Association of overweight with postoperative acute kidney injury among patients receiving orthotopic liver transplantation: an observational cohort study

Jian Zhou¹, Lin Lyu², Lin Zhu², Yongxin Liang², He Dong² and Haichen Chu²*

Abstract

Background: Acute kidney injury (AKI) is a common postoperative complication of orthotopic liver transplantation (OLT). So far, little attention has been paid on the association between overweight and AKI after OLT, and animal models or clinical studies have drawn conflicting conclusions. The objective of our study was to determine whether overweight (BMI [Body Mass Index] ≥ 25 kg/m²) is associated with an increased risk of AKI after OLT.

Methods: This retrospective cohort study included 244 patients receiving OLT in the Affiliated Hospital of Qingdao University between January 1, 2017, and August 29, 2019. Preoperative, intraoperative, and postoperative data were collected retrospectively. The primary outcome was the development of AKI as defined by Kidney Disease, Improving Global Outcome (KIDGO) staging system. Logistic regression analysis was used to determine the relationship between overweight and the occurrence of postoperative AKI. Data analysis was conducted from September to October 2019, revision in April 2020.

Results: Among 244 patients receiving OLT (mean [standard deviation] age, 54.1 [9.6] years; 84.0% male) identified, 163 patients (66.8%) developed postoperative AKI. Overweight (BMI ≥ 25 kg/m²) was associated with a higher rate of postoperative severe AKI (stage 2/3) compared with normal weight (BMI ≤ 25 kg/m²) (47.7% vs 28.7%; adjusted odds ratio [OR], 2.539; 95% confidence interval [CI], 1.389–4.642; P = 0.002). Furthermore, patients with obese were at even higher risk of postoperative severe AKI after controlling for confounding factors (adjusted OR: 3.705; 95% CI: 1.108–12.388; P = 0.033).

Conclusions: Overweight is independently associated with an increased risk of postoperative severe AKI among patients receiving OLT. The association of BMI with severe AKI after OLT is J-shaped.

Keywords: Overweight, Body mass index, BMI, Obesity paradox, Acute kidney injury, AKI, Orthotopic liver transplantation
Background
Acute kidney injury (AKI) is a common postoperative complication of orthotopic liver transplantation (OLT), with varying occurrence ranging from 5 to 94% [1]. This extreme variability may be due to the absence of a uniform definition of AKI and rapid improvement of surgical technique and perioperative management over the past few decades. Alternatively, the study subjects are diverse in this setting. Among patients who develop AKI after OLT, approximately 8–17% need renal replacement therapy (RRT) [2]. AKI after OLT was associated with immediate complications including volume overload, metabolic acidosis, and electrolyte disturbances [3] and an increased rate of inferior long-term outcomes such as mortality, graft loss, infection, chronic kidney disease, prolonged stay in the intensive care unit (ICU), and augmented hospital costs [1, 4].

A wide variety of mechanisms have been reported for the development of AKI after OLT, such as female gender, pre-existing diabetes mellitus (DM) [5], the severity of liver disease [6], preoperative renal dysfunction [7–9], perioperative events especially hepatic ischaemia reperfusion injury (IRI) [10, 11], toxicity of immunosuppressive therapy [2, 12, 13], and other miscellaneous etiologies. As the etiology of AKI after OLT are multifactorial and not well understood, timely preventive therapy or medical interventions performed during the initiation phase of AKI can minimize the extent of injury and promote renal recovery, therefore the cornerstone for reducing the development of AKI after OLT is early recognition of high-risk patients alongside active perioperative management.

Generally, overweight is exposed to numerous comorbidities and a higher risk of postoperative complications in patients receiving operation procedure especially OLT [14]. However, there is well-grounded body of work demonstrating that patients with overweight actually do better in the long term when compared with normal weight individuals in general surgery [15], and there is anecdotal evidence that patients with overweight might have a better physiological reserve to handle the stress of surgery in clinical practice. So far, little attention has been paid on the association between overweight and AKI after OLT, and animal models or clinical studies have drawn conflicting conclusions [16, 17].

The objective of our study was to determine the association between overweight (BMI [Body Mass Index] ≥ 25 kg/m²) and postoperative AKI in patients receiving OLT. Our secondary objective was to assess whether overweight was associated with other postoperative outcomes in patients receiving OLT.

Methods
Patient population and data source
This single-center retrospective observational study included all consecutive patients undergoing OLT from January 1, 2017, to August 29, 2019, in the Affiliated Hospital of Qingdao University, Qingdao, China. All surgical procedures were performed by dedicated surgical team with extensive experience in the field. We used electronic health records to extract all study variables and archived paper medical records to identify and correct the outlying values. Data retrieval was performed by 2 reviewers (Yongxin Liang and He Dong) who were blinded to data analysis and data accuracy was verified by reabstracting 10% (n = 25) of the study sample. Patients with important missing values (ie, serum creatinine) were excluded from the model. Patients were excluded if they had: (1) Patients under 18 years of age (N = 12); (2) Combined transplantation (N = 11); (3) RRT before OLT (N = 3); (4) kidney transplantation before OLT (N = 4); (5) re-transplantation (N = 7); and (6) Lack of important data (N = 60). Ultimately, 244 patients were enrolled in final analyses.

The study protocol was approved by the Institutional Review Board of the Ethics Committee of the Affiliated Hospital of Qingdao University with waiver of participant consent (protocol number: QYFY WZLL 25725). This study was also performed in accordance with the Declaration of Helsinki (2013) of the World Medical Association [18]. This study followed the Strengthening the Reporting of Observational Studies in Epidemiology (STROBE) reporting guideline [19].

Primary exposure
Overweight and obesity are defined by the World Health Organization as abnormal or excessive fat accumulation that presents a risk to health [20]. We use the body mass index (BMI), the weight (in kilograms) of a person divided by the square of his or her height (in metres), as a crude population measure of overweight. A person with a BMI equal to or more than 25 is considered overweight. A person with a BMI of 30 or more is generally considered obese. Data for BMI were not adjusted for ascites because the volume of ascites drained at the time of transplantation did not differ significantly between groups.

Primary outcome
Our primary outcome was the occurrence of AKI after OLT. According to Kidney Disease: Improving Global Outcomes (KDIGO) serum creatinine criteria [21], the AKI after OLT was defined by an increase in serum creatinine (SCr) of 0.3 mg/dL or greater within 48 h or an increase in SCr of 1.5 or greater times baseline (to convert creatinine level to micromoles per liter, multiply by 88.4) within 7 days or a urine volume less than 0.5 ml/kg/hour for 6 h. Baseline SCr measurements were defined as the most recent value of SCr before surgery. We did not use urine volume as a criterion because this was not available for all patients.
Additional characteristics
We also collected the demographic data and clinical features that might related to AKI of all enrolled patients. These variables included age, gender, personal history of smoking and alcoholism, assessment of comorbidities (hypertension, DM, chronic kidney disease [CKD], hepatitis B virus [HBV] infection, hepatitis C virus [HCV] infection, encephalopathy, and ascites), underlying liver diseases (hepatocellular carcinoma combined with viral hepatitis, hepatocellular carcinoma, viral hepatitis, alcohol-related liver disease and others [Budd Chiari syndrome, Wilson disease, autoimmune hepatitis, drug-induced hepatitis and cryptogenic hepatitis]), Child-Pugh score, Model for End-Stage Liver Disease (MELD) score, preoperative laboratory data (creatinine, sodium, lymphocyte, lactic acid, glucose, albumin, total bilirubin), intraoperative clinical data (estimated blood loss, received albumin, received packed red blood cell [RBC], total volume of transfusion, nadir mean blood pressure, requirement of vasopressors, anhepatic time, cold ischaemia time and duration of surgery), postoperative clinical outcomes (mechanical ventilation, RRT, hospital days and hospital mortality), peak serum aspartate transaminase (AST) within the first 24 h after OLT was used as a measure of IRI [3].

Statistical analysis
Categorical data were presented as number (percentage) and compared between two groups using the chi-square test or Fisher exact test, as appropriate. Quantitative data were tested for normality by the Kolmogorov-Smirnov test, and data were considered normally distributed if the P value was > 0.05. Normally distributed quantitative data were presented as mean ± standard deviation (SD) and compared by the independent-sample t test or ANOVA as appropriate, while nonnormally distributed quantitative data were expressed as median (interquartile range [IQR]) and compared by the Mann-Whitney U test or Kruskal-Wallis H test as appropriate. We stratified baseline demographics and clinical characteristics by AKI episode (yes/no) in order to select potential confounding factors to be included in multivariate analysis. We used multivariable logistic regression to determine the relationship between BMI and AKI after OLT with adjustment for covariates. Patients were assigned to one of four categories: underweight (BMI < 18.5), normal weight (18.5 ≤ BMI < 25), overweight (25 ≤ BMI < 30), or obese (BMI ≥ 30). Non-linear relationships were explored through a restricted cubic spline with four knots at the 5th, 35th, 65th, and 95th centiles. Three models were used, with model 1 being unadjusted; model 2 adjusting model 1 for a priori defined variables based on literature research and clinical relevance (female gender, CTP score, MELD score, pre-existing CKD, pre-existing DM, preoperative SCr, requirement of vasopressors, postoperative peak AST, Tacrolimus); model 3 adjusting model 2 for potential confounding factors based on univariate analyses (female gender, preoperative SCr, preoperative lactic acid, intraoperative RBC transfusion). All regression model results were reported as an odds ratio (OR) with an associated 95% confidence interval (CI). All analyses were performed using SPSS software (SPSS, version 23.0, IBM Corp., Chicago, IL, United States) and STATA software (STATA, version 12.0, Stata Corp LP, College Station, TX, United States).

Results
Study population characteristics
A total of 244 patients aged 18 years and older receiving OLT were included in the study (Fig. 1). The mean (SD) age was 54.1 (9.6) years, with 205 (84.0%) male and 39 (16.0%) female. At the time of surgery, 88 (36.1%) patients smoked tobacco and 94 (38.2%) had an alcohol consumption. The most common underlying liver disease was hepatocellular carcinoma combined with viral hepatitis (103 [42.2%]) followed by the virus-related hepatic cirrhosis (51 [20.9%]). In total, 163 (66.8%) patients developed postoperative AKI. Compared with patients who did not develop postoperative AKI, these patients had higher BMI (24.69 [22.49–27.12] vs 23.03 [20.81–25.55], P = 0.004), preoperative SCr (65.00 [50.00–78.00] vs 71.00 [64.00–87.50], P < 0.001) and intraoperative RBC transfusion (12.00 [6.00–14.00] vs 9.00 [4.00–13.50], P = 0.026), and a slightly higher proportion of female gender (31 [19.0%] vs 8 [9.9%, P = 0.066) and preoperative blood lactic acid (1.20 [1.00–1.60] vs 1.30 [1.10–1.75], P = 0.092), though not statistically significant (0.05 < P < 0.10). All demographic and clinical characteristics of the study patients are provided in Table 1.

Incidence of AKI
Overall, 163 patients (66.8%) were subsequently diagnosed with AKI after OLT (Table 2). Approximately, 65.6% (107/163) of AKI occurred on the first postoperative day (Fig. 2). Among those patients, 76 (31.3%) was classified as stage 1, 52 (21.3%) as stage 2 and 35 (14.3%) as stage 3 acute kidney injuries. The incidence of severe AKI (stage 2/3) was 0.0, 28.7, 47.7 and 50.5% in the underweight group, normal weight group, overweight group and obese group, respectively.

Overweight and postoperative AKI
We used a restricted cubic spline to flexibly model and visualize the non-linear relationship. After adjusted for confounding factors, the association between BMI and AKI after OLT were J-shaped (Fig. 3). In the primary analyses (model 1), there were no statistically significant differences in postoperative AKI among different BMI categories. However, overweight was associated with a higher incidence of postoperative severe AKI compared...
with normal weight (crude OR: 2.266; 95% CI, 1.290–3.980; \( P = 0.004 \)). This relationship between overweight and postoperative severe AKI remained robust (adjusted OR: 2.539; 95% CI, 1.389–4.642; \( P = 0.002 \)) in the fully adjusted model (model 3). Furthermore, when compared with normal weight, patients with obese were at even higher risk of postoperative severe AKI (adjusted OR: 3.705; 95% CI, 1.108–12.388; \( P = 0.033 \)), independent of traditional risk factors. (Table 3).

**Overweight and other perioperative outcomes**

There were no statistically significant differences in postoperative ventilation time, postoperative hospitalization, or hospital mortality among underweight, normal weight, overweight and obese groups (Table 2).

**Discussion**

In this retrospective cohort study at our center, 163 (66.8%) patients who received OLT were diagnosed with postoperative AKI. Among these 163 patients, 87 (53.4%) developed severe AKI (stage 2/3) and 12 (7.4%) required RRT during postoperative hospitalization. Overweight were associated with an increased risk of postoperative severe AKI and this relationship was independent of known confounding factors.

Postoperative AKI is a common complication following OLT. In keeping with previous studies [1], the overall incidence of postoperative AKI and postoperative severe AKI at our center were 66.8 and 35.6%, respectively. The slightly higher incidence of AKI may be due to relatively lower diagnostic threshold of KDIGO criteria. This lends further support to the notion that it is time to pay close attention to postoperative AKI, as even transient AKI is associated with worse patients and graft outcomes [22].

Our results suggest that overweight is associated with an increased risk for severe AKI after OLT. A couple of explanations can be given for this association. First, while the pathophysiology of postoperative AKI is multifactorial and may have a complex interplay of each other, a major factor may be hypoperfusions of kidney and subsequent systemic and local processes that can occur during surgery [23]. Specifically, adequacy fluid resuscitation of overweight especially obese patients creates additional challenges since it is still uncertain whether the dose based on actual body weight or by formula-based weight adjustment [17]. Overweight and obesity also make it more complicated and challenging to operative field exposure and certain surgical procedures such as clamping and unclamping the portal vein which is often followed by hemodynamic instability [4]. Next, as a component of metabolic syndrome, obesity is also a significant risk factor for cardiovascular disease, hypertension, DM, and CKD, which may provide a vulnerable physiological reserve to handle the stress hypoperfusion of kidney during surgery [24]. Then, overweight and obesity leads to glomerular
Table 1 Demographic and Clinical Characteristics of the 246 patients, Stratified by AKI

| Cohort Characteristics | AKI, No. (%) | No (n = 81) | P Value  |
|------------------------|-------------|-------------|----------|
| Age, mean (SD), y      | 54.8 (9.6)  | 53.2 (9.7)  | 0.281    |
| Female, No. (%)        | 31 (19.0)   | 8 (9.9)     | 0.066    |
| BMI, median (IQR), kg/m² | 24.7 (22.5–27.1) | 23.0 (20.8–25.6) | 0.004 |
| Personal history       |             |             |          |
| Smoking, No. (%)       | 59 (37.1)   | 29 (37.7)   | 0.934    |
| Alcoholism, No. (%)    | 59 (37.6)   | 35 (46.1)   | 0.216    |
| Comorbidities          |             |             |          |
| Hypertension, No. (%)  | 44 (27.0)   | 25 (30.9)   | 0.527    |
| Diabetes mellitus, No. (%) | 58 (35.6) | 28 (34.6) | 0.876 |
| Chronic kidney disease, No. (%) | 20 (12.3) | 10 (12.3) | 0.986 |
| HBV, No. (%)           | 78 (47.9)   | 47 (58.0)   | 0.134    |
| HCV, No. (%)           | 6 (3.7)     | 4 (4.9)     | 0.641    |
| Encephalopathy, No. (%)| 11 (6.7)    | 6 (7.4)     | 0.849    |
| Ascites, No. (%)       | 37 (22.7)   | 18 (22.2)   | 0.933    |
| Underlying liver disease |             |             | 0.356    |
| Hepatocellular carcinoma, No. (%) | 65 (39.9) | 38 (46.9) |          |
| Chronic hepatitis, No. (%) |             |             |          |
| Hepatocellular carcinoma, No. (%) | 36 (22.1) | 12 (14.8) |          |
| Viral hepatitis, No. (%) | 32 (19.6)  | 19 (23.5)   |          |
| Alcohol-related liver disease, No. (%) | 11 (6.7) | 7 (8.6) |          |
| Other, No. (%)         | 19 (11.7)   | 5 (6.2)     |          |
| GFR, median (IQR)      | 145.0 (108.4–186.3) | 134.5 (104.0–164.3) | 0.144 |
| Child-Pugh score, median (IQR) | 8.0 (7.0–9.0) | 8.0 (7.0–9.0) | 0.268 |
| MELD score, median (IQR) | 6.7 (0.5–12.5) | 7.2 (2.1–14.6) | 0.474 |

Preoperative laboratory data

|                     | AKI, No. (%) | No (n = 81) | P Value |
|---------------------|--------------|-------------|---------|
| Creatinine, median (IQR), mg/dL | 65.0 (50.0–78.0) | 71.0 (64.0–88.0) | < 0.001 |
| Sodium, median (IQR), mEq/L       | 140.0 (138.0–142.0) | 140.0 (137.0–142.0) | 0.865 |
| Albumin, median (IQR), g/dL       | 35.0 (30.6–39.7) | 35.3 (30.9–40.6) | 0.520 |
| Total bilirubin, median (IQR), umol/L | 49.1 (26.6–129.3) | 44.6 (20.9–397.0) | 0.568 |
| Glucose, median (IQR), mg/dL      | 5.0 (4.4–6.4) | 5.2 (4.6–6.2) | 0.282 |
| Lactic acid, median (IQR), mmol/L | 1.2 (1.0–1.6) | 1.3 (1.1–1.8) | 0.092 |

Surgery

|                     | AKI, No. (%) | No (n = 81) | P Value  |
|---------------------|--------------|-------------|----------|
| Estimated blood loss, median (IQR), L | 2.0 (1.0–2.5) | 1.5 (0.8–2.1) | 0.146 |
| RBC transfusion, median (IQR), units | 12.0 (6.0–14.0) | 9.0 (4.0–13.5) | 0.026 |
| Nadir MAP, median (IQR), mmHg         | 64.9 (61.7–68.5) | 64.9 (63.3–69.3) | 0.923 |

Requirement of vasopressors (large dose), No. (%)

|                     | AKI, No. (%) | No (n = 81) | P Value |
|---------------------|--------------|-------------|---------|
| Duration of surgery, median (IQR), min | 531.0 (468.0–572.0) | 531.0 (470.0–570.0) | 0.811 |
| Cold ischemia time, median (IQR), min | 382.0 (326.0–384.0) | 383.0 (325.0–387.0) | 0.628 |
| Anhepatic phase, median (IQR), min   | 60.0 (52.0–60.0) | 60.0 (56.0–64.0) | 0.128 |

Postoperative parameter

|                     | AKI, No. (%) | No (n = 81) | P Value |
|---------------------|--------------|-------------|---------|
| Peak AST, median (IQR), U/L | 526.0 (72.5–795.5) | 285.0 (79.0–742.0) | 0.805 |
| Tacrolimus, median (IQR), ng/mL | 9.00 (6.65–11.50) | 8.30 (6.30–12.10) | 0.657 |
| Postoperative hospitalization, median (IQR), day | 27.0 (21.0–30.0) | 25.0 (20.0–28.0) | 0.334 |
hyperfiltration, augmented urinary albumin loss, and deterioration of renal function related to a focal segmental glomerulosclerosis. The obesity-associated ultrastructural and functional changes are collectively called obesity-related glomerulopathy [17]. Besides, obesity induced a chronic mild inflammatory condition with increased adipokines, and evidence has shown that adiponectin may have a pivotal role in the pathogenesis of acute renal ischemia/reperfusion injury [25, 26]. Moreover, patients with increasing BMI showed a rise in markers of oxidative stress and endothelial dysfunction, which are stronger predictors of postoperative AKI than inflammatory markers [27]. As such it is plausible that overweight results in an increased risk for postoperative severe AKI.

An animal study reported that obese swine receiving the surgery procedure presented with attenuated kidney dysfunction after cardiac surgery, Sleeman and colleagues proposed that chronic hyper-inflammatory status induced by pre-operative obesity had ‘pre-conditioning’ effects against ischemia/reperfusion injury as a possible explanation of the phenomenon [28]. In the study of Ju Yeon Park, no significant difference was found in occurrence of AKI after OLT between underweight and normal weight patients [29]. So far, systemic reviews aiming to understand the relationships between obesity and postoperative AKI have not been thoroughly performed in patients undergoing OLT. A previous study suggested that patients with BMI > 25 should get more attention even in patients with normal preoperative renal function [30]. Another study included recipient BMI as a predictor in their prediction model for AKI after liver transplantation [31]. In the study of Hilmi, weight > 100 kg was associated with the development of AKI within 72 h [5]. In addition, overweight has been studied in general surgery and other clinical settings showing similar outcomes [32]. Our results support these findings and our study is unique by stratifying patients by different BMI categories and the J-shaped association of BMI and severe AKI after OLT. Together these results suggest that overweight is a strong modifiable risk factor of severe AKI after OLT.

Obesity paradox has been an area of intensive investigation in diverse clinical settings over the past few years. It refers to the phenomenon that being overweight and moderately obese was actually associated with superior clinical outcomes (compared with patients with normal and low BMI) [33]. For example, In the study of Moon H, patients in the overweight groups showed a higher risk of postoperative AKI, but these patients had a more favorable prognosis in terms of overall mortality after coronary artery bypass grafting [32]. In another study that was based on patients in the ICU, an increased BMI was highly associated with the development of AKI, but with a decreased in-hospital mortality [34]. With respect to OLT, an recent meta-analysis concluded that BMI > 30 increased post-transplant complications, though length of hospital or ICU were not increased as compared to controls [14]. However, our results showed no relationship between overweight and other immediate postoperative outcomes. Future study is required with wider populations, better markers combined with animal studies to fully elucidate the physiological mechanism of obesity paradox and underlying obesity-related comorbidities. In addition, we will continue to explore the impact of BMI on progression to CKD and mortality in OLT recipients with AKI.

### Table 1

Demographic and Clinical Characteristics of the 246 patients, Stratified by AKI (Continued)

| Cohort Characteristics | AKI, No. (%) | P Value |
|------------------------|-------------|---------|
|                        | Yes (n = 163) | No (n = 81) |
| Hospital mortality, No. (%) | 5 (3.1) | 2 (2.5) | 1.000 |
Strengths and limitations
This study examines whether overweight is associated with the risk for postoperative AKI and other postoperative outcomes in patients receiving OLT. We were able to examine a cohort of patients who had a OLT at our center and we controlled for various confounders, including patient and surgery factors. In addition, we used restricted cubic spline to analyze the non-linear relationship of BMI and severe AKI after OLT, and it turned out to be J-shaped.

This study has several limitations, including but not limited to its retrospective nature. The occurrence of AKI was identified using SCr. Although it is the most widely used marker for renal function in clinical practice, it is likely that SCr overestimates preoperative renal function and delayed diagnosis and underestimate the severity of postoperative AKI, as patients awaiting OLT tend to have a reduced creatinine production compared with healthy subjects and fluid accumulation might mask the increase in SCr [3]. We mitigated this limitation by defining the baseline SCr from most recent SCr measurement made 3 to 90 days prior to operation. The incidence of postoperative AKI in our cohort (66.8%) is consistent with previous reports. A further limitation was our definition of overweight, defined as BMI ≥ 25

Fig. 2 AKI incidence and prevalence during immediate postoperative time

Fig. 3 Non-linear relationship between BMI and AKI after OLT. Legend: Cubic spline analysis showed J-shaped association of BMI and postoperative severe AKI after adjusting for confounding factors
kg/m². BMI per se has limitations since it cannot represent body composition especially in scenarios of sarcopenic obesity and oedema or ascites [14]. We did not take into account the influence of a correction of BMI according to the volume of ascites because ascites drained at the time of OLT did not differ significantly between groups in our study. Also, potential unmeasured confounding remains from variables not measured in our study (e.g., donor factors, use of angiotensin-converting enzyme inhibitors, reperfusion syndrome). Future studies should clarify whether these variables are related to the occurrence and severity of AKI after OLT.

Conclusions
Overweight is associated with an increased risk of postoperative severe AKI among patients receiving OLT. Overweight is an independent risk factor of postoperative severe AKI among patients receiving OLT after adjusting for confounding factors. There were no statistically significant differences in postoperative outcomes among different BMI categories.

Abbreviations
AKI: Acute kidney injury; OLT: Orthotopic liver transplantation; RRT: Renal replacement therapy; IR: Hepatic ischaemia reperfusion injury; BMI: Body mass index; STROBE: Strengthening the Reporting of Observational Studies in Epidemiology; KIDGO: Kidney Disease, Improving Global Outcome; SCr: Serum creatinine; DM: Diabetes mellitus; CKD: Chronic kidney disease; HBV: Hepatitis B virus; HCV: Hepatitis C virus; MELD: Model for End-Stage Liver Disease; RBC: Red blood cell; AST: Aspartate transaminase; SD: Standard deviation; IQR: Interquartile range; OR: Odds ratio; CI: Confidence interval

Authors’ contributions
JZ, HCC, and LZ contributed to study conception and design; JZ, HCC, YXL, and HD collected the data; JZ, LL, L2, analyzed and interpreted the data; HCC reviewed the statistical methods and analysis; JZ wrote the manuscript; HCC, LL, LZ made the critical revision of the article. All authors read and approved the final manuscript.

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Availability of data and materials
The datasets used and analysed during the current study are available from the corresponding author on reasonable request.

Ethics approval and consent to participate
The study protocol was approved by the Institutional Review Board of the Ethics Committee of the Affiliated Hospital of Qingdao University with waiver of participant consent (protocol number: QYFY WZLL 25725).

Consent for publication
Not applicable.

Competing interests
The authors declare that they have no competing interests.

Author details
1Department of Anesthesiology, Qingdao University Medical College, Qingdao, China. 2Department of Anesthesiology, The Affiliated Hospital of Qingdao University, No. 59, Haier Road, Qingdao 266100, Shandong Province, China.

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Table 3 Association of Overweight with Postoperative AKI

| Outcome | Model 1 | | | Model 2 | | | Model 3 | | |
|---------|---------|----------------|----------------|---------|----------------|----------------|---------|----------------|----------------|
|         | OR (95%CI) | P Value | OR (95%CI) | P Value | OR (95%CI) | P Value |
| AKI     |         |         |         |         |         |         |
| Normal weight | 1 (reference) | | 1 (reference) | | 1 (reference) | |
| Underweight | 0.360 (0.083–1.570) | 0.174 | 0.240 (0.051–1.140) | 0.073 | 0.219 (0.045–1.063) | 0.059 |
| Overweight | 1.745 (0.962–3.168) | 0.067 | 1.820 (0.985–3.363) | 0.056 | 1.781 (0.962–3.297) | 0.066 |
| Obese | 2.200 (0.586–8.260) | 0.243 | 2.522 (0.633–10.052) | 0.190 | 2.427 (0.608–9.684) | 0.209 |
| Severe AKI (stage 2/3) |         |         |         |         |         |         |
| Normal weight | 1 (reference) | | 1 (reference) | | 1 (reference) | |
| Underweight | – | | – | | – | |
| Overweight | 2.266 (1.290–3.980) | 0.004 | 2.560 (1.401–4.678) | 0.002 | 2.539 (1.389–4.642) | 0.002 |
| Obese | 2.487 (0.818–7.559) | 0.108 | 3.741 (1.119–12.510) | 0.032 | 3.705 (1.108–12.388) | 0.033 |

Note: - not available
Model 1 was unadjusted
Model 2 was adjusted for a priori defined variables based on literature research and clinical relevance (female gender, CTP score, MELD score, pre-existing CKD, pre-existing DM, preoperative SCr, requirement of vasopressors, postoperative peak AST, Tacrolimus)
Model 3 was adjusted for the same variables as model 2 and for potential confounding factors based on univariate analyses (female gender, preoperative lactic acid, intraoperative RBC transfusion)
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