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**RESEARCH ARTICLE** 

# Prevalence of hypertension and its determinants in Ethiopia: A systematic review and meta-analysis

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## Abstract

## Introduction

Hypertension is a major public health problem globally and it is a leading cause of death and disability in developing countries. This review aims to estimate the pooled prevalence of hypertension and its determinants in Ethiopia.

## Methods

A systematic literature search was conducted at the electronic databases (PubMed, Hinari, and Google Scholar) to locate potential studies. Heterogeneity between studies checked using Cochrane Q test statistics and I<sup>2</sup> test statistics and small study effect were checked using Egger's statistical test at 5% significance level. Sensitivity analysis was checked. A random-effects model was employed to estimate the pooled prevalence of hypertension and its determinants in Ethiopia.

## Results

In this review, 38 studies that are conducted in Ethiopia and fulfilled the inclusion criteria with a total number of 51,427 study participants were reviewed. The overall pooled prevalence of hypertension in the country was 21.81% (95% CI: 19.20–24.42,  $I^2 = 98.35\%$ ). The result of the review also showed that the point of prevalence was higher among males (23.21%) than females (19.62%). When we see the pervasiveness of hypertension from provincial perspective; the highest prevalence of hypertension was observed in Addis Ababa (25.35%) and the lowest was in Tigray region (15.36%). In meta-regression analysis as the mean age increases by one year, the likelihood of developing hypertension increases by a factor of 0.58 times ( $\beta = 0.58$ , 95% CI: 0.31–0.86,  $R^2 = 36.67$ ). Male sex (OR = 1.29, 95% CI: 1.03–1.61,  $I^2 = 81.35\%$ ), age > 35 years (OR = 3.59, 95% CI: 2.57–5.02,  $I^2 = 93.48\%$ ), overweight and/or obese (OR = 3.34, 95% CI: 2.12–5.26,  $I^2 = 95.41\%$ ), khat chewing (OR = 1.42, 95% CI:  $I^2 = 62\%$ ), alcohol consumption (OR = 1.50, 95% CI: 1.21–1.85,  $I^2 = 64\%$ ), family history of hypertension (OR = 2.56, 95% CI: 1.64–3.99,  $I^2 = 83.28\%$ ), and

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family history of diabetes mellitus (OR = 3.69, 95% CI: 1.85-7.59,  $I^2 = 89.9\%$ ) are significantly associated with hypertension.

## Conclusion

Hypertension is becoming a major public health problem in Ethiopia. Nearly two out of ten individuals who are older than 18 years living with hypertension. Sex, age, overweight and/ or obese, khat chewing, alcohol consumption, and family history of hypertension and diabetes mellitus are statistically significant determinant factors for hypertension in Ethiopia. Primary attention should be given for behavioral risk factors to tackle the alarming increase of hypertension in Ethiopia.

## Introduction

Globally, more than 1.13 billion people living with hypertension, of this two-thirds living in Low and Middle-Income Countries (LMICs) [1]. In the globe, by the end of 2025 1.56 billion people will live with hypertension [2]. In Africa, 46% of adults whose age is older than 25 years and above living with hypertension [3]. The prevalence of hypertension in Africa has raised from 19.7% in 1990 to 30.8% in 2010 [4]. One in every five people live with hypertension in LMICs and studies showed that 3 out of 4 people in these countries will live with hypertension by the end of 2025 [5]. Besides, 74.7 million people living with hypertension in Sub-Saharan Africa, and it will rise to 125.5 million by the end of 2025 [6]. These trends have been strongly linked with lifestyle changes such as an increase in smoking tobacco use, excessive alcohol consumption, and physical inactivity [7, 8]. To tackle the burden of hypertension, the Pan-African Society of Cardiology (PASCAR) identified 10 action points to be implemented by African ministers to achieve a 25% decline by the end of 2025 [7, 9].

In Ethiopia, non-communicable diseases account for 39% of all causes of mortality of which, cardiovascular disease accounts for 16% [10]. On the other hand, hypertension constitutes the majority (62.3%) of all the causes of cardiovascular-related morbidity and mortality [11]. This is because high blood pressure increases the risk of life-threatening complications on vital organs like heart, blood vessels, brain, and kidney which leads to premature mortality and disability [12].

In 2015, a systematic review and meta-analysis was conducted in Ethiopia [13]; but this study did not identify the pooled effects of factors affecting the prevalence of hypertension. Besides, there are several studies published after the previous review. Therefore, this systematic review and meta-analysis gives updated pooled prevalence and factors affecting the prevalence of hypertension in Ethiopia.

Moreover, documenting the updated pooled prevalence and its determinants of hypertension will help to achieve the action plan of the Pan-African Society of Cardiology and global targets regarding hypertension. Therefore, the objective of this systematic review and metaanalysis is to synthesize updated pooled prevalence and its determinants of hypertension in Ethiopia. The finding of this review will show the trends of hypertension in Ethiopia and that can be used for health planners, policymakers, and for the community itself to curve the alarming rise of hypertension in Ethiopia.

## Methods

#### Study setting and search strategies

Ethiopia is found in the horn of Africa and has nine administrative regional states and two city administrations. Potential studies were identified using electronic databases (PubMed/MED-LINE, Hinari, Google scholar) and google search. Besides, unpublished theses were also reviewed out from some research centers and library sources. The sources are reviewed limited to English language and studies published after 01/01/2000. The task of searching sources was carried out from all stated electronic databases performed on October/24/2019. All included studies defined hypertension as Systolic Blood Pressure (SBP)  $\geq$  140 mmHg and/or a Diastolic Blood Pressure (DBP)  $\geq$  90 mmHg or known hypertensive patients on treatment. The search MeSH headings were hypertension and synonyms for hypertension were used. The synonyms of hypertension are "blood pressure, high", "and blood pressures, high", "high blood pressure", and "high blood pressures". Finally, the search combination used as; "Hypertension" OR "Blood Pressure, High" OR "High Blood Pressure" OR "High Blood Pressures" OR "Blood Pressures, High" AND Ethiopia (S1 Table).

## **Eligibility criteria**

We used CoCoPop (Condition, Context, and Population) approach for prevalence studies to declare inclusion and exclusion criteria.

#### Inclusion criteria and exclusion criteria

Studies conducted on the prevalence and/or associated factors of hypertension in Ethiopia were included. Besides, all full-text articles written in English language (with response rate > 85%), with participants older than 18 years and published after January 01/2000 are included for this review. Studies conducted on pregnancy-induced hypertension, for the reason that has no prevalence report on hypertension, and hypertension prevalence reports on other comorbidities excluded for this review.

#### Measurement of the outcome variable

The primary outcome of interest for this review was to estimate the pooled prevalence of hypertension and its determinants. Potentials of extracted factors from each study considered as an independent factor for hypertension.

#### Study selection and data collection

All the studies reviewed through different electronic databases were combined, exported, and managed using Endnote version X9.2 (Thomson Reuters, Philadelphia, PA, USA) software. All duplicate studies were removed and full-text studies downloaded using Endnote software and manually. The eligibility of each study was completely assessed independently by two reviewers (SA. &YA.). Exaggerated differences in the results of the two reviewers narrowed through discussion and other reviewer members (ST. & DA.).

#### Assessment of the quality of the individual studies

The quality of the studies assessed using the validated modified version of a quality assessment tool for prevalence studies [14]. Two reviewers (SA. & YA.) were independently assessed to check the quality of the included studies. The problem of subjectivities between the two reviewers was solved through discussion and other review teams (ST. & DA.). The quality

assessment tool has nine-questions. Based on the score of the quality assessment tool the highest score had the minimum risk of bias. Overall scores range from (0-3), (4-6), and (7-9), which are declared low, moderate, and high risk of bias respectively [14].

#### Data extraction and management

All-important parameters extracted from each study were reviewed by two authors (SA. &YA.) independently using Microsoft Excel. The discrepancies between the two authors managed through discussion and/or the other authors (ST. & DA.). The data extraction format was prepared using the assistance of the Joanna Briggs Institute (JBI) data extraction tool for prevalence studies. For each study, authors, years of publication, study design, sample size, the prevalence of hypertension with their standard error, and determinant factors effect size with their standard error were extracted.

#### Statistical analysis

The extracted data were exported to STATA/MP version 16.0 software for analysis. The pooled prevalence of hypertension and its determinants analyzed by the random effects model using DerSimonian-Laird model weight [15]. Heterogeneity in meta-analysis is mostly inevitable due to differences in study quality, its sample size, method, and different outcome measurements across studies [16, 17]. Statistically, significant heterogeneity was checked by Cochrane Q-test and I<sup>2</sup> statistics [18]. To minimize the variance of estimated points between primary studies, a subgroup analysis was carried out in reference to the regions, age categories, and residence. Besides, a sensitivity analysis was also conducted to determine the influence of single studies on the pooled estimates. Univariate meta-regression conducted using year of publication, the mean age of the respondent from primary studies, sample size, and region using random effects model. Publication bias (small study effect) checked using graphically and Egger's statistical test [19]. Statistically significant Egger's test (P-value < 0.05) indicates that the presence of a small study effect and handled by non-parametric trim and fill analysis using the random effects model [20].

## Results

#### Study selection and identification

Of the 784 studies reviewed, 336 were excluded, because they were duplications. By reading their titles and abstracts, 406 studies excluded as they were irrelevant for this review. Again, five studies excluded, because of the outcome not reported, inadequate sample size, and lack of full text. Finally, 38 potential studies have been included for qualitative and quantitative synthesis influences as summarized in the PRISMA flow diagram [21] (Fig 1).

#### Characteristics of included studies

Among the included studies, 20 (52.60%) studies published after 2016. All the included studies were cross-sectional surveys, of which 27 community based, six health facility-based, and five studies were institutional-based (Schools, College, Bank. . .). Overall, a total number of 51,427 study participants who are older than 18 years included for this review. The minimum and maximum sample sizes were 306 and 9788 respectively [22, 23]. A minimum of (7.47%) and maximum of (41.90%) prevalence of hypertension were reported from the studies conducted in the Oromia region [24, 25]. Five regions and two city administrations (Addis Ababa and Dire Dawa) were represented for this review. Seven from Amhara Region [26–32], eight from Oromia Region [24, 25, 33–38], six from South National and Nationalities of People's Region

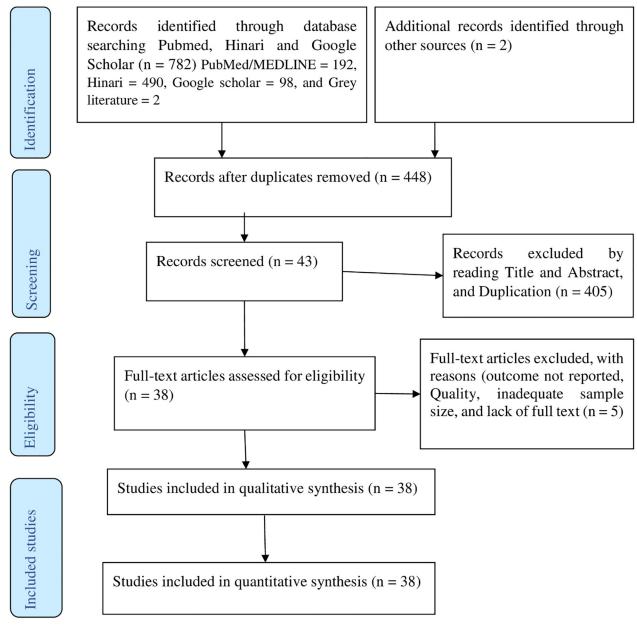


Fig 1. PRISMA flow diagram of article selection for systematic review and meta-analysis of the prevalence of hypertension and its determinants in Ethiopia.

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(SNNPR) [22, 39–43], four from Tigray Region [44–47], three from Somali Region [48–50], eight from Addis Ababa [51–58], two from Dire Dawa [37] and one national study in Ethiopia [23] were included. No studies reviewed from Gambela, Afar, Benishangul Gumez, and Harari Regional states of Ethiopia (Table 1).

## The pooled prevalence of hypertension in Ethiopia

In random effects model, the pooled prevalence of hypertension in Ethiopia was 21.81 (95% CI = 19.20-24.42); significant heterogeneity observed among studies ( $I^2 = 98.4$ , P-

	Characteristics of the inc	inded studies and the	ii prevalence of ii	ypertension in 1			
S. No	Author	Publication year	Region	Sample size	Response rate (%)	Prevalence of hypertension	Quality score
1	Zekewos et al. [42]	2019	SNNPR	425	-	21.80	1
2	Kiber et al. [30]	2019	Amhara	456	95.6	12.50	2
3	Shukuri et al. [25]	2019	Oromia	401	96	41.90	1
4	Abebe et al. [ <u>56</u> ]	2019	Addis Ababa	487	100	34.70	0
5	Roba et al. [ <u>59</u> ]	2019	Dire Dawa	872	96.5	24.40	3
6	Belachew et al. [27]	2018	Amhara	308	100	27.30	0
7	Gebreyes et al. [23]	2018	National	9788	95.4	18.05	0
8	Bayray et al. [46]	2018	Tigray	1523	99.7	15.90	0
9	Tesfaye et al. [38]	2018	Oromia	648	97	14.2	0
10	Esaiyas et al. [40]	2018	SNNPR	620	99.6	19.70	0
11	Bekele et al. [51]	2018	Addis Ababa	758	100	15.90	0
12	Asfaw et al. [41]	2018	SNNPR	524	99.8	30.00	0
13	Mara et al. [39]	2018	SNNPR	346	97.4	23.00	0
14	Neba et al. [50]	2017	Somali	548	100	21.90	0
15	Demisse et al. [28]	2017	Amhara	3057	94.8	27.40	0
16	Asresahegn et al. [48]	2017	Somali	487	98.9	28.30	0
17	Birhanu Tolera [57]	2017	Addis Ababa	401	98.5	14.00	1
18	Seifu et al. [49]	2017	Somali	330	100	13.30	1
19	Gebrihet et al. [45]	2017	Tigray	521	96	16.50	0
20	Fikadu et al. [52]	2016	Addis Ababa	1866	100	21.00	0
21	Tadele et al. [22]	2016	SNNPR	306	95.9	27.80	3
22	Abdissa et al. [54]	2015	Addis Ababa	2716	100	24.90	1
23	Anteneh et al. [32]	2015	Amhara	678	99.6	25.10	1
24	Asresahegn et al. [34]	2015	Oromia	830	100	36.40	3
25	Angaw et al. [55]	2015	Addis Ababa	629	96	27.30	0
26	Birlew et al. [24]	2015	Oromia	4055	90.7	7.47	2
27	Abebe et al. [26]	2015	Amhara	2141	97.3	27.90	1
28	Bissa et al. [35]	2014	Oromia	701	96.02	21.30	1
29	Zikru et al. [47]	2014	Tigray	709	99.7	11.00	3
30	Mengistu et al. [44]	2014	Tigray	1183	100	18.10	1
31	Tadesse et al. [31]	2014	Amhara	610	100	7.70	2
32	Helelo et al. [43]	2014	SNNPR	518	96.6	22.40	2
33	Gudina et al.[36]	2014	Oromia	396	93.8	16.92	2
34	Gudina et al. [33]	2013	Oromia	734	100	13.20	0
35	Nshisso et al. [53]	2012	Addis Ababa	2153	100	19.10	1
36	Awoke et al. [29]	2012	Amhara	679	97.6	28.30	0
37	Muluneh et al. [37]	2012	Oromia	3223	-	9.30	2
38	Tesfaye et al. [58]	2009	Addis Ababa	648	93.2	14.20	1

Table 1. Characteristics of the included studies and their prevalence of hypertension in Ethiopia, 2019.

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value < 0.001). The highest weight among studies observed from the studies conducted by Muluneh et al. [37], Gebreyes et al. [23], and Birlew et al. [24] (Fig 2). Among 23 studies in the random effects model, the pooled prevalence of hypertension among males were 23.21 (95% CI:18.86–27.57) (Fig 3) with statistically significant heterogeneity ( $I^2 = 97.5\%$ , P-value < 0.001). Besides, the overall pooled prevalence of hypertension among females were 19.62 (95% CI: 16.26–22.97) (Fig 4); heterogeneity ( $I^2 = 96.08\%$ , P-value < 0.001). Egger's

Study (Authors, Year)			ffect Size ith 95% CI		Weight (%)
Abebe et al., 2015	-	27.90 [	26.00,	29.80]	2.69
Abebe et al., 2019		34.70 [	30.47,	38.93]	2.55
Awoke et al., 2012		28.30 [	24.91,	31.69]	2.61
Angaw et al., 2015		27.30 [	23.82,	30.78]	2.60
Asresahegn et al. , 2015		36.40 [	33.13,	39.67]	2.62
Asfaw et al., 2018		30.00 [	26.08,	33.92]	2.57
Mara et al., 2018		23.00 [	18.57,	27.43]	2.53
Anteneh et al. , 2015	-=-	25.10 [	21.84,	28.36]	2.62
Asresahegn et al., 2017		28.30 [	24.30,	32.30]	2.57
Bekele et al., 2018		32.30 [	28.97,	35.63]	2.61
Bayray et al, 2018	-	15.90 [	14.06,	17.74]	2.69
Belachew et al, 2018		27.30 [	22.32,	32.28]	2.48
Gudina et al., 2014		16.92 [	13.23,	20.61]	2.59
Demisse et al. , 2017		27.40 [	25.82,	28.98]	
Gebrihet et al, 2017		16.50 [	13.31,	19.69]	2.62
Esaiyas et al, 2018		19.70 [	16.57,	22.83]	2.63
Gebreyes et al, 2018		18.05 [	17.29,	18.81]	
Tesfaye et al, 2018	-	14.20 [	11.51,	16.89]	
Gudina et al., 2013	-	13.20 [	10.75,	15.65]	
Helelo et al. , 2014		22.40 [	18.81,	25.99]	2.60
Kiber et al., 2019	-	12.50 [	9.46,	15.54]	
Roba et al., 2019		24.40 [	21.55,	27.25]	
Mengistu, 2014	-	18.10 [	15.91,	20.29]	2.68
Muluneh et al., 2012		9.30 [	8.30,	10.30]	
Seifu et al. , 2017		13.30 [	9.64,	16.96]	2.59
Shukuri et al. , 2019		41.90 [	37.07,	46.73]	2.50
Zekewos et al. , 2019	+	21.80 [	19.97,	23.63]	2.69
Tadesse et al., 2014	<b>.</b>	7.70 [	5.58,	9.82]	2.68
Birhanu tolera, 2017		14.00 [	10.60,	17.40]	2.61
Nshisso et al. , 2012	-	19.10 [	17.44,	20.76]	2.70
Fikadu et al, 2016		21.00 [	19.15,	22.85]	2.69
Abdissa et al. , 2015		24.90 [	23.27,	26.53]	2.70
Bissa et al., 2014	-	21.30 [	18.27,	24.33]	2.63
Tesfaye et al, 2009		30.28 [	28.71,	31.85]	2.70
Naba et al, 2017		21.90 [		25.36]	2.61
Tadele et al, 2016		27.80 [	22.78,	32.82]	2.48
Zikru et al, 2014	<b>.</b>	11.00 [	8.70,	13.30]	2.67
Birlew et al., 2015		7.47 [	6.66,	8.28]	2.72
Overall		21.81 [	19 20	24 421	
Heterogeneity: $\tau^2 = 65.01$ , $I^2 = 98.35\%$ , $H^2 = 60.62$ Test of $\theta_i = \theta_j$ : Q(37) = 2243.06, p < 0.001 Test of $\theta = 0$ : z = 16.37, p < 0.001	5 10 15 20 25 30 35 40 45	-	10.20,	£7,72]	
	5 10 15 20 20 50 55 40 45				

Random-effects DerSimonian-Laird model

Fig 2. Pooled prevalence of hypertension age greater than 18 years in Ethiopia.

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Study (Authors, Year)		Effect Size with 95% Cl	Weight (%)
Abebe et al., 2015	-	26.30 [ 23.54, 29.06]	4.48
Awoke et al., 2012	-	26.00 [ 21.22, 30.78]	4.33
Asresahegn et al. , 2015		38.10 [ 32.19, 44.01]	4.21
Asfaw et al., 2018	-	26.40 [ 21.21, 31.59]	4.29
Mara et al., 2018		24.00 [ 16.48, 31.52]	4.02
Bekele et al., 2018		36.40 [ 31.10, 41.70]	4.28
Bayray et al, 2018	-	20.00 [ 17.34, 22.66]	4.49
Belachew et al, 2018		32.70 [ 25.27, 40.13]	4.03
Gudina et al., 2014	-	13.10 [ 9.05, 17.15]	4.39
Esaiyas et al, 2018		24.50 [ 20.15, 28.85]	4.37
Gebreyes et al, 2018	<b>.</b>	15.50 [ 12.77, 18.23]	4.48
Gudina et al., 2013		15.30 [ 11.28, 19.32]	4.39
Helelo et al. , 2014		26.20 [ 20.95, 31.45]	4.28
Kiber et al., 2019		5.80 [ 3.11, 8.49]	4.49
Roba et al., 2019	-	25.30 [ 20.26, 30.34]	4.30
Mengistu, 2014	- <b>-</b>	21.00 [ 17.21, 24.79]	4.41
Tadesse et al., 2014		9.25 [ 6.58, 11.92]	4.49
Fikadu et al, 2016		23.30 [ 20.83, 25.77]	4.50
Abdissa et al. , 2015	-	30.23 [ 27.56, 32.90]	4.49
Bissa et al., 2014	-	22.50 [ 18.58, 26.42]	4.40
Naba et al, 2017		22.00 [ 16.91, 27.09]	4.30
Tadele et al, 2016		49.00 [ 41.66, 56.34]	4.04
Birlew et al., 2015		6.40 [ 5.31, 7.49]	4.55
Overall	▲	23.21 [ 18.86, 27.57]	
Heterogeneity: $T^2 = 108.20$ , $I^2 = 97.54\%$ , $H^2 = 40.64$			
Test of $\theta_i = \theta_i$ : Q(22) = 894.17, p < 0.001			
Test of $\theta = 0$ : z = 10.44, p < 0.001			
	0 5 10 15 20 25 30 35 40 45 50 55 6	0	
Random-effects DerSimonian-Laird model			

Fig 3. Pooled prevalence of hypertension among males in Ethiopia, 2019.

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statistical test evidenced that has no publication bias among the included studies ( $\beta$  = -0.615, P-value = 0.91).

## Handling heterogeneity

Significant heterogeneity observed from random effects model pooled estimate. To handle this heterogeneity sensitivity analysis, subgroup analysis, and meta-regression analysis were performed.

## Sensitivity analysis

From the random effects model, there are no studies that excessively influence the overall pooled estimate of hypertension (S1 Fig).

Study (Authors, Year)		Effect Size with 95% Cl	Weight (%)
Abebe et al., 2015		29.30 [ 26.68, 31.92]	4.66
Awoke et al., 2012		30.30 [ 25.53, 35.07]	4.37
Asresahegn et al. , 2015		28.50 [ 15.01, 41.99]	2.70
Asfaw et al., 2018		34.00 [ 28.09, 39.91]	4.17
Mara et al., 2018		22.50 [ 17.01, 27.99]	4.25
Bekele et al., 2018		29.40 [ 25.15, 33.65]	4.45
Bayray et al, 2018	. <b></b>	10.40 [ 8.06, 12.74]	4.68
Belachew et al, 2018		21.90 [ 15.39, 28.41]	4.06
Gudina et al., 2014		24.80 [ 17.35, 32.25]	3.88
Esaiyas et al, 2018		12.00 [ 7.83, 16.17]	4.46
Gebreyes et al, 2018	-	16.30 [ 14.11, 18.49]	4.70
Gudina et al., 2013		11.70 [ 8.65, 14.75]	4.61
Helelo et al. , 2014		18.40 [ 13.93, 22.87]	4.42
Kiber et al., 2019		18.10 [ 11.65, 24.55]	4.07
Roba et al., 2019		10.25 [ 7.79, 12.71]	4.67
Mengistu, 2014	-	16.40 [ 13.53, 19.27]	4.63
Tadesse et al., 2014	- <b></b>	3.20 [ 0.44, 5.96]	4.64
Fikadu et al, 2016	- E	17.50 [ 14.77, 20.23]	4.65
Abdissa et al. , 2015	-	21.17 [ 19.15, 23.19]	4.71
Bissa et al., 2014		19.62 [ 14.84, 24.40]	4.37
Naba et al, 2017		21.80 [ 17.08, 26.52]	4.38
Tadele et al, 2016		- 36.00 [ 27.68, 44.32]	3.70
Birlew et al., 2015		8.50 [ 7.31, 9.69]	4.77
Overall	+	19.62 [ 16.26, 22.97]	
Heterogeneity: $\tau^2 = 61.09$ , $I^2 = 96.08\%$ , $H^2 = 25.51$			
Test of $\theta_i = \theta_i$ : Q(22) = 561.26, p < 0.001			
Test of $\theta = 0$ : z = 11.46, p < 0.001			
,	0 5 10 15 20 25 30 35 40	¬ 45	
Random-effects DerSimonian-Laird model			

Fig 4. Pooled prevalence of hypertension among females in Ethiopia, 2019.

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## Subgroup analysis

Even though subgroup analysis was carried out across the administrative regions of the country, age category, and residence as the source of heterogeneity was not handled. In the subgroup analysis, the highest prevalence of hypertension observed in Addis Ababa (25.35%) followed by Southern Nations Nationalities and People's Region (23.83%); whereas the lowest prevalence was in Tigray regional state of Ethiopia (15.36%). The pooled prevalence of hypertension (27%) was higher in the age category which is older than 35 years. Also, the highest prevalence of hypertension was observed in urban inhabitants (22.85%) (Table 2).

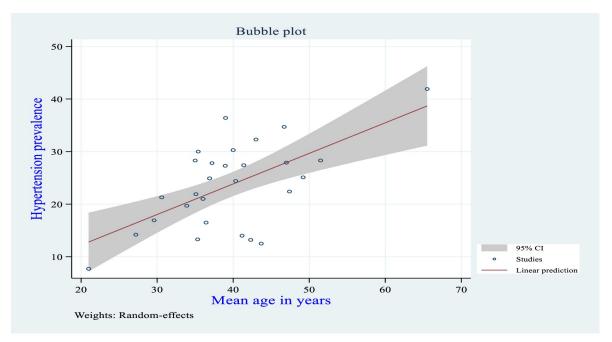
Variables		Included studies	Sample size	Prevalence (95%CI)	Heterogeneity (I <sup>2</sup> , p-value)
By region	Tigray	4	3936	15.36 (12.33-18.39)	85.4%, < 0.001
	Amhara	7	7929	22.27 (15.44-29.11)	98.1%, < 0.001
	Oromia	8	10988	19.83 (14.09-25.28)	98.7%, < 0.001
	SNNPR	6	2739	23.83 (20.93-26.72)	77.0%, < 0.001
	Addis Ababa	8	9658	25.35 (21.25-29.45)	96.3%, < 0.001
	Somali	3	1365	21.14 (12.86-29.42)	93.3%, < 0.001
By age category	> 18 years	25	38360	19.92 (24.28-29.56)	98.4%, < 0.001
	> 25 years	7	8304	24.37 (19.84-28.89)	95.4%, < 0.001
	> 30 years	2	1196	23.86 (21.22-26.49)	15.9%, 0.275
	> 35years	3	2888	26.92 (24.28-29.56)	53.7%, 0.115
By residence	Rural	5	10814	18.45 (12.41-24.48)	99.03, < 0.001
	Urban	28	26554	22.85 (20.34-25.36)	95.91, < 0.001
	Both urban and rural	5	14059	18.45 (12.41-24.48)	98.19, < 0.001

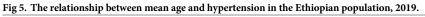
Table 2. Sub-group pooled prevalence of hypertension in Ethiopia, 2019 (n = 38).

https://doi.org/10.1371/journal.pone.0244642.t002

#### Meta-regression

Univariate meta-regression analysis revealed that the mean age and region were statistically significant with hypertension. As the mean age increased by one year, the likelihood of developing hypertension increases by a factor of 0.58 times ( $\beta = 0.58$ , 95% CI: 0.31–0.86); with a total proportion of hypertension explained by the covariate mean age by 36.67% (adjusted R<sup>2</sup> = 36.67). The linear relationship between mean age and hypertension was presented as shown in Fig 5 below. Besides, the pooled prevalence of hypertension was higher in the capital city of





https://doi.org/10.1371/journal.pone.0244642.g005

Study level variables		Adjusted R <sup>2</sup>	Standard error	Coefficients (95% CI)
Mean age		36.67	0.14	0.58 (0.31–0.86) *
Publication year		00	0.57	0.58(-0.54-1.69)
Sample size		00	0.0008	0.00072 (-0.0023–0.0009)
Regions	Tigray	1	1	1
	Amhara		4.59	6.88 (-2.13-15.89)
	Addis Ababa	20	4.48	10.01(1.22–18.80) *
	Oromia		4.49	4.39 (-4.40–13.19)
	SNNPR		4.76	8.67 (-0.65–18.00)
	Somali		5.64	5.77 (-5.30-16.54)
	Dire Dawa		8.21	9.03 (-7.07–25.12)

#### Table 3. Univariate meta-regression analysis results for the prevalence of hypertension in Ethiopia, 2019.

#### NB:

\* = Statistically significant at 5% level, CI = Confidence Interval.

https://doi.org/10.1371/journal.pone.0244642.t003

Addis Ababa, Ethiopia as compared to Tigray regional state of Ethiopia ( $\beta$  = 10.01, 95% CI: 1.22–18.80) (Table 3).

#### Factors associated with hypertension

As summarized in <u>Table 4</u>, sex, age, Body Mass Index (BMI), chat chewing, alcohol consumption, and family history of hypertension and diabetes mellitus were statistically significant factors for hypertension.

#### Table 4. Summary of the pooled effects of factors associated with hypertension in Ethiopia, 2019.

Variables		OR (95% CI)	Heterogeneity (I <sup>2</sup> , P-value)	Egger's P-value	Total studies	Sample size
Sex	Female	1				
	Male	1.29 (1.03–1.61) *	81.35%, < 0.001	0.544	15	19957
Age	< 35 years	1	1			
	> 35 years	3.59 (2.57-5.02) *	93.48%, < 0.001	0.487	15	27365
BMI	Normal	1				
	Underweight	0.68 (0.30-1.56)	94.00%, < 0.001	0.229	16	
	Overweight and /or obese	3.34 (2.12–5.26) *	95.41%, < 0.001	0.176	18	13383
Khat chewing	No	1	1			
	Yes	1.42 (1.10–1.85) *	62.2%, 0.005	0.267	10	8687
Smoking	No	1	1			
	Yes	1.55 (1.00-2.38)	67.56%, 0.002	0.873	10	9556
Alcohol drinking	No	1	1			
	Yes	1.50 (1.21–1.85) *	64.0%, 0.001	0.005	14	12988
Physical activity	Active	1	1			
	Inactive	1.24 (0.83-1.85)	91.28%, < 0.001	0.0002	15	
Family history of HTN	No	1	1			
	Yes	2.56 (1.64-3.99) *	83.28%, < 0.001	0.016	11	5918
Family history of DM	No	1	1			
	Yes	3.69 (1.85–7.59) *	89.93%, < 0.001	0.4707	9	14660

#### NB:

\* = Statistically significant at 5% level, OR = Odds Ratio, CI = Confidence Interval.

https://doi.org/10.1371/journal.pone.0244642.t004

Study (Authors, Year)		exp(ES) with 95% CI	Weight (%)
Abebe et al., 2015		0.86 [ 0.71, 1.04]	9.19
Asfaw et al., 2018		1.92 [ 1.14, 3.23]	6.41
Asresahegn et al., 2017		2.40 [ 1.32, 4.36]	5.76
Bekele et al., 2018		1.37 [ 1.02, 1.85]	8.36
Bayray et al, 2018	-	2.06 [ 1.49, 2.84]	8.17
Demisse et al. , 2017	•	- 1.42 [ 0.12, 17.14]	0.74
Gudina et al., 2013	-	1.35 [ 0.88, 2.07]	7.23
Helelo et al. , 2014		2.03 [ 1.05, 3.93]	5.29
Kiber et al., 2019		0.27 [ 0.14, 0.54]	5.04
Roba et al., 2019		1.39 [ 0.88, 2.20]	6.96
Tadesse et al., 2014		3.12 [ 1.16, 8.38]	3.36
Fikadu et al, 2016	-	1.43 [ 1.13, 1.80]	8.90
Abdissa et al. , 2015		1.40 [ 1.18, 1.67]	9.30
Tadele et al, 2016		0.97 [ 0.59, 1.60]	6.60
Birlew et al., 2015	-	0.74 [ 0.57, 0.96]	8.69
Overall	•	1.29 [ 1.03, 1.61]	
Heterogeneity: $\tau^2 = 0.13$ , $I^2 = 81.35\%$ , $H^2 = 5.36$			
Test of $\theta_i = \theta_i$ : Q(14) = 75.05, p < 0.001			
Test of $\theta$ = 0: z = 2.25, p = 0.02			
and server and the server and the server - Solida Bab	0.12 1/4 1/2 1 2 4 8 17	7. <b>14</b>	
Random-effects DerSimonian-Laird model			
Fig 6 The association between males and hypertension			

Fig 6. The association between males and hypertension.

https://doi.org/10.1371/journal.pone.0244642.g006

Fifteen [22, 24, 26, 28, 30, 31, 33, 41, 43, 46, 48, 51, 52, 54, 59] studies were included to identify the association between sex and hypertension. Five of these studies [22, 26, 28, 33, 59] had no statistically significant association between sex and hypertension. From random effects model estimate, the pooled odds of developing hypertension among males were 29% more likely to develop hypertension than females (OR = 1.29, 95% CI: 1.03–1.61); with statistically significant heterogeneity between studies ( $I^2 = 81.3\%$ , P-value < 0.001) (Fig 6). Egger's test indicates that no small study effect (P-value = 0.544) and in random effects model there was no single study that excessively influences the pooled estimate of hypertension (S2 Fig).

The pooled effect of age has a significant association with hypertension. From fifteen [23, 24, 28, 30, 35, 40, 41, 46, 48, 50, 51, 52, 54–56] studies only one [24] study had no significant association between age and hypertension. The pooled odds of developing hypertension among individuals older than 35 years was 3.59 times higher than age younger than 35 years (OR = 3.59, 95% CI: 2.57–5.02) (Fig 7); with statistically significant heterogeneity among studies (I<sup>2</sup> = 93.5%, P-value < 0.001). There is no small study effect (P-value = 0.485) and in random effects model, there was no single study excessively influence the pooled estimate of effect size (S3 Fig).

A total of eighteen [22, 25, 27–29, 31, 32, 35, 40, 43, 45–47, 49–51, 55, 56] studies included to estimate the association between BMI and hypertension. The results of the test statistics indicate that significant heterogeneity was observed between studies ( $I^2 = 95.41\%$ , P-value < 0.001). Egger's test evidenced that there was no publication bias (P-value = 0.176).

Ctudy (Authors Veer)	0	35 years		Control		Odds Ratio	Weight
Study (Authors, Year)	HTN	Non-HTN	HIN	Non-HTN		with 95% CI	(%)
Abebe et al., 2019	144	190	25	128		3.88 [ 2.40, 6.27]	6.49
Angaw et al., 2015	113	164	59	293		3.42 [ 2.37, 4.95]	6.86
Asfaw et al., 2018	138	284	19	83		2.12 [ 1.24, 3.64]	6.27
Asresahegn et al., 2017	78	124	60	225		2.36 [ 1.58, 3.52]	6.76
Bekele et al., 2018	231	361	14	152		6.95 [ 3.92, 12.31]	6.14
Bayray et al, 2018	49	60	18	269		12.20 [ 6.64, 22.43]	6.00
Demisse et al. , 2017	667	1,181	170	1,708	-	5.67 [ 4.72, 6.82]	7.32
Esaiyas et al, 2018	70	165	45	289		2.72 [ 1.79, 4.15]	6.69
Gebreyes et al, 2018	286	846	83	552		2.25 [ 1.72, 2.94]	7.14
Kiber et al., 2019	50	282	7	117		2.96 [ 1.31, 6.73]	5.17
Fikadu et al, 2016	304	554	116	930	-	4.40 [ 3.47, 5.58]	7.21
Abdissa et al. , 2015	498	792	180	1,246	-	4.35 [ 3.59, 5.28]	7.30
Bissa et al., 2014	100	81	49	470		11.84 [ 7.82, 17.94]	6.71
Naba et al, 2017	55	137	65	291		1.80 [ 1.19, 2.72]	6.72
Birlew et al., 2015	189	2,298	114	1,454 -	-	1.05 [ 0.82, 1.34]	7.20
Overall					•	3.59 [ 2.57, 5.02]	
Heterogeneity: $\tau^2 = 0.39$	9. $ ^2 = 9$	3.48%. H <sup>2</sup>	= 15.	34			
Test of $\theta_i = \theta_j$ : Q(14) = 2							
Test of $\theta$ = 0: z = 7.48,							
				-	1 2 4 8 16		
Random-effects DerSimonian-Laird	model						

https://doi.org/10.1371/journal.pone.0244642.g007

Again, from random effects model, no individual studies excessively influence the pooled estimate of the effect size (S4 Fig). From the random effects model pooled estimate, the likelihood of developing hypertension among overweight and/or obese individuals was 3.34 times higher than the normal-weight individuals (OR = 3.34, 95% CI: 2.12–5.26) (Fig 8).

The pooled effects between khat chewing and hypertension was assessed using ten studies [22, 24, 33, 35, 36, 46, 48, 49, 55, 57]. Among the included studies, six [22, 24, 33, 36, 38, 48] of them reported that khat chewing has not a statistically significant association with hypertension. Based on Egger's test there was no publication bias (P-value = 0.498). Besides, from random effects model there was no single study that excessively influences the pooled effect size (S5 Fig). Khat chewers have 42% more likelihood to develop hypertension than non-khat chewers (OR = 1.42, 95% CI: 1.10–1.85) (Fig 9), with moderate heterogeneity ( $I^2 = 62.2\%$ , Pvalue = 0.005).

The association between alcohol consumption and hypertension was assessed using 14 studies [22, 24, 26, 27, 30, 34-36, 38, 55-57, 59]. Moderate heterogeneity was also observed from the random effects model ( $I^2 = 64.04\%$ ) and there is no evidence of a single study that affects the pooled effects size in the sensitivity analysis (S6 Fig). Egger's test evidenced that small study effect (P-value = 0.001). After non-parametric trim and fill analysis (Fig 10), alcohol consumption had a negative effect on hypertension. From the random-effects trim and fill analysis, alcohol drinkers were more likely to develop hypertension by half as compared to non-drinkers (OR = 1.50, 95% CI: 1.21-1.85).

Fig 7. Forest plot for the association between age and hypertension.

	0	&/or Obese		ontrol				dds Ratio	Weight
Study (Authors, Year)	HTN	Non-HTN	HTN	Non-HTN			wit	h 95% Cl	(%)
Abebe et al., 2019	83	92	79	182		-	2.08 [	1.40, 3.09	5.69
Awoke et al., 2012	81	129	101	298		-	1.85 [	1.30, 2.65	] 5.74
Angaw et al., 2015	105	67	60	341		-	8.91 [	5.90, 13.44	] 5.68
Anteneh et al., 2015	59	65	100	402		-	3.65 [	2.41, 5.53	] 5.67
Bekele et al., 2018	166	190	79	323			3.57 [	2.59, 4.93	] 5.78
Bayray et al, 2018	108	317	129	74	-		0.20 [	0.14, 0.28	5.74
Belachew et al, 2018	16	14	60	186			3.54 [	1.63, 7.68	] 5.08
Demisse et al. , 2017	284	473	485	1,411			1.75 [	1.46, 2.09	] 5.90
Gebrihet et al, 2017	51	31	33	364			- 18.15 [	10.25, 32.12	5.44
Esaiyas et al, 2018	65	127	51	323		-	3.24 [	2.13, 4.93	5.67
Helelo et al., 2014	73	53	42	330			10.82 [	6.71, 17.45	5.59
Seifu et al., 2017	25	95	15	77			1.35 [	0.67, 2.74	5.20
Shukuri et al., 2019	67	26	91	181			5.13 [	3.05, 8.61	] 5.53
Tadesse et al., 2014	11	17	27	337			8.08 [	3.44, 18.97	] 4.92
Bissa et al., 2014	60	72	86	433		-	4.20 [	2.77, 6.34	5.67
Naba et al, 2017	58	93	61	294		-	3.01 [	1.96, 4.61	] 5.65
Tadele et al, 2016	49	50	36	171			4.66 [	2.73, 7.93	] 5.50
Zikru et al, 2014	34	118	40	409		-	2.95 [	1.79, 4.86	] 5.55
Overall						+	3.34 [	2.12, 5.26	]
Heterogeneity: $\tau^2 = 0$			= 21.7	9					
Test of $\theta_i = \theta_j$ : Q(17)									
Test of $\theta = 0$ : $z = 5.2$	21, p < 0.00	1					-		

1/4

4

1

16

Random-effects DerSimonian-Laird model

Fig 8. The association between body mass index and hypertension.

https://doi.org/10.1371/journal.pone.0244642.g008

Study (Authors, Year)		exp(ES) Weight with 95% CI (%)
Angaw et al., 2015		1.62 [ 1.04, 2.53] 12.71
Asresahegn et al., 2017		1.32 [ 0.88, 1.97] 13.61
Gudina et al., 2014		1.60 [ 0.90, 2.84] 10.22
Tesfaye et al, 2018		— 4.56 [ 0.61, 34.11] 1.55
Gudina et al., 2013		0.50 [ 0.24, 1.03] 7.84
Seifu et al. , 2017	-	1.98 [ 1.47, 2.68] 15.88
Birhanu tolera, 2017		- 6.90 [ 1.44, 33.07] 2.44
Bissa et al., 2014		1.50 [ 1.07, 2.11] 14.98
Tadele et al, 2016		1.30 [ 0.37, 4.55] 3.57
Birlew et al., 2015	-	1.07 [ 0.84, 1.36] 17.19
Overall	•	1.42 [ 1.10, 1.85]
Heterogeneity: $\tau^2 = 0.09$ , $I^2 = 62.24\%$ , $H^2 = 2.65$		
Test of $\theta_i = \theta_i$ : Q(9) = 23.83, p < 0.001		
Test of $\theta$ = 0: z = 2.65, p = 0.01		
	1/4 1 4 16	
Random-effects DerSimonian-Laird model		

Fig 9. The association between khat chewing and hypertension.

https://doi.org/10.1371/journal.pone.0244642.g009

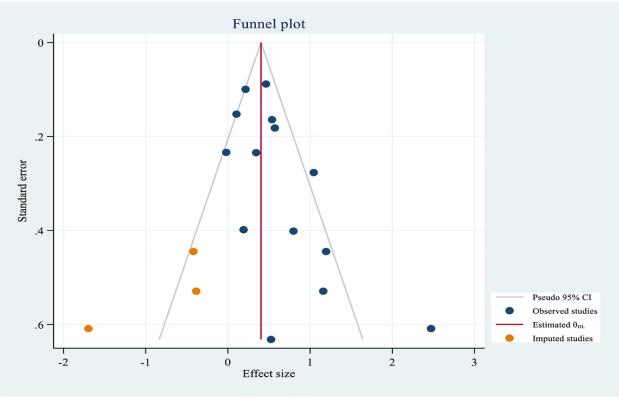


Fig 10. Trim and fill analysis funnel plot for alcohol consumption.

https://doi.org/10.1371/journal.pone.0244642.g010

A total of fifteen studies [22, 25, 26, 28, 29, 32, 33, 35, 41, 43, 45, 48, 49, 56, 59] were included to determine the association between physical activity and hypertension; of them, four studies had no statistically significant association with hypertension. From random effects model estimate, significant heterogeneity observed ( $I^2 = 91.3\%$ , P-value < 0.001). Egger's test indicates that evidence of publication bias (P-value = 0.002). After non-parametric trim and fill analysis, physical exercise and hypertension has no significant association (OR = 1.24, 95% CI: 0.83–1.85).

As the results of eleven studies [22, 25, 29, 30, 33, 34, 43, 48–50, 55], family history of hypertension and hypertension had statistically significant association. The random effects model evidenced that statistically significant heterogeneity across studies ( $I^2 = 83.3\%$ , Pvalue < 0.001). From the sensitivity analysis random effects model estimate there is no single study that excessively influences pooled effect size (S7 Fig). Egger's test showed that the presence of a small study effect (P-value = 0.016). After non-parametric trim and fill analysis pooled estimate (Fig 11), the pooled odds of developing hypertension among individuals who had a family history of hypertension were 2.56 times higher than their counterparts (OR = 2.56, 95% CI:1.64–3.99).

Furthermore, the association between the family of diabetes mellitus and hypertension was identified using nine studies [22, 23, 29, 32, 33, 48, 55, 56, 59]; among them, four studies [22, 23, 48, 59] showed that there is no statistically significant association between family history of diabetes mellitus and hypertension. The random effects model estimate showed that statistically significant heterogeneity between studies ( $I^2 = 89.9\%$ , P-value < 0.001) and Egger's test

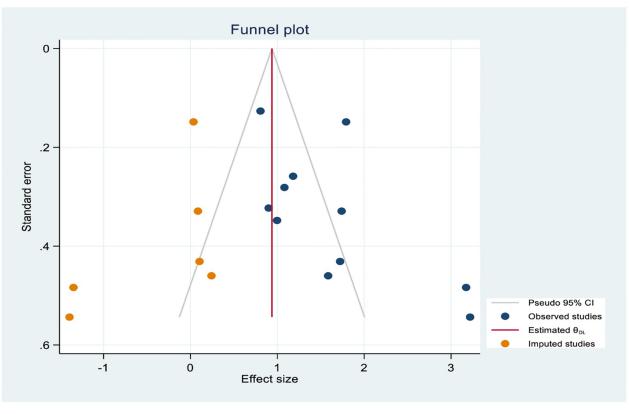


Fig 11. Trim and fill analysis funnel plot for a family history of hypertension.

https://doi.org/10.1371/journal.pone.0244642.g011

showed that there is no publication bias (P-value = 0.47). From random effects model sensitivity analysis, there is no single study that excessively affects the pooled effect size (S8 Fig). Form random effects model estimate individuals who had a family history of diabetes mellitus are 3.69 times more likely to develop hypertension than the reference category (OR = 3.69, 95% CI: 1.85–7.59) (Fig 12) (Table 4).

#### Discussion

Non-communicable diseases are becoming a double burden of public health problem in developing countries [60]; besides hypertension prevalence is rising in developing countries in contrast to developed nations [61]. This systematic review and meta-analysis will give the update pooled estimates of hypertension in Ethiopia which gives invaluable information to policy-makers, health planners, and the community itself.

This systematic review and meta-analysis revealed that the pooled prevalence of hypertension in Ethiopia was 21.81% (95% CI: 19.20–24.42), which was consistent with a study conducted in rural communities of Sub-Saharan Africa (22%), Kenya (22.8%), and a meta-analysis from Vietnam (21.1%) [62–64]. However, the finding of this meta-analysis was lower than the previous meta-analysis reports in LMICs (32.3%), among adults in Africa (57.0%), a metaanalysis study on undiagnosed hypertension in Sub-Saharan Africa (30%), Nigeria (28.9%), India (29.8%), Pakistan (26.34%), and a study in Nepal (25.1%) [65–71]. The prevalence of hypertension in this review was higher than a study conducted a previous systematic review in Ethiopia and a study conducted in Ghana [13, 72]. The possible reason for this discrepancy

Study (Authors, Year)		exp(ES) with 95% CI	Weight (%)
		with 5570 Of	(70)
Abebe et al., 2019		– 15.47 [ 7.76, 30.86]	11.67
Awoke et al., 2012		4.15 [ 1.77, 9.72]	11.00
Angaw et al., 2015		13.56 [ 6.91, 26.60]	11.73
Anteneh et al., 2015		2.78 [ 1.63, 4.73]	12.24
Asresahegn et al., 2017		3.90 [ 0.88, 17.20]	8.22
Gebreyes et al, 2018		1.25 [ 0.91, 1.72]	12.84
Gudina et al., 2013		4.13 [ 1.47, 11.63]	10.20
Roba et al., 2019		1.22 [ 0.73, 2.03]	12.31
Tadele et al, 2016		2.50 [ 0.81, 7.71]	9.79
Overall		3.69 [ 1.85, 7.39]	
Heterogeneity: $\tau^2 = 0.95$ , $I^2 = 89.93\%$ , $H^2 = 9.93$			
Test of $\theta_i = \theta_i$ : Q(8) = 79.47, p < 0.001			
Test of $\theta$ = 0: z = 3.69, p < 0.00		_	
	1 2 4 8 16		

Random-effects DerSimonian-Laird model

Fig 12. The association between family history of diabetes mellitus and hypertension.

https://doi.org/10.1371/journal.pone.0244642.g012

might be the time of the study, the age group of the population studied, the diagnosis criteria for hypertension, and the study setting.

From subgroup analysis by region, the highest prevalence of hypertension (25%) was observed in the capital city of Ethiopia, Addis Ababa. This is similar to subgroup analysis by the residence which is the prevalence of hypertension (23%) was higher in urban inhabitants. The possible justification might be, urbanization may be linked to low physical activity, consumption of unhealthy diet and stress which may again leads to the high burden of non-communicable diseases [73–76].

This review also identifies the determinant factors of hypertension. In random effects model pooled estimate, sex, age, body mass index, khat chewing, alcohol consumption, family history of hypertension, and family history of diabetes mellitus were significantly associated with hypertension.

From the random effects model estimate, the pooled odds of developing hypertension among males were 29% higher than females. This finding was similar with the studies conducted in Nepal, Varanasi India, Burkina Faso, Debrecen city of Hungary, and a meta-analysis study from Vietnam [62, 69, 77–80], whereas it is not similar to a study conducted at Uganda [81]. The possible reason might be males were more vulnerable to behavioral risk factors for hypertension.

The pooled effect of age greater than 35 years was 3.6 times higher than age less than 35 years to develop hypertension, which is similar to the community-based studies conducted in Uganda, Nepal, Benin, Varanasi city India, and another city of New Delhi, India [79–83]. As well, from meta-regression analysis showed that mean age and hypertension had a direct linear relationship. Age is one of the non-modifiable risk factors for hypertension. As a result, this is the fact that cardiovascular system is strongly affected by ageing; besides, ageing causes the structural and functional changes in the blood vessels that may lead to cardiovascular morbid-ity and mortality [84].

This review also evidenced that individuals being overweight and/or obese were venerable to hypertension. The likelihood of developing hypertension among overweight and/or obese individuals were three times higher than normal in their body mass index. This finding is similar to the previous studies conducted in different countries [63, 77–79, 81]. Besides, a study conducted in Japan evidenced that as 1 kg/m<sup>2</sup> increase in body mass index increases the odds of developing hypertension by 23% among males and 35% among females [85]. This study strengths the fact that high body mass index increases blood cholesterol level which leads to hypertension [86]. Furthermore, this review evidenced that khat (Catha edulis) chewers were 42% more likely to develop hypertension than their counterparts which was similar to the studies conducted in Ethiopia, Yemen, and a meta-analysis study from Ethiopia [87–90]. Khat contains chemicals cathinone, cathine, and amphetamine. Cathinone is structurally related to amphetamine which increases levels of dopamine in the brain by acting on the catecholamin-ergic synapses [91] and increase blood pressure and heart rate [92, 93].

The pooled estimates of alcohol drinking and hypertension were statistically significant in random effects model estimate with moderate heterogeneity between studies. The odds of developing hypertension among drinkers were higher than by half as compared to non-drinkers. This finding was similar to the studies done in North American and France [94–96]. Another study evidenced that consuming three or more drinks of alcohol per day which approximately doubles the risk of developing hypertension [97]. Alcohol consumption affects the central nervous system which enhances cardiac output and has an effect on peripheral vascular effects [98].

Furthermore, family history of hypertension was a potential determinant factor for hypertension. Individuals who had a family history of hypertension have almost five times more chance to develop hypertension than individuals who had no family history of hypertension. This finding was similar to the previous studies conducted in China, Sri Lanka, and Mexico [99–101]. In addition, individuals who had a family history of diabetes mellitus were 3.7 times more likely to develop hypertension as compared to their counterparts. These factors are nonmodifiable risk factors for hypertension. The possible association of family history of hypertension and diabetes mellitus with hypertension might be close blood relatives might have the same genes which may predispose to hypertension. Besides, close blood relatives might have experience of common behavioral practices that may predispose to hypertension.

This study follows some strengths and limitations. Our review adds considerable knowledge of the updated prevalence of hypertension in Ethiopia. All included studies use the same definition to declare hypertension. Subgroup analysis was performed to minimize statistical heterogeneity. Multiple factors were also included to identify the significant factors for hypertension. However, substantial statistically significant heterogeneity was observed across studies which undermine the pooled estimate of hypertension suggests that chance could be responsible for between-study variability. Sub-group analysis could not identify the source of heterogeneity. Though, meta-regression analysis suggested that mean age and region explain some source of heterogeneity.

## **Conclusions and recommendations**

In conclusion, hypertension is becoming a major public health problem in Ethiopia. Nearly two out of ten individuals who are older than 18 years in Ethiopia are living with hypertension. The highest prevalence of hypertension was observed in Addis Ababa and the lowest was in Tigray region. Sex, age, overweight and/or obesity, chat chewing, alcohol consumption, family history of hypertension and family history of diabetes mellitus were statistically significant factors for hypertension. Based on the finding of this review, we recommend that health planners,

policymakers, and the community itself should give prior attention to behavioral risk factors such as chat chewing, alcohol drinking and sedentary lifestyle.

## Supporting information

S1 Table. Studies search strategies and entry terms from different electronic databases on the prevalence and determinants of hypertension. (DOCX)

**S1** Fig. Sensitivity analysis plot for the pooled prevalence of hypertension. (TIF)

**S2** Fig. Assessment of sensitivity analysis plot for factor sex. (TIF)

**S3 Fig. Assessment of sensitivity analysis plot for the factor age.** (TIF)

S4 Fig. Assessment of sensitivity analysis plot for factor among obese and/or overweight. (TIF)

**S5 Fig. Assessment of sensitivity analysis plot for factor Khat Chewing.** (TIF)

**S6 Fig.** Assessment of sensitivity analysis plot for factor alcohol consumption. (TIF)

S7 Fig. Assessment of sensitivity analysis plot for factor family history of hypertension. (TIF)

**S8 Fig.** Assessment of sensitivity analysis plot for factor alcohol consumption. (TIF)

**S1 Checklist.** (DOC)

S1 File.

(XLSX)

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