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Letters to the editor

Comment on “Development of a prediction index for persistent acute kidney injury following orthotopic liver transplant”

1 Dear Editor:

2
3 Toro-Cisneros et al. [1] recently conducted a study aimed at creat-
Q4 ing a prediction index for persistent acute kidney injury (pAKI) fol-
5 lowing orthotopic liver transplant (OLT). The research sought to
6 identify key preoperative, intraoperative, and postoperative factors
7 that could forecast the onset of pAKI, a significant complication
8 known to worsen outcomes after liver transplantation. Several fac-
9 tors were found to be strongly associated with an increased risk of
10 pAKI, including male gender, the need for reoperation, transfusion of
11 more than six units of red blood cells, anhepatic time exceeding 50
12 min, and the use of norepinephrine at dosages greater than 0.25 mg/
13 kg/min. The authors created a prediction model and scoring system
14 utilizing these risk factors to identify patients at an elevated risk for
15 pAKI following OLT. The model demonstrated robust discriminatory
16 capability, with the area under the receiver operating characteristic
17 curve (AUC) indicating its efficacy in predicting pAKI within the study
18 cohort. However, several issues merit further exploration.

19 First, as described in the study [1], “**Baseline SCr was defined as**
20 **the mean SCr value corresponding to 6 months before hospitaliza-**
21 **tion or the minimum SCr value obtained during hospitalization if**
22 **the previous values were not available.**” The method employed to
23 define baseline serum creatinine (SCr) introduces considerable bias
24 and may result in inaccuracies in evaluating kidney function prior to
25 transplantation. In certain cases, baseline serum creatinine (SCr) may
26 be established using values from six months preceding hospitaliza-
27 tion, potentially failing to accurately represent the patient’s true renal
28 function prior to orthotopic liver transplantation (OLT), particularly if
29 renal function has fluctuated during that interval. Alternatively, for
30 certain patients, baseline serum creatinine (SCr) may be established
31 using SCr values from the three days preceding hospitalization, which
32 would more accurately represent the patient’s renal function imme-
33 diately prior to transplantation. The variation in baseline definition
34 may affect the accuracy of the study’s risk predictions and their clini-
35 cal implications.

36 Second, while the study [1] highlights the long-term effects of
37 pAKI on kidney function, the 5-year follow-up period may be inade-
38 quate to comprehensively assess the full range of chronic kidney dis-
39 ease (CKD) progression. Chronic kidney disease progresses through
40 various stages over an extended timeframe, and the 5-year

assessment period may overlook earlier or later stages of CKD devel- 41
opment that are essential for comprehending long-term renal out- 42
comes. Furthermore, alterations in renal function over time can be 43
affected by other factors beyond pAKI, including the kind and length 44
of immunosuppressive treatment, pre-existing comorbidities, and 45
subsequent problems that may occur following transplantation. The 46
study inadequately considered these parameters, thereby restricting 47
the generalizability and precision of the findings related to CKD 48
development post-transplantation. 49

Third, a notable difference in surgical bleeding was observed 50
between the persistent AKI and no AKI groups (3 [IQR 1.6–6] liters 51
vs. 2 [IQR 1.2–3] liters, $p < 0.01$; Table 1) [1], suggesting a possible 52
link between surgical bleeding and the onset of persistent AKI. None- 53
theless, this clinically significant variable was conspicuously missing 54
from both the bivariate and multivariate logistic regression analyses 55
investigating pAKI risk factors (Table 2). The omission signifies a 56
notable methodological limitation, given that surgical bleeding is a 57
recognized risk factor for postoperative acute kidney injury (AKI) 58
through various mechanisms, including hypotension, hemodilution, 59
and transfusion-related effects. The exclusion of this essential predic- 60
tor from the final model may have introduced selection bias and 61
potentially compromised the integrity of the proposed risk index. 62
Future investigations should thoroughly evaluate the independent 63
impact of surgical blood loss on the risk of pAKI, while considering 64
potential confounding variables. 65

Uncited references

[2,3].

Declaration of competing interest

None.

References

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