



Editorial: Transplant Oncology of Liver Malignancies

Cheng-Maw Ho*

Department of Surgery, National Taiwan University Hospital and College of Medicine, Taipei, Taiwan

Keywords: transplant oncology, liver transplantation, hepatocellular carcinoma, malignancy, risk factors

Editorial on the Research Topic

Transplant Oncology of Liver Malignancies

The evolving field of transplant oncology targets at oncological issues related to potential transplant candidates or recipients. For the Research Topic: Transplant oncology of liver malignancies, we narrowed the field down to liver malignancies of which progress had been actively accumulating. Liver transplantation sits in the center of transplant oncology (1) because, compared to other solid organ transplants, it can achieve long-term survival in transplant candidates with malignancies, mainly hepatocellular carcinoma (HCC) (2). However, by definition, transplant oncology can expand to all transplant fields.

Novel risk factors and biomarkers of malignancies that specific and unique to patients in the transplant setting, such as cross-match (3) and graft size (Kim et al.), can be evaluated under this theme. Kim et al. suggested that positive graft weight change during organ solution perfusion indicates poor prognosis in live donor liver transplantation (LDLT) for HCC. These factors may not carry a weight as heavy as the traditional oncological risk factors but could potentially fine-tune the oncological outcomes in transplant setting. Therefore, evidence derived from non-transplant setting is of paramount importance in caring transplant candidates or recipients. For example, preoperative assessment or prediction of microvascular invasion is of significance in selecting a reasonable surgical strategy to prolong patient survival [Zhang et al., (4)].

As the expansion of eligible criteria to liver transplantation for HCC, the outcomes of liver transplantation for primary, recurrent, or down-staged HCCs might be different. Downstaging of HCC to reduce post liver transplant recurrence is another hotspot for investigation [Bhatti et al., (5)]. Bhatti et al. reported their data that LDLT without prior downstaging could reach comparable survival for T2-T4a HCC with lower AFP. Further refining the eligible criteria enable these transplant candidates with HCC (once “outsiders”) to enjoy the benefit of liver transplantation.

Therapeutic role of liver transplantation in liver malignancies other than HCC (cholangiocarcinoma, sarcomas, metastases from neuroendocrine tumors, or colorectal cancer) remains to be precisely defined and requests more real-world data for validation (Finotti et al., Houben et al.).

Because almost all oncological treatments for transplant recipients are off-label use, investigator-initiated trials and real-world experiences are needed to provide solid evidence. Meanwhile, precisely harnessing immunotherapy in the setting of transplant oncology is a vital issue (6).

Looking forward, with the advance of oncology, organ transplantation could become a bridge to further oncological treatment for those who receive limited onco-therapeutic options due to the presence of organ failure. Inversely, development in transplant oncology may enhance precision immunological manipulation of oncological patients without transplant.

OPEN ACCESS

Edited and reviewed by:

Yinghong Shi,
Fudan University, China

*Correspondence:

Cheng-Maw Ho
miningho@ntu.edu.tw

Specialty section:

This article was submitted to
Surgical Oncology,
a section of the journal
Frontiers in Surgery

Received: 08 November 2021

Accepted: 14 December 2021

Published: 05 January 2022

Citation:

Ho C-M (2022) Editorial: Transplant
Oncology of Liver Malignancies.
Front. Surg. 8:811223.
doi: 10.3389/fsurg.2021.811223

I would like to express special thanks to my co-editors: Ye Xin Koh and Nicolas Syn; and reviewers for their excellent work to accomplish this Research Topic collection.

REFERENCES

1. Sapisochin G, Hibi T, Ghobrial M, Man K. The ILTS Consensus Conference on transplant oncology: Setting the stage. *Transplantation*. (2020) 104:1119–20. doi: 10.1097/TP.0000000000003175
2. Mazzaferro V, Regalia E, Doci R, Andreola S, Pulvirenti A, Bozzetti F, et al. Liver transplantation for the treatment of small hepatocellular carcinomas in patients with cirrhosis. *N Engl J Med*. (1996) 334:693–9. doi: 10.1056/NEJM199603143341104
3. Ho CM, Hu RH, Wu YM, Ho MC, Lee PH. Cross-match as an immuno-oncological risk factor for hepatocellular carcinoma recurrence and inferior survival after living donor liver transplantation: a call for further investigation. *Clin Med Insights Oncol*. (2020) 14:1179554920968774. doi: 10.1177/1179554920968774
4. Ho CM, Wu CY, Lee PH, Lai HS, Ho MC, Wu YM, et al. Analysis of the risk factors of untransplantable recurrence after primary curative resection for patients with hepatocellular carcinoma. *Ann Surg Oncol*. (2013) 20:2526–33. doi: 10.1245/s10434-013-2940-7
5. Ho CM, Lee CH, Lee MC, Zhang JF, Chen CH, Wang JY, et al. Survival after treatable hepatocellular carcinoma recurrence in liver recipients: a nationwide cohort analysis. *Front Oncol*. (2021) 10:616094. doi: 10.3389/fonc.2020.616094
6. Ho CM, Chen HL, Hu RH, Lee PH. Harnessing immunotherapy for liver recipients with hepatocellular carcinoma: a review from a transplant oncology perspective. *Ther Adv Med Oncol*. (2019) 11:1758835919843463. doi: 10.1177/1758835919843463

AUTHOR CONTRIBUTIONS

The author confirms being the sole contributor of this work and has approved it for publication.

Conflict of Interest: The author declares that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

Publisher's Note: All claims expressed in this article are solely those of the authors and do not necessarily represent those of their affiliated organizations, or those of the publisher, the editors and the reviewers. Any product that may be evaluated in this article, or claim that may be made by its manufacturer, is not guaranteed or endorsed by the publisher.

Copyright © 2022 Ho. This is an open-access article distributed under the terms of the Creative Commons Attribution License (CC BY). The use, distribution or reproduction in other forums is permitted, provided the original author(s) and the copyright owner(s) are credited and that the original publication in this journal is cited, in accordance with accepted academic practice. No use, distribution or reproduction is permitted which does not comply with these terms.