REVIEW

Physical Activity and the Risk of Liver Cancer: A Systematic Review and Meta-Analysis of Prospective Studies and a Bias Analysis

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Abstract

Background: Physical inactivity is an established risk factor for several cancers of the digestive system and female reproductive organs, but the evidence for liver cancers is less conclusive.

Methods: The aim of this study was to synthesize prospective observational studies on the association of physical activity and liver cancer risk by means of a systematic review and meta-analysis. We searched Medline, Embase, and Scopus from inception to January 2019 for prospective studies investigating the association of physical activity and liver cancer risk. We calculated mean hazard ratios (HRs) and 95% confidence intervals (CIs) using a random-effects model. We quantified the extent to which an unmeasured confounder or an unaccounted selection variable could shift the mean hazard ratio to the null.

Results: Fourteen prospective studies, including 6,440 liver cancers, were included in the systematic review and meta-analysis. The mean hazard ratio for high compared with low physical activity was 0.75 (95% CI = 0.63 to 0.89; 95% prediction interval = 0.52 to 1.07; $I^2 = 64.2\%$). We estimated that 67.6% (95% CI = 56.6% to 78.5%) of all true effect estimates would have a hazard ratio less than 0.8. Bias analysis suggested that an unobserved confounder would have to be associated with a 1.99-fold increase in the risk of physical activity or liver cancer to explain away the observed mean hazard ratio. An unaccounted for selection variable would have to be related to exposure and endpoint with a relative risk of 1.58 to explain away the mean hazard ratio.

Conclusions: Physical activity is inversely related to the risk of liver cancer. Further studies with objectively measured physical activity and quasi-experimental designs addressing confounding are needed.

Worldwide, liver cancer was the fourth leading cause of cancer death in 2015 after lung, colorectal, and stomach cancer (1). The most common type of primary liver cancer is hepatocellular carcinoma. The burden of liver cancer varies markedly by sex and geographic region (2). Major risk factors include infections (hepatitis B virus, hepatitis C virus), aflatoxins, metabolic factors (diabetes, excess body fatness), and behavioral factors (alcohol consumption, tobacco) (3,4). In the past several decades, we have witnessed an increase in the incidence of liver cancers in Western countries (1,5). Obesity and physical inactivity have emerged as risk factors for a number of cancers of the digestive system and female reproductive organs (6,7). Based on current knowledge, factors related to energy balance, encompassed by

physical activity and obesogenic effects of diet, are estimated to account for 10%–15% of the overall cancer burden (8). A previous systematic review and meta-analysis found an inverse association between total physical activity and liver cancer risk (9). However, that systematic review and meta-analysis (9) was not comprehensive, because it did not include a recent multinational cohort study (10).

Nonetheless, as stated by the World Cancer Research Fund International (7) and an umbrella review (6), the existing evidence is inconclusive to support a beneficial effect of physical activity for lowering the risk of liver cancer. Moreover, meta-analysis of observational studies might exaggerate bias because of confounding by increasing statistical power and reducing

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random variability (11–15). In particular, few existing prospective studies on the association between physical activity and liver cancer risk were adjusted for all relevant confounding factors, including hepatitis infection (10,16). In addition, observational studies can be subject to selection bias, whereby the parameter being estimated differs from a causal effect in the total population because of differential restriction of the study population (17). Therefore, bias analysis is particularly important in meta-analyses of observational studies, where a central role is to assess the current quality of evidence. We conducted a systematic review of the literature and summarized eligible prospective studies on the association of physical activity and liver cancer risk and performed bias analyses to examine the robustness to unobserved confounding and selection bias.

Methods

Literature Search

We conducted this systematic review and meta-analysis following the Meta-analysis of Observational Studies in Epidemiology (18) guideline. Two investigators (SEB, SS) searched Medline, Embase, and Scopus from database inception to January 23, 2019, for studies that investigated the association between physical activity and the risk of liver cancer. We tailored search strategies to each database and used controlled vocabulary and search filters where available, or Boolean search methods and free-text terms. Full search terms are provided in the [Supplementary Boxes 1–3 \(available online\)](#). We restricted the search to human studies. No date or language restrictions were used. References of relevant publications were handsearched for additional articles.

Study Selection

Two authors (SEB, SS) independently performed the search, read the full texts of all identified articles to determine whether each study met the eligibility criteria, and resolved discrepancies by discussion. Included studies fulfilled the following criteria: 1) prospective design (ie, cohort, nested case control, case cohort, follow-up of randomized controlled trial); 2) physical activity used as an exposure variable; 3) liver cancer incidence or liver cancer mortality as outcome; 4) performance among general (healthy) populations older than 18 years without prevalent cancers at baseline; and 5) provision of hazard ratios (HRs) or relative risks (RRs) estimates with 95% confidence intervals (CIs). Studies were excluded if they measured muscle strength, physical fitness or cardiorespiratory fitness but not physical activity. If multiple studies reported on the same dataset, we included the one with the largest number of cases or longer duration of follow-up, more detailed report of physical activity, and better control of confounding variables. We considered the following to represent a minimal set of adjustment variables: age, sex, educational attainment, smoking, alcohol consumption, and coffee intake (10,16,19).

Data Extraction and Evaluation of Study Quality

Data were extracted using a standardized data extraction form. Two authors (SB and SS) independently checked the data. For each of the eligible studies, the following information was extracted: first author's name, publication year, study name, population, country, age at baseline, sex, number of participants and cases, cancer site, type and intensity of physical activity,

comparisons made, hazard ratios or relative risks and 95% confidence intervals from multivariable adjusted models, and confounding factors (Table 1). Because liver cancer is relatively rare by the end of follow-up, hazard ratios were treated as estimators of relative risks (26–28). The hazard ratio and 95% confidence interval were inverted for one study (25) that used the most active group as the reference category. A within-study inverse-variance weighted fixed-effect model was performed to obtain an overall estimate when hazard ratios were separately reported for men and women (23,25).

We used the Newcastle-Ottawa Scale (29) to assess the quality of included studies consisting of representativeness of the cohort, whether the nonexposed participants were drawn from the same population as the exposed, ascertainment of the exposure, whether the outcome of interest was absent at the study onset, comparability (conditional exchangeability) of the exposed and unexposed (that is, if adjustment for the minimal adjustment set, as defined above, was made), ascertainment of the outcome, whether the length of the follow-up was long enough (at least 5 years) for the outcome to occur, and the completeness of the follow-up (loss to follow-up <20%). A minimum score of 2 can be awarded for comparability and a maximum of 1 can be given for each of the remaining items. A study can achieve a maximum possible quality score of 9.

Statistical Analysis

We used the Hartung-Knapp-Sidik-Jonkman random-effects meta-analysis approach [whose advantages were demonstrated previously (30,31)] to combined study-specific log hazard ratio, with the empirical Bayes estimator for the between-study variance (τ^2) (30), for estimation of a mean hazard ratio, with corresponding 95% confidence intervals and 95% prediction intervals (32). The 95% confidence interval from a random-effects model contains highly probable values for the mean hazard ratio (32,33). The 95% prediction interval estimates where the true hazard ratio is to be expected for 95% of studies in similar settings in the future (32,33). We reported the percentage of total variation due to heterogeneity (I^2); values greater than 50% to 75% are considered moderate to large (33). Cochran Q statistic was used to test for between-study heterogeneity. Heterogeneity was further examined by estimating the proportion of studies in the population with evidence for a clinically meaningful effect (using thresholds for a mean HR of ≤ 0.7 , ≤ 0.8 , ≤ 0.9) and the estimated proportion of studies with harmful effects (ie, HR > 1) (14,32). Random-effects meta-analysis was additionally stratified by study quality [Newcastle-Ottawa Scale (29) < 6 vs ≥ 6 points], endpoint (incident liver cancer vs liver cancer mortality), type of physical activity (total physical activity, leisure-time physical activity, vigorous physical activity, sports participation), sex (as some studies included only male or female participants), and geographic region (Asia, Europe, United States). Because there were few studies in these strata, we did not perform meta-regression to identify sources of heterogeneity. All statistical tests were two-sided, and P values less than 0.05 were considered as statistically significant.

Bias Analysis

We performed bias analysis to examine possible effects due to inclusion of small studies, selective publication of positive findings, and sensitivity to unobserved confounding and selection bias. We evaluated publication bias and small study effects

Table 1. Characteristics of prospective studies of physical activity and liver cancer

Author, year (reference)	Study, age range at baseline, follow-up	Study population (country)	Outcome, No. cases	Definition of physical activity	Comparison	Study quality score*	HR or RR (95% CI)	Controlled covariates	Included in primary or secondary analysis
Baumeister et al., 2019 (10)	European Prospective Investigation into Cancer and Nutrition Study 438 948 men and women 25–70 y Median follow-up: 14.9 y	General population (Denmark, France, Germany, Greece, Italy, the Netherlands, Norway, Spain, Sweden, and the United Kingdom)	Incident liver cancer, 275	Total physical activity	Active vs inactive	9	0.55 (0.38 to 0.80)	Sex, age, education, smoking, baseline alcohol, lifetime alcohol, coffee intake	Primary analysis
Moore et al., 2016 (20)	National Institutes of Health–AARP Diet and Health Study 507 826 men and women 50–71 y Mean follow-up: 11.0 y	American Association of Retired Persons members (United States)	Incident liver cancer, 619	Moderate and vigorous leisure-time physical activity	90th to 10th percentile of study-specific physical activity measure	9	0.52 (0.42 to 0.65)	Age, sex, smoking, alcohol, education, race	Primary analysis
	Breast Cancer Detection Demonstration Project 37 228 women 39–93 y Median follow-up: 9 y	Participants of screening program (United States)	Incident liver cancer, 28	Moderate and vigorous leisure-time physical activity	90th to 10th percentile of study-specific physical activity measure	6	1.58 (0.55 to 4.57)	Age, sex, smoking, alcohol, education, race	Primary analysis
	Cohort of Swedish men 40 919 men 45–80 y Median follow-up: 10 y	General population (Sweden)	Incident liver cancer, 51	Moderate and vigorous leisure-time physical activity	90th to 10th percentile of study-specific physical activity measure	8	0.42 (0.19 to 0.94)	Age, sex, smoking, alcohol, education, race	Primary analysis
	Cancer Prevention Study II 154 425 men and women 50–74 y Median follow-up: 14.2 y	General population (United States)	Incident liver cancer, 181	Moderate and vigorous leisure-time physical activity	90th to 10th percentile of study-specific physical activity measure	9	1.03 (0.68 to 1.54)	Age, sex, smoking, alcohol, education, race	Primary analysis

(continued)

Table 1. (continued)

Author, year (reference)	Study, age range at baseline, follow-up	Study population (country)	Outcome, No. cases	Definition of physical activity	Comparison	Study quality score*	HR or RR (95% CI)	Controlled covariates	Included in primary or secondary analysis
Arem et al., 2014 (21)	Iowa Women's Health Study 37 584 women 52–70 y Median follow-up: 20 y	General population (United States)	Incident liver cancer, 50	Moderate and vigorous leisure-time physical activity	90th to 10th percentile of study-specific physical activity measure	6	1.19 (0.54 to 2.63)	Age, sex, smoking, alcohol, education, race	Primary analysis
	Physician's Health Study I and II 27 890 men 40–87 y Median follow-up: 20 y	Health professionals (United States)	Incident liver cancer, 43	Moderate and vigorous leisure-time physical activity	90th to 10th percentile of study-specific physical activity measure	8	0.76 (0.33 to 1.78)	Age, sex, smoking, alcohol, education, race	Primary analysis
	Prostate, Lung, Colorectal and Ovarian Cancer Screening Trial 60 200 men and women 52–77 y Mean follow-up: 10 y	Patients at medical centers (United States)	Incident liver cancer, 71	Moderate and vigorous leisure-time physical activity	90th to 10th percentile of study-specific physical activity measure	5	0.63 (0.33 to 1.22)	Age, sex, smoking, alcohol, education, race	Primary analysis
	Swedish Mammography Cohort, 33 006 women 48–83 y Mean follow-up: 10 y	General population (Sweden)	Incident liver cancer, 21	Moderate and vigorous leisure-time physical activity	90th to 10th percentile of study-specific physical activity measure	8	2.45 (0.69 to 8.73)	Age, sex, smoking, alcohol, education, race	Primary analysis
	Women's Health Study 39 414 women 45–74 y Median follow-up: 10.1 y	Health professionals (United States)	Incident liver cancer, 21	Moderate and vigorous leisure-time physical activity	90th to 10th percentile of study-specific physical activity measure	8	0.64 (0.19 to 2.13)	Age, sex, smoking, alcohol, education, race	Primary analysis
	National Institutes of Health–AARP Diet and Health Study 293 511 men and women 50–71 y Median follow-up: 12.1 y	American Association of Retired Persons members (United States)	Liver cancer mortality, 397	Moderate and vigorous leisure-time physical activity	>7 h per wk vs never or rare	9	0.71 (0.52 to 0.98)	Sex, age, body mass index, education, race, alcohol, healthy eating score, calories, marriage status, diabetes, smoking	Subgroup analysis by endpoint

(continued)

Table 1. (continued)

Author, year (reference)	Study, age range at baseline, follow-up	Study population (country)	Outcome, No. cases	Definition of physical activity	Comparison	Study quality score*	HR or RR (95% CI)	Controlled covariates	Included in primary or secondary analysis
Wen et al., 2011 (22)	MJ Health Management Institution 416 175 men and women 20–79 y Mean follow-up: 8.4 y	Health checkpoint population (Taiwan)	Incident liver cancer, 1,676	Leisure-time physical activity	≥16.5 vs ≤3.75 metabolic equivalents hours per week	5	0.81 (0.71 to 0.92)	Age, sex, smoking, alcohol, fasting blood glucose, systolic blood pressure, total cholesterol, body mass index, diabetes	Primary analysis
Inoue et al., 2008 (23)	Japanese Public Health Center-based Prospective Study 79 771 men and women 45–74 y Mean follow-up: 7.5 y	Population-based and participants health checkpoint programs (Japan)	Incident liver cancer, 263	Total physical activity	Highest vs lowest quartile of metabolic equivalents per day	6	0.60 (0.41 to 0.89)	Sex, age, area, energy intake, diabetes, smoking, alcohol, body mass index	Primary analysis
Yun et al., 2008 (24)	National Health Insurance Corporation 444 963 men >40 y Mean follow-up: 6 y	Health checkpoint population (Korea)	Incident liver cancer, 2,676	Vigorous physical activity	≥5 times per wk at ≥30 min per session vs ≤4 times per wk at <30 min per session	6	0.88 (0.81 to 0.95)	Age, sex, dietary preference, alcohol, body mass index, employment, fasting blood glucose	Primary analysis
Suzuki 2007 (25)	Japanese Collaborative Cohort Study for Evaluation of Cancer Risk 69 752 men and women 40–79 y Median follow-up: 12 y	Health checkpoint population (Japan)	Liver cancer mortality, 465	Sports participation	>3 vs <1 h per wk	5	0.81 (0.71 to 0.92)	Age, sex, area	Primary analysis

*Study quality score: Newcastle-Ottawa Scale. CI = confidence interval; HR = hazard ratio; RR = relative risk.

(funnel plot asymmetry) using the regression-based tests proposed by Egger and Debray (34), the trim-and-fill method (35), and the Copas selection model (35). We conducted adjustment for small study bias using the Rücker regression-based shrinkage estimator (35). Random-effects meta-analysis of observational studies can produce biased estimates of pooled effect sizes if the synthesized individual studies are subject to unmeasured confounding or selection bias (11–13). We used sensitivity analyses for unmeasured confounding in a random-effects meta-analysis. Specially, we quantified the magnitude of unmeasured confounding capable of reducing the mean hazard ratio to the null (15). This approach to sensitivity analysis is a meta-analytic extension of the E-Value, a popular metric that quantifies the minimum strength of association that an unmeasured confounder would need to have with the exposure and the outcome on the relative risk scale to fully account for an observed exposure-outcome association, above and beyond the measured covariates (26). Given the heterogeneity of the random-effects model estimate, we additionally reported the unobserved confounding strength required to nullify 10% to 50% of studies with true hazard ratio larger than a meaningful scientific threshold (ie, $HR < 0.9$) (15). We used an upper bounding factor approach to selection bias to determine the strength of selection on the relative risk scale that would be necessary to explain away the mean hazard ratio (17). The statistical software R [version 3.5.1, Foundation for Statistical Computing, Vienna, Austria; packages E-Value (15), meta, metafor, meta-misc (34), and metasens (35)] was used.

Results

Systematic Review

Our literature search identified a total of 400 publications (Figure 1). After removal of duplicate publications, 172 studies remained for title and abstract screening, of which 22 were potentially relevant for full-text review. Among the 22 reviewed publications, studies (with possible overlap) were excluded from the primary meta-analysis on physical activity and liver cancer if the exposure variable was not physical activity ($n = 6$) (36–41), or if the outcome was not liver cancer ($n = 1$) (19), or the effect estimates for physical activity were not shown ($n = 4$) (36,37,41,42). Additionally, studies were excluded if the study design was not prospective ($n = 5$) (43–47) or if the article had a shorter follow-up or fewer events than another publication from the same study ($n = 5$) (16,41,48–50).

This left a total of 14 eligible studies from six publications (10,20,22–25). Taken together, those studies included 2.39 million individuals and 6,440 liver cancer events or deaths. Characteristics of included studies are shown in Table 1. Self-reported physical activity, measured at baseline, served as exposure variable in all studies. The mean follow-up time was 11.6 years (range = 6 to 20). The age of included participants ranged from 20 to 93 years (median = 45) at baseline examination. Among the 14 included studies, eight were conducted in the United States, two in Europe, and four in Asian countries. The quality of the studies was moderate to high for 11 studies and low for three studies.

Meta-Analysis for the Association Between Physical Activity and Liver Cancer Risk

The random-effects meta-analysis showed that physical activity is inversely associated with liver cancer risk. The mean

hazard ratio for liver cancer risk, comparing high and low levels of physical activity, was 0.75 (95% CI = 0.63 to 0.89) (Figure 2). There was moderate heterogeneity between studies (Cochran Q $P = .001$; $I^2 = 64.2\%$). Based on the point and τ^2 estimates from the random-effects model, we estimated that 30.7% (95% CI = 19.0% to 42.4%), 67.6% (95% CI = 56.6% to 78.5%), and 90.3% (95% CI = 76.0% to 99.1%) of all true effect estimates will have mean hazard ratios below 0.7, 0.8, and 0.9, respectively. We further estimated that 2% (95% CI = 0.13% to 8.3%) of all studies have hazard ratios larger than 1.0. The 95% prediction interval for the mean hazard ratio of 0.75 was 0.52 to 1.07; this additionally suggests that, with high probability, a new hazard ratio drawn from the population will be less than 1. When the meta-analysis was stratified by study quality, endpoint (incidence, mortality), type of physical activity, sex, and region, we found difference with regard to type of physical activity (Cochran Q $P < .001$) (Table 2). Studies using total physical activity as the exposure variable revealed the largest hazard ratio, and a study that used the participation in sports as the exposure revealed the smallest hazard ratio.

Bias Analysis

There was little evidence for publication bias using analyses of funnel plot asymmetry, as indicated by Egger test ($P = .46$) and Debray test ($P = .23$) (Supplementary Table 1 and Figure 1, available online). We used the trim-and-fill method, Copas selection model, Rücker's shrinkage procedure, and the leave-one-out procedure to examine small study bias (Supplementary Table 2 and Figure 2, available online). The trim-and-fill method mirrored one of the cohorts included in a pooled analysis of nine US and European cohorts (20). The mean hazard ratios and 95% confidence intervals from these bias analyses were in agreement with our primary finding. Of note, when the NIH-AARP Diet and Health Study was excluded in leave-one-out analysis, heterogeneity was substantially reduced ($I^2 = 37.2\%$, $\tau^2 = 0.006$), and the mean hazard ratio was 0.81 (95% CI = 0.71 to 0.92; prediction interval = 0.66 to 0.98).

In the bias analysis for unobserved confounding, for an unmeasured confounder to explain the mean hazard ratio of 0.75, an unobserved confounder would have to be associated with a 1.99-fold increase in the risk of physical activity or liver cancer, above and beyond the measured confounders. For an unmeasured confounder to bring the upper confidence limit of 0.89 for this mean hazard ratio above 1.0, the unmeasured confounder would have to be related to physical activity and liver cancer risk with a relative risk of 1.49 above and beyond the measured confounders. Unobserved confounder strengths with relative risk of 2.22, 2.03, 1.90, 1.78, and 1.68, respectively, would be necessary to shift 10%, 20%, 30%, 40%, and 50% of all true studies with hazard ratio less than 0.9 to 1.0 (Supplementary Figure 3, available online). By calculating the upper bounding factor $\sqrt{1/0.75} + \sqrt{1/0.75 - 1/0.75} = 1.58$, we come to the conclusion that unaccounted for selection variables would have to be related to physical activity and liver cancer with a relative risk of 1.58 to produce an observed hazard ratio of 0.75 if the true hazard ratio were equal to 1.0.

Discussion

The results of this meta-analysis, including 6,440 liver cancer cases from 14 prospective observational studies, indicated that high vs low physical activity reduces the risk of liver cancer by

Figure 1. Flowchart of systematic review and meta-analysis selection.

25%. There was moderate heterogeneity between studies, but we estimated that 90% of all true hazard ratios would be smaller than 0.9 and 68% of all true hazard ratio would be smaller than 0.8.

To our knowledge, one previous systematic review and meta-analysis (9) has summarized available prospective cohorts on the association of physical activity and liver cancer risk. This meta-analysis produced a mean relative risk of 0.75 (95% CI=0.66 to 0.86; $I^2=0\%$) for the association of high vs low total physical activity and liver cancer risk, although differences in the evaluation of individual studies exist. By comparison, a pooled analysis from 10 cohorts reported a mean hazard ratio of 0.73 (95% CI=0.55 to 0.98; $I^2=55\%$) (20). Of note, the meta-analysis (9) and the pooled analysis (20) did not include the most recent pan-European cohort study on physical activity and liver cancer risk (10).

A concern of our meta-analysis includes the potential for uncontrolled confounding. Although most estimates in the individual studies were adjusted for relevant confounders, meta-analysis of observational associations can be biased because of residual confounding. Most notably, chronic infections with hepatitis virus B and C are strong liver cancer risk factors and account for more than 70% of liver cancers worldwide (3,4). Very few data from prospective settings are available from populations with high hepatocellular carcinoma incidence and a greater contribution of chronic hepatitis infection (10). Accordingly, because of the potential for residual confounding by unmeasured confounders in several of the individual studies, we performed sensitivity analysis to quantify the extent to which associations across studies were robust to unmeasured confounding. We found that a modestly strong unobserved confounder, with a 1.99-fold increase of the risk of the exposure or

Figure 2. Forest plot from random-effects meta-analysis of prospective studies on the risk of liver cancer comparing high and low levels of physical activity. Liver cancer was defined as HCC (hepatocellular carcinoma, C22.0) and/or IHBC (intrahepatic bile duct cancer, C22.1). Study-specific hazard ratios (HR) are represented by black diamonds (with their 95% confidence interval [CI] as error bars). hazard ratios were combined using a Hartung-Knapp-Sidik-Jonkman random-effects model, yielding a mean hazard ratio and its 95% confidence interval and 95% prediction interval. The dotted line represents the mean hazard ratio. Two-sided P value for between-study heterogeneity based on Cochran Q statistic. AARP = National Institutes of Health-AARP Diet and Health Study; BCDDP = Breast Cancer Detection Demonstration Project; COSM = Cohort of Swedish Men; CPSII = Cancer Prevention Study II; EPIC = European Prospective Investigation into Cancer and Nutrition Study; IWHS = Iowa Women's Health Study; JCCSECR = Japanese Collaborative Cohort Study for Evaluation of Cancer Risk; JPHCPS = Japanese Public Health Center-based Prospective Study; MJHMI = MJ Health Management Institution; NHIC = National Health Insurance Corporation; PHS = Physician's Health Study I and II; PLCO = Prostate, Lung, Colorectal and Ovarian Cancer Screening Trial; SMC = Swedish Mammography Cohort; WHS = Women's Health Study.

Table 2. Subgroup meta-analyses of physical activity and liver cancer*

Subgroup	No. of studies	HR (95% CI)†	I ² , %	τ ²	P _{h1} ‡
Study quality					.42
NOS ≥6	11	0.73 (0.57 to 0.94)	72.0	0.032	
NOS <6	3	0.80 (0.70 to 0.92)	0	0	
Endpoint					0.45
Incident liver cancer	13	0.74 (0.61 to 0.90)	67.0	0.023	
Liver cancer mortality	3	0.80 (0.68; 0.94)	0	0	
Type of physical activity					<.001
Total physical activity	2	0.57 (0.44 to 0.75)	0	0	
Leisure-time physical activity	10	0.75 (0.67 to 0.83)	61.5	0.0364	
Vigorous physical activity	1	0.88 (0.81, 0.95)	0	0	
Sports participation	1	0.81 (0.61 to 1.07)	0	0	
Sex					.26
Men§	5	0.86 (0.74 to 1.00)	29.0	<0.001	
Women	6	0.87 (0.50 to 1.52)	28.6	0.031	
Men and women¶	5	0.68 (0.48 to 0.97)	76.2	0.032	
Geographic region					.56
Asia	4	0.85 (0.80 to 0.91)	31.1	<0.001	
Europe	3	0.62 (0.43 to 0.88)	79.6	0.096	
United States	7	0.64 (0.54 to 0.75)	53.5	0.052	

*Liver cancer was defined as HCC (hepatocellular carcinoma, C22.0) and/or IHBC (intrahepatic bile duct cancer, C22.1). CI = confidence interval; HR = hazard ratio; NOS = Newcastle-Ottawa Scale; I² = percentage of total variance explained by τ²; τ² = between-study variance.

†Hazard ratios were combined using a Hartung-Knapp-Sidik-Jonkman random-effects model.

‡Two-sided P value was calculated using Q test for subgroup differences.

§Primary study included only men or reported HR for men.

||Primary study included only women or reported HR for women.

¶Primary study included men and women and reported one HR for men and women.

outcome, would suffice to explain away the mean hazard ratio of 0.75. A confounder strength of a 1.49-fold increased risk of exposure and outcome would suffice to render the estimate statistically nonsignificant. Individuals with hepatitis B virus and hepatitis C virus infections show a 15- to 20-fold increase in the risk of hepatocellular carcinoma compared with seronegative individuals (51), indicating the potential for unobserved confounding by hepatitis virus infection to account for the observed association between physical activity and liver cancer risk. Selection bias with a 1.58-fold increased risk for unaccounted for selection variables would be sufficient to shift the observed hazard ratio to the null. Thus, a potential role for chronic hepatitis infection and other unobserved confounders and selection factors (such as lack of access to medical care or adverse socioeconomic conditions associated with cohort participation) remains to bias the observed association of physical activity and liver cancer risk.

Limitations of our meta-analysis include that individual studies relied on self-reported measures of physical activity, which may introduce measurement error and potentially biased effect estimates. Self-reported measures of activity may be affected by mood states, social desirability, or cognitive biases that also may affect liver cancer liability. However, a majority of the studies used physical activity questionnaires that underwent rigorous validation against objective measures of energy expenditure and cardiorespiratory fitness. A further concern is that assessments of type, duration, and intensity of physical activity differed somewhat by study. In addition, not all studies assessed moderate and vigorous physical activities separately. Because the majority of the included studies reported risk estimates for high vs low levels of physical activity, we could not examine intensities of physical activity required to reduce liver cancer risk using dose-response meta-analysis. Finally, only one included study (10) reported effect estimates on types of hepatobiliary cancers, which precluded analyses by cancer subtype.

A large number of experimental and observational studies suggest that the most important mechanism by which physical activity affects gastrointestinal cancer occurrence is lowering body weight (visceral fatness in particular) (52,53). The exact biologic mechanisms underlying the association between adiposity and liver cancer are not fully established, but main mechanisms include dysregulation of adipose tissue-derived inflammation (immune-related and other), alterations in insulin signaling, growth hormone pathways, and sex hormone metabolism (53–55).

In conclusion, this quantitative synthesis of all available prospective studies suggests that physical activity reduces the risk of liver cancer. This study leveraged quantitative bias analysis to test how much unobserved confounding and selection bias would suffice to bias effect estimates. Moderately strong unobserved confounding and selection bias suffice to explain away the observed inverse association between physical activity and liver cancer risk. Studies with more detailed and objectively measured physical activity assessed at multiple time points throughout the life course, and adjustment for hepatitis infection, are needed to provide additional evidence on this association. Recent developments in quasi-experimental designs (56,57) offer the opportunity to further reduce potential confounding bias. Stronger evidence for causal associations is of great importance because few modifiable factors for preventing liver cancer are known.

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