

Research Paper

# Prognostic and Clinicopathological Significance of lncRNA MVIH in Cancer Patients

Yi Zhang<sup>1\*</sup>, Shibu Lin<sup>2\*</sup>, Xianjin Yang<sup>1✉</sup>, Xu Zhang<sup>1✉</sup>

1. Department of General Surgery, the First People's Hospital of Neijiang, Neijiang 641000, Sichuan Province, P. R. China.

2. Department of Hepatobiliary Surgery, The First Affiliated Hospital of Hainan, Medical College, Haikou 570102, Hainan Province, P. R. China

\*These authors contributed equally to this work.

✉ Corresponding authors: Xianjin Yang, E-mail: 2439015338@qq.com, Tel +86 138 9059 3536; Xu Zhang, E-mail: 1990486976@qq.com, Tel +86 139 8143 2513

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## Abstract

**Objectives:** lncRNA associated with microvascular invasion in hepatocellular carcinoma (lncRNA MVIH) is a newly discovered long non-coding RNA that aberrantly up-regulates and holds prognostic value in various tumors. The aim of the review and meta-analysis is to assess its prognostic potential and functions in malignant tumors.

**Materials and methods:** PubMed, PMC, EMBASE, Web of Science, Cochrane Library, China National Knowledge Infrastructure (CNKI), and Wanfang Database were carefully searched for articles published as of Jan. 1, 2018. Hazard ratios (HRs) or odds ratios (ORs) with 95% confidence intervals (95% CIs) were calculated to demonstrate prognostic value of lncRNA MVIH using Stata 12.0 software.

**Results:** Six original studies including 830 cancer patients were ultimately enrolled in this meta-analysis. The pooled results showed that elevated lncRNA MVIH expression was significantly correlated with unfavorable OS (HR=2.17, 95% CI: 1.58-2.76,  $p < 0.001$ ) and RFS/DFS/PFS (HR=2.21, 95% CI: 1.54-2.87,  $p < 0.001$ ) in cancer patients. Additionally, high expression of lncRNA MVIH was related to positive lymph node metastasis (OR = 3.04, 95% CI: 1.37-6.75,  $p = 0.006$ ) and advanced clinical stage (OR = 2.52, 95% CI: 1.68-3.79,  $p < 0.001$ ).

**Conclusions:** lncRNA MVIH over-expression was associated with poor clinical outcomes in multiple cancer types. More studies are warranted to verify and strengthen the clinical value of lncRNA MVIH in human cancers.

Key words: MVIH, lncRNA, cancer, prognosis, meta-analysis

## Introduction

In recent years, long non-coding RNAs have drawn increasingly attention and interest [1-3]. By definition, it is one class of non-coding RNA molecules with more than 200 nucleotides in length but no open reading frame [4-5]. In the past several decades, it has been considered as the "junk" of genome. But now, lncRNAs have been rising stars and serving pivotal roles in a variety of biological processes, such as gene regulation, posttranscriptional regulation, and intercellular signaling [6-9]. Importantly, many lncRNAs have been identified as being cancer-specific, which could exert carcinogenic or anti-tumor effects in tumor occurrence and

progression [10-13]. Moreover, the abnormal expression of lncRNAs was also reported to be correlated with clinical features and cancer prognosis [14-15]. lncRNAs are currently believed to be utilized as novel therapeutic targets and biomarkers that could aid cancer treatment and prognostication.

Microvascular invasion in hepatocellular carcinoma (MVIH) is a newly found lncRNA. It was first reported to be up-regulated in hepatocellular carcinoma and participated in the process of microvascular invasion [16]. Remarkably, it could serve as a prognostic predictor for patients after hepatectomy [16]. Subsequently, lncRNA MVIH has

been reported to be involved in tumor progression, and is suggestive of prognostic potential in multiple human cancers [17-19]. For example, Nie et al [17] reported that lncRNA MVIH was up-regulated in tissues of non-small cell lung cancer, and increased MVIH expression levels could promote cell proliferation and invasion, whereas high lncRNA MVIH levels were associated with malignant biological behavior (such as lymph node metastasis) and poor prognosis. In breast cancer, Lei et al [18] found that elevated MVIH expression levels could promote cell proliferation and the cell cycle, while high lncRNA MVIH was an unfavorable prognostic factor and was associated with Ki67 expression but not with lymph node metastasis. It should be taken into consideration that the individual study consisted of a relatively small sample and some discrepancies existed in these studies for clinical outcomes. In addition, still no study gives a compressive assessment of prognostic values of lncRNA MVIH in human cancers. Therefore, we carried out this systematic review and meta-analysis to clarify the predictive value of lncRNA MVIH in various cancer patients.

## Material and Methods

### Search Strategy

Two reviewers (Zhang Y and Lin SB) performed the literature retrieval, independently. The following electronic databases were screened: PubMed, PMC, EMBASE, Web of Science, Cochrane Library, China National Knowledge Infrastructure (CNKI), and Wanfang Database. The latest search was up to Jan. 1, 2018. The following keywords were used for literature searching: "lncRNA associated with microvascular invasion in hepatocellular carcinoma" OR "lncRNA MVIH" OR "MVIH" OR "Microvascular invasion in hepatocellular carcinoma." In order to supplement our literature search, the reference lists of relevant articles were observed for additional eligible studies.

### Inclusion and Exclusion Criteria

Eligible articles were identified based on the following inclusion criteria: 1) The expression of lncRNA MVIH was detected in any human tumors; 2) Correlation of lncRNA MVIH with prognosis was reported; 3) Sufficient data were available for calculating the hazard ratio (HR) with 95% CI. 4) Patients were categorized as high or low group according to lncRNA MVIH expression level. Articles were excluded when they did not cover the points above.

### Data Extraction and Quality Assessment

Two investigators (Zhang Y and Lin SB) independently extracted data from identified studies

according to unified form. Extracted data elements included the following records: 1) Name of first author, publication year, country of origin, study design; cancer type, sample size, expression pattern, tumor stage, criterion of high expression, detection method, follow-up time, outcome measures, analysis type; 2) Hazard ratio (HR) with 95% CI for cancer prognosis; 3) Patient number for high expression and low expression, lymph node metastasis (LNM), distant metastasis (DM), microvascular invasion (MVI), and tumor-node-metastasis (TNM) stage.

If a study reported the data in multivariate analysis and univariate analysis, the hazard ratio (HR) with the corresponding 95% CI was directly extracted from multivariate analysis. Those indirectly reported HRs and 95% CIs were extracted from graphical survival plots using Engauge Digitizer version 4.1 (<https://sourceforge.net/projects/digitizer/>).

The Newcastle-Ottawa quality assessment scale (NOS) with score range from 0-9 was applied to assess the quality of all included studies. A high quality study was identified as having a score of  $\geq 6$ .

### Statistical Analysis

In this meta-analysis, Stata version 12.0 (Stata Corporation, College Station, TX, USA) was used to calculate all statistical analyses. Higgins  $I^2$  statistics and Cochran's Q-test were applied to assess the heterogeneity among studies. When the percentage of  $I^2$  was greater than 50% or  $P_{het}$  less than 0.05, a random-effects model was used. Otherwise, a fixed-effects model was applied to the analyses. Begg's and Egger's test were utilized to detect the publication bias. In order to evaluate the robustness of the results, sensitivity analysis was performed by omitting the study one by one to assess the effects on the pooled results. A  $p$ -value  $< 0.05$  was considered to be statistically significant.

## Results

### Study selection and Characteristics

Initially, there were a total of 220 articles retrieved from a number of online databases after duplications removed. Following, 29 articles were left when 191 articles were excluded after titles and abstracts were screened for the following reasons: reviews, no association between lncRNA MVIH expression and human cancers, no clinical outcomes. After further screening of the remaining 29 studies, 23 were excluded due to lack of survival data, unrelated to specific prognosis. Finally, six available studies were deemed applicable to the meta-analysis [16-21]. The detailed steps for screening eligible articles is presented in **Figure 1**.

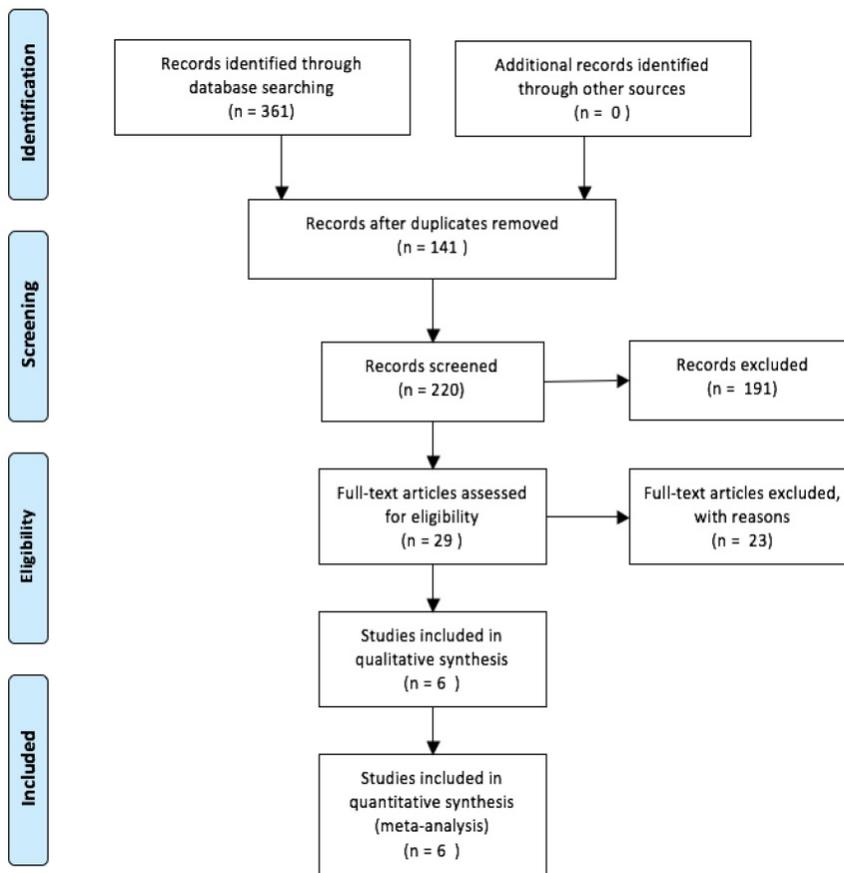
All six studies were published from 2012 to 2017. They were all identified as high-quality. Those studies enrolled a total of 830 cancer patients, with a mean subject size of 138.3, ranging from 42 to 250. All studies measured the expression of lncRNA MVIH in tissue specimens by quantitative real-time polymerase chain reaction (qRT-PCR) and reported the relationship between lncRNA MVIH expression and cancer prognosis.

Six studies reported patients' OS, and one focused on RFS, DFS, and PFS, respectively. Four studies estimated the relationship between lncRNA MVIH and TNM stage, and four focused on LNM, while one focused on MVI and DM, respectively. The cancers investigated consisted of hepatocellular carcinoma (HCC), non-small cell lung cancer (NSCLC), breast cancer (BC), gastric cancer (GC), glioma, and colorectal carcinoma (CRC). All included studies were retrospective, conducted in China, and focused on an Asian population. **Table 1 and Table 2** summarize the main information and data of all included studies.

## Results of the Meta-analysis

### MVIH Expression and OS

Six clinical studies that comprised a total of 830 cancer patients were conducted to analyze the OS; a



**Figure 1.** The steps for screening eligible articles.

fixed-effects model was utilized for no significant heterogeneity ( $I^2=0.0\%$ ,  $P_{\text{het}}=0.823$ ). From the pooled result, we found that increased lncRNA MVIH expression was correlated with poorer OS in cancer patients (HR=2.17, 95% CI: 1.58-2.76; **Figure 2A**). In the stratified analysis, we found that elevated MVIH expression could be a prognostic biomarker for patients with gastrointestinal (GI) cancers (HR: 2.14, 95% CI: 1.45-2.84,  $p < 0.001$ ) or patients with non-gastrointestinal cancers (HR: 2.25, 95% CI: 1.15-3.35,  $p < 0.001$ ). In the subgroup analyses by follow-up time, MVIH could act as a prognostic factor in groups with follow-up time  $< 5$  years (HR: 1.87, 95% CI: 1.03-2.70,  $p < 0.001$ ) or groups with follow-up time  $\geq 5$  years (HR: 2.48, 95% CI: 1.65-3.31,  $p < 0.001$ ). We also found that a high expression of MVIH could act as an unfavorable independent prognostic factor for OS based on multivariate analysis (HR: 2.31, 95% CI: 1.54-3.08,  $p < 0.001$ ). Stratification analyses for all subgroups are presented in detail in **Figure 2B-2D**.

### MVIH Expression and DFS/RFS/PFS

A total of 3 studies including 617 patients investigated the prognostic value of MVIH on cancer progression or recurrence, with a pooled HR of 2.21 (95% CI: 1.54-2.87;  $p < 0.001$ , **Figure 3**). No heterogeneity was observed among the studies, a fixed-effects model was performed to the calculation ( $I^2=0.0\%$ ,  $P_{\text{het}}=0.712$ ). Among them, one focused on the prognostic value of MVIH expression on cancer relapse (HR=1.99, 95% CI: 1.31-3.00), one reported that patients with high MVIH expression levels had poor disease-free survival (HR=2.54, 95% CI: 1.06-6.12). A final one explored the relationship between MVIH and tumor progression (HR=2.57, 95% CI: 1.64-4.04).

### MVIH Expression and Clinicopathological Characteristics

Four studies comprising of 488 cancer patients reported the correlation of MVIH with lymph node metastasis (LNM) in multiple tumors. A random-effect model was used for obvious heterogeneity observed across studies ( $I^2=66.4\%$ ,  $P_{\text{het}}=0.030$ ), the pooled results showed that patients with high MVIH expression in tumor tissues

preferentially metastasize to the lymph nodes (OR = 3.04, 95% CI: 1.37-6.75;  $p = 0.006$ , **Figure 4A**).

Furthermore, four studies with 536 patients declared the relationship between MVIH expression levels and clinical stage, with a fixed-effects model on

account of no obvious heterogeneity ( $I^2=0.0\%$ ,  $P_{het} = 0.681$ ). The combined results indicated that the patients with high MVIH expression have a more advanced stage (III/IV) (OR = 2.52, 95% CI: 1.68-3.79;  $p < 0.001$ , **Figure 4B**).

**Table 1** Main characteristics of all studies included in the meta-analysis.

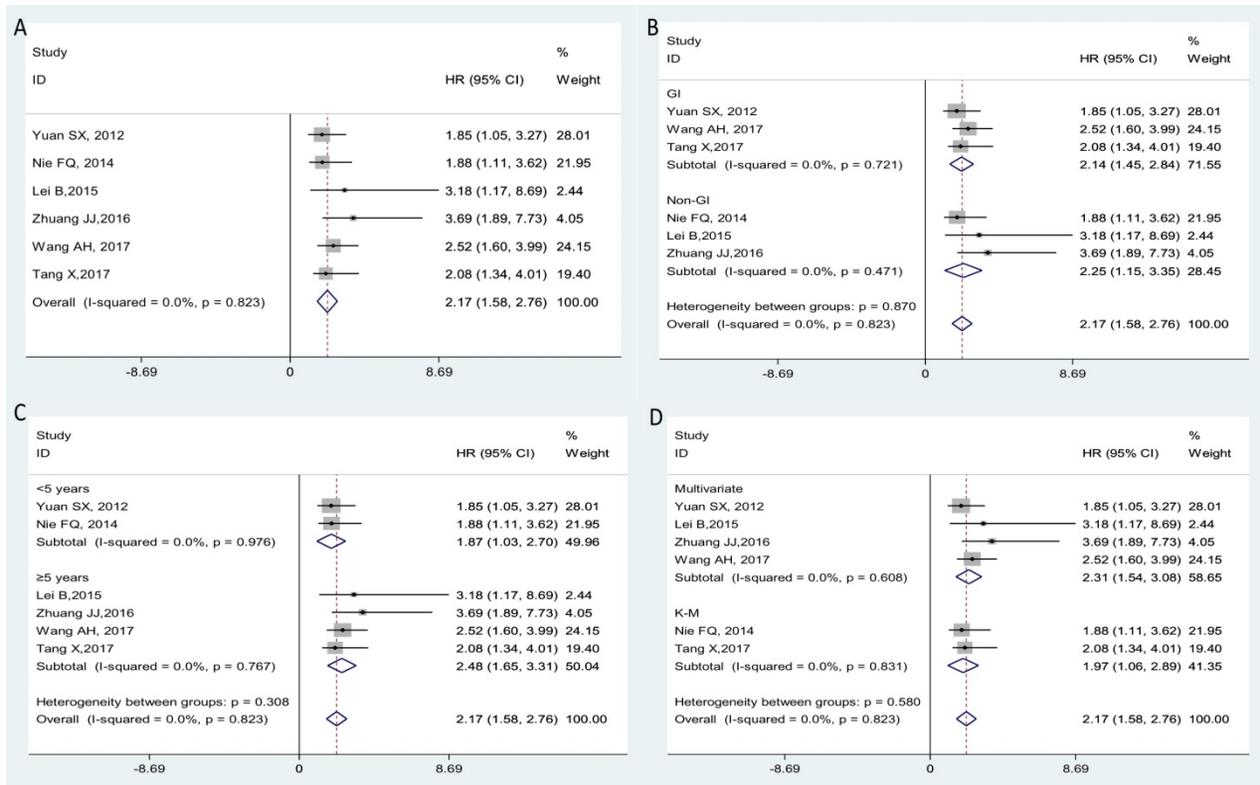
First author, Year	Country	Cancer type	Study design	Sample size	HE (%)	Stage	Criterion of HE	Follow-up	MVIH assay	Detected sample	Outcome measures	NOS score
Yuan SX, 2012 [16]	China	HCC	R	215	50.20%	I-III	at or above the 50th percentile	< 5 years	qRT-PCR	Tissue	OS, RFS	8
Nie FQ, 2014 [17]	China	NSCLC	R	42	50.00%	I-IIIa	MVIH expression ratio $\geq$ mean ratio(2.12-fold)	< 5 years	qRT-PCR	Tissue	OS	6
Lei B, 2015 [18]	China	BC	R	250	50.00%	NR	according to the mean value	$\geq$ 5 years	qRT-PCR	Tissue	OS, DFS	9
Zhuang JJ, 2016 [19]	China	Glioma	R	127	48.80%	I-IV	based on the median expression levels	$\geq$ 5 years	qRT-PCR	Tissue	OS	7
Tang X, 2016 [21]	China	CRC	R	44	77.27%	I-IV	according to the mean value	$\geq$ 5 years	qRT-PCR	Tissue	OS	6
Wang AH, 2017 [20]	China	GC	R	152	44.10%	I-IV	MVIH cut-off point (3.85)	$\geq$ 5 years	qRT-PCR	Tissue	OS, PFS	7

Abbreviations: R, retrospective; HCC, hepatocellular carcinoma; NSCLC, non-small cell lung cancer; BC, breast cancer; GC, gastric cancer; CRC, colorectal carcinoma; NR, not reported; HE, high expression; OS, over survival; DFS, disease-free survival; RFS, recurrence-free survival; PFS, progression-free survival; qRT-PCR, quantitative real-time polymerase chain reaction.

**Table 2** HRs and 95% CIs of cancer prognosis and progression associated with MVIH expression in all included studies.

First author, Year	Country	Cancer type	Case number of MVIH expression										HR(95%CI) for OS	HR(95%CI) for RFS/DFS/PFS
			High	High with LNM	High with DM	High with MVI	High with II/IV	Low	Low with LNM	Low with DM	Low with MVI	Low with II/IV		
Yuan SX, 2012 [16]	China	HCC	108	NR	NR	40	22	107	NR	NR	23	13	1.85(1.05-3.27)#	1.99(1.31-3.00)#
Nie FQ, 2014 [17]	China	NSCLC	21	14	NR	NR	11	21	7	NR	NR	6	1.88(1.11-3.62)*	NR
Lei B, 2015 [18]	China	BC	125	77	NR	NR	NR	125	58	NR	NR	NR	3.18(1.17-8.69)#	2.54(1.06-6.12)#
Zhuang JJ, 2016 [19]	China	Glioma	62	NR	NR	NR	41	65	NR	NR	NR	29	3.69(1.89-7.73)#	NR
Tang X, 2017 [21]	China	CRC	34	29	NR	NR	NR	10	1	NR	NR	NR	2.08(1.34-4.01)*	NR
Wang AH, 2017 [20]	China	GC	67	32	4	NR	25	85	27	3	NR	12	2.52(1.60-3.99)#	2.57(1.64-4.04)#

Abbreviations: HCC, hepatocellular carcinoma; NSCLC, non-small cell lung cancer; BC, breast cancer; GC, gastric cancer; CRC, colorectal carcinoma; OS, over survival; DFS, disease-free survival; RFS, recurrence-free survival; PFS, progression-free survival; NR, not reported; LNM, lymph nodes metastasis; DM, distant metastasis; MVI, microvascular invasion; \*HR calculated from survival curves; #HR directly from multivariate analysis.



**Figure 2.** Subgroup analysis for OS. Over-all (A); stratified by the cancer type (B), by follow-up time (C) and by analysis type (D).

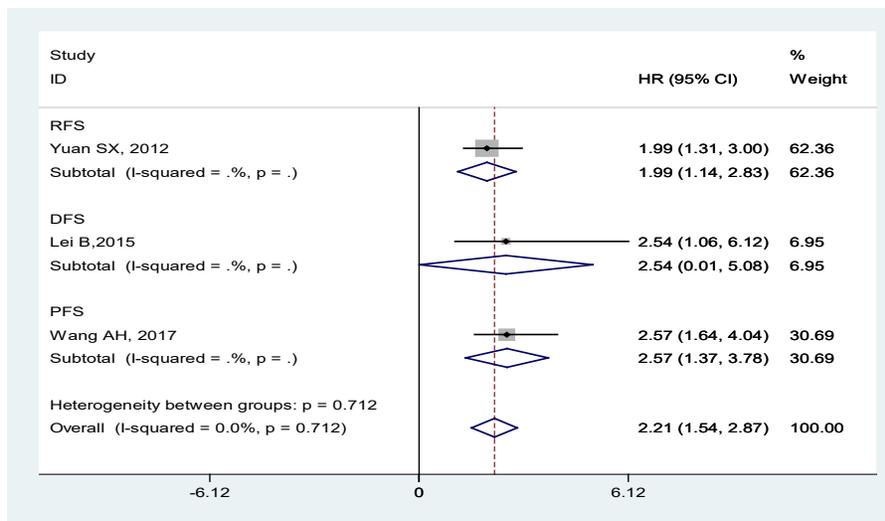


Figure 3. Meta-analysis of the pooled HRs of DFS/RFS/PFS of patients with high MVIH expression.

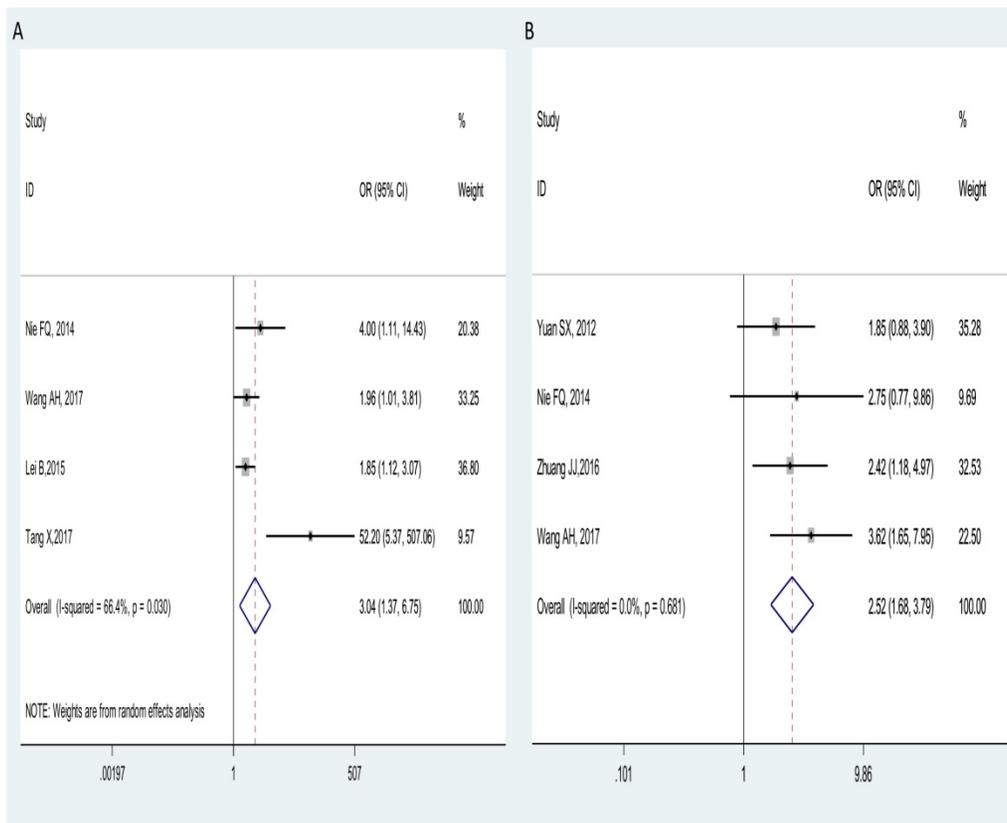


Figure 4. Forest plots of combined analyses associated with MVIH expression. (A): LNM; (B): clinical stage.

The studies by Yuan et al [16] and Wang et al [20] conducted the microvascular invasion (MVI) and distant metastasis (DM) analysis of cancer patients, respectively. Those data suggested that the overexpression of MVIH was associated with frequent MVI ( $p=0.016$ ), but was unable to accurately predict the occurrence of DM ( $p=0.476$ ).

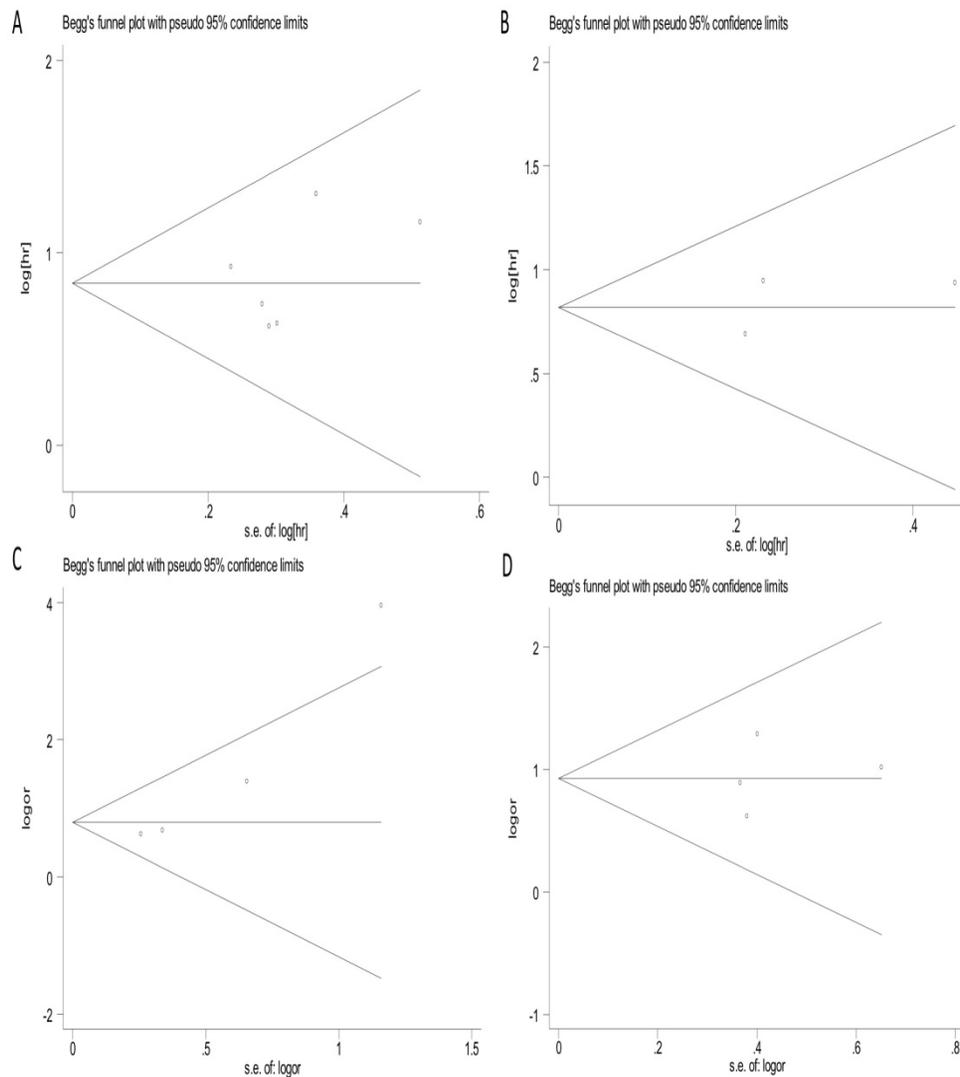
**Publication Bias**

Funnel plots and Begg’s and Egger’s test were

used to evaluate the publication bias in this meta-analysis. As shown in Figures 5A-5D, the Begg’s funnel plots with pseudo 95% CIs were symmetric. The test results also indicated no significant publication bias in the meta-analysis.

**Sensitivity Analysis**

As illustrated in Figures 6A-6D, the analyzed results suggested that our results were comparatively credible and stable.

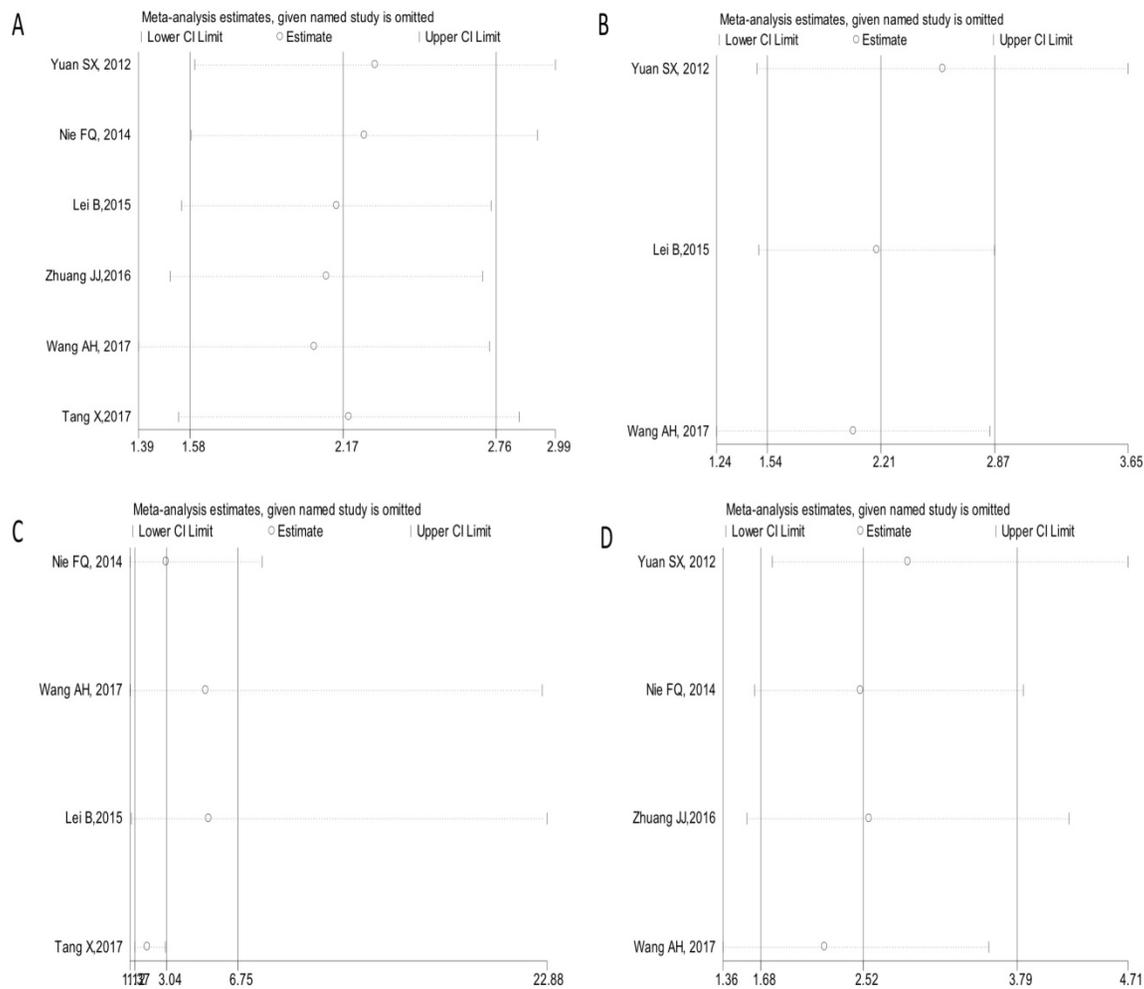


**Figure 5.** Begg's funnel plots of publication bias test. (A) OS; (B) RFS/DFS/PFS; (C) LNM; (D) clinical stage.

## Discussion

MVIH is a novel identified lncRNA transcript, the majority of the MVIH sequence was located within the intron of the RPS24 gene, but both the MVIH and RPS24 genes were independently transcribed [16, 22]. As a cancer-associated lncRNA, MVIH was discovered in several cancers and showed carcinogenic roles in tumor progression. In hepatocellular cancer, MVIH could activate tumor-inducing angiogenesis by inhibiting the secretion of phosphor-glycerate kinase 1, while MVIH expression was inversely correlated with cancer prognosis. Therefore, deregulation of lncRNA MVIH could be utilized as a potential target for new adjuvant therapies and serves as a predictor for poor recurrence-free survival after hepatectomy [16]. Furthermore, Shi et al [23] found mechanistically MVIH could promote cell growth and inhibit cell

apoptosis of HCC via inhibiting miR-199a expression. Finally, Cheng et al [24] reported that ARID1A could interact with lncRNA MVIH and ARID1A suppressed HCC cells proliferation and migration by inhibiting MVIH. In breast cancer, the knockdown of MVIH expression by siRNA could suppress cell proliferation and cell cycle, while increasing cell apoptosis, whereas up-regulated MVIH expression showed the converse [18]. MVIH also displayed oncogenic functions in NSCLC, high levels of MVIH could partly promote cell proliferation and invasion via regulating MMP2 and MMP9 protein expression *in vitro* [17]. MVIH also showed the same oncogenic effect in glioma cell migration and invasion [19]. Although MVIH has been discovered in several different kinds of cancers, the exact functional roles of it in carcinogenesis remain unclear and need to be further elucidated.



**Figure 6** Sensitivity analysis under specific model. (A): for OS; (B): for RFS/DFS/PFS; (C): for LNM; (D): for clinical stage.

As we all know, this is the first meta-analysis that elaborated the clinical value of lncRNA MVIH in human cancers. We demonstrated that the high expression of lncRNA MVIH was significantly related with unsatisfactory clinical outcomes; the patients with high MVIH level in tissues had a worse OS when compared with those with low MVIH expression (HR=2.17, 95% CI: 1.58-2.76), and the prognostic value of MVIH on cancer progression or recurrence was also revealed (HR=2.21, 95% CI: 1.54-2.87). Furthermore, the clinical role of MVIH expression was also exhibited in regard to LNM (OR = 3.04, 95% CI: 1.37-6.75) and tumor stages (OR = 2.52, 95% CI: 1.68-3.79). In this meta-analysis, no significant heterogeneity was observed when we carried out OS and DFS/RFS/PFS analysis of MVIH, as well as comparison for TNM stage. The results from the sensitivity analysis and publication bias evaluation also further indicated our analyses were credible and stable.

Although this meta-analysis was conducted with standard procedure and rigorous statistics, our

conclusion still has several limitations that should be clearly interpreted. First of all, the total sample size was relatively small and the number of enrolled cancer types still needs to be enlarged. Secondly, the quantity of included studies was not sufficient for a more comprehensive analysis for other prognostic indicators and clinic-pathological features, such as DFS, DM or MVI. Thirdly, all included were retrospective researches, and only a Chinese population was applicable in this meta-analysis, potentially minimizing the analyzing value to some extent and affecting the broader applicability of our conclusions. Fourthly, studies with positive results are more easily published than negative results, which might cause potential publication bias and some missing data. At last, the cut-off value of high lncRNA MVIH expression is needed to be unified before its clinical application. Regarding these deficiencies, the role of lncRNA MVIH in multiple human carcinomas might be exceedingly evaluated. Therefore, further multi-center studies that are well-designed and at a

large-scale are needed to verify the function of lncRNA MVIH in various cancers.

To summarize, the present meta-analysis highlighted the clinical effect of lncRNA MVIH in predicting prognosis, metastasis, and clinical stage in multiple cancers. To better evaluate the prognostic role of lncRNA MVIH expression in human cancers, higher quality studies with larger sample sizes are required for a more complete confirmation.

## Competing Interests

The authors have declared that no competing interest exists.

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