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Manuscript NO: 34551

Manuscript Type: Case Report

Heart failure after liver transplant - diagnostic and therapeutic challenge:
Case series

Manish Tandon, Sunaina Tejpal Karna, Chandra Kant Pandey, Ravindra Chaturvedi

Institutional review board statement: The study was reviewed and approved by the Institutional Review Board of Institute of Liver and Biliary Sciences Institutional Review Board vide letter no. F.25/5/107/ILBS/AC/2016/11252/511 dated 01/05/2017.

Name: Manish Tandon

Signature:  Date: 3/07/2017



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No F.25/5/107/ILBS/AC/2016/11252/511

Dated - 01/05/2017

Communication of Decision of the Institutional Ethics Committee(IEC) / Institutional Review Board(IRB)

IEC/2017/50/MA02

Protocol Title: "Retrospective analysis of patients operated upon for Liver Transplant and had post transplant Heart Failure".

Principal Investigator: Dr. Manish Tandon, Associate Professor-Anesthesia

Name & Address of Institution: INSTITUTE OF LIVER AND BILIARY SCIENCES, D-1, VASANT KUNJ, NEW DELHI

☒ New review ☐ Revised review ☐ Expedited review

Date of review (D/M/Y): 15/04/2017

Date of previous review, if revised application: NA

Decision of the IEC/ IRB:

☐ Recommended ☒ Recommended with minor modifications
☐ Revision to be submitted again ☐ Rejected

Suggestions/ Reasons/ Remarks: Comments : (Minor Modifications)

- Proforma for data collection needs to be submitted.
- Timeline for data collection needs to be mentioned in the protocol.

Remarks: The members who were present and participated in the voting are as follows:

- | | |
|-----------------------------|------------------|
| 1. Dr. A.K. Agarwal | Chairman |
| 2. Dr. Veena Malhotra | Co-Chairman |
| 3. Dr. Sanjeev Sachdeva | Member |
| 4. Dr. Akhil C. Banerjee | Member |
| 5. Sh. Sanjay Poddar | Member |
| 6. Dr. Hanuman Prasad Yadav | Member |
| 7. Dr. Nirupama Trehanpati | Member |
| 8. Dr. Amar Mukund | Member |
| 9. Dr. Rajeev Khanna | Member |
| 10. Mrs. Vibhuti Sharma | Member |
| 11. Dr. Ekta Gupta | Member Secretary |

Please note *

1. Inform IEC/IRB immediately in case of any adverse events and serious adverse events.
2. Inform IEC/IRB in case of any change of study procedure, site and investigator
3. This permission is only for period mentioned above. Annual report to be submitted to IEC/IRB.
4. Members of IEC/IRB have right to monitor the trial with prior intimation.

(Dr. Ekta Gupta)

Member Secretary, IRB/IEC, ILBS

Member Secretary, IRB/IEC, ILBS