

Antacid exposure and immunotherapy outcomes among patients with advanced hepatocellular carcinoma

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Abstract

Background: Antibiotic exposure has been associated with worse outcomes with immune checkpoint inhibitors (ICIs) in cancer patients, likely due to disruption of the gut microbiome. Other commonly prescribed medications, such as proton pump inhibitors (PPIs) and histamine-2-receptor antagonists (H2RAs), are also known to disrupt the microbiome, but data on their association with ICI outcomes are conflicting.

Methods: We conducted a retrospective, multicenter, international cohort study including 314 hepatocellular carcinoma (HCC) patients treated with ICIs from 2017 to 2019 to assess the association between PPI or H2RA exposure (up to 30 days before ICI) and overall survival. Secondary outcomes included overall response rate (ORR) and development of any treatment-related adverse events (AEs).

Results: Baseline PPI/H2RA exposure was not associated with overall survival in univariable (HR 1.01, 95% CI 0.75–1.35) or multivariable analysis (HR 0.98, 95% CI 0.71–1.36). Baseline PPI/H2RA exposure was not associated with either ORR (OR 1.32, 95% CI 0.66–2.65) or AEs (OR 1.07, 95% CI 0.54–2.12) in multivariable analysis.

Conclusions: Our results suggest that exposure to PPI/H2RA prior to ICIs does not adversely affect outcomes in HCC patients.

Keywords: antacid, hepatocellular carcinoma, immunotherapy, proton pump inhibitor

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Introduction

There is growing interest in potential interactions between commonly prescribed medications, such as antibiotics, proton pump inhibitors (PPIs), steroids, and opioids, and cancer immunotherapy.¹ Antibiotic exposure has been repeatedly linked to worse immunotherapy outcomes.^{2–8} Experimental evidence suggests that this association is mediated through the gut microbiome, which is disrupted by antibiotics.^{5,9–12} Several other commonly prescribed medication classes, including antacids such as PPIs and histamine-2-receptor antagonists (H2RAs), are also known

to alter the gut microbiome,^{13–16} and recent reports have linked PPI usage to worse immunotherapy outcomes in lung and bladder cancer.^{7,17,18}

Immune checkpoint inhibitors (ICIs) have become an important component of hepatocellular carcinoma (HCC) management. ICI agents were initially approved by the US Food and Drug Administration in 2017 for second-line treatment of advanced HCC on the basis of promising overall response rates (ORRs) of 20% in early phase studies.^{19,20} More recently, the combination of atezolizumab and bevacizumab became the first

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immunotherapy regimen to improve overall survival (OS) over sorafenib in advanced HCC in a phase III trial and has become standard of care.²¹ However, ORRs remain limited, underlining the need to identify markers and mediators of ICI resistance and response.

Given the increasing use of ICIs in HCC management and the lack of published data addressing the effect of antacids on outcomes in this setting, we conducted this observational study to test the associations of PPI/H2RA exposure before ICI treatment with survival and response.

Methods

Study population

The study population comprised HCC patients treated with immunotherapy between 2017 and 2019 at 11 tertiary referral centers in the United States, Europe, and Asia. An earlier version of the cohort with fewer sites was described previously.²² Included patients had a diagnosis of HCC in accordance with American Association for the study of Liver Disease²³ and European Association for the Study of the Liver²⁴ guidelines, received systemic ICI therapy (either monotherapy or in combination), and had measurable disease according to RECIST 1.1²⁵ criteria at ICI commencement. Baseline antacid data were not available from two US sites, both of which were excluded ($N=102$). An additional eight patients were excluded due to missing baseline antacid data.

Clinical variables were obtained through manual review of electronic medical records by investigators at each site using a standardized data collection form, including specific fields for concomitant medications such as antacids. Antacids were defined as PPIs (omeprazole, lansoprazole, dexlansoprazole, esomeprazole, pantoprazole, rabeprazole) or H2RAs (famotidine, ranitidine, cimetidine, nizatidine). Data were censored on 20 February 2020. Disease staging was conducted by computerized tomography or magnetic resonance imaging prior to ICI initiation and at approximately 9-week intervals during treatment.

This study was conducted in accordance with the Declaration of Helsinki. The Institutional Review Board (IRB) at Imperial College London acted as the central IRB, whose review was accepted by all participating institutions' IRBs (Ref. 17/WA/0161/R18009) (Supplemental Table 1). The

central IRB determined that this research involved no greater than minimal risk and approved a waiver for informed consent.

Study design

This was a retrospective cohort study with a primary outcome of OS, measured from the date of initiation of ICI to the date of death from any cause or last follow-up. Secondary outcomes were ORR, defined as the proportion of patients with a best response of either complete response (CR) or partial response (PR) by RECIST v1.1 criteria; disease control rate (DCR), defined as CR, PR, or stable disease (SD) by RECIST v1.1 criteria; and the development of treatment-related adverse events (AEs) of any grade. AEs were identified from clinical notes in conjunction with laboratory and radiographic evidence. Events were deemed treatment-related based on known side-effect profiles of ICI drugs and the judgment of the treating physician, with study investigators validating the association during chart review. AEs were graded following the National Cancer Institute Common Terminology Criteria for Adverse Events *versus* 5.0. All outcome data were obtained from the electronic medical records of the individual institutions.

The primary predictor in the analysis was baseline antacid (H2RA or PPI) exposure within 30 days prior to ICI initiation. Exposure was defined as an active prescription in the medical record per clinical notes or prescription records. Data were also collected on antacid exposures concurrent with ICI treatment; that is, antacid exposures between the dates of ICI initiation and ICI cessation. Baseline H2RA and PPI exposure were also considered separately as secondary predictors. Additional clinically relevant covariates were age, sex, alpha-fetoprotein (AFP) >400 ng/ml, presence of cirrhosis (clinically or radiologically diagnosed), Barcelona Clinic Liver Cancer (BCLC) stage (A–B *versus* C–D), Child–Turcotte Pugh class, Eastern Cooperative Oncology Group (ECOG) performance status (0 *versus* ≥1), and antibiotic exposure within 30 days prior to ICI.

Statistical analysis

Descriptive statistics were summarized using medians and interquartile ranges (IQRs) for continuous variables, and frequency and proportions for categorical variables. The Wilcoxon rank-sum test and Fisher's exact test were used to compare

Table 1. Baseline characteristics, by baseline antacid exposure.

Variable	No antacid exposure (N=204)	Antacid exposure (N=110)	p value	All patients (N=314)
Age (years)	65.6 (59.1–72.6)	67.9 (58.2–70.7)	0.77	66 (58.7–71.6)
Male	161 (78.9%)	87 (79.1%)	1	248 (79%)
USA	119 (58.3%)	35 (31.8%)	<0.001	154 (49%)
Europe	23 (11.3%)	38 (34.5%)	<0.001	61 (19.4%)
Asia	62 (30.4%)	37 (33.6%)	0.634	99 (31.5%)
PD-1 monotherapy	186 (91.2%)	85 (77.3%)	<0.001	271 (86.3%)
PD-1/CTLA-4 Combination	8 (3.9%)	13 (11.8%)	0.016	21 (6.7%)
PD-1/TKI Combination	10 (4.9%)	12 (10.9%)	0.063	22 (7%)
First-line ICI	93 (45.6%)	44 (40%)	0.404	137 (43.6%)
Second-line ICI	102 (50%)	56 (50.9%)	0.906	158 (50.3%)
Third-line or later	9 (4.4%)	10 (9.1%)	0.135	19 (6.1%)
Prior sorafenib	110 (53.9%)	65 (59.1%)	0.406	175 (55.7%)
Prior local therapy	182 (89.2%)	95 (86.4%)	0.467	277 (88.2%)
Cirrhosis	147 (72.1%)	78 (70.9%)	0.896	225 (71.7%)
Chronic hepatitis B	57 (27.9%)	31 (28.2%)	1	88 (28%)
Chronic hepatitis C	76 (37.3%)	42 (38.2%)	0.903	118 (37.6%)
HBV/HCV co-infection	6 (2.9%)	2 (1.8%)	0.718	8 (2.5%)
Non-viral liver disease	65 (31.9%)	35 (31.8%)	1	100 (31.8%)
Alcoholic liver disease	35 (17.2%)	23 (20.9%)	0.447	58 (18.5%)
Non-alcoholic steatohepatitis	21 (10.3%)	9 (8.2%)	0.688	30 (9.6%)
Other liver disease	7 (3.4%)	4 (3.6%)	1	11 (3.5%)
ECOG \geq 1	88 (43.1%)	57 (51.8%)	0.155	145 (46.2%)
BCLC A	5 (2.5%)	0 (0%)	0.167	5 (1.6%)
BCLC B	51 (25%)	30 (27.3%)	0.686	81 (25.8%)
BCLC C	146 (71.6%)	77 (70%)	0.795	223 (71%)
BCLC D	2 (1%)	3 (2.7%)	0.348	5 (1.6%)
Child–Pugh A	137 (67.2%)	78 (70.9%)	0.441	215 (68.5%)
Child–Pugh B	58 (28.4%)	27 (24.5%)	0.507	85 (27.1%)
Child–Pugh C	7 (3.4%)	3 (2.7%)	1	10 (3.2%)
Portal venous thrombosis	68 (33.3%)	40 (36.4%)	0.535	108 (34.4%)
Extrahepatic metastasis	103 (50.5%)	59 (53.6%)	0.637	162 (51.6%)

(continued)

Table 1. (continued)

Variable	No antacid exposure (N= 204)	Antacid exposure (N= 110)	p value	All patients (N= 314)
No measurable intrahepatic disease	26 (12.7%)	9 (8.2%)	0.261	35 (11.1%)
Multifocal (≥ 3) intrahepatic nodules	115 (56.4%)	59 (53.6%)	0.632	174 (55.4%)
Maximum diameter of largest lesion (cm)	6.15 (3–12.5)	6 (3.6–10.1)	0.663	6 (3.2–11.2)
Alpha-fetoprotein (ng/ml)	183.3 (10.9–2563.5)	223.65 (14.25–4615.5)	0.408	183.3 (12.4–2917.7)
Baseline PPI only [†]	NA	85 (78%)	NA	85 (27.1%)
Baseline H2RA only	NA	17 (15.6%)	NA	17 (5.4%)
Baseline PPI and H2RA exposure	NA	7 (6.4%)	NA	7 (2.2%)
No concurrent antacid [‡]	147 (72.4%)	10 (9.3%)	<0.001	157 (50.5%)
Concurrent PPI only	34 (16.7%)	73 (67.6%)	<0.001	107 (34.4%)
Concurrent H2RA only	9 (4.4%)	19 (17.6%)	<0.001	28 (9%)
Concurrent PPI and H2RA exposure	13 (6.4%)	6 (5.6%)	1	19 (6.1%)
Baseline antibiotic exposure	14 (6.9%)	26 (23.6%)	<0.001	40 (12.7%)
Baseline steroid exposure	4 (2%)	10 (9.1%)	0.006	14 (4.5%)

Data are presented as medians and interquartile ranges or counts and proportions.

[†]One patient had baseline antacid exposure, but the type of antacid was not documented.

[‡]Two patients with baseline antacid exposure had missing data on concurrent antacid exposure; one patient without baseline antacid exposure had missing data on concurrent antacid exposure.

BCLC, Barcelona Clinic Liver Cancer; CTLA-4, cytotoxic T-lymphocyte-associated protein 4; ECOG, Eastern Cooperative Oncology Group; H2RA, histamine-2-receptor antagonist; HBV, hepatitis B virus; HCV, hepatitis C virus; ICI, immune checkpoint inhibitor; PD-1, programmed cell death protein 1; PPI, proton pump inhibitor; TKI, tyrosine kinase inhibitor.

the distribution of continuous and categorical variables between antacid exposures, respectively. AFP values were missing in 17 patients and were imputed as the median value. Median OS was determined using the Kaplan–Meier method, and Kaplan–Meier survival curves were compared using the log-rank test.

Univariable and multivariable Cox proportional hazards models were used to test the association of the predictors with OS. Some covariates for the multivariable model were selected *a priori*, based on the number of events in the dataset and existing literature. These covariates were: age, sex, geographic region, AFP > 400 ng/ml, BCLC (A–B versus C–D), and prior antibiotics, in addition to the primary predictor. Additional covariates significant in the univariable regression

analysis were added to the multivariable model, provided they were not redundant with covariates already being included, for example, BCLC stage incorporates Child–Pugh score and performance status. The proportional hazards assumption was tested and the final model was stratified by geographic region; that is, separate baseline hazard functions were estimated for each geographic region. There was no evidence of multicollinearity with variance inflation factors of all predictors < 5. A summary of each model was presented using hazard ratios (HRs) and 95% confidence intervals (CIs). Subgroup analyses were conducted using a model including the interaction term between antacid exposure and the grouping variable. The HR and 95% CI for antacid exposure in each subgroup was presented, along with the *p* value of the interaction term.

Univariable and multivariable logistic regression models were used to test the association of predictors with the secondary outcomes ORR, DCR, and AE. A similar set of covariates were selected for the multivariable models *a priori*, with additional covariates added based on significance in univariable analysis, absence of collinearity, and available degrees of freedom. A summary of each model was presented using odds ratios (OR) and 95% CIs.

All analyses were conducted in R version 4.0.0 (Vienna, Austria).

Results

Baseline characteristics

The cohort included 314 patients with HCC treated with ICIs between 2017 and 2019 at centers in Asia ($N=99$, 31.5%), the United States ($N=154$, 49%), and Europe ($N=61$, 19.4%). Clinical characteristics of the cohort are reported in Table 1.

The cohort was predominantly male ($N=248$, 79%), with a median age of 66 years (IQR 59–72). At baseline, 225 (71.7%) patients had clinical or radiographic evidence of cirrhosis. The most common underlying causes of liver disease were hepatitis C virus ($N=118$, 37.6%) and hepatitis B virus ($N=88$, 28%), followed by alcoholic liver disease ($N=58$, 18.5%) and non-alcoholic steatohepatitis ($N=30$, 9.6%). Liver function was preserved (Child–Pugh A) in most patients ($N=215$, 68.5%).

Most patients ($N=223$, 71%) met criteria for BCLC stage C HCC at the time of ICI initiation. The majority of patients were treated with anti-PD-1 monotherapy ($N=271$, 86.3%). Half of the patients ($N=158$, 50.3%) had received one prior systemic treatment, whereas 43.6% ($N=137$) were naïve to systemic treatment. Most patients ($N=277$, 88.2%) had also had prior local therapy, most commonly surgical resection ($N=103$, 32.8%).

Treatment outcomes

There were 190 deaths (60.5%) during a median follow-up of 9.2 months (IQR 4.0–16.1). Based on best radiographic response, there were 123 patients with SD (39.2%), 31 with PRs (9.9%), and 21 with CRs (6.7%), resulting in a DCR of 55.7% and an ORR of 16.6%. Median OS in the whole cohort was 12.3 months (95% CI 9.9–15.7). The median duration of ICI treatment was 3.7 months (IQR 1.9–8.6). The most common

reason for treatment discontinuation was progressive disease ($N=178$, 56.7%). Treatment-related AEs developed in 33.8% of patients ($N=106$), with 15.6% of patients ($N=49$) experiencing AEs of grade 2 or higher. The most common AEs involved the liver ($N=41$, 13.1%) and skin ($N=33$, 10.5%).

Antacid exposures

Baseline exposure to antacids, either a PPI or a H2RA, within 30 days prior to ICI treatment was present in 35% of patients ($N=110$) (Table 1). Most baseline exposures were to PPIs ($N=85$, 27.1%), but some were exposed to H2RAs ($N=17$, 5.4%), or both ($N=7$, 2.2%). Among those with baseline antacid exposure, 90.9% ($N=100$) also had antacid exposure concurrent with their ICI treatment.

Additional details on duration and indications for antacid usage were available for 224 patients, 76 of whom had baseline antacid exposure and 117 of whom had concurrent antacid exposure (Supplemental Tables 2 and 3). Most baseline antacids were prescribed for more than 4 weeks (77.6%), with the most common indications being acid reflux or dyspepsia (43.4%) and procedures or other prophylactic reasons (48.7%). Most concurrent antacid exposures originated from prescriptions beginning before ICI treatment (63.6%) and the most common indications were also acid reflux or dyspepsia (37.9%) and procedures or other prophylaxis (37.1%).

There were baseline differences between the patients with or without antacid exposures (Table 1). Antacid-exposed patients were more commonly from European sites (34.5% of exposed *versus* 11.3% of unexposed, $p<0.001$) and less commonly from US sites (31.8% of exposed *versus* 58.3% of unexposed, $p<0.001$). Antacid-exposed patients were less likely to have been treated with anti-PD-1 monotherapy (77.3% of exposed *versus* 91.2% of unexposed, $p<0.001$), but more likely to have been treated with a combination of anti-PD-1 and anti-CTLA-4 agents (11.8% of exposed, 3.9% of unexposed, $p=0.02$). Antacid-exposed patients were also more likely to have been exposed to antibiotics (23.6% of exposed *versus* 6.9% of unexposed, $p<0.001$) or steroids (9.1% of exposed *versus* 2.0% of unexposed, $p=0.006$) in the 30 days prior to ICI initiation.

Table 2. Univariable and multivariable Cox proportional hazards models for overall survival.

Variable	Univariable HR (95% CI)	p value	Multivariable HR* (95% CI)	p value
Age (years)	0.997 (0.99–1.01)	0.642	1.00 (0.99–1.01)	0.958
Male	1.07 (0.75–1.54)	0.701	1.12 (0.77–1.61)	0.560
Geographic region				
Europe versus USA	0.97 (0.67–1.41)	0.866	–	–
Asia versus USA	0.83 (0.60–1.17)	0.292	–	–
Immunotherapy treatment				
PD-1/CTLA-4 versus PD-1	0.78 (0.44–1.37)	0.380	–	–
PD-1/TKI versus PD-1	0.80 (0.45–1.41)	0.443	–	–
Second-line or later	1.15 (0.86–1.54)	0.339	–	–
Cirrhosis	1.06 (0.77–1.45)	0.740	–	–
Liver disease				
HCV versus HBV	0.85 (0.60–1.22)	0.388	–	–
HBV/HCV versus HBV	0.16 (0.02–1.19)	0.074	–	–
Non-viral versus HBV	1.09 (0.76–1.56)	0.650	–	–
ECOG ≥ 1	1.55 (1.15–2.09)	0.004	–	–
BCLC C/D	1.45 (1.04–2.03)	0.030	1.22 (0.85–1.75)	0.276
Child–Pugh B/C	1.94 (1.43–2.62)	<0.001	–	–
Portal venous thrombosis	1.73 (1.29–2.32)	<0.001	1.51 (1.10–2.08)	0.011
Extrahepatic metastasis	1.06 (0.80–1.42)	0.673	–	–
≥ 3 intrahepatic nodules	1.69 (1.26–2.26)	<0.001	1.83 (1.34–2.50)	<0.001
Maximum diameter of largest lesion (cm)	0.998 (0.99–1.00)	0.503	–	–
Alpha-fetoprotein >400 ng/ml	1.39 (1.04–1.85)	0.025	1.31 (0.96–1.79)	0.084
Baseline antacid exposure	1.01 (0.75–1.35)	0.971	0.98 (0.71–1.36)	0.909
Baseline PPI exposure	1.14 (0.84–1.54)	0.409	–	–
Baseline H2RA exposure	0.58 (0.31–1.10)	0.095	–	–
Baseline antibiotic exposure	1.38 (0.92–2.06)	0.122	1.23 (0.78–1.94)	0.370

*Stratified by geographic region.

BCLC, Barcelona Clinic Liver Cancer; CI, confidence interval; CTLA-4, cytotoxic T-lymphocyte-associated protein 4; ECOG, Eastern Cooperative Oncology Group; H2RA, histamine-2-receptor antagonist; HBV, hepatitis B virus; HCV, hepatitis C virus; HR, hazard ratio; PD-1, programmed cell death protein 1; PPI, proton pump inhibitor; TKI, tyrosine kinase inhibitor.

Association of baseline antacid use with immunotherapy outcomes

In univariable analysis, baseline antacid exposure was not associated with OS (HR 1.01, 95% CI 0.75–1.35), nor was baseline PPI exposure (HR 1.14, 95% CI 0.84–1.54) or H2RA exposure

(0.58, 95% CI 0.31–1.10) (Table 2). Factors associated with OS were: ECOG PS > 0 (HR 1.55, 95% CI 1.15–2.09), BCLC C or D (HR 1.45, 95% CI 1.04–2.03), Child–Pugh class B or C (HR 1.94, 95% CI 1.43–2.62), portal venous thrombosis (HR 1.73, 95% CI 1.29–2.32),

Table 3. Univariable and multivariable logistic regression for overall response.

Variable	Univariable OR (95% CI)	<i>p</i> value	Multivariable OR (95% CI)	<i>p</i> value
Age (years)	0.99 [0.97–1.02]	0.459	0.99 [0.97–1.02]	0.534
Male	0.95 [0.45–1.97]	0.883	0.88 [0.41–1.85]	0.730
Geographic region				
Europe <i>versus</i> USA	0.63 [0.26–1.53]	0.306	0.53 [0.20–1.44]	0.213
Asia <i>versus</i> USA	1.28 [0.66–2.48]	0.462	1.16 [0.58–2.35]	0.676
Immunotherapy treatment				
PD-1/CTLA-4 <i>versus</i> PD-1	1.90 [0.70–5.17]	0.209	–	–
PD-1/TKI <i>versus</i> PD-1	0.53 [0.12–2.36]	0.403	–	–
Second line or later	0.83 [0.45–1.51]	0.537	–	–
Cirrhosis	1.74 [0.83–3.67]	0.142	–	–
Liver disease				
HCV <i>versus</i> HBV	0.94 [0.46–1.9]	0.852	–	–
HBV/HCV <i>versus</i> HBV	0.77 [0.08–6.99]	0.812	–	–
Non-viral <i>versus</i> HBV	0.56 [0.25–1.25]	0.159	–	–
ECOG ≥ 1	1.08 [0.59–1.97]	0.799	–	–
BCLC C/D	0.83 [0.43–1.59]	0.567	–	–
Child–Pugh B/C	1.33 [0.70–2.51]	0.383	–	–
Portal venous thrombosis	1.04 [0.55–1.95]	0.905	–	–
Extrahepatic metastasis	0.85 [0.47–1.55]	0.602	–	–
≥ 3 intrahepatic nodules	0.66 [0.36–1.22]	0.185	–	–
Maximum diameter of largest lesion (cm)	1.00 [0.99–1.01]	0.994	–	–
Alpha-fetoprotein >400 ng/ml	0.95 [0.51–1.74]	0.858	–	–
Baseline antacid exposure	1.06 [0.57–1.98]	0.847	1.32 [0.66–2.65]	0.429
Baseline PPI exposure	1.15 [0.60–2.18]	0.676	–	–
Baseline H2RA exposure	1.10 [0.36–3.42]	0.865	–	–
Baseline antibiotic exposure	0.67 [0.25–1.82]	0.435	0.60 [0.20–1.74]	0.343
BCLC, Barcelona Clinic Liver Cancer; CI, confidence interval; CTLA-4, cytotoxic T-lymphocyte-associated protein 4; ECOG, Eastern Cooperative Oncology Group; H2RA, histamine-2-receptor antagonist; HBV, hepatitis B virus; HCV, hepatitis C virus; OR, odds ratio; PD-1, programmed cell death protein 1; PPI, proton pump inhibitor; TKI, tyrosine kinase inhibitor.				

multifocal (≥ 3) intrahepatic nodules (HR 1.69, 95% CI 1.26–2.26), and AFP > 400 ng/ml (HR 1.39, 95% CI 1.04–1.85).

In multivariable analysis, baseline antacid exposure remained unassociated with OS (HR 0.98, 95% CI 0.71–1.36) (Table 2). Significant

independent predictors of survival in this model were multifocal intrahepatic disease (HR 1.83, 95% CI 1.34–2.50) and portal venous thrombosis (HR 1.51, 95% CI 1.10–2.08).

In subgroup analyses, there were trends towards heterogeneous effects of antacid exposure based

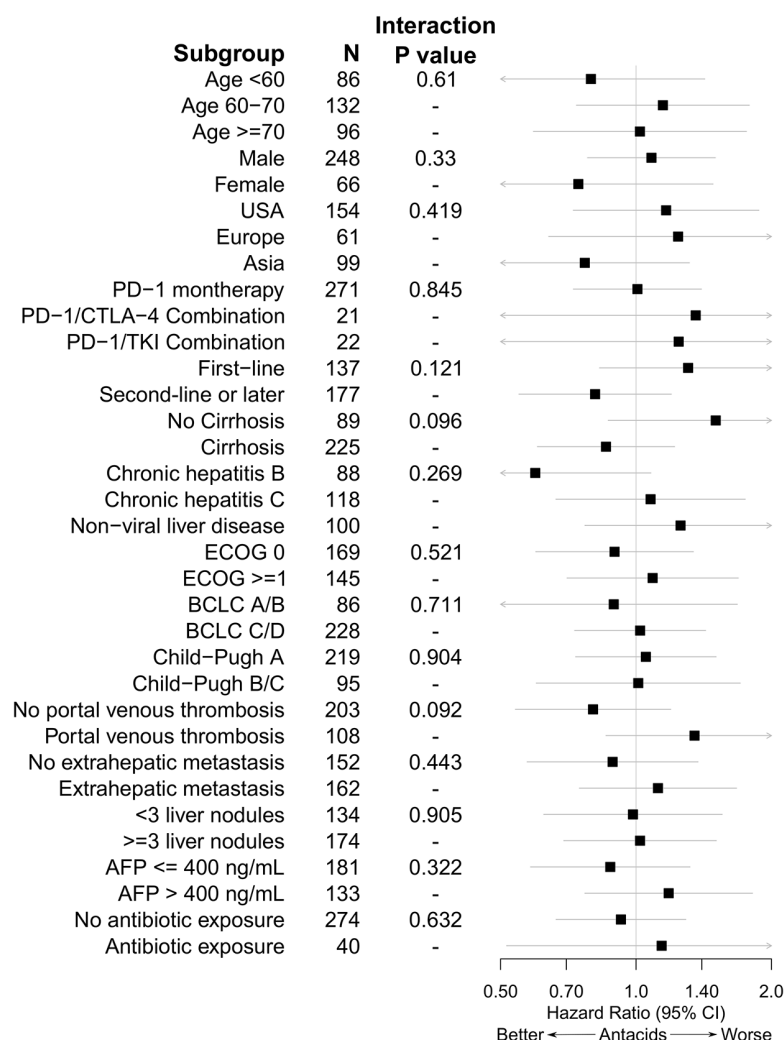


Figure 1. Cox proportional hazards models for overall survival taking into account interactions between baseline antacid exposure and subgroup membership. Hazard ratios and 95% confidence intervals for antacid exposure are presented along with *p* values for the interaction term in each model.

on cirrhosis status (interaction *p* = 0.096) and the presence or absence of portal venous thrombosis (interaction *p* = 0.092) (Figure 1).

To control more strictly for possible confounding by baseline antibiotic exposure, we repeated the OS analysis while excluding the 40 patients who had received baseline antibiotics. Baseline antacid use was not associated with OS in univariable analysis (HR 0.92, 95% CI 0.66–1.29) nor multivariable analysis (HR 0.92, 95% CI 0.63–1.34) after excluding antibiotic-exposed patients.

Baseline antacid exposure was not associated with either ORR (univariable OR 1.06, 95% CI 0.57–1.98;

multivariable OR 1.32, 95% CI 0.66–2.65) or DCR (univariable OR 1.56, 95% CI 0.95–2.57; multivariable OR 1.34, 95% CI 0.75–2.39). There were no significant associations with ORR in either univariable or multivariable analysis (Table 3). DCR was associated with AFP > 400 ng/mL (OR 0.57, 95% CI 0.36–0.9), European sites (OR 3.97, 95% CI 1.91–8.24), and chronic hepatitis C (OR 2.04, 95% CI 1.14–3.66) in univariable analysis. AFP > 400 ng/mL (OR 0.55, 95% CI 0.33–0.92) and European sites (OR 3.94, 95% CI 1.73–9.03) remained independently associated with DCR in multivariable analysis (Table 4).

Baseline antacid exposure was associated with development of any AE (OR 1.85, 95% CI 1.14–3.05) in univariable analysis, but not multivariable analysis (OR 1.07, 95% CI 0.54–2.12) (Table 5). Predictors independently associated with development of any AE were European site (OR 8.33, 95% CI 2.64–26.4) and maximum diameter of the largest nodule in centimeters (OR 1.05, 95% CI 1.03–1.07). Compared with anti-PD-1 monotherapy, combination treatments were associated with AEs in univariable (PD-1/CTLA-4 OR 5.1, 95% CI 1.98–13.1; PD-1/TKI 8.17, 95% CI 2.89–23.1) but not multivariable analysis (PD-1/CTLA-4 OR 1.08, 95% CI 0.29–4.06; PD-1/TKI OR 1.68, 95% CI 0.41–6.89).

Discussion

In this multicenter retrospective cohort study of patients with advanced HCC and treated with ICI therapy, there was no association between OS and exposure to PPIs or H2RAs within 30 days prior to ICI initiation. Furthermore, there were no significant associations between antacid exposure and secondary outcomes such as OR, disease control, or AEs.

Previous observational studies of antacid exposure and immunotherapy outcomes have yielded mixed results. Several single-center retrospective studies focusing on patients with non-small cell lung cancer (NSCLC) or melanoma have failed to find an association between PPI use and immunotherapy outcome.^{26–29} Two of these studies were able to detect significant associations between immunotherapy outcome and antibiotic use, but not PPI exposure.^{27,29} Another study using a combination of retrospective and early phase clinical trial data found significant associations between outcomes and antibiotic exposure, but not PPI exposure.⁵

Table 4. Univariable and multivariable logistic regression for disease control rate.

Variable	Univariable OR (95% CI)	p value	Multivariable OR (95% CI)	p value
Age (years)	1.02 (1.00–1.05)	0.028	1.01 (0.99–1.03)	0.472
Male	0.98 (0.55–1.74)	0.930	1.16 (0.62–2.18)	0.648
Geographic region				
Europe <i>versus</i> USA	3.97 (1.91–8.24)	<0.001	3.94 (1.73–9.03)	0.001
Asia <i>versus</i> USA	1.2 (0.70–2.04)	0.509	1.40 (0.75–2.60)	0.296
Immunotherapy treatment				
PD-1/CTLA-4 <i>versus</i> PD-1	2.02 (0.76–5.37)	0.160	–	–
PD-1/TKI <i>versus</i> PD-1*	–	–	–	–
Second line or later	1.08 (0.68–1.73)	0.741	–	–
Cirrhosis	1.02 (0.60–1.71)	0.955	–	–
Liver disease				
HCV <i>versus</i> HBV	2.04 (1.14–3.66)	0.016	1.88 (0.95–3.71)	0.069
HBV/HCV <i>versus</i> HBV	5.26 (0.59–46.9)	0.138	8.25 (0.84–81.9)	0.070
Non-viral <i>versus</i> HBV	1.55 (0.85–2.81)	0.152	1.08 (0.54–2.16)	0.836
ECOG ≥ 1	0.89 (0.56–1.43)	0.637	–	–
BCLC C/D	0.88 (0.52–1.49)	0.635	0.93 (0.52–1.63)	0.790
Child–Pugh B/C	0.78 (0.47–1.29)	0.325	–	–
Portal venous thrombosis	0.94 (0.57–1.54)	0.802	–	–
Extrahepatic metastasis	1.07 (0.67–1.7)	0.790	–	–
≥ 3 Intrahepatic nodules	0.66 (0.41–1.06)	0.083	–	–
Maximum diameter of largest lesion (cm)	1.01 (0.997–1.02)	0.198	–	–
Alpha-fetoprotein >400 ng/ml	0.57 (0.36–0.92)	0.021	0.55 (0.33–0.92)	0.024
Baseline antacid exposure	1.56 (0.95–2.57)	0.079	1.34 (0.75–2.39)	0.332
Baseline PPI exposure	1.31 (0.79–2.2)	0.299	–	–
Baseline H2RA exposure	2.29 (0.82–6.44)	0.115	–	–
Baseline antibiotic exposure	0.72 (0.37–1.44)	0.355	0.63 (0.29–1.39)	0.252
*All 20 patients treated with PD-1/TKI attained disease control. BCLC, Barcelona Clinic Liver Cancer; CI, confidence interval; CTLA-4, cytotoxic T-lymphocyte-associated protein 4; ECOG, Eastern Cooperative Oncology Group; H2RA, histamine-2-receptor antagonist; HBV, hepatitis B virus; HCV, hepatitis C virus; OR, odds ratio; PD-1, programmed cell death protein 1; PPI, proton pump inhibitor; TKI, tyrosine kinase inhibitor.				

Conversely, *post hoc* analyses of phase II and III clinical trial data have found independent associations between PPI exposure and immunotherapy outcomes. Using data from the Checkmate 069 trial comparing ipilimumab *versus* ipilimumab with

nivolumab in melanoma, Homicsko *et al.*³⁰ found significantly reduced ORR, progression-free survival, and OS among PPI-exposed patients; these associations were maintained in multivariable analysis. Chalabi *et al.*⁷ and Hopkins *et al.*¹⁷ have also

Table 5. Univariable and multivariable logistic regression for any treatment-related adverse event.

Variable	Univariable OR (95% CI)	p value	Multivariable OR (95% CI)	p value
Age (years)	1.02 (0.995–1.04)	0.131	0.999 (0.97–1.03)	0.960
Male	0.66 (0.38–1.16)	0.149	0.69 (0.33–1.47)	0.341
Geographic region				
Europe <i>versus</i> USA	9.58 (4.84–18.9)	<0.001	8.33 (2.64–26.4)	<0.001
Asia <i>versus</i> USA	2.14 (1.21–3.81)	0.009	0.61 (0.24–1.55)	0.295
Immunotherapy treatment				
PD-1/CTLA-4 <i>versus</i> PD-1	5.10 (1.98–13.1)	<0.001	1.08 (0.29–4.06)	0.908
PD-1/TKI <i>versus</i> PD-1	8.17 (2.89–23.1)	<0.001	1.68 (0.41–6.89)	0.469
Second line or later	0.96 (0.60–1.53)	0.848	–	–
Cirrhosis	0.75 (0.45–1.26)	0.277	–	–
Liver disease				
HCV <i>versus</i> HBV	1.90 (1.02–3.51)	0.042	1.69 (0.74–3.86)	0.213
HBV/HCV <i>versus</i> HBV*	–	–	–	–
Non-viral <i>versus</i> HBV	2.30 (1.22–4.32)	0.010	1.37 (0.58–3.25)	0.475
ECOG \geq 1	0.55 (0.34–0.89)	0.014	–	–
BCLC C/D	0.88 (0.52–1.48)	0.634	0.70 (0.37–1.35)	0.288
Child–Pugh B/C	0.65 (0.38–1.09)	0.104	–	–
Portal venous thrombosis	0.92 (0.56–1.51)	0.732	–	–
Extrahepatic metastasis	1.23 (0.77–1.97)	0.384	–	–
\geq 3 intrahepatic nodules	0.81 (0.51–1.30)	0.386	–	–
Maximum diameter of largest lesion (cm)	1.03 (1.01–1.04)	<0.001	1.05 (1.03–1.07)	<0.001
Alpha-fetoprotein >400 ng/ml	1.40 (0.87–2.25)	0.161	1.15 (0.62–2.13)	0.650
Baseline antacid exposure	1.85 (1.14–3.01)	0.013	1.07 (0.54–2.12)	0.850
Baseline PPI exposure	1.84 (1.11–3.05)	0.018	–	–
Baseline H2RA exposure	0.63 (0.24–1.63)	0.338	–	–
Baseline antibiotic exposure	0.71 (0.34–1.48)	0.356	1.17 (0.44–3.1)	0.750
*No patients with HBV/HCV co-infection (N=8) developed treatment-related adverse events. BCLC, Barcelona Clinic Liver Cancer; CI, confidence interval; CTLA-4, cytotoxic T-lymphocyte-associated protein 4; ECOG, Eastern Cooperative Oncology Group; H2RA, histamine-2-receptor antagonist; HBV, hepatitis B virus; HCV, hepatitis C virus; OR, odds ratio; PD-1, programmed cell death protein 1; PPI, proton pump inhibitor; TKI, tyrosine kinase inhibitor.				

recently reported large analyses showing negative associations between PPI exposure and atezolizumab outcomes in lung and bladder cancer, respectively.

Chalabi *et al.*⁷ used data from the OAK and POPLAR trials of atezolizumab *versus* chemotherapy to examine the impact of antibiotic and PPI exposure in 1512 immunotherapy-treated

patients with NSCLC. They found that both antibiotic and PPI exposure were independently associated with reduced OS among atezolizumab-treated patients, but not chemotherapy-treated patients. However, the interaction between PPIs and treatment was not significant.

Hopkins *et al.*¹⁷ examined PPI use as a predictor of outcomes among 1360 bladder cancer patients treated with atezolizumab or chemotherapy in the IMvigor210 and IMvigor211 trials. They similarly found that PPI exposure was an independent predictor of worse OS and PFS in atezolizumab-treated but not chemotherapy-treated patients; in this case, the interaction between PPI and treatment was statistically significant. Overall, these analyses from clinical trials are the most compelling evidence thus far that PPI usage can influence immunotherapy outcome in solid tumors.

Unlike prior studies, our study focuses on patients with HCC. The question of antacid exposure and immunotherapy is particularly relevant in HCC for both clinical and mechanistic reasons. Patients with chronic liver disease and HCC are often prescribed PPIs, whether they may be clinically indicated or not.³¹ Our results suggest that those with indications for PPIs may use them prior to immunotherapy without adversely affecting outcomes.

Mechanistically, the negative findings of our study raise questions of whether the immune microenvironment or microbiome in HCC differs from lung and bladder cancer to the extent that medication exposures have less influence on immunotherapy outcomes. The liver has been recognized as a site of myeloid-derived suppressor cell accumulation,³² and liver metastases have been associated with worse immunotherapy outcomes in lung cancer.^{33,34} Both cirrhosis and HCC are associated with altered gut microbiota and the microbiome has been linked with progression of cirrhosis to HCC.^{35–38}

It is possible that the baseline dysbiosis of cirrhotic HCC patients limits the impact of medication-induced perturbations to the microbiome. Intriguingly, we found a suggestive but non-significant interaction ($p=0.096$) between antacid exposure and cirrhosis status with regards to OS. One recent study of eight HCC patients reported microbial differences between immunotherapy-responders and non-responders but excluded patients with advanced liver disease.³⁹ More microbiome data from immunotherapy-treated

patients are needed to understand the interactions between medication exposures, the microbiome, and immunotherapy outcomes in HCC.

Another mechanistic link between PPIs and immunotherapy outcomes may be through the pH of the tumor microenvironment. Pre-clinical studies have linked the acidity of the tumor microenvironment with T-cell anergy and immune escape.^{40,41} Experiments in mice have demonstrated that systemic PPIs can increase intratumoral pH,⁴² which can in turn improve response to immunotherapy.⁴¹ The net effect of PPIs on immunotherapy outcomes may depend on the balance between detrimental impacts on the microbiome *versus* beneficial changes in the tumor microenvironment.

The external validity of these findings is bolstered by the size and geographic diversity of this cohort. Unlike clinical trials, which have been mostly limited to patients with preserved liver function, this observational cohort reflects a real-world clinical population. In-depth characterization of each patient's tumor also allowed us to control for potential confounders such as performance status, liver function, disease stage, and antibiotic exposure.

This study has several limitations. Although our cohort size compares favorably with other retrospective studies examining this question,^{26–29} power may have been limited to detect significant effects. Our definition of antacid exposure did not incorporate duration or dose of antacid treatment, and medication adherence could not be assessed from the medical record alone. However, this definition and its limitations is consistent the definition of exposure used in prior studies, including the large studies of Chalabi *et al.*⁷ and Hopkins *et al.*¹⁷ Although we captured antibiotic and steroid use, we did not record use of other common medications which may also influence the microbiome or the immune system. Ultimately, this retrospective analysis was exploratory and the findings are hypothesis-generating. Untangling causal relationships between medications, the microbiome, and cancer immunotherapy will require larger retrospective clinical datasets, in-depth mechanistic studies, and randomized clinical trials, where feasible.

In conclusion, these observational data suggest that PPIs and H2RAs do not adversely impact ICI therapy outcomes in patients with advanced

HCC. These findings require validation in prospective cohorts and provide motivation for mechanistic studies to dissect the interactions of medication exposures and the microbiome in HCC as compared to other solid tumors.

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Author contributions

TJ and CA conceived and designed the study. SD, NP, TP, NN, PCL, CJL, HH, NN, SP, PF, MN, UK, AS, YHH, MK, LR, TM, and DJP made substantial contributions to the acquisition of data. TJ, UO, and JYL analyzed the data. TJ and CA interpreted the data. TJ drafted the manuscript. All authors reviewed the work for important intellectual content.

Conflict of interest statement

DJP received lecture fees from ViiV Healthcare and Bayer Healthcare and travel expenses from BMS and Bayer Healthcare; consulting fees for Mina Therapeutics, Eisai, Roche, and Astra Zeneca; received research funding (to institution) from MSD and BMS.

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Ethics approval and consent to participate

This study was conducted in accordance with the Declaration of Helsinki and was approved by the Institutional Review Board in each participating institution.

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Supplemental material

Supplemental material for this article is available online.

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