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Letter to the editor

## Life expectancy after liver transplantation for acute hepatic necrosis



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Acute hepatic necrosis (AHN) – also referred to as acute liver failure, fulminant hepatic failure, fulminant hepatic necrosis, and fulminant hepatitis – refers to severe and acute liver injury in a patient without cirrhosis or pre-existing liver disease. Without rapid correction, if possible, of the precipitating causes, short-term mortality is high. Liver transplantation in AHN is uncommon, representing only 3% of all such transplants.[1]. Long-term survival of those who received transplantation has apparently not been studied. In the present work we compute life expectancies, and also examine whether survival has improved since 2002.

The data and methods used here are the same as those in the prior studies.[2,3] Briefly, we analyzed de-identified data from the OPTN database, which contains information on 130,665 first time, single-organ liver transplants in the USA. We restricted attention to patients meeting three criteria: (1) reason for transplant given as AHN (etiology codes 4100 to 4110), (2) age 35 to

74 years, and (3) transplanted during the MELD era, calendar years 2002 to 2018. Within the overall group we examined various subsets with a sufficient number of patients: those whose AHN was due to drug toxicity (code 4100, “Drug”), Hepatitis Type C (4104), and Hepatitis Type B-HBSag+ (4102). Remaining causes were combined as “Other”.

Demographic and medical characteristics of the patients are given in Supplemental Table 1, and the multivariate Cox (proportional hazards regression) survival models in Supplemental Table 2.

The Drug and Type C groups had very similar adjusted long-term survival (hazard ratio [HR] = 0.98,  $P = 0.85$ ), and were combined in what follows. That the “Drug” group had a worse prognosis than those with Type B Hepatitis (multivariate HR = 1.41,  $P < 0.0001$ ) was expected, as such patients are more likely to have high (rather than low) grade necrosis,[4] which is correlated with survival[5] We are not aware of any obvious reason that the Type C group had similarly poor survival

**Table 1**  
Life expectancies based on the multivariate model of Supplemental Table 2.

Starting Time	Current Age	Males				Females			
		Drug / Type C	Type B	All Others	GP	Drug / Type C	Type B	All Others	GP
<b>From tx</b>	40	17	24	21	39	19	26	23	43
	50	14	20	18	30	16	22	20	33
	60	12	17	15	22	13	19	17	25
	70	10	14	13	15	11	16	14	17
<b>1-yr post</b>	41	18	25	22	38	21	27	25	42
	51	16	21	19	29	18	23	21	33
	61	14	18	16	21	15	20	18	24
	71	12	16	14	14	13	17	15	16
<b>5-yrs post</b>	45	16	22	20	34	19	25	22	38
	55	14	19	17	26	16	21	19	29
	65	12	16	14	18	13	17	16	21
	75	10	13	12	11	11	14	13	13

compared with Type B, though it may be noted that Hepatitis C in AHN does not often manifest in isolation[6] Over the 17-year study period, there was evidence of improved survival (HR = 0.96 per calendar year;  $P < 0.0001$ ). As in prior studies, the improvement appeared to be limited to the first few years after transplant.

The resulting life expectancies by age, sex, and subset are shown in Table 1 below. Overall, life expectancy was reduced from that of the general population (GP), and varied according to age, medical risk factors, and health status. Such information may prove helpful in medical decision-making regarding treatment for both liver disease and other medical conditions.

### Disclaimer

The data reported here have been supplied by the United Network for Organ Sharing as the contractor for the Organ Procurement and Transplantation Network. The interpretation and reporting of these data are the responsibility of the authors and in no way should be seen as an official policy of or interpretation by the OPTN or the U.S. Government.

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### Declaration of Competing Interest

None.

### Supplementary materials

Supplementary material associated with this article can be found, in the online version, at [doi:10.1016/j.liver.2021.100066](https://doi.org/10.1016/j.liver.2021.100066).

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