



Original article

Palliative care and end stage liver disease: A cohort analysis of palliative care use and factors associated with referral

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ABSTRACT

Introduction and Objectives: Prevalence and mortality of chronic liver disease have risen significantly. In end stage liver disease, the survival of patients is approximately two years. Despite the poor prognosis and high symptom burden of these patients, integration of palliative care is limited. We aim to assess associated factors and trends in palliative care use in recent years.

Materials and Methods: A Multicenter retrospective cohort of patients with end stage liver disease who suffered in-hospital mortality between 2017 and 2019. Information regarding patient demographics, hospital characteristics, comorbidities, etiology, decompensations, and interventions was collected. Two-sided tests and logistic regression analysis were used to identify factors associated with palliative care use.

Results: A total of 201 patients were analyzed, with a yearly increase in palliative care consultation: 26.7 % in 2017 to 38.3 % in 2019. Patients in palliative care were older (65.72 ± 11.70 vs. 62.10 ± 11.44 ; $p = 0.003$), had a lower Karnofsky functionality scale ($\chi=18.104$; $p = 0.000$) and had higher rates of hepatic encephalopathy (32.1 % vs. 17.4 %, $p = 0.007$) and hepatocarcinoma (61.7 % vs. 26.2 %; $p = 0.000$). No differences were found for Model for End-stage Liver Disease (19.28 ± 6.60 vs. 19.90 ± 5.78 ; $p = 0.507$) or Child-Pugh scores ($p = 0.739$). None of the patients who die in the intensive care unit receive palliative care (0 % vs 31.6 %; $p = 0.000$). Half of the palliative care consultations occurred 6,5 days before death.

Conclusions: Palliative care use differs based on demographics, disease complications, and severity. Despite its increasing implementation, palliative care intervention occurs late. Future investigations should identify approaches to achieve an earlier and concurrent care model.

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1. Introduction

The prevalence and mortality of chronic liver disease have risen significantly worldwide in the last decades [1,2]. Portugal shows this same trend, with death from liver disease ranking 8th and being the European country with the highest mortality from hepatocarcinoma [3]. Typically, chronic liver disease progresses from a compensated phase to the emergence of one or more liver related decompensations, such as ascites, hepatic encephalopathy, or gastrointestinal

bleeding, a phase designated as decompensated liver cirrhosis [4]. End stage liver disease (ESLD) is defined as advanced fibrosis of the liver with one or more liver related decompensations or liver related complications such as spontaneous bacterial peritonitis, hepatorenal syndrome, hepatopulmonary syndrome, or hepatocarcinoma [4]. The survival of patients with ESLD is approximately two years, and the appearance of hepatocarcinoma can accelerate the course of the disease and worsen the prognosis [4]. In ESLD, liver transplantation is the only curative treatment. However, some patients are not eligible, and the scarcity of organs determines that only a few patients have access to this therapeutic strategy [5].

Despite the poor prognosis associated with ESLD and the high symptom burden of these patients, studies show that the integration of palliative care in ESLD is reduced and restricted, usually after exclusion from the liver transplantation list or on the last days of life [6–8]. In addition to symptomatic control, palliative care teams facilitate timely discussion of the care plan and treatment goals [9], rarely discussed in these patients [10,11]. They are also important in

Abbreviations: ESLD, End Stage Liver Disease; ICD-9-CM, International Classification of Diseases, Ninth Revision, Clinical Modification; ACLF, Acute-on-chronic Liver Failure; MELD-Na, Model for End-Stage Liver Disease (MELD)-Na; BCLC, Barcelona Clinic Liver Cancer

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supporting caregivers with high physical and psychological burdens [12]. Nevertheless, several studies show patients' misperception of palliative care, which continues to be regarded as synonymous with end-of-life, the emphasis of assistant physicians on a curative approach, and the reduced number of physicians in palliative care, which act as barriers to the early integration of palliative care in ESLD [9,13,14].

The growing research [15,16], the recognition of scientific societies through the publication of recommendations on the topic [17] and studies showing the benefit of palliative care intervention in the symptomatic improvement of patients with ESLD, may explain the growing trend in referring patients with ESLD to palliative care [18–20]. However, with most studies taking place in the United States [21,22], the existing data in Europe are still limited [23]. A previous study in Portugal found that the rate of referral to palliative care was less than 8% [24].

The objectives of this study are to determine palliative care referral rates and patterns for patients who died with ESLD between 2017 and 2019 and to identify the clinical and patient factors associated with referral.

2. Materials and Methods

2.1. Study design and patients

We performed a retrospective cohort study of patients with ESLD who died in three Portuguese hospitals between January 1, 2017, and December 31, 2019. This convenience sample aims to represent different hospital types as it allows us to know the reality of a university center with a liver transplant unit, a district hospital with a very established liver center, and a local health unit with a established palliative care team. All of these hospitals simultaneously offer structured hepatology and palliative care services.

Patients were identified by screening the electronic medical record for the presence of the International Classification of Diseases, Ninth Revision, Clinical Modification (ICD-9-CM) codes for cirrhosis, chronic liver disease, and indicators of hepatic decompensation. Patients under 18 years old, patients who died from acute hepatic failure or due to liver transplant, and those with ESLD but who died from a cause unrelated to the evolution of ESLD were excluded. These criteria aimed to minimize selection bias and capture a broad scope of patients. The study was approved by each hospital's ethical commission.

2.2. Outcomes

The primary outcomes of interest were referral to palliative care and predictors of use of palliative care. Secondary outcomes included the time between referral and death.

2.3. Variables

Data were collected using the hospital's digital information system. We collected patients' characteristics that could impact outcomes: age, sex, Charlson Comorbidity Index [25] and Karnofsky Performance Scale [26]. Liver-specific variables included liver disease etiology, liver disease complications (ascites, spontaneous bacterial peritonitis, hepatic encephalopathy, gastrointestinal bleeding, hepatorenal syndrome, acute-on-chronic liver failure [ACLF] and hepatocarcinoma), and liver disease severity (Model for End-Stage Liver Disease (MELD)-Na and Child-Pugh scores) [27,28]. Hepatocarcinoma was classified according to the Barcelona Clinic Liver Cancer (BCLC) staging system [29].

2.4. Statistical analyses

Categorical variables were specified as counts and percentages, and continuous variables were specified using mean and standard deviations when normally distributed and as medians and ranges otherwise. Comparisons between categorical variables were performed with Pearson's chi-square test or Fisher's exact test where appropriate, and comparisons between continuous variables were carried out using the Student's t-test for normally distributed variables or the Wilcoxon rank-sum test otherwise. Two-sided tests were used and were considered statistically significant when p -values were < 0.05 . Logistic regression analysis was applied to identify factors associated with palliative care use, and a post hoc power analysis was carried out. Analyses were performed using the Statistical Package for the Social Sciences (SPSS) for Mac (version 26).

2.5. Ethical statement

The study protocol conforms to the ethical guidelines of the 1975 Declaration of Helsinki as reflected in a priori approval by the Ethics Committee of Centro Hospitalar Universitário do Porto (2021.351;284-DEFI/299-CE), Centro Hospitalar Trás-os-Montes e Alto Douro (3335/CE) and Unidade Local de Saúde de Matosinhos (131/CES/JAS). As a non-interventional study of deceased patients, the Ethics Committee waived written informed consent.

3. Results

3.1. General characterization of the sample

Between 2017 and 2019, 420 patients diagnosed with chronic liver disease died, of which 219 were excluded, leaving 201 patients for analysis (Fig. 1), granting a post hoc power analysis with a test power of 90% and a type I error of 5%.

The general characteristics of this population are found in Tables 1 and 2. The mean age was 63.18 ± 11.61 years, with 50% of patients being up to 63 years old and the majority of patients being male (74.5%; $n = 150$). On average, these patients had a Charlson index of 7.02 ± 2.94 , with a median of 7.00, and the majority had a Karnofsky functionality scale of 50–70 (67.7%; $n = 136$). There are some comparative differences for the female gender (Table 2).

Overall, the main cause of ESLD is alcohol (60.2%; $n = 121$), followed by hepatitis C virus infection (22.9%; $n = 46$). Regarding the type of decompensation, it was observed that overall, the main types were: ascites (21.6%; $n = 42$); hepatic encephalopathy (21.1%; $n = 41$) and gastrointestinal bleeding (17.5%; $n = 34$). The Meld-Na was 19.72 ± 6.02 , with a median of 19.00 and almost half had a Child-Pugh class B score (48.8%; $n = 98$). On average, patients had been diagnosed with chronic liver disease for 4.01 ± 3.23 years, with a median of 3 years, and the majority of patients (63.2%; $n = 127$) did not have hepatocarcinoma.

3.2. Evolution of referral to palliative care

Were evaluated by palliative care 29.9% ($n = 60$) of the patients, with an annual increase in the referral rate from 26.7% in 2017 to 38.3% in 2019, the results are presented in Fig. 2. Even though without statistical significance ($\chi^2=1.322$; $p = 0.516$).

3.3. Factors associated with referral to palliative care (Tables 3 and 4)

Patients evaluated by palliative care had a higher average age of 65.72 ± 11.70 vs. 62.10 ± 11.44 years ($T = 8.733$; $p = 0.003$) and were significantly female ($\chi^2=8.733$; $p = 0.003$), with more women assessed than theoretically expected. It should also be noted that these patients had a higher Charlson index 7.92 ± 2.87 vs. 6.63 ± 2.90



Fig. 1. Flow diagram showing the screening and inclusion process for the cohort.

Table 1
Characteristics of patients who died with advanced chronic liver disease between 2017 and 2019.

		n	%
Gender	Female	51	25.4
	Male	150	74.5
Karnofsky Score	0–40	19	9.5
	50–70	136	67.7
	80–100	46	22.9
Etiology of Chronic Liver Disease	Alcohol	121	60.2
	Metabolic dysfunction-associated fatty liver	2	1.0
	Other	18	9.0
	Hepatitis B Virus	14	7.0
Type of decompensation	Hepatitis C Virus	46	22.9
	Ascites	42	21.6
	Hepatic encephalopathy	41	21.1
	Upper gastrointestinal Bleeding	34	17.5
	Other	3	1.5
	Spontaneous bacterial peritonitis	30	15.5
	Hepato-renal syndrome	19	9.8
Child-Pugh Score	Acute-on-Chronic Liver Failure	25	12.9
	A	33	16.4
	B	98	48.8
	C	70	34.8
Hepatocarcinoma	Yes	74	36.8
	No	127	63.2
Place of Death	General Ward	156	77.6
	Intensive Care Unit	45	22.4
Assessment by Palliative Care	Yes	60	29.9
	No	141	70.1

Table 2
Characterization of age at death, Charlson index, length of illness and MELD-Na index by sex.

Variable	Total $\bar{X} \pm s$	Female $\bar{X} \pm s$	Male $\bar{X} \pm s$	T Test (p)
Age at death	63.18 ± 11.61	67.16 ± 12.91	61.83 ± 10.85	2.648 (0.010)
Charlson Index	7.02 ± 2.94	7.16 ± 2.91	6.97 ± 2.96	0.384 (0.701)
Length of illness	4.01 ± 3.23	4.96 ± 3.35	3.69 ± 3.14	2.462 (0.015)
MELD-Na Index	19.72 ± 6.02	20.00 ± 6.48	19.20 ± 5.88	0.388 (0.698)

$\bar{X} \pm s$ – mean ± standard deviation; T test (p) – Parametric test statistics (significance level) for comparing results between men and women.

($T = 2.871$; $p = 0.005$) and a lower Karnofsky functionality scale ($\chi = 18.104$; $p = 0.000$).

The majority of patients not assessed by palliative care (66.0 %; $n = 93$) had alcohol as the cause of ESLD ($\chi = 6.537$; $p = 0.012$). The type of decompensation was significantly associated with palliative care assessment ($\chi = 17.649$; $p = 0.007$), highlighting the high number of cases of ascites and hepatic encephalopathy in these patients. The presence of hepatocarcinoma was also significantly associated with the evaluation for palliative care ($\chi = 22.707$; $p = 0.000$). No patient who died in the intensive care unit was referred to palliative care assessment ($\chi = 26.673$; $p = 0.000$).

On average, MELD-Na values are identical between patients assessed by palliative care and those who were not: 19.28 ± 6.60 vs. 19.90 ± 5.78 ($T = -0.664$; $p = 0.507$). Also, the severity of the disease according to the Child-Pugh score was not significantly associated with the assessment for palliative care ($\chi = 0.606$; $p = 0.739$). The number of decompensations, number of emergency visits and number of hospitalizations presented, on average, higher results in patients assessed by palliative care, but without statistically significant differences.

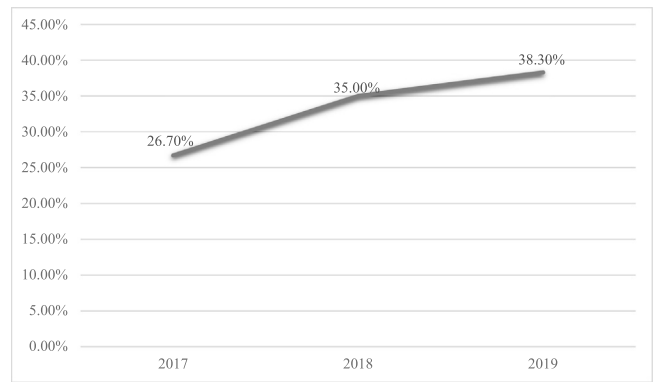


Fig. 2. Proportion of patients with end stage liver disease who received palliative care by year*

* $\chi = 1.322$; $p = 0.516$.

3.4. Multivariate analysis

To assess the significance of age, sex, Charlson index, disease duration, MELD-Na, alcohol consumption, presence of hepatocarcinoma and Karnofsky index on the probability of referral to palliative care, logistic regression was used using the enter method (Table 5). The sensitivity of the model is 89.4 % and the specificity is 48.3 %, revealing that the presence of hepatocarcinoma ($b_{HCC} = 1.256$; $X_{Wald}^2 = 6.725$; $p = 0.010$) and the Karnofsky index ($b_{Karnofsky} = -0.042$; $X_{Wald}^2 = 8.088$; $p = 0.004$) have a statistically significant effect on the logit of the probability of referral to palliative care. The absence of hepatocarcinoma reduces the probability of referral to palliative care by 71.5 % vs. 28.5 %. It is also clear that for each increase of one unit in the Karnofsky index, the probability of referral to palliative care decreases by 4.1 %.

3.5. Palliative care intervention time

The time between assessment for palliative care and the patient's death varied between a minimum of 1 day and a maximum of 196 days, with a median of 6.5 days. However, patients with hepatocarcinoma had, on average, a time between referral to palliative care and death, 30.11 ± 43.77 days, longer than patients without hepatocarcinoma, 9.09 ± 13.28 days ($T = 2.726$; $p = 0.009$). In fact, 50 % of patients with hepatocarcinoma died within 12 days of referral, and patients without hepatocarcinoma died within four days of referral.

4. Discussion

Our study shows that palliative care intervention differs based on factors such as the presence of hepatocarcinoma or the patient's functional state. We further confirm a major increase in the rate of referral to palliative care, which, however, continues to occur late. With the majority of studies using this methodology focusing on the United States [21,22], this is one of the few European studies of its kind [23,24] and the first in the country to use a multicenter strategy.

This study describes a population with similar characteristics to other studies, showing the weight of alcohol consumption in chronic liver disease [23,30], as well as of hepatitis C virus infection. The low number of cases of chronic liver disease caused by metabolic dysfunction-associated fatty liver disease is highlighted in comparison to other studies [23,30]. It is known that the etiologies of cirrhosis are changing over time and differ by region in the world, so in the near future, we believe we can see an increase in our population of chronic liver disease caused by metabolic dysfunction-associated fatty liver disease.

Table 3
Factors associated with referral to palliative care.

		Palliative care assessment		Total n (%)	χ (p)
		Yes n (%)	No n (%)		
Sex	Female	22 (36.7)	29 (20.6)	51 (25.4)	8.733 (0.003)
	Male	38 (63.3)	112 (79.4)	150 (74.6)	
Karnofsky Score	0-40	10 (16.7)	9 (13.3)	19 (9.5)	18.104 (0.000)
	50-70	47 (78.3)	89 (63.1)	136 (67.7)	
	80-100	3 (5.0)	43 (30.5)	46 (22.9)	
Chronic liver disease etiology	Alcohol	28 (46.1)	93 (66.0)	121 (60.2)	6.537 (0.012)
	Other	32 (53.3)	48 (34.0)	80 (39.8)	
Type of decompensation	Ascites	18 (32.1)	24 (17.4)	42 (21.6)	17.649 (0.007)
	Hepatic encephalopathy	17 (30.4)	24 (17.4)	41 (21.1)	
	Gastrointestinal bleeding	11 (19.6)	23 (16.7)	34 (17.5)	
	Other	0 (0.0)	3 (2.2)	3 (1.5)	
	Spontaneous bacterial peritonitis	5 (8.9)	25 (18.1)	30 (15.5)	
	Hepatorenal syndrome	3 (5.4)	16 (11.6)	19 (9.8)	
Child-Pugh	ACLF	2 (3.6)	23 (16.7)	25 (12.9)	0.606 (0.739)
	A	8 (13.3)*	25 (17.7)	33 (16.4)	
	B	30 (50.0)	68 (48.2)	98 (48.8)	
	C	22 (36.7)	48 (34.0)	70 (34.8)	
Hepatocarcinoma	Yes	61.7 %	26.2 %	36.8 %	22.707 (0.000)
	No	38.3 %	73.8 %	63.2 %	
Hospital with transplant unit	Yes	56.7 %	77.3 %	71.1 %	8.733 (0.003)
	No	43.3 %	22.7 %	28.9 %	
Location of death	Ward	60 (100.0)	96 (68.1)	156 (77.6)	26.673 (0.000)
	ICU	0 (0.0)	45 (31.6)	45 (22.4)	

χ (p) – Chi-square independence test statistic (significance level)

* elderly patients with hepatocarcinoma

Table 4

Analysis of age at death, Charlson index, length of illness, MELD-Na index, number of decompensations, emergencies, or hospitalizations when referring to palliative care.

Variable	Palliative care assessment $\bar{X} \pm s$	No palliative care assessment $\bar{X} \pm s$	T Test (p)
Age at death	65.72 ± 11.70	62.10 ± 11.44	2.038 (0.043)
Charlson Index	7.92 ± 2.87	6.63 ± 2.90	2.871 (0.005)
Length of illness	4.50 ± 3.64	3.80 ± 3.03	1.405 (0.161)
MELD-Na Index	19.28 ± 6.60	19.90 ± 5.78	-0.664 (0.507)
Number of decompensations	1.47 ± 1.31	1.26 ± 1.26	1.087 (0.282)
Number of emergencies	2.45 ± 2.06	1.91 ± 1.98	1.729 (0.085)
Number of hospitalizations	0.95 ± 1.08	0.69 ± 0.94	1.726 (0.086)

$\bar{X} \pm s$ – mean ± standard deviation; T (p) – Parametric test statistics (level of significance) for comparing results between patients considered for palliative care and those who were not.

Table 5

Multivariate analysis of factors associated with palliative care referral.

Variable	B (coefficient)	Wald	p	Exp(B)	Confidence interval 95%	
					Linf	Lsup
Age	0.004	0.050	0.822	1.004	0.969	1.041
Sex	-0.708	2.954	0.086	0.493	0.220	1.104
Charlson	-0.008	0.007	0.936	0.992	0.826	1.192
Length of illness	0.004	0.005	0.942	1.004	0.902	1.117
MELD-Na	0.015	0.204	0.652	1.015	0.952	1.082
Alcohol consumption	0.629	2.296	0.130	1.876	0.831	4.232
Hepatocarcinoma	-1.256	6.725	0.010	0.285	0.110	0.736
Transplant unit	-1.229	7.997	0.005	0.292	0.125	0.686
Karnofsky	-0.042	8.088	0.004	0.959	0.932	0.987
Constant	2.943	2.264	0.132	18.973		

This study confirms that palliative care intervention in ESLD varies according to the characteristics of the patient, the disease, and its complications. It also suggests that referral to palliative care is associated with being older and being female. However, in some studies, advanced age appears to be associated with referral to palliative care [21,31], this finding is not unanimous, with studies failing to prove

the influence on referral [30] or to show this relationship in the presence of younger individuals [32].

The results were also able to demonstrate that the existence of comorbidities or low functionality also seems to be related to greater referral to palliative care. In logistic regression, a statistically significant effect was documented on the probability of referral to palliative

care, with a reduction in the referral rate with an improvement in the patient's functionality. In fact, there have been publications in the literature suggesting the relevance of the Karnofsky Performance Scale in the mortality of patients with ESLD [27,33].

Similar to another study, the presence of liver disease caused by alcohol was associated with non-referral for palliative care [30]. But we have to remember that not only these patients, but also their caregivers present high rates of clinical depression and burden [34]. The types of decompensation, namely ascites and hepatic encephalopathy, were also associated with greater referral to palliative care. These findings have also been demonstrated in other studies, both for ascites [21,31], and for hepatic encephalopathy [32]. We believe that the symptomatic and functional impact that these complications have on the patient may favor the need to discuss a care plan and thus unlock referral to palliative care.

It is also confirmed what appears to be the most consensual data in many studies. The association between the presence of hepatocarcinoma and palliative care assessment generally induces a high referral rate [20,30]. In the logistic regression model presented, the absence of hepatocarcinoma reduces the probability of referral to palliative care by 71.5 %. We also verify the reduced implementation of palliative care in patients admitted to the intensive care unit [32,35]. In our study, 0 patients were referred to palliative care. There are studies showing the apparent influence of Child-Pugh and Meld-Na on referral to palliative care [31]. However, similar to another study [30], this finding was not observed in the sample examined.

This study shows an increasing prevalence in referral to palliative care when compared to initial studies [6,21,31,36]. There is an average percentage of 29.9 %, close to a study carried out in recent years [22]. The referral rate in 2019 (38.3 %) is already close to values published in recent studies [32,37]. It must be acknowledged that, being an analysis of deceased patients, there may be a greater referral to palliative care compared to patients who remained alive and that there are methodological differences between the different studies, as well as external factors, such as the hospital culture and the existence of structured hepatology and palliative care services, which can influence the rates described in the literature.

However, 50 % of patients in the sample died within 6.5 days after assessment by palliative care. In other words, we continue to have late referrals, as supported by data in other studies [11,21,22]. In fact, in a published questionnaire, 97 % of liver transplant team professionals continued to regard palliative care teams as end-of-life care [14]. There are several studies that show the benefit of palliative care intervention in improving the symptoms of patients with ACLD [18–20], as well as reducing disproportionate measures [38], reducing healthcare costs [39], and increasing the discussion of advance directives [38]. Being referral too often delayed until patients are approaching death, the psycho-social and therapeutic benefits of palliative care input may not be achievable. Therefore, more than meeting the referral rate, we will have to start working on timely and adequate referrals as representative of the quality of care.

This is one of only a few European studies examining palliative care referral in ESLD and is strengthened by its multicentric nature and considerable sample size. So, it is anticipated that these findings can be used as a baseline for comparison in future studies. There are some limitations, including the retrospective nature of the study. Findings may be limited by the support of clinical documentation, and, being a retrospective study, we were unable to gather information on patient satisfaction, symptom burden, or family's perspective. Finally, the geographic, demographic, and provider practice patterns may limit the generalizability of our findings to other centers.

The implementation of this work is intended to highlight the importance of the topic for the scientific community and reinforce the existing knowledge in the area of palliative care and ESLD, in line with what was suggested by Patel *et al.* [16] and Fricker *et al.* [40].

5. Conclusions

In this study, we demonstrate the increasing intervention of palliative care in patients with ESLD, especially taking into account the functional status or the presence of hepatocarcinoma. However, we continue to see late referrals. Future investigations should identify approaches to achieve an early and concurrent care model that incorporates palliative services into ESLD teams in order to improve patient satisfaction, quality of life, and symptom burden of patients with ESLD.

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Conflicts of interest

None.

Author contributions

Hugo Miguel Oliveira: Investigation, Conceptualization, Visualization, Data curation, Formal analysis, Software, Formal analysis, Writing – original draft, Writing – review & editing. Helena Pessegueiro Miranda: Conceptualization, Data curation, Formal analysis, Writing – original draft, Writing – review & editing. Francisca Rego: Conceptualization, Data curation, Formal analysis, Writing – original draft, Writing – review & editing. Rui Nunes: Conceptualization, Data curation, Formal analysis, Writing – original draft, Writing – review & editing.

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References

- [1] Sepanlou SG, Safiri S, Bisignano C, Ikuta KS, Merat S, Saberifirooz M, et al. The global, regional, and national burden of cirrhosis by cause in 195 countries and territories, 1990–2017: a systematic analysis for the Global Burden of Disease Study 2017. *Lancet Gastroenterol Hepatol* 2020;5(3):245–66. [https://doi.org/10.1016/S2468-1253\(19\)30349-8](https://doi.org/10.1016/S2468-1253(19)30349-8).
- [2] Moon AM, Singal AG, Tapper EB. Contemporary epidemiology of chronic liver disease and cirrhosis. *Clin Gastroenterol Hepatol* 2020;18(12):2650–66. <https://doi.org/10.1016/j.cgh.2019.07.060>.
- [3] Da Rocha MC, Marinho RT, Rodrigues T. Mortality Associated with hepatobiliary disease in Portugal between 2006 and 2012. *GE Port J Gastroenterol* 2018;25(3):123–31. <https://doi.org/10.1159/000484868>.
- [4] D'Amico G, Garcia-Tsao G, Pagliaro L. Natural history and prognostic indicators of survival in cirrhosis: a systematic review of 118 studies. *J Hepatol* 2006;44(1):217–31. <https://doi.org/10.1016/j.jhep.2005.10.013>.
- [5] Terrault NA, Francoz C, Berenguer M, Charlton M, Heimbach J. Liver transplantation 2023: status report, current and future challenges. *Clin Gastroenterol Hepatol* 2023;21(8):2150–66. <https://doi.org/10.1016/j.cgh.2023.04.005>.
- [6] Poonja Z, Brisebois A, Van Zanten SV, Tandon P, Meeberg G, Karvellas CJ. Patients with cirrhosis and denied liver transplants rarely receive adequate palliative care or appropriate management. *Clin Gastroenterol Hepatol* 2014;12(4):692–8. <https://doi.org/10.1016/j.cgh.2013.08.027>.
- [7] Kathalia P, Smith A, Lai JC. Underutilization of palliative care services in the liver transplant population. *World J Transplant* 2016;6(3):594. <https://doi.org/10.5500/wjt.v6.i3.594>.
- [8] Peng JK, Hepgul N, Higginson IJ, Gao W. Symptom prevalence and quality of life of patients with end-stage liver disease: a systematic review and meta-analysis. *Palliat Med* 2019;33(1):24–36. <https://doi.org/10.1177/0269216318807051>.
- [9] Verma M, Tapper EB, Singal AG, Navarro V. Nonhospice palliative care within the treatment of end-stage liver disease. *Hepatology* 2020;71(6):2149–59. <https://doi.org/10.1002/hep.31226>.

- [10] Sprange A, Ismond KP, Hjartarson E, Chavda S, Carbonneau M, Kowalczewski J, et al. Advance care planning preferences and readiness in cirrhosis: a prospective assessment of patient perceptions and knowledge. *J Palliat Med* 2020;23(4):552–7. <https://doi.org/10.1089/jpm.2019.0244>.
- [11] Najafian N, Sack JS, Delisle AM, Jakab S. Advance care planning for patients with cirrhosis in a structured inpatient/outpatient hepatology program. *J Palliat Med* 2019;22(11):1445–8. <https://doi.org/10.1089/jpm.2018.0261>.
- [12] Roth K, Lynn J, Zhong Z, Borum M, Dawson NV. Dying with end stage liver disease with cirrhosis: insights from support. *J Am Geriatr Soc* 2000;48(S1):S122–30. <https://doi.org/10.1111/j.1532-5415.2000.tb03121.x>.
- [13] Ufere NN, Donlan J, Waldman L, Patel A, Dienstag JL, Friedman LS, et al. Physicians' perspectives on palliative care for patients with end-stage liver disease: a national survey study. *Liver Transplant* 2019;25(6):859–69. <https://doi.org/10.1002/lt.25469>.
- [14] Beck KR, Pantilat SZ, O'Riordan DL, Peters MG. Use of palliative care consultation for patients with end-stage liver disease: survey of liver transplant service providers. *J Palliat Med* 2016;19(8):836–41. <https://doi.org/10.1089/jpm.2016.0002>.
- [15] Verma S, Hingwala J, Low JTS, Patel AA, Verma M, Bremner S, et al. Palliative clinical trials in advanced chronic liver disease: challenges and opportunities. *J Hepatol* 2023. <https://doi.org/10.1016/j.jhep.2023.06.018>.
- [16] Patel AA, Woodrell C, Ufere NN, Hansen L, Tandon P, Verma M, et al. Developing priorities for palliative care research in advanced liver disease: a multidisciplinary approach. *Hepatol Commun* 2021;5(9):1469–80. <https://doi.org/10.1002/hep4.1743>.
- [17] Rogal SS, Hansen L, Patel A, Ufere NN, Verma M, Woodrell CD, et al. AASLD practice guidance: palliative care and symptom-based management in decompensated cirrhosis. *Hepatology* 2022;76(3):819–53. <https://doi.org/10.1002/hep.32378>.
- [18] Baumann AJ, Wheeler DS, James M, Turner R, Siegel A, Navarro VJ. Benefit of early palliative care intervention in end-stage liver disease patients awaiting liver transplantation. *J Pain Symptom Manag* 2015;50(6):882–6. <https://doi.org/10.1016/j.jpainsymman.2015.07.014>.
- [19] Verma M, Navarro V. Palliative care for end-stage liver disease population. *J Palliat Med* 2017;20(1):4. <https://doi.org/10.1089/jpm.2016.0375>.
- [20] Rush B, Walley KR, Celi LA, Rajoriya N, Brahmamania M. Palliative care access for hospitalized patients with end-stage liver disease across the United States. *Hepatology* 2017;66(5):1585–91. <https://doi.org/10.1002/hep.29297>.
- [21] Holden JH, Shamseddeen H, Johnson AW, Byriel B, Subramoney K, Cheng YW, et al. Palliative care and hospice referrals in patients with decompensated cirrhosis: what factors are important? *J Palliat Med* 2020;23(8):1066–75. <https://doi.org/10.1089/jpm.2019.0501>.
- [22] Ufere NN, Halford JL, Caldwell J, Jang MY, Bhatt S, Donlan J, et al. Health care utilization and end-of-life care outcomes for patients with decompensated cirrhosis based on transplant candidacy. *J Pain Symptom Manag* 2020;59(3):590–8. <https://doi.org/10.1016/j.jpainsymman.2019.10.016>.
- [23] Woodland H, Buchanan RM, Pring A, Dancox M, McCune A, Forbes K, et al. Inequity in end-of-life care for patients with chronic liver disease in England. *Liver Int* 2023(July):1–11. <https://doi.org/10.1111/liv.15684>.
- [24] Carvalho JR, Vasconcelos M, Marques da Costa P, Marinho RT, Fatela N, Raimundo M, et al. Identifying palliative care needs in a Portuguese liver unit. *Liver Int* 2018;38(11):1982–7. <https://doi.org/10.1111/liv.13865>.
- [25] Charlson ME, Pompei P, Ales KL, MacKenzie CR. A new method of classifying prognostic comorbidity in longitudinal studies: development and validation. *J Chronic Dis* 1987;40(5):373–83. [https://doi.org/10.1016/0021-9681\(87\)90171-8](https://doi.org/10.1016/0021-9681(87)90171-8).
- [26] Crooks V, Waller S, Smith T, Hahn TJ. The use of the Karnofsky performance scale in determining outcomes and risk in geriatric outpatients. *J Gerontol* 1991 Jul;46(4):M139–44. <https://doi.org/10.1093/geronj/46.4.m139>.
- [27] Kamath PS, Wiesner RH, Malinchoc M, Kremers W, Therneau TM, Kosberg CL, et al. A model to predict survival in patients with end-stage liver disease. *Hepatology* 2001;33(2):464–70. <https://doi.org/10.1053/jhep.2001.22172>.
- [28] Pugh RN, Murray-Lyon IM, Dawson JL, Pietroni MC, Williams R. Transection of the oesophagus for bleeding oesophageal varices. *Br J Surg* 1973;60(8):646–9. <https://doi.org/10.1002/bjs.1800600817>.
- [29] Forner A, Reig M, Bruix J. Hepatocellular carcinoma. *Lancet* 2018;391(10127):1301–14. [https://doi.org/10.1016/S0140-6736\(18\)30010-2](https://doi.org/10.1016/S0140-6736(18)30010-2).
- [30] Chen H, Johnston A, Palmer A, Mickenbecker M, O'Sullivan T, Clark P. Too little, too late: palliation and end-stage liver disease. *J Gastroenterol Hepatol* 2021;1–4 (Australia). <https://doi.org/10.1111/jgh.15499>.
- [31] O'Leary JG, Tandon P, Reddy KR, Biggins SW, Wong F, Kamath PS, et al. Underutilization of hospice in inpatients with cirrhosis: the NACSELD experience. *Dig Dis Sci* 2020;65(9):2571–9. <https://doi.org/10.1007/s10620-020-06168-8>.
- [32] Kieffer SF, Tanaka T, Ogilvie AC, Gilbertson-White S, Hagiwara Y. Palliative care and end-of-life outcomes in patients considered for liver transplantation: a single-center experience in the US midwest. *Am J Hosp Palliat Med* 2022;1–9. <https://doi.org/10.1177/10499091221142841>.
- [33] Tandon P, Reddy KR, O'Leary JG, Garcia-Tsao G, Abraldes JG, Wong F, et al. A Karnofsky performance status–based score predicts death after hospital discharge in patients with cirrhosis. *Hepatology* 2017;65(1):217–24. <https://doi.org/10.1002/hep.28900>.
- [34] Brisebois A, Ismond KP, Carbonneau M, Kowalczewski J, Tandon P. Advance Care Planning (ACP) for specialists managing cirrhosis: a focus on patient-centered care. *Hepatology* 2018;67(5):2025–40. <https://doi.org/10.1002/hep.29731>.
- [35] Oud L. Patterns of palliative care utilization among patients with end stage liver disease during end-of-life hospitalizations: a population-level analysis. *J Crit Care* 2018;48:290–5. <https://doi.org/10.1016/j.jcrrc.2018.09.012>.
- [36] Plunkett A, Mortimore M, Good P. Palliative care in cirrhosis with decompensation. *Intern Med J* 2019;49(7):904–8. <https://doi.org/10.1111/imj.14348>.
- [37] Sohal A, Chaudhry H, Sharma R, Dhillon N, Kohli I, Singla P, et al. Recent trends in palliative care utilization in patients with decompensated liver disease: 2016–2020 National Analysis. *J Palliat Med* 2024;27(3):335–44. <https://doi.org/10.1089/jpm.2023.0367>.
- [38] Kearney A, Tiwari N, Cullen O, Legg A, Arbi I, Douglas C, et al. Improving palliative and supportive care in advanced cirrhosis: the HepatoCare model of integrated collaborative care. *Intern Med J* 2023;1–9. <https://doi.org/10.1111/imj.16248>.
- [39] Patel AA, Walling AM, May FP, Saab S, Wenger N. Palliative care and health care utilization for patients with end-stage liver disease at the end of life. *Clin Gastroenterol Hepatol* 2017;15(10):1612–1619.e4. <https://doi.org/10.1016/j.cgh.2017.01.030>.
- [40] Fricker ZP, Serper M. Current knowledge, barriers to implementation, and future directions in palliative care for end-stage liver disease. *Liver Transplant* 2019;25(5):787–96. <https://doi.org/10.1002/lt.25434>.