

Clinicopathological significance and prognostic role of p-STAT3 in patients with hepatocellular carcinoma

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Background and aim: Constitutive activation of STAT3 through its phosphorylation (p-STAT3) plays a key role in the development and progression of various cancers. However, the relationship between p-STAT3 expression and the clinicopathological features and prognostic value in patients with hepatocellular carcinoma (HCC) remains controversial. We conducted a meta-analysis to evaluate the role of p-STAT3 in HCC.

Methods: The PubMed, Cochrane Library, Web of Science, EMBASE, Chinese CNKI, and Chinese Wanfang databases were searched using the appropriate terms to find the relevant studies on p-STAT3 and HCC. The relationship between p-STAT3 expression and clinicopathological characteristics and prognostic value was established. Pool odds ratios (ORs) and hazard ratios (HRs) with 95% CIs were calculated using the STATA 14.2 software.

Results: The eight articles included in this meta-analysis comprised 752 patients. Expression of p-STAT3 was associated with incidence, age, liver cirrhosis, tumor size, vascular invasion, and TNM stage of HCC, but it was not related to gender, alpha-fetoprotein (AFP), hepatitis B surface antigen (HBsAg), number of tumors, and tumor differentiation. Additionally, the expression of p-STAT3 was related to a poor 3- and 5-year overall survival rate and disease-free survival rate.

Conclusion: Expression of p-STAT3 was associated with the incidence, age, liver cirrhosis, tumor size, vascular invasion, and TNM stage. Thus, p-STAT3 can be a reliable prognostic biomarker for HCC. Further high-quality studies with larger numbers of patients are needed.

Keywords: p-STAT3, hepatocellular carcinoma, prognosis, meta-analysis

Background

Hepatocellular carcinoma (HCC) is prone to lead to intrahepatic metastasis, and the prognosis is poor. More than half of the 660,000 new cases of HCC worldwide each year occur in China.^{1,2} The effective method for the treatment of HCC is a comprehensive treatment based on surgical resection. However, occult onset of HCC can occur and make it difficult to diagnose in the early stage, which leads to many patients with HCC being diagnosed in the middle or late stage. As a result, only 20%–25% of patients with liver cancer are amenable to surgery and the surgical resection rate is very low. In addition, postoperative recurrence and metastasis are common, so the long-term survival rate of patients with HCC remains low.^{3–5} Recently, studies on the mechanisms of recurrence and metastasis of liver cancer at the molecular level have been reported. Nevertheless, a better understanding of the pathogenesis of HCC is needed for the early prediction of its occurrence and prognosis. Thus, the identification of reliable biomarkers for HCC would be valuable.

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Recently, multiple studies have shown that STAT3, an important member of the STAT protein family, is involved in the development and progression of tumors. The constitutive activation of STAT3 through its phosphorylation is thought to be involved in the development, invasion, and metastasis of malignant tumors.⁶ Sustained levels of phosphorylated STAT3 (p-STAT3) have been found in gastric cancer,⁷ pancreatic cancer,⁸ prostate cancer,⁹ breast cancer,¹⁰ colorectal cancer,¹¹ melanoma,¹² and other malignant tumors. In addition, p-STAT3 has been recognized as an oncoprotein. In particular, p-STAT3 expression has been found to be associated with clinicopathological characteristics and prognosis of patients with HCC, but controversy remains. For instance, Wu et al¹³ indicated that the expression of p-STAT3 is only related to TNM stage but not to other aspects including age, gender, and cirrhosis. Zhang et al¹⁴ reported associations of p-STAT3 expression with TNM stage and metastasis. Mano et al¹⁵ found that p-STAT3 expression was correlated with tumor size, level of alpha-fetoprotein (AFP), and intrahepatic metastasis, but not with tumor differentiation or vascular invasion. Yang et al¹⁶ found that p-STAT3 expression was associated with intratumor microvessel density in HCC and p-STAT3 was a significant predictor of overall survival (OS) of patients with HCC ($P=0.036$).

Two meta-analyses assessed the relationship between STAT3 and prognosis of digestive or various solid tumors. One study¹⁷ reported that STAT3 expression suggested a poor 3- and 5-year OS rate in solid tumors. However, this meta-analysis did not assess the relationship between p-STAT3 specifically and prognosis and instead included data of p-STAT3 and STAT3 combined. Moreover, there were little data on the association of p-STAT3 with the clinicopathological features of HCC. The other study¹⁸ explored the relationship between p-STAT3 expression and prognosis of patients with digestive tumors. In this meta-analysis, there were few articles for HCC and no data were included on the correlation between p-STAT3 expression and the 3- and 5-year survival and clinicopathological features of patients with HCC.

These deficiencies prompted the present meta-analysis so as to more thoroughly evaluate the relationship between the expression of p-STAT3 and the clinicopathological features and prognosis of patients with HCC.

Methods

Search strategy and study selection

The PubMed, Cochrane Library, Web of Science, EMBASE, Chinese CNKI, and Chinese Wanfang databases were searched up to September 15, 2017, with no limit on the

earlier date. The keywords used to perform the search were “p-STAT3” or “phosphorylated signal transducer and activator of transcription 3” and “hepatocellular carcinoma” or “hepatic tumor” or “liver tumor” or “hepatic cancer” or “liver cancer” or “HCC.”

The inclusion criteria for studies on HCC and p-STAT3 were as follows: 1) the expression of p-STAT3 in patients with HCC was measured by immunohistochemistry (IHC) analysis; 2) the results included the survival information (OS and disease-free survival [DFS] rates) and clinicopathological characteristics; 3) hazard ratios (HRs) for OS or DFS can be calculated for the survival curves; and 4) in case of repeated studies, the most recent report is included in the meta-analysis.

The exclusion criteria for studies were as follows: 1) case reports, reviews, editorials, letters, conference abstracts, or animal trials and 2) repeated studies based on the same databases.

Data extraction and quality assessment

All eligible articles were independently screened in the same way by two independent reviewers (CJ Liang and YC Xu). The following data were extracted from eligible studies: first author, publication year, country or region, case numbers, clinicopathological characteristics, 3- and 5-year OS rates, and DFS rate (Table 1). The Newcastle-Ottawa Scale (NOS) criteria were used to assess the quality of the studies. A study was considered to be a low-quality or high-quality study when the NOS score was <6 or ≥ 6 , respectively.

Statistical analyses

The STATA 14.2 (StataCorp LP, College Station, TX, USA) and Engauge Digitizer 10.0 software were used to calculate all statistical data or extract the survival data from Kaplan–Meier curves. Pooled odds ratios (ORs) were used to assess the relationship between p-STAT3 expression and clinicopathological features (gender [male vs female], age [>50 vs ≤ 50 years], hepatitis B surface antigen [HbsAg] [yes vs no], AFP [>400 vs ≤ 400], liver cirrhosis [yes vs no], tumor size [≤ 5 vs >5 cm], vascular invasion [yes vs no], number of tumors [single vs multiple], tumor differentiation [low vs high + moderate], and TNM stage [I + II vs III + IV]). Pooled HR with 95% CI was used to evaluate the association between p-STAT3 expression and 3- and 5-year OS and DFS. Heterogeneity was assessed by the I^2 test and Q test if the $I^2 < 50\%$ fixed-effects model was used, otherwise the random-effects model was chosen. Funnel plots and Egger's tests were used to assess the potential publication bias. The source of heterogeneity was determined by sensitivity

and subgroup analyses. If the pooled estimated OR or HR (95% CI) did not overlap 1, the effect of the expression of p-STAT3 on the clinicopathological features and survival was considered significant.

Results

Identification of eligible studies, study characteristics, and quality assessment

A total of 780 potentially relevant studies were selected from the PubMed, Web of Science, Cochrane Library, EMBASE, CNKI, and Chinese Wanfang databases. Duplicated studies, non-HCC studies and nonoriginal articles, comprising 717 articles, were excluded (Figure 1). The remaining 63 articles were further evaluated by reading the abstracts, of which 28 articles were excluded as they were non-p-STAT3-related human studies or did not analyze tumor tissues. The remaining 35 studies were assessed by reading the full text. Of them, 27 studies were excluded because of lack of clinicopathological data. Ultimately, eight^{13–16,19–22} eligible studies involving 752 patients with HCC were included in this meta-analysis. The studies were published between 2007 and 2014, and the sample size ranged from 68 to 138 patients. Six studies were from mainland China, one study was from Japan, and one study was from Taiwan. Five of these studies were published in English, and the others were published in Chinese (Table 1). All eight studies were high quality.

Association between p-STAT3 in HCC and clinicopathological features

The association between p-STAT3 expression and clinicopathological features of patients with HCC was evaluated. As shown in Table 2 and Figure 2A–K, p-STAT3 was overexpressed in HCC tissue compared to adjacent nontumor tissue (OR =6.27, 95% CI =2.85, 13.79, $P=0.000$, $I^2=76.5%$, random effect) and p-STAT3 expression was associated with age (OR =1.45, 95% CI =1.01, 1.49, $P=0.044$, $I^2=0$, fixed effect), liver cirrhosis (OR =1.56, 95% CI =1.01, 2.43, $P=0.046$, $I^2=0$, fixed effect), tumor size (OR =0.56, 95% CI =0.40, 0.78, $P=0.001$, $I^2=45.2%$, fixed effect), vascular invasion (OR =3.43, 95% CI =2.13, 5.52, $P=0.000$, $I^2=34.2%$, fixed effect), and TNM stage (OR =0.36, 95% CI =0.19, 0.70, $P=0.042$, $I^2=66.1%$, random effect). However, p-STAT3 expression was not significantly related to gender (OR =0.91, 95% CI =0.63, 1.31, $P=0.618$, $I^2=0$, fixed effect), HBsAg (OR =1.11, 95% CI =0.75, 1.65, $P=0.597$, $I^2=0$, fixed effect), AFP (OR =1.40, 95% CI =0.92, 2.14, $P=0.121$, $I^2=0$, fixed effect), number of tumors (OR =0.64, 95% CI =0.35, 1.18, $P=0.155$, $I^2=0$, fixed effect), and tumor differentiation

(OR =1.81, 95% CI =0.79, 4.15, $P=0.161$, $I^2=69.3%$, random effect). The results of the subgroups analysis based on p-STAT3 expression, tumor differentiation, and TNM stage by sample size and NOS score are presented in Table 3. p-STAT3 was overexpressed in studies with a small sample size ($n \leq 100$, OR =7.70, 95% CI =2.82, 21.00, $P=0.003$, $I^2=78.1%$, random effect) or those ranked with a low NOS score (NOS ≤ 7 , OR =5.46, 95% CI =2.64, 11.31, $P=0.000$, $I^2=65.7%$, random effect). However, p-STAT3 expression was not obviously significant in studies with large sample size ($n > 100$, OR =5.78, 95% CI =0.56, 59.51, $P=0.140$, $I^2=65.2%$, random effect) or studies ranked as high quality (NOS > 7 , OR =20.14, 95% CI =0.26, 595.43, $P < 0.001$, $I^2=87.4%$, random effect). Heterogeneity was evident in each subgroup. Also, p-STAT3 expression was related to tumor differentiation in a subgroup of studies with a large sample size ($n > 100$, OR =1.78, 95% CI =1.15, 3.69, $P=0.015$, $I^2=0$, random effect). However, p-STAT3 expression was not related to differentiation in other subgroups and heterogeneity was evident in the subgroups of studies with a small sample size and low NOS score subgroup ($n \leq 100$, $I^2=85%$ and NOS ≤ 7 , $I^2=82.8%$, respectively). Additionally, heterogeneity was significant for each TNM stage group. The sources of the heterogeneities could not be identified because of the limitations of sample size and NOS score. Despite this, p-STAT3 expression was related to poor clinicopathological features. It is conceivable that p-STAT3 plays an oncogenic role in HCC.

Association between p-STAT3 expression in HCC and 3- and 5-year OS and DFS

Six studies comprising 610 patients were analyzed to determine the relationship between p-STAT3 expression and the 3-year OS (HR =1.69, 95% CI 1.07, 2.31; $I^2=0.0$, $P < 0.001$). Five studies comprising 542 patients were evaluated to establish the correlation between p-STAT3 and the 5-year OS (HR =1.67, 95% CI 1.18, 2.15; $I^2=0.0$, $P < 0.001$). Three studies comprising 352 patients were examined to find the relationship between p-STAT3 expression and DFS (HR =1.94, 95% CI 1.42, 2.47; $I^2=42.0%$, $P < 0.001$). These results (Figure 3) indicated that there is a correlation between the p-STAT3 expression and poor prognosis of patients with HCC. Additionally, these results reinforce the view that p-STAT3 may be a reliable prognostic biomarker for HCC.

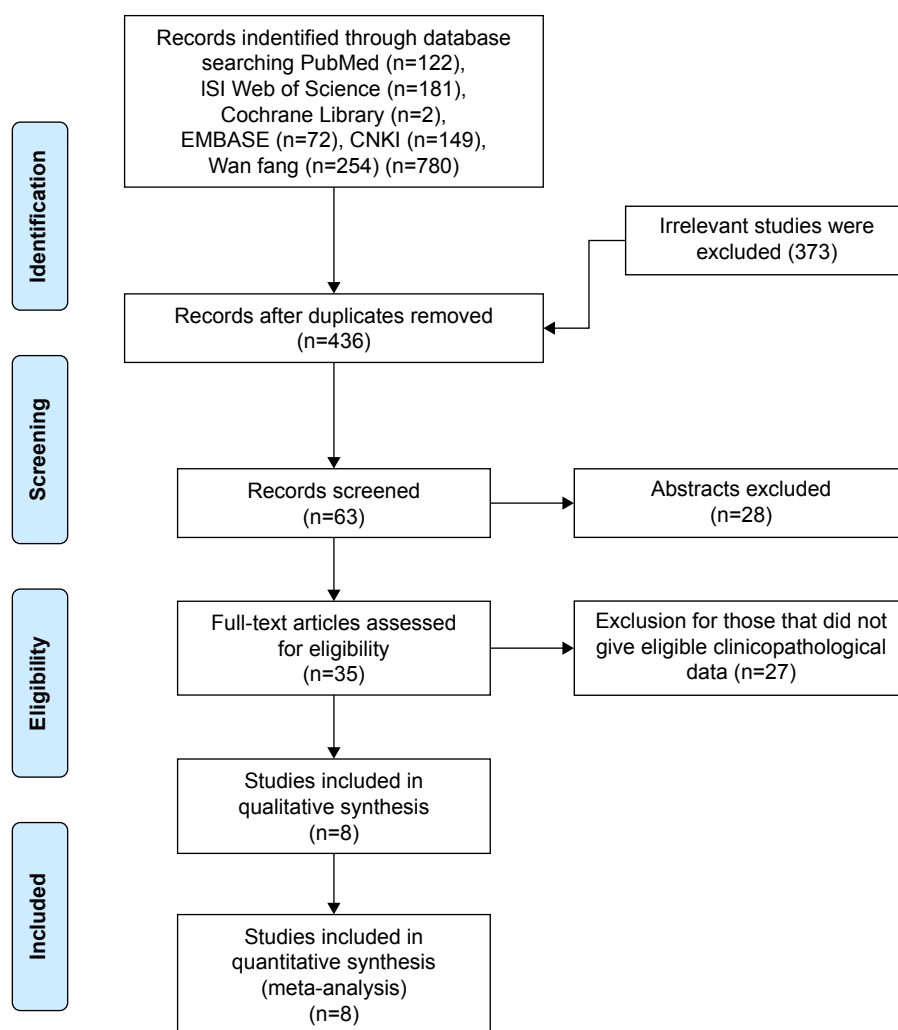
Publications bias

No publication biases were evident for p-STAT3 expression ($P=0.06$), gender ($P=0.711$), age ($P=0.230$), HBsAg

Table 1 Characteristics of studies included in the meta-analysis

Studies	Year	Country	Kind	Number of patients	Laboratory method	Expression, cancer (±) and control (±)	Gender, male (±) and female (±)	Age (years), >50 (±) and ≤50 (±)	HBsAg, yes (±) and no (±)
Yang et al ¹⁶	2007	Taiwan	HCC	69	IHC	50/19 14/55	38/14 12/5	NA	30/12 20/7
Wu et al ¹²	2011	China	HCC	138	IHC	75/63 35/75	60/55 15/8	51/35 24/28	60/51 15/12
Wu et al ²²	2011	China	HCC	113	IHC	NA	55/37 13/8	45/22 23/23	58/34 10/11
Zhang et al ¹⁴	2012	China	HCC	100	IHC	58/42 0/10	48/32 10/10	28/20 30/22	47/38 11/4
Zepeng et al ¹⁹	2013	China	HCC	90	IHC	46/44 0/90	35/30 11/14	25/23 21/21	NA
Mano et al ¹⁵	2013	Japan	HCC	101	IHC	NA	26/55 10/10	NA	8/14 28/51
Tao et al ²⁰	2014	China	HCC	73	IHC	38/35 20/53	31/30 7/5	NA	32/26 6/9
Lin et al ²¹	2014	China	HCC	68	IHC	46/22 18/50	38/18 8/4	17/6 29/16	41/18 5/4

Abbreviations: AFP, alpha-fetoprotein; HBsAg, hepatitis B surface antigen; HCC, hepatocellular carcinoma; IHC, immunohistochemistry; DFS, disease-free survival; NA, not applicable; OS, overall survival.

**Figure 1** Flow diagram depicting the study selection.

Liver cirrhosis, yes (\pm) and no (\pm)	AFP, >400 (\pm) and \leq 400 (\pm)	Tumor size (cm), \leq 5 (\pm) and >5 (\pm)	Tumor number, single (\pm) and multiple (\pm)	Vascular invasion, yes (\pm) and no (\pm)	Differentiation, low (\pm) and high and moderate (\pm)	TNM stage, I, II (\pm) and III, IV (\pm)	Survival information	Quality score
36/12	25/8	35/10	NA	17/5	3/5	37/13	N	6
14/7	18/8	15/9		33/14	47/14	13/6		
69/56	NA	13/28	NA	NA	2/3	50/56	OS RFS	8
6/7		62/35			73/60	25/7		
63/38	NA	17/15	NA	NA	2/0	47/39	OS DFS	7
5/7		51/30			66/45	21/6		
38/22	32/19	29/26	NA	NA	24/8	35/25	OS	8
20/20	26/23	29/16			34/34	23/17		
NA	36/27	27/32	29/32	14/1	10/10	24/40	OS	8
	10/17	19/12	17/12	32/43	36/34	22/4		
14/22	NA	NA	NA	24/30	16/19	NA	OS DFS	8
22/43				12/35	20/46			
NA	12/14	12/20	28/39	23/8	11/3	11/19	N	7
	26/21	26/15	10/6	15/27	27/32	27/16		
NA	32/13	18/10	36/17	31/8	31/3	16/17	OS	7
	14/9	28/12	10/5	15/14	15/19	30/5		

($P=0.462$), AFP ($P=1$), liver cirrhosis ($P=0.806$), tumor size ($P=0.707$), number of tumors ($P=0.296$), vascular invasion ($P=0.452$), tumor differentiation ($P=0.368$), TNM stage ($P=0.230$), 3-year OS ($P=0.268$), 5-year OS ($P=0.368$), and DFS ($P=1$) according to the Begg's test (Figures 3 and 4).

Discussion

STAT3 becomes activated in response to interleukin-6 (IL-6) and other cytokines and functions as an acute-phase factor response factor. The *STAT3* gene is located in human chromosome 17. The 4815 bp gene harbors 24 exons that include STAT3 α , β , and γ .^{23,24} STAT3 is involved in early embryonic development and differentiation of bone marrow

cells, regulation of cell growth and differentiation, proliferation, malignant transformation, and apoptosis inhibition, and other important physiological functions.²⁵⁻²⁷

Recent experimental studies revealed the key role of STAT3 in the development of tumors. STAT3 is activated by a variety of carcinogens, including viral infections, ionizing radiation, growth factors, oncogenes, and inflammatory mediators.²⁸ Constitutive activation of STAT3 occurs most often in tumor cells and generally does not occur in normal cells. p-STAT3 is overexpressed in a variety of tumor tissues and cells. Such overexpression is associated with clinicopathological features and prognosis of patients with various cancers, including HCC. However, the relationship

Table 2 p-STAT3 clinicopathological features for hepatocellular carcinoma

Heterogeneity							
Clinicopathological features	Number of studies	Number of patients	Pooled OR (95% CI)	PHet	I^2 (%)	P-value	Model used
Expression (positive:negative)	6	958	6.27 (2.85, 13.79)	0.001	76.5	<0.001	Random
Gender (male:female)	8	752	0.91 (0.63, 1.31)	0.673	0.0	0.618	Fixed
Age (years) (>50: \leq 50)	5	509	1.45 (1.01, 1.49)	0.704	0.0	0.044	Fixed
HBsAg (yes:no)	7	662	1.11 (0.75, 1.65)	0.568	0.0	0.597	Fixed
AFP (>400: \leq 400)	5	390	1.40 (0.92, 2.14)	0.529	0.0	0.121	Fixed
Liver cirrhosis (yes:no)	5	521	1.56 (1.01, 2.43)	0.942	0.0	0.046	Fixed
Tumor size (cm) (\leq 5:>5)	7	651	0.56 (0.40, 0.78)	0.090	45.2	0.001	Fixed
Vascular invasion (yes:no)	5	401	3.43 (2.13, 5.52)	0.193	34.2	<0.001	Fixed
Tumor number (single:multiple)	3	231	0.64 (0.35, 1.18)	0.568	0.0	0.155	Fixed
Differentiation (low:H + M)	8	752	1.81 (0.79, 4.15)	0.002	69.3	0.161	Random
TNM stage (I + II:III + IV)	7	651	0.36 (0.19, 0.70)	0.007	66.1	0.003	Random

Notes: Random, random-effects model; fixed, fixed-effects model; H + M, high + moderate level of differentiation.

Abbreviations: AFP, alpha-fetoprotein; CI, confidence interval; HBsAg, hepatitis B surface antigen; OR, odds ratio; PHet, probability of heterogeneity.

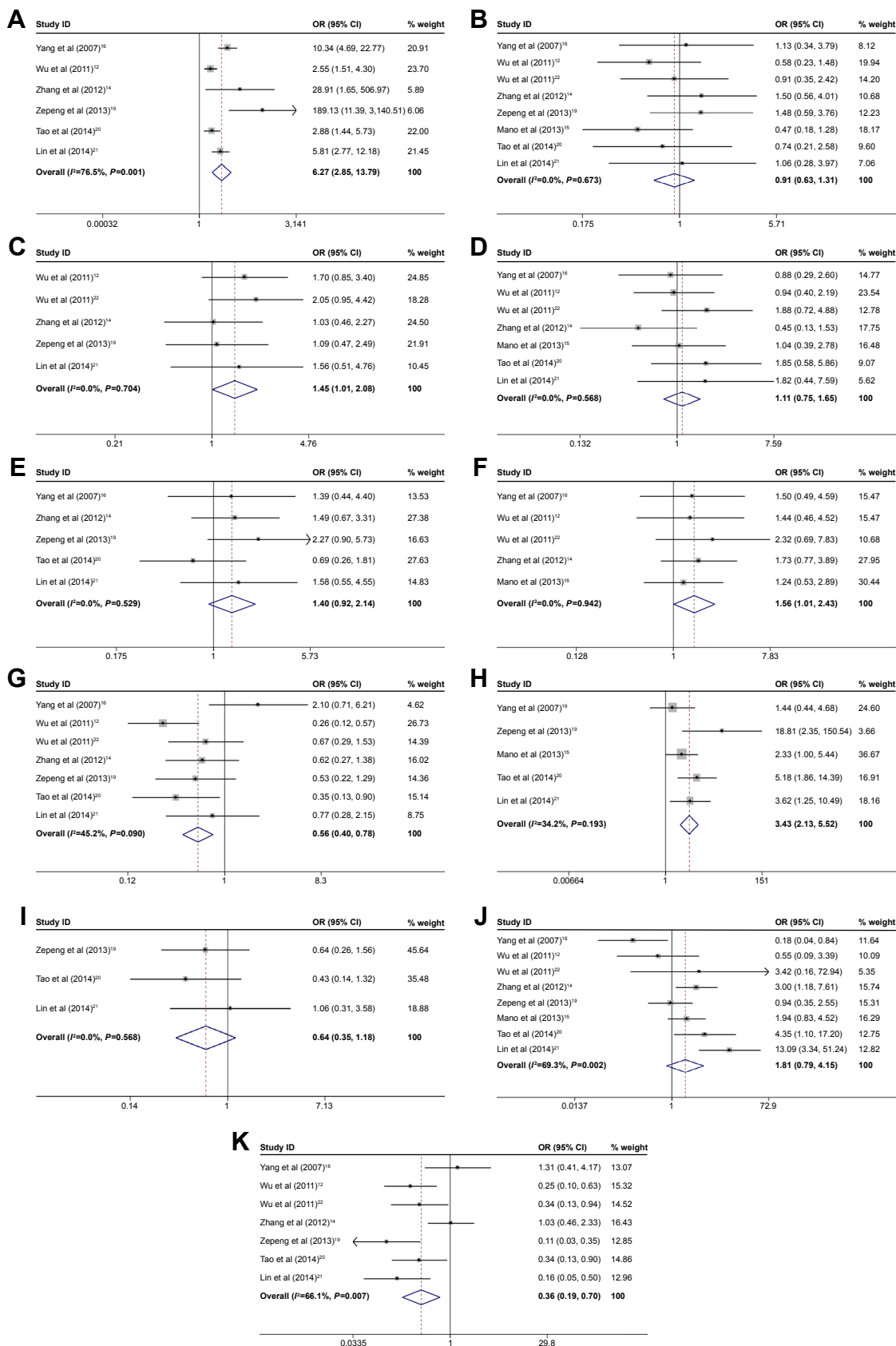


Figure 2 Forest plot of studies evaluating the association between p-STAT3 expression and clinicopathological features including expression (A), gender (B), age (C), HBsAg (D), liver cirrhosis (E), AFP (F), tumor size (G), number of tumors (H), vascular invasion (I), tumor differentiation (J), and TNM stage (K).

Note: Weights are from random-effects analysis.

Abbreviations: AFP, alpha-fetoprotein; HBsAg, hepatitis B surface antigen; OR, odds ratio.

Table 3 Subgroup analysis of expression, differentiation, and TNM stage

Subgroups	Number of studies	Number of patients	Pooled OR (95% CI)	PHet	I ² (%)	P-value
Expression						
Sample size						
≤ 100	4	600	7.70 (2.82–21.00)	0.003	78.1	<0.001
> 100	2	358	5.78 (0.56–59.51)	0.090	65.2	0.140
NOS score						
≤ 7	3	153	5.46 (2.64–11.31)	0.054	65.7	<0.001
> 7	3	437	20.14 (0.26–595.43)	<0.001	87.4	0.082
Differentiation						
Sample size						
≤ 100	4	300	1.78 (0.33–9.54)	<0.001	85.0	0.500
> 100	4	452	2.06 (1.15–3.69)	0.426	0.0	0.015
NOS score						
≤ 7	4	323	2.41 (0.32–18.17)	0.001	82.8	0.393
> 7	4	429	1.59 (0.84–2.99)	0.227	30.9	0.151
TNM stage						
Sample size						
≤ 100	3	300	0.30 (0.11–0.84)	0.016	71.1	0.022
> 100	4	351	0.46 (0.19–1.12)	0.055	65.6	0.088
NOS score						
≤ 7	4	323	0.39 (0.18–0.87)	0.082	55.3	0.021
> 7	3	328	0.32 (0.09–1.17)	0.004	81.8	0.086

Abbreviations: NOS, Newcastle-Ottawa Scale; OR, odds ratio; PHet, probability of heterogeneity.

between p-STAT3 expression and the clinicopathological features and prognosis of patients with HCC is unclear and contentious.

The current meta-analysis of eight studies comprising 752 HCC patients revealed the association of the expression of p-STAT3 with HCC incidence, age, liver cirrhosis, tumor size, vascular invasion, and TNM stage. These findings are evidence of the involvement of p-STAT3 in the development and progression of HCC. Furthermore, p-STAT3 expression was related to a poor 3- and 5-year OS rate as well as the DFS rate, suggesting that p-STAT3 could be a reliable biomarker for the prognosis of patients with HCC.

STAT3 appears to be involved in the tumor inflammatory response, tumor cell transformation, survival, proliferation, angiogenesis, invasion, and metastasis. For instance, Miyatsuka et al²⁹ suggested that the activation of STAT3 in pancreatic cells upregulates the expression of pancreatic-duodenal homeobox 1 (PDX1), leading to the transformation of pancreatic acinar cells to ductal cells, which is the basis of pancreatic cancer. Their results suggested that activation of STAT3 promotes tumor cell transformation. Also, Fan et al³⁰ reported that the expressions of the proapoptotic factors BCL2L1 and BIRC5 were downregulated when STAT3 expression was downregulated in colon cancer cells. Li et al³¹ found that silencing STAT3 can significantly downregulate BCL2 and lead to increased glioma cell apoptosis, suggesting

that STAT3 can protect tumor cells from apoptosis. Masuda et al³² showed that p-STAT3 inhibits the expression of cyclin D1, which contributes to the proliferation of head and neck squamous cell carcinoma. In addition, p-STAT3 expression has been positively correlated to cyclin D1 expression, while the activation of epidermal growth factor receptor in glioma cells can cause excessive activation of STAT3, and the resulting increased expression of cyclin D1 drives the accelerated proliferation of tumor cells.³³ Moreover, STAT3 was found to promote invasion and metastasis of tumor cells by upregulating matrix metalloproteinases (MMP1, MMP2 and MMP9) and overexpression of p-STAT3 was closely related to tumor angiogenesis.^{22,34} Furthermore, p-STAT3 participates in the interaction between tumor cells and the immune system, which leads to the inhibition of the release of some inflammatory mediators, such as tumor necrosis factor (TNF) and nitric oxide (NO), which in turn contributes to a weakened inflammatory response and promotion of the escape of tumor cells from T-cell-mediated killing.^{35,36}

This meta-analysis confirms that p-STAT3 is associated with clinicopathological features and may be a specific biomarker for the prognosis of patients with HCC. Nevertheless, these conclusions must remain tentative due to the many limitations of this meta-analysis. First, most of the included studies were conducted in China, so the results may not be generalizable to non-Chinese or non-Asian individuals. More

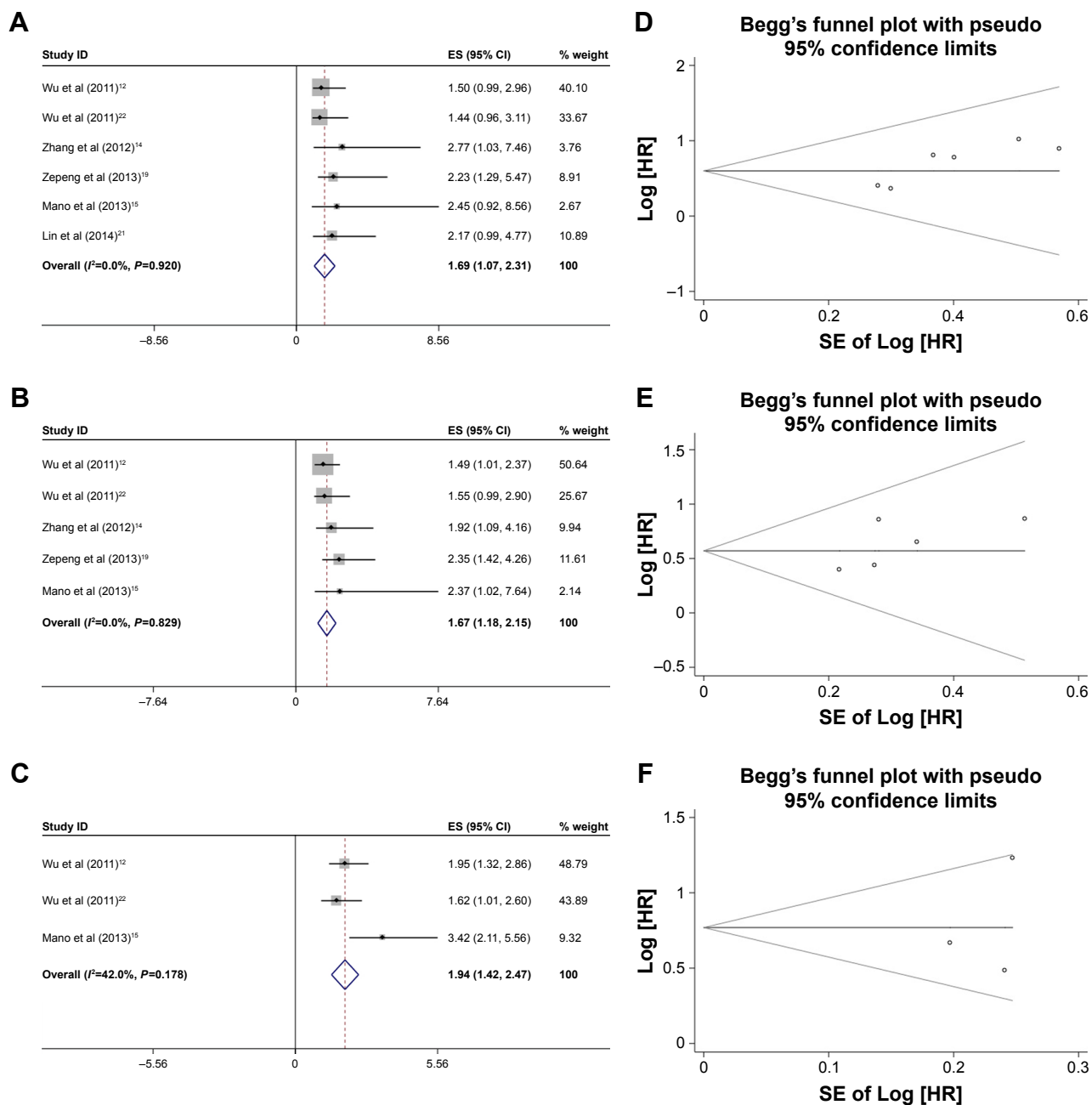


Figure 3 Pooled analysis for the association between p-STAT3 expression and the 3-year OS rate (**A**: Forest plot, **D**: Begg's publication bias plot), the 5-year OS rate (**B**: Forest plot, **E**: Begg's publication bias plot), and the DFS rate (**C**: Forest plot, **F**: Begg's publication bias plot).

Abbreviations: DFS, disease-free survival; HR, hazard ratio; OS, overall survival.

studies will be needed to confirm or refute the present findings in other races and ethnicities. Second, although subgroups' analysis involved sample size and NOS score, there was heterogeneity present and the source of such heterogeneity was unclear. A possible explanation is the experimental methods used in the included studies. For instance, IHC analysis, which was used in all studies, is a semiquantitative research method that can be influenced by the use of different antibodies and cutoff values. Careful inspection of the articles

determined that the source of the p-STAT3 antibody differed between some of the studies (some purchased from Santa Cruz Biotechnology Inc., Dallas, TX, USA, and others from Cell Signaling Technology CST, Danvers, MA, USA). Furthermore, the antibody dilutions used in the IHC procedures were different (1:200, 1:150, 1:100, and 1:50 in the studies that reported the dilution). Additionally, the positive cutoff values differed (19%, 25%, or 50%). These differences likely influenced the outcome of the decisions. High-quality studies

involving large numbers of individuals that utilize the same antibody, antibody dilution, and cutoff value are needed to further support the present findings. Finally, HRs were obtained using the Engauge Digitizer 10.0 software, which may have caused some offset.

Conclusion

Despite the limitations, the results are still significant. The expression of p-STAT3 was associated with HCC incidence, as well as age, liver cirrhosis, tumor size, vascular invasion, and TNM stage in patients with HCC. Additionally,

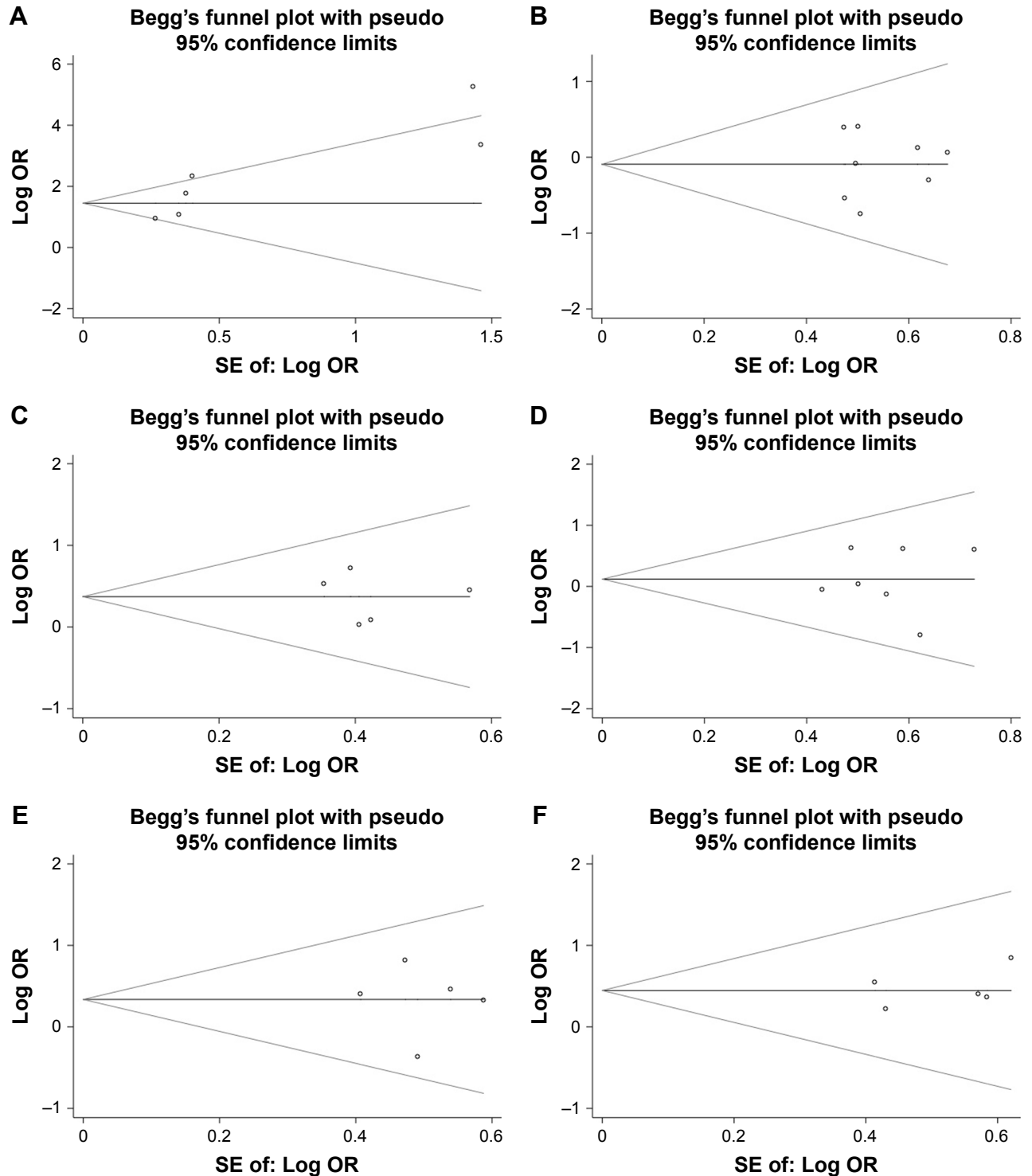


Figure 4 (Continued)

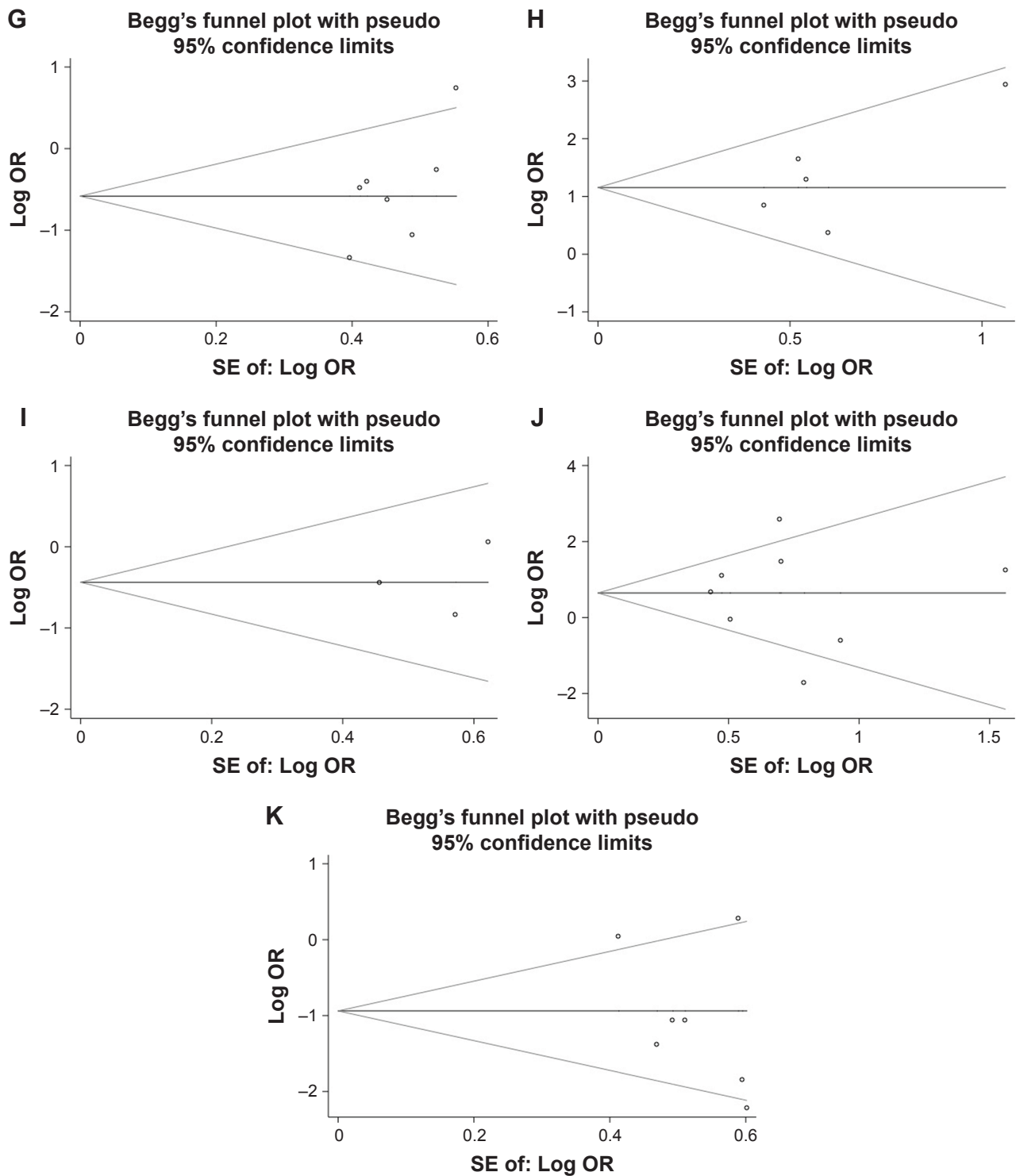


Figure 4 Begg's publication bias plot for the p-STAT3-related studies including expression (A), gender (B), age (C), HBsAg (D), liver cirrhosis (E), AFP (F), tumor size (G), number of tumors (H), vascular invasion (I), differentiation (J), and TNM stage (K).

Abbreviations: AFP, alpha-fetoprotein; HBsAg, hepatitis B surface antigen; OR, odds ratio.

p-STAT3 expression correlated with poor 3- and 5-year OS and DFS. The collective findings implicate p-STAT3 as a reliable biomarker for the prognosis of patients with HCC, pending the results of high-quality studies involving larger numbers of individuals.

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Disclosure

The authors report no conflicts of interest in this work.

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