

## Review Article

# Does tea consumption correlate to risk of fracture? A meta-analysis

An Yan<sup>1\*</sup>, Hao-Hua Zhang<sup>2\*</sup>, Shang-Quan Wang<sup>1</sup>, Yong Zhao<sup>1</sup>

<sup>1</sup>Department of Comprehensive Orthopedics, Wangjing Hospital of China Academy of Chinese Medical Sciences, Beijing 100102, China; <sup>2</sup>Department of Orthopaedics, Beijing Jishuitan Hospital, The Fourth Clinical College of Peking University, Beijing 100035, China. \*Equal contributors.

Received January 11, 2015; Accepted March 17, 2015; Epub June 15, 2015; Published June 30, 2015

**Abstract:** Fractures are important causes of healthy damage and economic loss nowadays. The conclusions of observational studies on tea consumption and fracture risk are still inconsistent. The objective of this meta-analysis is to determine the effect of tea drinking on the risk of fractures. In this study, a comprehensive literature search was conducted in Pubmed, Embase and reference lists of the relevant articles. Observational studies that reported an estimate of the association between tea drinking and incidence of fractures were included. A meta-analysis was conducted by the STATA software. The results indicated that a total of 9 studies involving 147,950 individuals that examined the association between tea consumption and risk of fractures were included in this meta-analysis. The odds ratios (ORs) with 95% confidence intervals (CIs) were pooled using a random-effects model. The pooled OR of 9 observational studies for the tea consumption on risk of fracture was 0.89 (95% CI, 0.78-1.04). In the subgroup analyses, no significant association was detected in neither cohort studies (n = 3; OR, 0.97; 95% CI, 0.89-1.06) nor case-control studies (n = 6; OR, 0.91; 95% CI, 0.70-1.19), respectively. Because of the varied and limited data of the included studies, we are not able to conduct the dose-response meta-analysis. In conclusion, tea consumption might be not associated with the risk of fractures. The following large-sample and well-designed studies are required to confirm the existing conclusions.

**Keywords:** Fracture risk, tea consumption, correlation, meta-analysis

## Introduction

Fractures are important causes of healthy damage and economic loss in the whole world. Several risk factors have been detected including the lower body mass density (BMD) [1] and the increased body mass index (BMI). Besides, the tobacco exposure [2], coffee intake [3], calcium and vitamin D intake [4] and history of fractures are also considered to be associated with the risk of fractures. The realization of the relationship between these modifiable factors and incidence of fractures would provide more effective strategy for the fracture prevention in the future.

Tea is the most popular beverage in the world. Several kinds of antioxidants, including flavonoid, catechin, thearubigin and theaflavin, exist in the tea. Nowadays, the tea drinking behavior is considered to be associated with the devel-

opment and progression of different kinds of diseases. Tea consumption is reported to be a protective factor of the cardiovascular diseases [5], Parkinson's disease [6] and several kinds of cancers [7, 8]. What's more, habitual tea drinking is reported to have a positive effect on the BMD [9, 10], which may thus prevent the incidence of fracture (osteoporotic fracture, in particular). However, tea is a beverage which contains caffeine as well. It was reported that the caffeine could decrease the BMD and subsequently increase the risk of fractures [11]. The contradictory effects of tea drinking make its role on the risk of fractures indistinct. Several epidemiological studies have researched the relationship between tea consumption and incidence of fracture; however, no accordant conclusion was obtained by now [10, 12]. Meta-analysis is a useful statistical tool to pool the relevant studies together and gain a more powerful conclusion. For instance, meta-analysis

## Correlation between tea consumption and fracture risk

was used to detect the risk factors of lung cancer [13], breast cancer [14], type 2 diabetes [15] and thyroid lesions [16]. The purposes of this study were: firstly, quantify the effect of tea consumption on the incidence of fractures by meta-analyzing the existing observational studies and provide quantitative and high-level evidence and secondly, provide some suggestions for the following advanced studies.

### Methods

#### *Search strategy and inclusion criteria*

This current meta-analysis was conducted according to the PRISMA guidelines and MOOSE guidelines [17, 18]. We searched two databases (Pubmed and EMBASE) with the key words “tea”, “green tea”, “black tea”, “flavonoid”, “catechin”, “thearubigin”, “polyphenol”, “theaflavin” in combine with “fracture” through July 8, 2013. No language or any other restrictions were set in the search strategy. The additional articles were obtained through consulting the reference lists of the relevant studies. When different articles from the same dataset were obtained, only the most recent study with available data was included in the meta-analysis. The contact with the corresponding author of certain article was conducted when more data was required.

A study would be included in this meta-analysis when it met the following inclusion criteria: (1) the association between tea consumption and risk of fractures was evaluated; (2) a cohort or case-control study design was obtained; (3) the value of relative risk (RR), odds ratio (OR) with 95% confidence intervals (CI) or enough data to calculate them were reported.

A study would be also excluded in this meta-analysis when it met the following inclusion criteria: (1) the fractures which were caused by the other reasons, such as injury, trauma, and etc. (2) The patients who consumed the tea only for a few days or months, or with intervals.

#### *Data extraction and assessment of study quality*

The data of each included article was extracted by two reviewers independently and any discrepancy was checked again and resolved through discussion. The following data was

extracted from each article: the name of the first author, publishing year, study design, study site, age and gender of participants, type and amount of cases, adjustments of the confounding factors, and the OR/RR value with 95% CI.

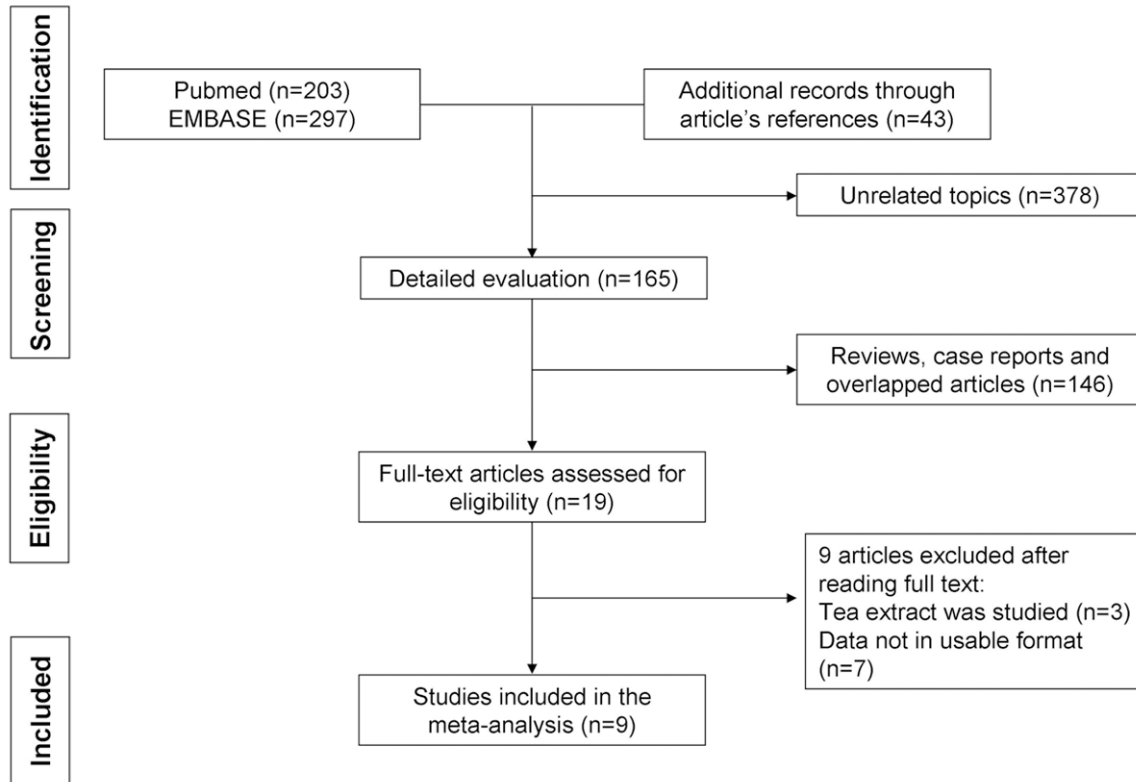
The study quality was assessed by two reviewers back to back and any disagreement was discussed with the third reviewer. All the included studies are cohort or case-control studies. Considering the observational study design of the included studies, the Newcastle-Ottawa Scale (NOS) was obtained to assess the methodological quality of the included studies [19]. It assessed the selection, comparability and exposure of a case-control study, while the selection, comparability and outcome of a cohort study. A maximum of 9 stars was scored for a study and the study with over 6 stars would be regarded as relative high quality.

#### *Statistical methods for the meta-analysis*

Both OR and RR were reported in the included studies and the OR was obtained to approximate RR in this meta-analysis. When both crude and adjusted OR/RR values were offered in the article, only the adjusted ones were adopted for the meta-analysis. A random-effects model was obtained to estimate the pooled effects, thus it would provide a more conservative conclusion. The effects of tea drinking on the incidence of fractures were measured with the OR with 95% CI. The heterogeneity among the included studies was detected by both  $\chi^2$  and  $I^2$  tests. When  $P$  for the heterogeneity  $< 0.1$  and  $I^2 > 50\%$ , the heterogeneity would be considered statistically significant. The source of the statistically significant heterogeneity was explored by both removing the included studies one by one and conducting the subgroup analysis. A cumulative meta-analysis was conducted as well through sorting all the studies by the publishing year.

We examined a potential nonlinear dose-response relationship between tea consumption and fracture risk using restricted cubic splines with three knots at percentiles 25%, 50%, and 75% of the distribution. A two-stage random-effects dose-response meta-analysis tested was adopted for the detection of the potential nonlinear relation. A  $P$  value for nonlinearity was calculated by testing the null hypothesis that the coefficient of the second spline is equal to 0 [20, 21].

## Correlation between tea consumption and fracture risk



**Figure 1.** Flow diagram showing the identification of relevant studies in the meta-analysis. A total of 500 articles (203 from Pubmed and 297 from EMBASE) were identified from the electronic database searching. Besides, 43 more records were identified through consulting the citations of the relevant reviews and articles. A total of 165 records were detailed evaluated and 19 full-text were assessed for eligibility after removing 146 reviews, case reports and overlapped articles. Subsequently 3 articles in which tea extract was studied and 7 ones in which the data not in usable format were excluding from the inclusion and in final, a total of 9 articles were included for the quantitative synthesis.

The sensitivity analyses were conducted to detect the robustness of the conclusions by two independent methods. Firstly, we conducted a one-way sensitivity analysis in which the included study was excluded one by one. Secondly, we excluded the studies with a relative lower methodological quality and assess the effect of tea consumption and risk of fracture. The potential publication bias was estimated by visually evaluation of funnel plot, Bgger's test and Egger's test concurrently [22, 23]. The Stata software package (version 11.0; Stata Corp., College Station, TX) was obtained to conduct the meta-analysis.

### Results

#### *Identification and selection of studies*

A total of 500 articles (203 from Pubmed and 297 from EMBASE) were identified from the

electronic database searching. Besides, 43 more records were identified through consulting the reference lists of the relevant reviews and articles. After excluding 378 articles with unrelated topics, a total of 165 records were detailed evaluated. Among the 165 articles, 19 full-texts were assessed for eligibility after removing 146 articles (reviews, case reports and overlapped articles). Subsequently 3 articles in which tea extract was studied and 7 ones in which the data not in usable format were excluding from the inclusion and in final, a total of 9 studies were included for the quantitative synthesis [24-33] (**Figure 1**).

#### *Study characteristics and quality*

A total of 17,965 cases and 147,950 individuals in 9 observational studies were indemnified in this current meta-analysis. All the detailed characteristics of each included study were

## Correlation between tea consumption and fracture risk

**Table 1.** Baseline of the included studies

Author	Year	Study design	Site	Age (year)	Gender	Type and No. of fractures	No. participants	Categories of tea consumption	Adjustment
Kreiger N [17]	1992	Case-control	Canada	50-84	Female	Hip fracture, 102 Wrist fracture, 154	533	2 categories, < 3 cup/day, ≥ 3 cup/day	Age, Quetelet index, ovariectomy, estrogens, replacement, cigarette, smoking
Johnell O [18]	1995	Case-control	Southern Europe	> 50	Female	Hip fracture, 2086	5618	4 categories, never, 1-2 cup/day, 3 cup/day, ≥ 4 cup/day	Age, center and BMI
Tavani A [19]	1995	Case-control	Italy	19-74	Female	Hip fracture, 279	1340	2 categories, no drinking, drinking	Age, education, body mass index, smoking status, total alcohol consumption, calcium intake, menopausal status, estrogen replacement therapy use.
Chen Z [20]	2003	Cohort	USA	50-79	Female	Hip fracture, 386 forearm/wrist fracture, 1809	91465	4 categories, < 1 cup/day, 1 cup/day, 2-3 cup/day, ≥ 4 cup/day	Age, BMI, ethnicity, hormone replacement therapy use, smoking, years since menopause, fracture history, osteoporosis drug use, walking, soy milk consumption, coffee drinking, depression
Hallstr D [21]	2010	Case-control	India	65.2 ± 15.1	M:F 86:114	Hip fracture, 100	200	2 categories, ≤ 1 cup/day, > 1 cup/day	Age, BMI, agility, eat pander, eat fish, active persons, take calcium supplements.
Jha RM [22]	1999	Case-control	Southern Europe	50-100	Male	Hip fracture, 730	1862	4 categories, never, 1-2 cup/day, 3 cup/day, ≥ 4 cup/day	Age, center and BMI
Kanis J [23]	2013	Case-control	China	71 ± 7	M:F 148:433	Hip fracture and femoral neck fracture, 396	581	2 categories, No drinking, drinking	NA
Zeng FF [24]	2011	Cohort	China	> 90	M:F 226:467	Osteoporotic fracture, 72	703	2 categories, Former drinking, current drinking	Age, gender, sleep habits, educational levels, religion habits, temperament.
Tomata Y [25]	2012	Cohort	Japan	> 65	M:F 6176: 7812	Hip fracture, 55	13988	4 categories, never, 1-2 cup/day, 3 cup/day, ≥ 4 cup/day	NA

## Correlation between tea consumption and fracture risk

**Table 2.** Quality assessment of included studies

Author	Study design	Quality assessment criteria			Overall quality
		Selection	Comparability	Outcome/exposure	
Kreiger N [17]	Case-control	***	**	***	8
Johnell O [18]	Case-control	***	*	**	6
Tavani A [19]	Case-control	***	*	**	7
Chen Z [20]	Cohort	***	*	**	6
HallströmH [21]	Cohort	****	**	***	9
Jha RM [22]	Case-control	**	**	*	5
Kanis J [23]	Case-control	***	**	***	8
Zeng FF [24]	Case-control	***	**	**	7
Tomata Y [25]	Cohort	***	***	**	8

The study quality is evaluated by Downs and Black score and Newcastle-Ottawa Scale (NOS). The NOS were obtained to assess the selection, comparability and exposure of the case-control study, while the selection, comparability and outcome for the cohort study. \*: one point; \*\*: two points; \*\*\*: three points.

presented in **Table 1**. The included studies were published between 1992 and 2013. A total of 3 cohort studies and 6 case-control studies were identified in this current meta-analysis. Among all the included studies, 4 studies were in European, 2 in Americas and 3 in Asia. The age, gender distribution, categories of tea consumption, and adjustments of confounding factors were demonstrated in **Table 1**.

The NOS were obtained to assess the selection, comparability and exposure of the case-control study, while the selection, comparability and outcome of the cohort study. All the scores of each part in the evaluation of all the studies were displayed in **Table 2**. Eight in nine studies were in relative high quality (over 6 stars) and the mean NOS score was 6.90 stars (standard deviation: 1.37).

### *Tea consumption and risk of fractures*

**Figure 2** showed the pooled estimation of the tea consumption and risk of fractures in a random-effects model. In this meta-analysis, tea intake was not associated the incidence of fractures (OR, 0.89; 95% CI, 0.78-1.04). No significant association was detected in neither cohort studies (n = 3; OR, 0.97; 95% CI, 0.89-1.06) nor case-control studies (n = 6; OR, 0.91; 95% CI, 0.70-1.19). When the geographical distributions of the included studies were considered, the studies that conducted in the Europe (n = 4; OR, 0.827; 95% CI, 0.61-1.13), the

Americas (n = 2; OR, 1.01; 95% CI, 0.88-1.1) and the Asia (n = 3; OR, 0.921; 95% CI, 0.68-1.25) showed no statistically significant results. In the two sexual subgroups, no significant association was in neither male group (n = 2; OR, 0.818; 95% CI, 0.62-1.09) nor female group (n = 6, OR, 0.961; 95% CI, 0.77-1.20). When stratified by the fracture types, no significant association between tea consumption and hip fracture (n = 8; OR, 0.897; 95% CI, 0.74-1.09) nor humeral fracture (n = 2; OR, 0.943; 95% CI, 0.71-

1.25) was detected. All the results of the subgroup analysis were presented in **Table 3**.

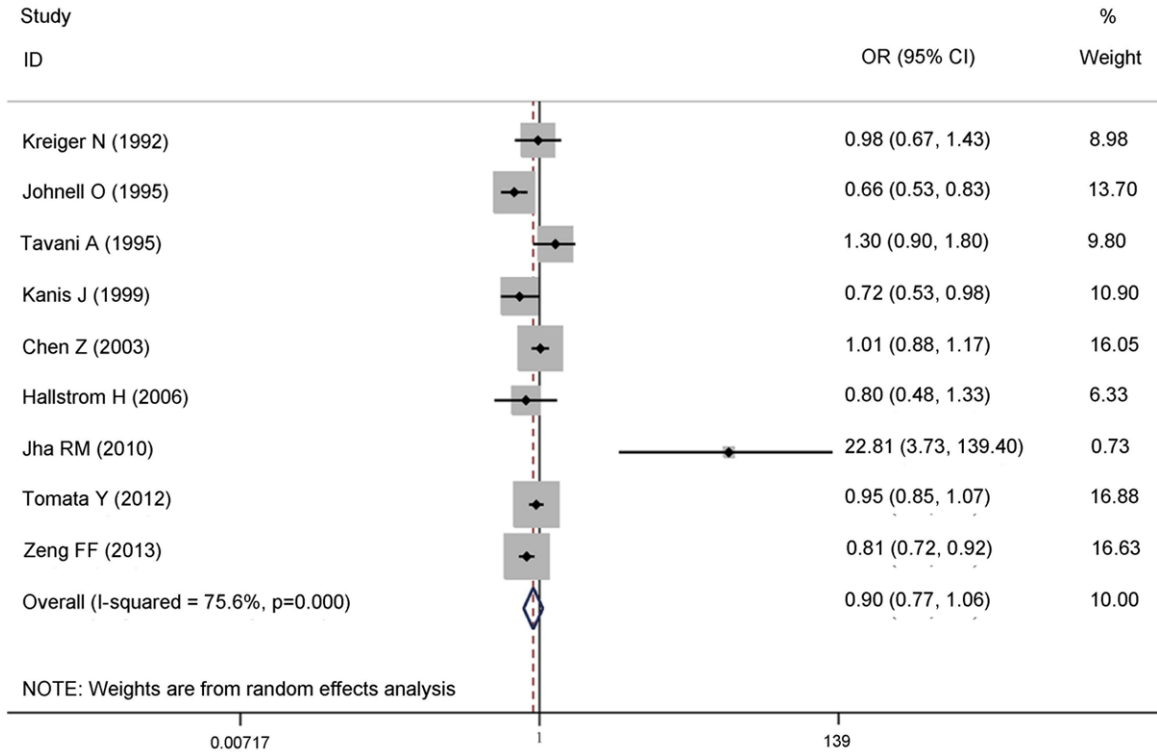
A cumulative meta-analysis was conducted as well. The cumulative effect was estimated after sorting the included studies by the publishing date. **Figure 3** demonstrated the forest plot of the cumulative effect of tea intake and risk of fractures. It could be detected that, from 1992 to 2013, the tea consumption was inversely associated with the risk of fractures; however, the association was not statistically significant constantly.

### *Heterogeneity and sensitivity analysis*

The heterogeneity was significant when all the 9 studies were pooled in the meta-analysis ( $I^2$ , 73.6%;  $P < 0.001$ ). We tried to explore the source by excluding the included studies one by one and re-count the heterogeneity; however, no single article could explain the source of the heterogeneity. The subgroup analysis by study design, study site, gender and fracture type demonstrated no satisfactory results in exploring the source of heterogeneity.

A one-way sensitivity analysis was conducted and **Figure 3** showed the results. No significant influence was detected after any single article excluded from the meta-analysis. Besides, we just included the articles with a relative high quality (over 6 stars NOS score in the meta-analysis); however, no significant association

## Correlation between tea consumption and fracture risk



**Figure 2.** Forest plot of risk estimates of the association between tea drinking and risk of fracture. In a random-effects meta-analysis, tea intake was not associated the incidence of fracture (OR, 0.89; 95% CI, 0.78-1.04). A significant heterogeneity was observed when all the 9 studies were pooled ( $I^2$ , 73.6%;  $P < 0.001$ ).

**Table 3.** Subgroup analysis of tea consumption and fracture risk with combined OR

	Subgroups	No. of studies	Summary effect		Heterogeneity		
			OR	(95% CI)	<i>p</i> Value	$I^2$ , %	<i>p</i> Value
Study design	Cohort	3	0.97	0.89-1.06	0.486	1.05	0.591
	Case-control	6	0.91	0.70-1.19	0.499	80.1	< 0.001
	Europe	4	0.827	0.61-1.13	0.232	72.4	0.012
Site	Americas	2	1.01	0.88-1.1	0.52	0	0.857
	Asia	3	0.921	0.68-1.25	1.41	82.2	0.001
Gender	Male	2	0.818	0.62-1.09	0.168	29.9	0.232
	Female	6	0.961	0.77-1.20	0.726	70	0.005
Fracture type	Hip fracture	8	0.897	0.74-1.09	0.276	75.8	< 0.001
	Humeral fracture	2	0.943	0.71-1.25	0.679	0	0.889

OR: odds ratio. The results with statistical significance were marked in bold.

was detected neither (OR, 0.88; 95% CI, 0.78-1.00). A significant heterogeneity should be noted as well ( $I^2 = 65.9%$ ,  $P = 0.005$ ).

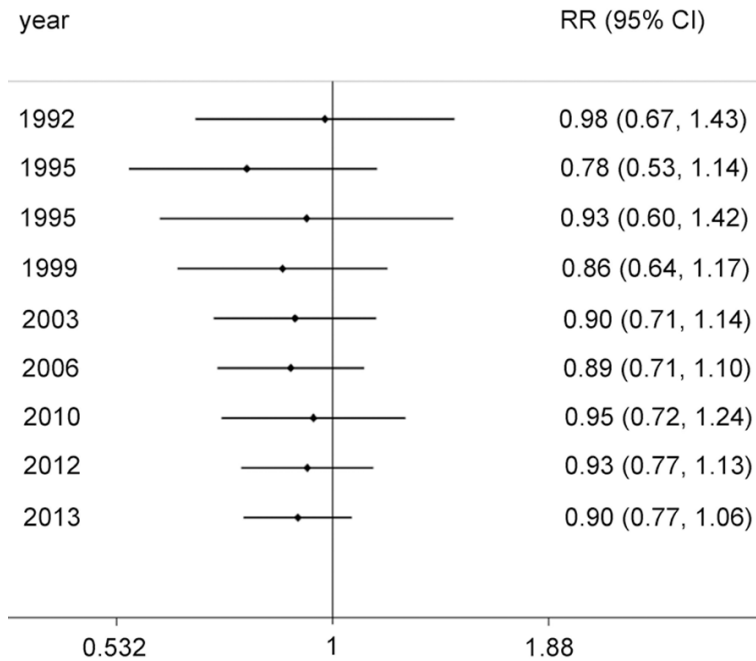
### Dose-response meta-analysis

Then, we assessed the dose-response relationship between tea consumption and the risk of

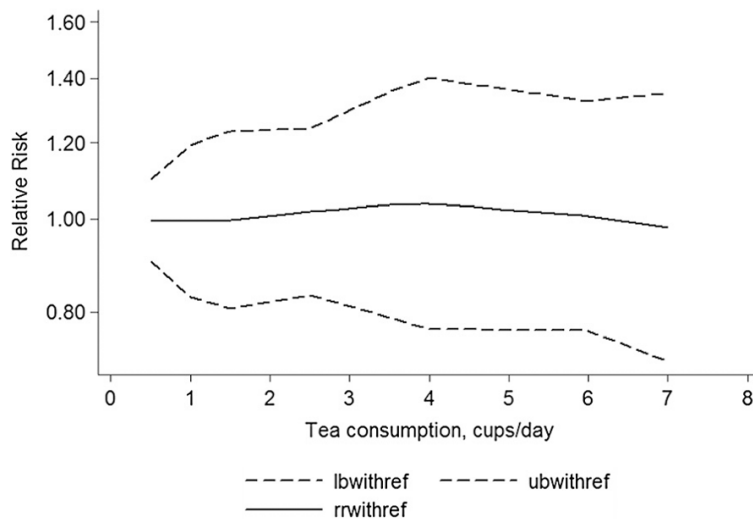
fracture progression. We found no obvious evidence of statistical significance from linearity ( $P = 0.983$ ). **Figure 4** demonstrated the dose-response relationship between tea consumption and risk of fracture.

Both Begg's graph and Egger's test were obtained to detect the potential publishing

## Correlation between tea consumption and fracture risk



**Figure 3.** Forest plot of the cumulative meta-analysis. From 1992 to 2013, the tea consumption was inversely associated with the risk of fracture, however, the association was not statistical significant constantly.



**Figure 4.** Dose-response relationship between tea intake and fracture risk.

bias. No significant publication bias was found in this current meta-analysis (funnel plot, symmetrical; Begg's test,  $P = 0.474$ ; Egger's test,  $P = 0.627$ ).

### Discussion

In this current meta-analysis, a total of 9 studies with 147,950 individuals were included. The

quantitative synthesis of these observational studies demonstrated that tea consumption showed neither protective nor harmful effect on the incidence of fractures. Meanwhile, the subgroup analysis by the study designs, study sites, genders and fracture types displayed no statistical significant results. The cumulative meta-analysis showed that an inverse but not significant association between tea intake and risk of fractures since 1992. No significant association was detected in the dose-response meta-analysis. Fractures are producing more and more physical illness and economic loss in the whole world. The studies on the detection of the risk factors of the fractures would provide us better knowledge on the prevention of the incidence of fractures. Several lifestyles and dietary factors, such as the tobacco exposure, coffee intake, calcium and vitamin D intake, were associated with the risk of fracture as mentioned above. Tea is regarded as the most frequently consumed beverage and the protective or harmful effects on the health are substantial public issues. As a modifiable factor, the understanding of the relationship between tea drinking and risk of fractures would provide potentially effective preventative measures of the fractures.

The association between tea consumption and BMD has been studied in several studies. However, PA collected the physical and lifestyle data from 62 postmenopausal women and the intake of tea was reported to be associated with increased BMD measurements [34]. Hegarty VM et al. compared the BMD measurements of 1,134 tea drinkers and 122 non-tea drinkers; and higher BMD was detected in the

## Correlation between tea consumption and fracture risk

tea drinkers [35]. However, as a beverage that contains caffeine, tea drinking was also considered to be risk factors of bone loss. In animal models, high doses of caffeine could induce calcium loss and influence the normal development of bone [36]. Nowadays tea, regarded as a beverage with multiple nutrients and caffeine, is a positive factor in the bone health [37].

Several epidemiological studies were also conducted to study the association and risk of fractures directly. A cross-sectional study in Sichuan China observed the association between habits of smoking, alcohol consumption, tea consumption and exercise and osteoporotic fracture among very old people. Among all the lifestyles, tea consumption was reported to be unrelated with the incidence of osteoporotic fracture [32]. Besides, the observational studies, including cohort and case-control studies, were conducted as well to detect the etiological effect of tea consumption on the incidence of fractures. In this current meta-analysis, we studied the association between tea drinking and risk of fractures. Through pooling the published observational studies together, we found that tea consumption was not associated with the risk of fractures. This result was similar with the most included studies. Among all the included studies, only a case-control study demonstrated that tea drinking was a risk factor of the incidence of fracture [29]. Through the comparison of 100 cases and 100 age and sex matched controls, tea drinkers was reported to have a higher risk of hip fractures (OR, 22.8; 95% CI, 3.73-139.43). The ethnic and regional differences might explain part of the result differences. However, the relative smaller amount of participants would result in more unsteadiness of the result. Contrarily, three case-control studies demonstrated that tea consumption produced a protective effect on the incidence of fracture [25, 30, 31]. However, these studies might not be powerful enough to influence the general conclusion in this meta-analysis.

In the subgroup analysis by the study designs, no association was detected between tea consumption and risk of fractures in cohort study group and case-control study group. The cohort study design could avoid the potential recall bias and would provide more credible conclusions. In our meta-analysis, the results of the

subgroups from both study designs were accordant and thus the conclusion was quite substantial. Besides, the sensitivity analysis when the studies with high quality were included showed a slight but not significant result (OR, 0.88; 95% CI, 0.78-1.00). Moreover, the cumulative meta-analysis showed that the association between tea drinking and incidence of fracture was not statistically significant; however, a potential trend that indicated the protective effect from tea consumption couldn't be ignored.

The data about the tea or tea extract on the bone metabolism is abundant. The tea contains several kinds of antioxidants, including flavan-3-ols, flavanols and flavones, and these polyphenol produced multiple effects on the bone metabolism. At a low concentration, polyphenol would increase osteoblast proliferation, while at a high concentration; polyphenol would inhibit its proliferation inversely [37]. Oka Y et al. reported that polyphenols, particularly epigallocatechin-3-gallate (EGCG), which were extracted from the green tea, could inhibit MMPs expression and activity. Meanwhile the tea polyphenols could inhibit osteoclast formation and differentiation in vitro [38]. Choi EM reported that treatment with catechin increased alkaline phosphatase activity which could indicate the activity of osteoblastic and reduced osteoblastic apoptosis induced by the tumor necrosis factor (TNF) [39]. Oxidative stress is regarded as a pivotal physiopathological process in the age-related bone loss [40]. Das AS et al. [41] showed that the black tea extract would protect the mononuclear cells from the oxidative stress and thus slowed the progression of bone loss. Apart from the polyphenol, the other contents in tea, such as fluorine and caffeine, also play important roles in the bone health [37, 42].

To our best knowledge, this is the first meta-analysis involving the relationship between tea consumption and risk of fracture. There are some strengths of this work. Firstly, a comprehensive literature search and advanced consulting the relevant references make available included in this meta-analysis. Secondly, the included studies are mainly of high quality and the sensitivity analysis showed that the results are quite tough, and thus the conclusions are credible.



However, there are some limitations could not be ignored in this study. Firstly, most of the studies followed a case-control study design, and therefore there were recall and selection bias which are inherent to retrospective studies. Even through the subgroup analyses by the study designs were conducted, the efficiency was limited by the absence of enough cohort studies. Secondly, tea is mainly grouped into green tea and black tea. The functional contents and effects were not identical for their two tea subtypes and thus the potential bias should be noted. The following epidemiological might need to take this into consideration.

In conclusion, tea consumption might not be associated with the risk of fracture. However, these conclusions should be considered with cautions because the inherent limitations in this meta-analysis. More large-sample and well-designed studies are required to confirm the existing conclusions or detect the covered significant association in this meta-analysis.

### Acknowledgements

This study was granted by the project of publication of series books (No. Z131110001913-154).

### Disclosure of conflict of interest

None.

**Address correspondence to:** Dr. Yong Zhao, Department of Comprehensive Orthopedics, Wangjing hospital, Huajiadi Road, Chaoyang District, Beijing 100102, China. Tel: +86-010-84739952; Fax: +86-010-84739952; E-mail: yongzhaobj@yeah.net

### References

- [1] Liu C, Kuang L, Wang L, Tian J. Management of combination fractures of the atlas and axis: a report of four cases and literature review. *Int J Clin Exp Med* 2014; 7: 2074-2080.
- [2] Johnston P, Gurusamy KS, Parker MJ. Smoking and hip fracture; a study of 3617 cases. *Injury* 2006; 37: 152-156.
- [3] Jokinen H, Pulkkinen P, Korpelainen J, Heikkinen J, Keinanen-Kiukaanniemi S, Jamsa T, Korpelainen R. Risk factors for cervical and trochanteric hip fractures in elderly women: a population-based 10-year follow-up study. *Calcif Tissue Int* 2010; 87: 44-51.
- [4] Moyer VA, US Preventive Services Task Force. Vitamin D and calcium supplementation to prevent fractures in adults: U.S. Preventive Services Task Force recommendation statement. *Ann Intern Med* 2013; 158: 691-696.
- [5] Yang CS, Hong J. Prevention of chronic diseases by tea: possible mechanisms and human relevance. *Annu Rev Nutr* 2013; 33: 161-181.
- [6] Kandinov B, Giladi N, Korczyn AD. Smoking and tea consumption delay onset of Parkinson's disease. *Parkinsonism Relat Disord* 2009; 15: 41-46.
- [7] Beliveau R, Gingras D. Green tea: prevention and treatment of cancer by nutraceuticals. *Lancet* 2004; 364: 1021-1022.
- [8] Yang CS, Wang X, Lu G, Picinich SC. Cancer prevention by tea: animal studies, molecular mechanisms and human relevance. *Nat Rev Cancer* 2009; 9: 429-439.
- [9] Hamdi Kara I, Aydin S, Gemalmaz A, Aktürk Z, Yaman H, Bozdemir N, Kurdak H, Sitmapinar K, Devran Sencar I, Başak O, Akdeniz M, İşildar H, Burgut E, Özcan S, Akça U, Dağdeviren N, Ungan M. Habitual tea drinking and bone mineral density in postmenopausal Turkish women: investigation of prevalence of postmenopausal osteoporosis in Turkey (IPPOT Study). *Int J Vitam Nutr Res* 2007; 77: 389-397.
- [10] Shen CL, Yeh JK, Cao JJ, Chyu MC, Wang JS. Green tea and bone health: evidence from laboratory studies. *Pharmacol Res* 2011; 64: 155-161.
- [11] Song QH, Zhang QH, Xu RM, Ma M, Zhao XP, Shen GQ, Guo YH, Wang Y. Effect of Tai-Chi exercise on lower limb muscle strength, bone mineral density and balance function of elderly women. *Int J Clin Exp Med* 2014; 7: 1569-1576.
- [12] Chang JR, Koo E, Agron E, Hallak J, Clemons T, Azar D, Sperduto RD, Ferris FL, Chew EY, Age-Related Eye Disease Study Group. Risk factors associated with incident cataracts and cataract surgery in the Age-related Eye Disease Study (AREDS): AREDS report number 32. *Ophthalmology* 2011; 118: 2113-2119.
- [13] Xu Z, Yu L, Zhang X. Association between the hOGG1 Ser326Cys polymorphism and lung cancer susceptibility: a meta-analysis based on 22,475 subjects. *Diagn Pathol* 2013; 8: 144.
- [14] Qin X, Peng Q, Qin A, Chen Z, Lin L, Deng Y, Xie L, Xu J, Li H, Li T, Li S, Zao J. Association of COMT Val158Met polymorphism and breast cancer risk: an updated meta-analysis. *Diagn Pathol* 2012; 7: 136.
- [15] Wang L, Teng Z, Cai S, Wang D, Zhao X, Yu K. The association between the PPARgamma2 Pro12Ala polymorphism and nephropathy susceptibility in type 2 diabetes: a meta-analysis based on 9,176 subjects. *Diagn Pathol* 2013; 8: 118.

## Correlation between tea consumption and fracture risk

- [16] de Matos LL, Del Giglio AB, Matsubayashi CO, de Lima FM, Del Giglio A, da Silva Pinhal MA. Expression of CK-19, galectin-3 and HBME-1 in the differentiation of thyroid lesions: systematic review and diagnostic meta-analysis. *Diagn Pathol* 2012; 7: 97.
- [17] Moher D, Liberati A, Tetzlaff J, Altman DG. Preferred reporting items for systematic reviews and meta-analyses: the PRISMA statement. *BMJ* 2009; 339: b2535.
- [18] Stroup DF, Berlin JA, Morton SC, Olkin I, Williamson GD, Rennie D, Moher D, Becker BJ, Sipe TA, Thacker SB. Meta-analysis of observational studies in epidemiology: a proposal for reporting. Meta-analysis of Observational Studies in Epidemiology (MOOSE) group. *JAMA* 2000; 283: 2008-2012.
- [19] Stang A. Critical evaluation of the Newcastle-Ottawa scale for the assessment of the quality of nonrandomized studies in meta-analyses. *Eur J Epidemiol* 2010; 25: 603-605.
- [20] Larsson SC, Orsini N, Wolk A. Vitamin B6 and risk of colorectal cancer: a meta-analysis of prospective studies. *JAMA* 2010; 303: 1077-1083.
- [21] Harrell FE Jr, Lee KL, Pollock BG. Regression models in clinical studies: determining relationships between predictors and response. *J Natl Cancer Inst* 1988; 80: 1198-1202.
- [22] Begg CB, Mazumdar M. Operating characteristics of a rank correlation test for publication bias. *Biometrics* 1994; 50: 1088-1101.
- [23] Egger M, Davey Smith G, Schneider M, Minder C. Bias in meta-analysis detected by a simple, graphical test. *BMJ* 1997; 315: 629-634.
- [24] Kreiger N, Gross A, Hunter G. Dietary factors and fracture in postmenopausal women: a case-control study. *Int J Epidemiol* 1992; 21: 953-958.
- [25] Johnell O, Gullberg B, Kanis JA, Allander E, Elffors L, Dequeker J, Dilsen G, Gennari C, Vaz LA, Lyritis G, Mazzuoli G, Miravet L, Passeri M, Cano PR, Rapado A, Ribot C. Risk factors for hip fracture in European women: the MEDOS Study. *Mediterranean Osteoporosis Study. J Bone Miner Res* 1995; 10: 1802-1815.
- [26] Tavani A, Negri E, La Vecchia C. Coffee intake and risk of hip fracture in women in northern Italy. *Prev Med* 1995; 24: 396-400.
- [27] Chen Z, Pettinger MB, Ritenbaugh C, LaCroix AZ, Robbins J, Caan BJ, Barad DH, Hakim IA. Habitual tea consumption and risk of osteoporosis: a prospective study in the women's health initiative observational cohort. *Am J Epidemiol* 2003; 158: 772-781.
- [28] Hallstrom H, Wolk A, Glynn A, Michaelsson K. Coffee, tea and caffeine consumption in relation to osteoporotic fracture risk in a cohort of Swedish women. *Osteoporos Int* 2006; 17: 1055-1064.
- [29] Jha RM, Mithal A, Malhotra N, Brown EM. Pilot case-control investigation of risk factors for hip fractures in the urban Indian population. *BMC Musculoskelet Disord* 2010; 11: 49.
- [30] Kanis J, Johnell O, Gullberg B, Allander E, Elffors L, Ranstam J, Dequeker J, Dilsen G, Gennari C, Vaz AL, Lyritis G, Mazzuoli G, Miravet L, Passeri M, Perez Cano R, Rapado A, Ribot C. Risk factors for hip fracture in men from southern Europe: the MEDOS study. *Mediterranean Osteoporosis Study. Osteoporos Int* 1999; 9: 45-54.
- [31] Zeng FF, Wu BH, Fan F, Xie HL, Xue WQ, Zhu HL, Chen YM. Dietary patterns and the risk of hip fractures in elderly chinese: a matched case-control study. *J Clin Endocrinol Metab* 2013; 98: 2347-2355.
- [32] Wang L, Liu C, Zhao QH, Tian JW. Classification and surgical management for the axis fracture complicated with adjacent segment instability. *Int J Clin Exp Med* 2014; 7: 657-664.
- [33] Tomata Y, Kakizaki M, Nakaya N, Tsuboya T, Sone T, Kuriyama S, Hozawa A, Tsuji I. Green tea consumption and the risk of incident functional disability in elderly Japanese: the Ohsaki Cohort 2006 Study. *Am J Clin Nutr* 2012; 95: 732-739.
- [34] Hoover PA, Webber CE, Beaumont LF, Blake JM. Postmenopausal bone mineral density: relationship to calcium intake, calcium absorption, residual estrogen, body composition, and physical activity. *Can J Physiol Pharmacol* 1996; 74: 911-917.
- [35] Hegarty VM, May HM, Khaw KT. Tea drinking and bone mineral density in older women. *Am J Clin Nutr* 2000; 71: 1003-1007.
- [36] Huang TH, Yang RS, Hsieh SS, Liu SH. Effects of caffeine and exercise on the development of bone: a densitometric and histomorphometric study in young Wistar rats. *Bone* 2002; 30: 293-299.
- [37] Dew TP, Day AJ, Morgan MR. Bone mineral density, polyphenols and caffeine: a reassessment. *Nutr Res Rev* 2007; 20: 89-105.
- [38] Oka Y, Iwai S, Amano H, Irie Y, Yatomi K, Ryu K, Yamada S, Inagaki K, Oguchi K. Tea polyphenols inhibit rat osteoclast formation and differentiation. *J Pharmacol Sci* 2012; 118: 55-64.
- [39] Choi EM, Hwang JK. Effects of (+)-catechin on the function of osteoblastic cells. *Biol Pharm Bull* 2003; 26: 523-526.
- [40] Banfi G, Iorio EL, Corsi MM. Oxidative stress, free radicals and bone remodeling. *Clin Chem Lab Med* 2008; 46: 1550-1555.
- [41] Das AS, Mukherjee M, Das D, Mitra C. Protective action of aqueous black tea (*Camellia si-*

## Correlation between tea consumption and fracture risk

nensis) extract (BTE) against ovariectomy-induced oxidative stress of mononuclear cells and its associated progression of bone loss. *Phytother Res* 2009; 23: 1287-1294.

[42] Aslan C, Melikoglu C, Ocal I, Saglam G, Sutcu R, Hosnuter M. Effect of epigallocatechin gal-

late on ischemia-reperfusion injury: an experimental study in a rat epigastric island flap. *Int J Clin Exp Med* 2014; 7: 57-66.