

RESULTS

After reviewing 46,476 unique patients from the MIMIC-III database and excluding those with readmission, age <16 years, and ventilation duration <48 hours, 2951 patients from the MICU and SICU were enrolled (figure 1). In the eICU-CRD, of the 139,226 unique patients, 6236 patients in the MICU and SICU were included after the exclusion of patients aged <16 years, those receiving invasive ventilation for less than 48 hours and that missing hospital discharge information (figure 1). The in-hospital mortality of MV patients was 31.53% in the MIMIC database and 28.62% in the eICU-CRD. Patients who died were older, had lower weight, higher OASIS and SOFA scores, higher lactate, and lower MAP (online supplemental eTable 1).

The original cohort baseline in the MIMIC database showed that patients who underwent TTE on the first day of MV had a more severe status in terms of the SOFA score, MAP, pH, pO2 and pCO2 (table 1); however, there was no statistically significant difference between the early TTE group and the control group in the eICU-CRD (table 1).

Primary outcomes

Univariate logistic regression analysis results of in-hospital mortality are shown in online supplemental eTable 2, and the details of multiple logistic regression are shown in online supplemental eTable 3. The clinical features of TTE during different time periods and no TTE at all are shown in online supplemental eTable 4. Then, we used PSM to standardise the differences between the early TTE group and the control group (online supplemental eTables 5 and 6). All covariates were balanced in the PSM cohort (online supplemental eFigures 3 and 4). The adjusted OR (MIMIC: OR 0.78, 95% CI (0.65 to 0.94), (p=0.01); eICU-CRD: OR 0.76, 95% CI (0.67 to 0.86), (p<0.01)) showed that the early use of TTE was beneficial to improve the in-hospital mortality of patients undergoing MV (figure 2).

Secondary outcomes

After PSM, early TTE was also associated with 30-day mortality in the MIMIC database (OR 0.71, 95% CI (0.57 to 0.88), (p=0.001) (online supplemental eFigure 5). Since the eICU-CRD only contains in-hospital mortality data, we had no way to assess the association between early TTE and 30-day mortality in the eICU-CRD cohort. In addition, we found that those who had early TTE had both more ventilation-free days (but there was no statistical difference in MIMIC, p=0.051) and more vasopressor-free days in 30 days than the control group (online supplemental eTable 7), which might be related to the management changes brought by early TTE, including a less amount of intravenous fluid (only in the eICU-CRD: 2156.88 (3635.99) vs. 1470 (2670.69), (p<0.001), the use of vasoactive drugs and the ventilator setting parameters (only in the MIMIC database). However, this conclusion should be regarded cautiously as far as the generalisation of results is concerned.

Sensitivity studies

We performed some sensitivity analyses, as summarised in figure 2 (in-hospital mortality) and online supplemental eFigure 5 (30-day mortality). We analysed all three cohorts, including the matched cohort, the original cohort with missing data, and the cohort after imputation, and found similar results: in-hospital mortality and 30-day mortality were improved in mechanically ventilated patients undergoing early TTE. In addition, early TTE was more conducive to the prognosis of patients than late TTE and non-TTE (online supplemental eFigure 6),
Table 1  Baseline characteristics of the original cohort

| Variables       | MIMIC cohort                  | eICU-CRD cohort                  |
|-----------------|-------------------------------|----------------------------------|
|                 | Control group | Early TTE | SMD | Control group | Early TTE | SMD |
| N               | 2028            | 903       | 4114 | 2122          |           |     |
| Sex (%)         | 0.004           |           | 0.061|               |           |     |
| F               | 976 (48.1)      | 433 (48.0)| 1803 (43.8) | 994 (46.8) |           |     |
| M               | 1052 (51.9)     | 470 (52.0)| 2311 (56.2) | 1128 (53.2) |           |     |
| Age (mean (SD)) | 62.33 (15.53)   | 62.43 (16.44)| 0.007| 62.04 (15.47) | 61.55 (15.92)| 0.031|
| Race (%)        | 0.07            |           | 0.131|               |           |     |
| Black           | 164 (8.1)       | 79 (8.7)  | 397 (8.6) | 227 (10.7)  |           |     |
| Hispanic        | 53 (2.6)        | 27 (3.0)  | 119 (2.9) | 35 (1.6)    |           |     |
| Other           | 436 (21.5)      | 170 (18.8)| 215 (5.2) | 161 (7.6)   |           |     |
| White           | 1375 (67.8)     | 627 (69.4)| 3383 (82.2)| 1699 (80.1)|           |     |
| Weight (mean (SD)) | 79.32 (20.37) | 81.82 (21.01)| 0.121| 83.62 (22.17)| 83.07 (22.70)| 0.024|
| HR (mean (SD))  | 90.71 (17.20)   | 92.18 (18.89)| 0.081| 94.73 (17.31)| 93.93 (17.09)| 0.046|
| MAP (mean (SD)) | 79.34 (11.33)   | 77.81 (11.45)| 0.134| 81.22 (12.06)| 82.38 (12.36)| 0.095|
| COPD (%)        | 0.031           |           | 0.102|               |           |     |
| CHF (%)         | 0.334           |           | 0.026|               |           |     |
| Asthma (%)      | 0.033           |           | 0.087|               |           |     |
| CKD (%)         | 0.148           |           | 0.065|               |           |     |
| Hypertension (%)| 0.11            |           | 0.035|               |           |     |
| ARDS (%)        | 0.2             |           | 0.091|               |           |     |
| Sepsis (%)      | 0.201           |           | 0.403|               |           |     |
| SOFA (mean (SD))| 5.94 (3.88)     | 7.62 (3.95)| 0.431| 6.25 (3.65)  | 6.02 (3.37) | 0.067|
| OASIS (mean (SD))| 38.25 (7.97)   | 40.91 (8.07)| 0.333| 33.27 (10.41)| 32.58 (10.66)| 0.065|
| Hb (mean (SD))  | 10.79 (2.05)    | 10.87 (2.15)| 0.039| 11.74 (2.69) | 11.53 (2.66)| 0.079|
| WBC (mean (SD)) | 12.08 (5.75)    | 12.49 (6.21)| 0.069| 12.77 (5.92) | 12.69 (5.86)| 0.013|
| pH (mean (SD))  | 7.36 (0.10)     | 7.33 (0.12)| 0.337| 7.33 (0.10)  | 7.34 (0.10) | 0.095|
| BUN (mean (SD)) | 29.67 (24.98)   | 34.32 (25.78)| 0.183| 30.12 (24.71)| 28.91 (23.17)| 0.051|
| pO2 (mean (SD)) | 142.68 (73.12)  | 129.50 (73.80)| 0.179| 120.42 (61.26)| 120.28 (58.24)| 0.002|
| Lactate (mean (SD)) | 2.05 (0.95) | 2.21 (1.10)| 0.155| 3.07 (2.52)  | 2.86 (2.34) | 0.09 |

Data are reported as the mean (SD) or no./total (%).
All data were extracted in the first 24 hours of ICU admission.
ARDS, acute respiratory distress syndrome; BUN, blood urea nitrogen; CHF, congestive heart failure; CKD, chronic kidney disease; COPD, chronic obstructive pulmonary disorder; Hb, haemoglobin; HR, heart rate; MAP, mean arterial pressure; OASIS, Oxford Acute Severity of Illness Score; SMD, standardised mean difference; SOFA, sequential organ failure assessment; WBC, white blood cell.
and the subgroup analysis also showed that early TTE was beneficial in improving in-hospital mortality in the ARDS cohort (in the MIMIC sample only) and the sepsis cohort (online supplemental eTable 8). The clinical characteristics of ARDS and sepsis patients are shown in online supplemental eTable 9.

**DISCUSSION**

In our study, the results provided evidence to support our hypotheses. After we adjusted for important confounding factors through PSM analysis, the results showed that early TTE was associated with a reduced risk of in-hospital mortality and 30-day mortality in these patients. The above results were verified by the data from the MIMIC III and eICU-CRD databases. We also tested several hypotheses to account for the mortality benefit. It may be that TTE caused a change in management and improved the patient’s prognosis, but based on current research, we cannot be completely sure.

We consider the following possible reasons for the improvement in in-hospital mortality and 30-day mortality with echocardiography: The MV affects haemodynamic as well as the respiratory system. The effect of ventilation on the haemodynamic of the heart is mainly due to changes in pleural pressure (Ppl), and Ppl affects both the inflow of the right ventricle (RV) and the outflow of the left ventricle (LV). While reducing the venous return, resulting in abnormal filling of the RV, it will also increase the resistance of the pulmonary vascular vessels and affect the outflow function of the RV. While affecting the RV, it also affects the function of the LV. In addition, in the case of inappropriate tidal volume, blood flow is mostly blocked due to transpulmonary pressure (TP) rising to a level that exceeds pulmonary artery pressure, resulting in a substantial right ventricular afterload. These effects lead to acute cardiac strain and functional and even organic lesions. Intuitive assessment of cardiac function can help clinicians adjust ventilator settings to minimise the occurrence of cardiovascular dysfunction while maintaining ventilation.

TTE is a valuable tool for monitoring haemodynamic bedside; it can easily provide dynamic haemodynamic parameters for assessing the type of shock that cannot be determined by clinical examination and information about heart-lung interactions in mechanically ventilated patients.

The afterload of RV is mainly composed of pulmonary blood flow resistance and vascular wall tension, and right ventricular ejection is tightly coupled with the pressure and compliance of the pulmonary artery. MV-induced pulmonary artery pressure variation (changes of TP and Ppl transmitted to the pulmonary blood vessels), as well as cardiac compression caused by increased lung volume, eventually leading to right heart dysfunction. In addition, due to the influence of positive pressure, the size of the inferior vena cava (IVC) will also alter, which affects the venous return. As shown in online supplemental eTable 7, early after PSM, the early TTE group had less fluid administered. Therefore, TTE may have helped directly (assessing fluid responsiveness by monitoring the RV pressure and stroke volume, as well as IVC) or indirectly (estimation of high LV filling pressure and diastolic dysfunction). The latter is associated with higher mortality in patients with sepsis.

Recently, similar to our study, Dessap et al found that early identification of right ventricular dysfunction based on acute cor pulmonale may allow the intensivist to design an intervention that might improve acute respiratory distress syndrome patient mortality, and Feng et al also showed that early transthoracic echocardiography could improve the prognosis of sepsis patients by changing their management. These findings increase the possibility that TTE can provide physicians with useful information in the ICU. In addition, echocardiography is non-invasive and provides reliable information in most cases. Thus, although the effect of echocardiography on patient outcomes has not yet been established, experts recommend that patients undergo MV for echocardiography. They suggested that echocardiography should be performed early in the course of management to quickly obtain information on ventricular dimensions and function and to assess changes in cardiac output in response to therapy. Our findings provide evidence for the early use of echocardiography in mechanically ventilated patients and the possibility that TTE can be better applied in the ICU.

The present analysis had some limitations. First, this was a retrospective registry study based on two large
CONCLUSION

To the best of our knowledge, this is the first report to determine the clinical value of TTE for patients undergoing MV. Early application of echocardiography during MV may be beneficial for the prognosis of patients. However, further prospective, multicentre, randomised controlled studies are needed to validate our results.

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Contributors XSY designed this study, collected and analysed data, and drafted the manuscript; XWZ designed this study, collected data, and drafted the manuscript; ZGC collected, compiled and analysed the data; WX and HJ analysed the data, interpreted the results and reviewed the manuscript; JS and WJC contributed with the study design and interpreted results; LLP and ZDF interpreted the data, reviewed the manuscript; YJP collected and compiled data; JPY designed and supervised this study and obtained funding, and is responsible for the overall content as guarantor. All authors read and approved the final manuscript.

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Competing interests None declared.

Patient consent for publication Not applicable.

Ethics approval Our study was in accordance with the ethical standards of the Declaration of Helsinki and was approved by the ethics review board of MIT and BIDMC (researchers certification number 1605699, record id 27752407). MMIC-III and eICU-CRD were retrospectives with lack of patient intervention, and all patients' data were de-identified; thus individual patient informed consent was not required.

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Data availability statement Data are available in a public, open access repository. Data are available upon reasonable request. All data relevant to the study are included in the article or uploaded as supplementary information. The datasets used during the current study are available from the corresponding author on reasonable request at wmpianjingye@126.com. However, reanalysis of the full data for other use requires approval by eICU-CRD (https://www.physionet.org/content/eicu-crd/2.0/) and the MMIC-III Institute (https://mimic.mit.edu/).

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