Poster

Health-related quality of life and stigma related to chronic hepatitis B: a systematic literature review

Poster No. PO-1725

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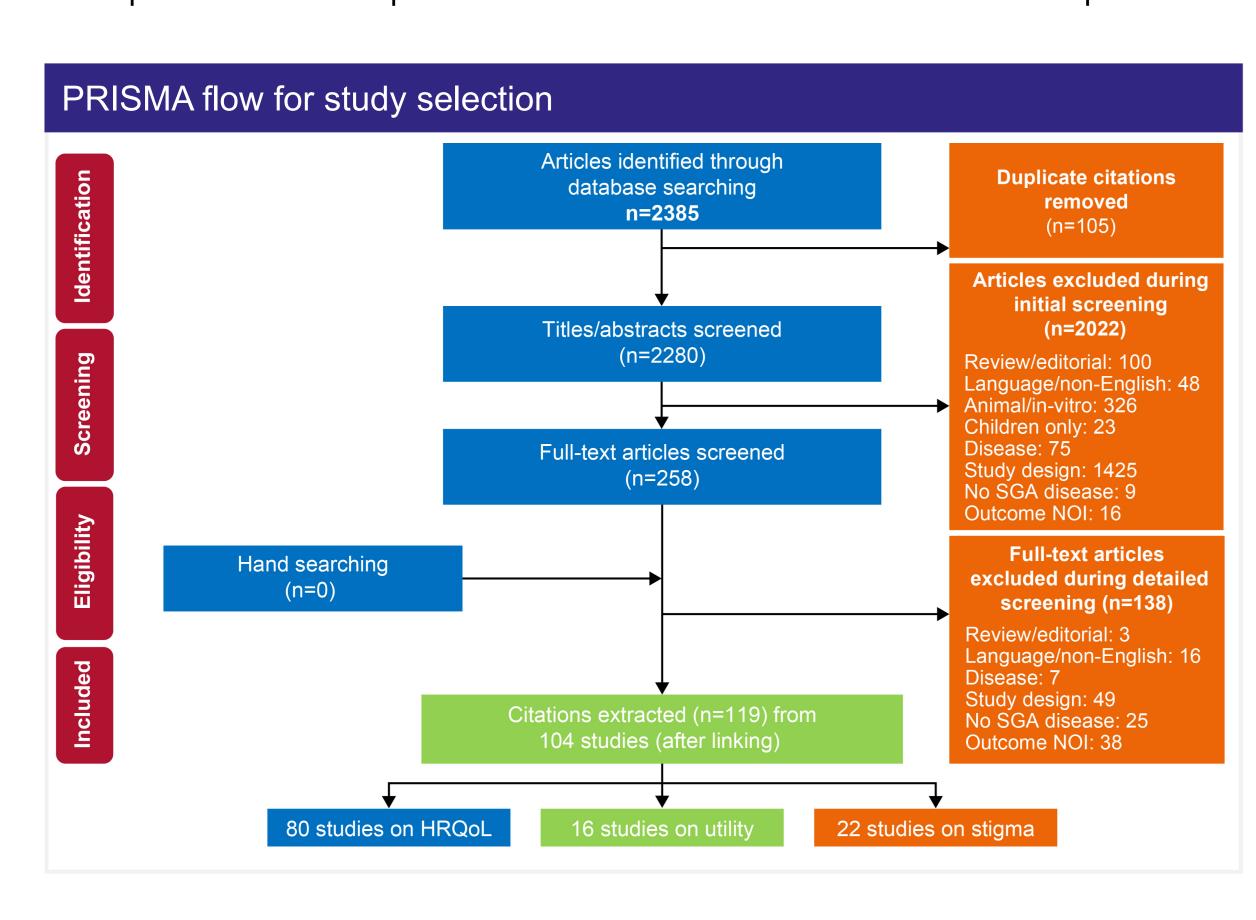
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Background

- The adverse impact of CHB infection on patients' HRQoL and patients' assessments of their health status (health utilities) has been investigated in numerous studies with diverse methodologies
- This is the first SLR to comprehensively identify and assess the published evidence on the HRQoL, utilities and stigma experienced by patients with CHB

Methods

- Data sources: Embase, MEDLINE, MEDLINE-In-Process and Cochrane Central from January 2004 to February 2020. Conference abstracts from 2017 to 2019
- Data extraction and quality assessment: data were extracted into a predefined data extraction form. Quality/risk of bias of included studies was assessed using the NICE critical appraisal checklist for RCTs and the Downs & Black (1998) checklist for nonrandomised/observational/single-arm studies^{1,2}
- Flow of studies through the review process: conducted and reported in accordance with PRISMA guidelines³
- Representative examples of data from these studies are shown in this poster



Assessment tools included:

Generic/nondisease-specific tools, e.g. • SF-36 (n=36)

• EQ-5D (n=11)

• World Health Organization Quality of Life Instrument, Short Form (WHOQOL-BREF, n=7)

Liver-disease-specific tools, e.g.

Chronic Liver Disease Questionnaire (CLDQ, n=16)

Hepatitis-specific tools, e.g.

- Hepatitis B Quality of Life (HBQOL, n=3)
- Hepatitis Quality of Life Questionnaire (HQLQ, n=2)

Generic PRO instruments assessing:

- Depression
- Anxiety Fatigue

References

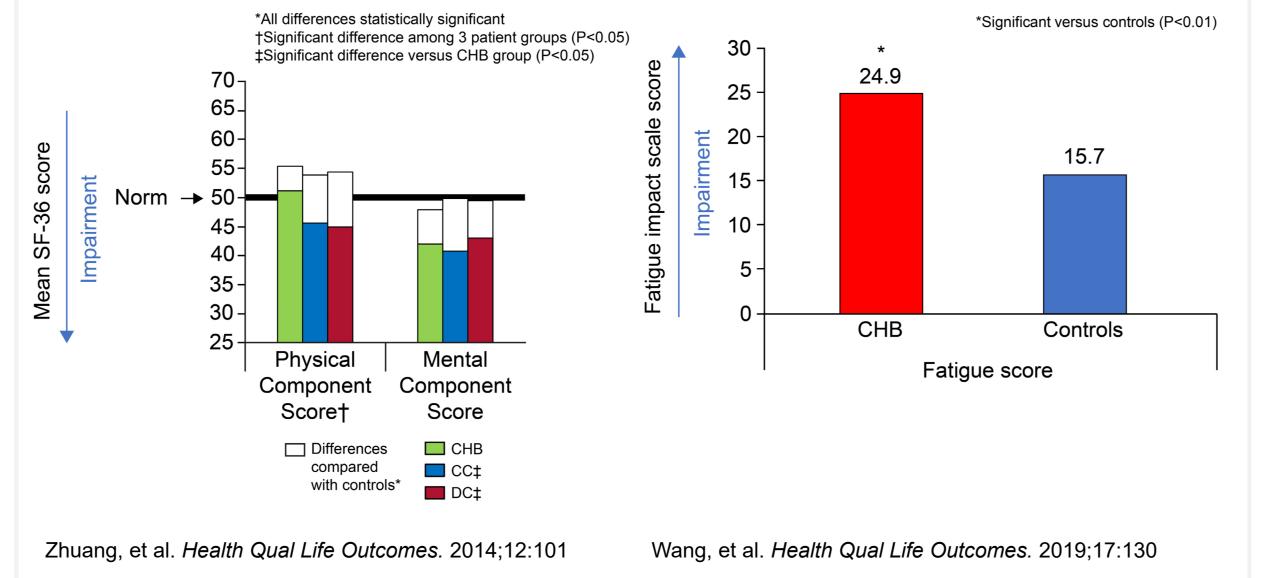
1. National Institute for Health and Care Excellence (NICE). *Specification for manufacturer/sponsor submission of* evidence. 2015. Available at: http://www.nice.org.uk/About/What-we-do/Our-Programmes/NICE-guidance/NICEtechnology-appraisal-guidance. Accessed 2 June 2021.

2. Downs SH, Black N. J Epidemiol Community Health. 1998;52:377-384.

3. Liberati A, et al. BMJ. 2009;339:b2700.

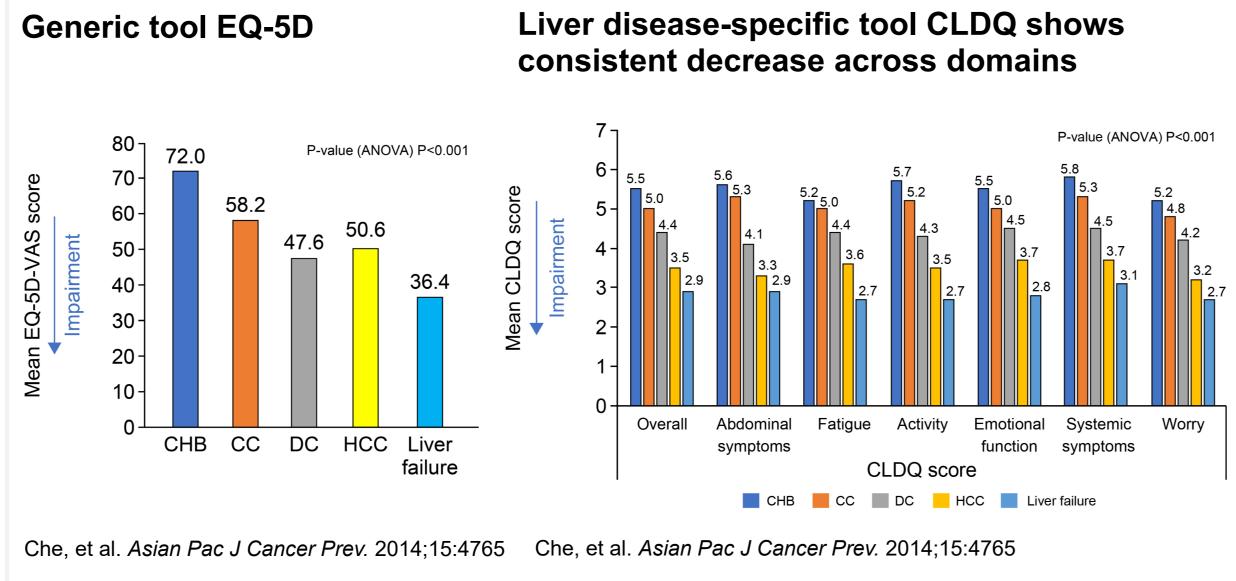
Results





In addition to seeing consistent impact on mental health and fatigue domains, domains sometimes impacted included general health, vitality and physical health

HRQoL declined significantly with worsening disease severity, from CHB to liver failure

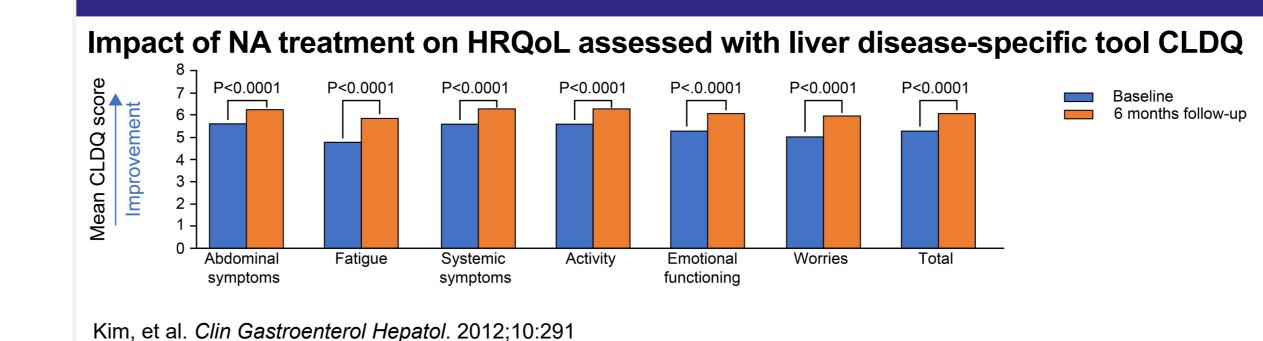


ANOVA used to compare mean scores among the five disease groups

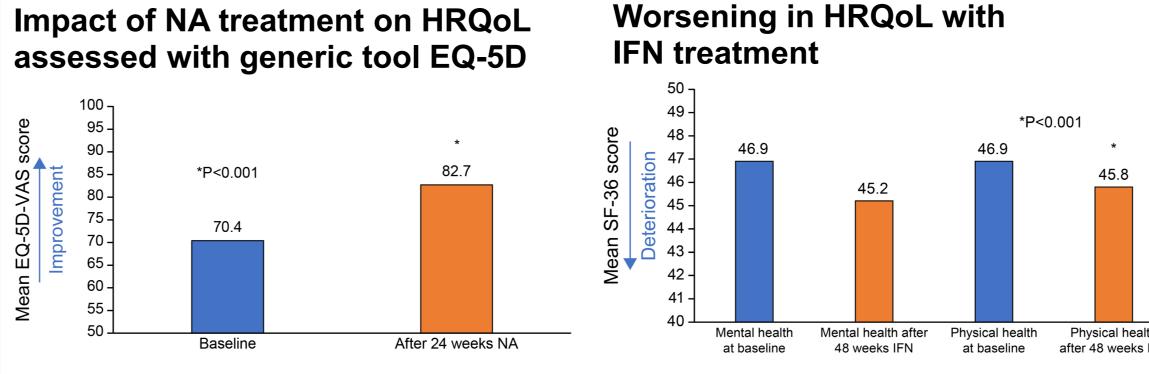
Disclosures

HUV showed a similar trend to declining HRQoL with worsening disease severity: HUVs ranged from 0.75 to 0.81 in asymptomatic carriers, 0.56 to 0.89 in compensated cirrhosis, down to 0.30 to 0.85 in decompensated cirrhosis and 0.32 to 0.85 in hepatocellular carcinoma. HUVs ranged from 0.54 to 0.63 in the first year after liver transplant, increasing to 0.62 to 0.86 in subsequent years after transplant. The ranges provided are lowest and highest values from different studies

NA therapy improved HRQoL in some studies, which correlated with viral suppression. In particular, improvements were seen in physical health, with less consistency in impact on other domains. IFNs were associated with worse HRQoL across all domains

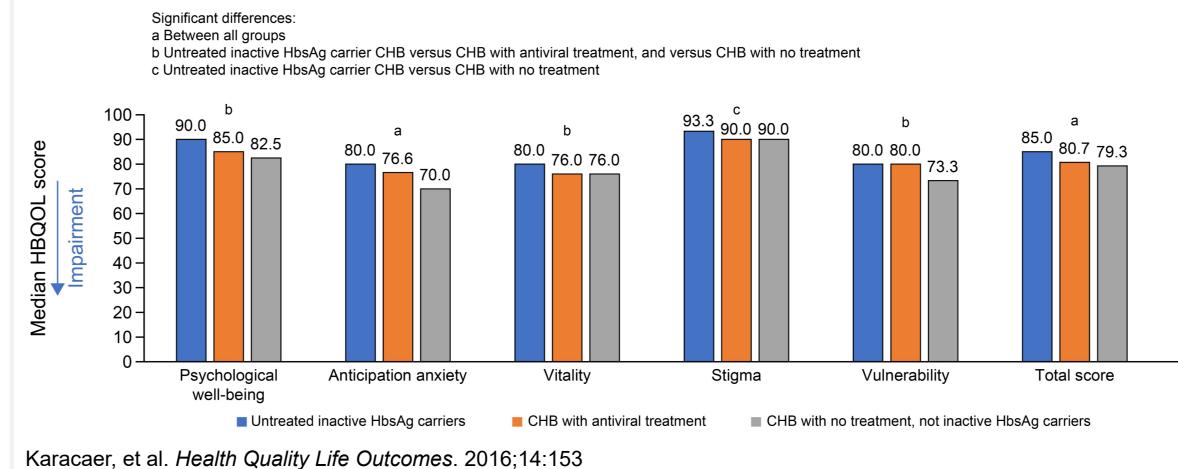


Kim, et al. Clin Gastroenterol Hepatol. 2012;10:291

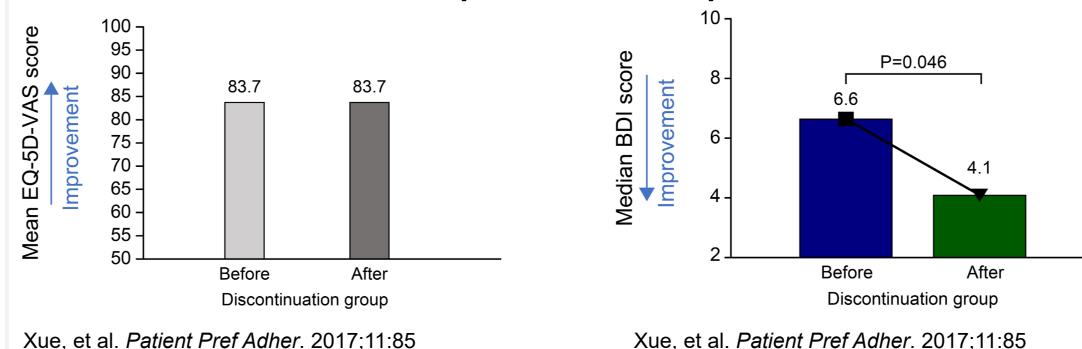


Marcellin, et al. Liver Int. 2008;28:477

Patients receiving and eligible for treatment have better HRQoL than patients eligible but untreated. However, HRQoL is highest in untreated inactive carriers







Conclusion

- Patients with CHB have impaired HRQoL, particularly in fatigue, mental health, emotional and social functioning, with no consistent improvement in these domains with current treatments
- The impact of interventions on HRQoL in clinical practice and trials should be further evaluated with diseasespecific instruments like HBQOL that focus on psychological well-being and stigma
- Stigma has been shown to have a major impact on patients with CHB infection. Research is needed on how best to reduce stigma and its effects

Stigma was shown to have a major impact on people living with CHB



- Knowledge gaps Fear of discrimination
- Level of education

Marital relationships

- Beliefs related to CHB (e.g. caused by a curse)
- Public health messages

Cultural constraints

 Access to health services Behaviors of health personnel

Socio-cultural & community factors

contributing to CHB-related stigma

Barriers to engagement

 Willingness to screen Employment

(CHB and caregivers)

- Seeking care
- Diagnosis acceptance Interpersonal relationships

Assessment tools

BDI: self-reported measure of depression severity, involving 21 items, each rated on a scale from 0 to 3. Global depression score is calculated from the mean score CLDQ: liver disease-specific questionnaire, including 29 items divided into six domains: fatigue,

abdominal pain, motional function, systemic symptoms, activity, and worry. Summary scores for each domain range from 1 (most impairment) to 7 (least impairment). Overall CLDQ score is calculated from the mean score

Impact and consequence of CHB-related stigma

EQ-5D visual analogue scale (VAS): records patient's self-rated health as a vertical line, with 0 representing worst and 100 the best possible health

HBQOL: hepatitis B-specific tool for assessment of HRQoL. Subscales include psychological well-being, anticipation anxiety, vitality, stigma, transmissibility, and vulnerability, each using Likert type scoring from 1 to 5

SF-36: generic tool with eight domains: physical functioning, role physical, bodily pain, general health, vitality, social functioning, role emotional, and mental health, on a range of 1 to 100. Two summary scores summarise physical and mental health-related components of SF-36

Abbreviations

BDI, Beck Depression Inventory; CC, compensated cirrhosis; CHB, chronic hepatitis B; DC, decompensated cirrhosis; HBV, hepatitis B virus; HCC, hepatocellular carcinoma; HRQoL, health-related quality of life; HUV, health utility value; IFN, interferon; NA, nucelos(t)ide analogue; NOI, not of interest; PRISMA, Preferred Reporting Items for Systematic Reviews and Metaanalyses; RCT, randomised clinical trial; SGA, subgroup data availability; SLR, systematic literature review.

• Coauthor conflict of interest: Stuart Kendrick, Lee Evitt: employees of GlaxoSmithKline Research and Development, UK, Ltd and hold GlaxoSmithKline stocks/shares. Rishabh Pandey, Ruchika Mittal, Kajal Thapa: employees of Parexel International. Parexel International received funding from GlaxoSmithKline to conduct this research.

• Editorial support (in the form of writing assistance, including preparation of the draft poster under the direction and guidance of the authors, collating and incorporating authors' comments for each draft, assembling tables and figures, grammatical editing, and referencing) was provided by Bill Wolvey, BSc, of Parexel International and was funded by GlaxoSmithKline.

- This study was funded by GlaxoSmithKline (Study 209774).
- On behalf of all authors, a video recording of this poster was prepared by Vera Gielen, who did not receive any payment for this recording.
- The presenting author, Vera Gielen, declares the following real or perceived conflicts of interest during the last 3 years in relation to this presentation: employee of GlaxoSmithKline Research and Development, UK, Ltd and holds GlaxoSmithKline stocks/shares.

Presented at The International Liver Congress™ 2021 (European Association for the Study of the Liver), Virtual, 23–26 June 2021