

BMJ Open Diagnostic accuracy of adrenal imaging for subtype diagnosis in primary aldosteronism: systematic review and meta-analysis

Yaqiong Zhou , Dan Wang, Licheng Jiang, Fei Ran, Sichao Chen, Peng Zhou, Peijian Wang

To cite: Zhou Y, Wang D, Jiang L, *et al.* Diagnostic accuracy of adrenal imaging for subtype diagnosis in primary aldosteronism: systematic review and meta-analysis. *BMJ Open* 2020;**10**:e038489. doi:10.1136/bmjopen-2020-038489

► Prepublication history and additional material for this paper is available online. To view these files, please visit the journal online (<http://dx.doi.org/10.1136/bmjopen-2020-038489>).

Received 12 March 2020

Revised 28 July 2020

Accepted 02 December 2020



© Author(s) (or their employer(s)) 2020. Re-use permitted under CC BY-NC. No commercial re-use. See rights and permissions. Published by BMJ.

Cardiology, Clinical Medical College and The First Affiliated Hospital of Chengdu Medical College, Chengdu, Sichuan, China

Correspondence to

Professor Peijian Wang; wpjmed@aliyun.com and Professor Peng Zhou; ap216@163.com

ABSTRACT

Objectives Accurate subtype classification in primary aldosteronism (PA) is critical in assessing the optimal treatment options. This study aimed to evaluate the diagnostic accuracy of adrenal imaging for unilateral PA classification.

Methods Systematic searches of PubMed, EMBASE and the Cochrane databases were performed from 1 January 2000 to 1 February 2020, for all studies that used CT or MRI in determining unilateral PA and validated the results against invasive adrenal vein sampling (AVS). Summary diagnostic accuracies were assessed using a bivariate random-effects model. Subgroup analyses, meta-regression and sensitivity analysis were performed to explore the possible sources of heterogeneity.

Result A total of 25 studies, involving a total of 4669 subjects, were identified. The overall analysis revealed a pooled sensitivity of 68% (95% CI: 61% to 74%) and specificity of 57% (95% CI 50% to 65%) for CT/MRI in identifying unilateral PA. Sensitivity was higher in the contrast-enhanced (CT) group versus the traditional CT group (77% (95% CI 66% to 85%) vs 58% (95% CI 50% to 66%). Subgroup analysis stratified by screening test for PA showed that the sensitivity of the aldosterone-to-renin ratio (ARR) group was higher than that of the non-ARR group (78% (95% CI 69% to 84%) vs 66% (95% CI 58% to 72%)). The diagnostic accuracy of PA patients aged ≤40 years was reported in four studies, and the overall sensitivity was 71%, with 79% specificity. Meta-regression revealed a significant impact of sample size on sensitivity and of age and study quality on specificity.

Conclusion CT/MRI is not a reliable alternative to invasive AVS without excellent sensitivity or specificity for correctly identifying unilateral PA. Even in young patients (≤40 years), 21% of patients would have undergone unnecessary adrenalectomy based on imaging results alone.

INTRODUCTION

Primary aldosteronism (PA) is one of the most common causes of endocrine hypertension, with a prevalence of approximately 20% in patients with resistant hypertension, 10% in those with severe hypertension and 6% in those with uncomplicated hypertension.¹

Strengths and limitations of this study

- This study is the first meta-analysis to synthesise the evidence regarding the diagnostic value of adrenal imaging for primary aldosteronism classification and demonstrated that CT/MRI is not a reliable alternative to invasive adrenal vein sampling (AVS) even in young patients (≤40 years).
- The main methodological limitations of this systematic review and meta-analysis are the exclusion of unpublished high-quality trials and foreign-language publications.
- Another potential limitation is that we encountered different AVS methods and large variation in the lateralisation criteria, which might also have affected the results for diagnostic accuracy.

Accumulating clinical and epidemiological evidence suggests that PA amplifies cardiovascular and cerebrovascular complications beyond essential hypertension prior to treatment, even after controlling elevated blood pressure.^{2,3} However, patients with unilateral resected PA have slightly better risk profiles than matched essential hypertensive patients. Patients with bilateral PA whose plasma renin activity is not suppressed have the same risk profiles as essential hypertensive patients; those whose renin activity remains suppressed have fourfold higher-risk profiles than controls, and titration of mineralocorticoid receptor antagonist therapy to raise renin might reduce this excess risk.^{4,5} Accordingly, early diagnosis and specific treatment of affected patients are key steps for the reversal of target-organ damage and prevention of cardiovascular and cerebrovascular events.

Selection of the most appropriate therapeutic strategy for patients with PA requires a distinction between unilateral and bilateral forms of PA. The former requires a unilateral adrenalectomy, mainly entailing

aldosterone-producing adenoma and, less commonly, unilateral adrenal hyperplasia. In contrast, the latter, also known as idiopathic hyperaldosteronism, is optimally treated with target medical therapy.³ Regarding the differentiation of the unilateral and bilateral subtypes, all current clinical practice guidelines recommend adrenal vein sampling (AVS) as the standard procedure for subtype diagnosis.^{6,7} However, several shortcomings of AVS have been reported, such as its technical challenges, invasive nature, poorly standardised procedures, high cost and lack of availability. Thus, it is urgent to explore alternative diagnostic methods without sacrificing accuracy.

Adrenal imaging with CT or MRI is recommended as the first step for subtype classification given the ease of performance and relative accessibility.⁸ By now, numerous studies have evaluated the diagnostic performance of CT/MRI in subtype diagnosis of PA, but the results have been inconsistent. Moreover, all these studies were limited by small sample sizes in a single centre, which limited the credibility of the results. In this context, systematic reviews and meta-analyses have the benefit of increasing the sample size, generating more precise results, which have been widely applied in clinical studies.⁹ In 2009, one systematic review reported that CT/MR-based diagnoses were discordant with AVS results in 37.8% of PA patients.¹⁰ However, the conclusions may not be reliable because of the potential for bias and concerns regarding the comparability of the included studies. Moreover, several additional studies were reported after this systematic review. We, thus, performed a comprehensive meta-analysis of all the available studies to evaluate the diagnostic value of adrenal imaging (CT/MRI) for subtype classification of PA.

METHOD

Patient and public involvement

Patient and public involvement

Search strategy

The study followed the guidelines specified in the Preferred Reporting Items for Systematic Reviews and Meta-Analyses Protocols.¹¹ We searched the PubMed, EMBASE and Cochrane Library databases from 1 January 2000 to 1 February 2020, using the following terms in combination, as both Medical Subject Headings (MeSH) or Emtree terms and text words: “primary aldosteronism”, “adrenal vein sampling” and “hyperaldosteronism”. The electronic search strategy for PubMed is shown in online supplemental table S1. To reflect modern practice, we decided to limit the publication date to after 1 January 2000. We searched articles published in English, and the references of relevant studies were also searched. All studies were carefully examined to exclude overlapping or potential duplicate data.

Eligibility criteria

We included a study if: (1) it used CT or MRI as a diagnostic test for PA subtyping; (2) it used AVS as the standard of reference. Successful AVS should be determined by calculating the selectivity index (SI), defined as the adrenal/peripheral vein cortisol ratio. Unilateral PA should be determined by calculating the lateralisation index (LI), defined as the aldosterone/cortisol ratio between the dominant and the non-dominant adrenal gland and (3) absolute numbers of true-positive, true-negative, false-positive and false-negative results were provided or could be derived. Identified studies had to be independent. In the case of multiple reports on the same population or subpopulation, the most recent or comprehensive information was used.

Data extraction and quality assessment

Data extraction from the eligible studies was performed by two independent investigators (YZ and PW) using a standardised data extraction form. The form included the following characteristics of each trial: first author's name and year of publication; study population characteristics, including sample size, geographical location, mean age and sex; diagnostic criteria characteristics, including screening test and confirmatory test for PA; AVS characteristics, including with/without adrenocorticotropic hormone (ACTH) stimulation, SI and LI; diagnostic test characteristics, including imaging methodology and whether contrast was administered. Differences between reviewers were resolved by discussion and consensus when necessary.

The methodological quality of the identified studies was assessed by two independent reviewers (YZ and PW) using the modified Quality Assessment of Diagnostic Accuracy Studies-2 (QUADAS-2) criteria. If a study was judged as ‘low’ on all domains relating to bias or applicability, then it was judged to be a high-quality study. If a study was judged to be ‘high’ and/or ‘unclear’ in more than one domain, then it was judged as a low-quality study. If a study was judged to be ‘unclear’ in one domain, it was considered an unclear-quality study.¹² Discrepancies were resolved by discussion and consensus.

Statistical analysis

Measures of diagnostic accuracy are reported as point estimates with 95% CIs. Sensitivity, specificity, the positive likelihood ratio (+LR) and the negative likelihood ratio (−LR) were modelled based on the true-positive, true-negative, false-positive and false-negative rates for each trial.¹³ The ratio of +LR to −LR was combined in a single global accuracy measure, the diagnostic OR (DOR). Summary sensitivity, specificity, +LRs, −LRs and DORs were assessed using a bivariate random-effects model. The approach assumes bivariate normal distributions for the logit transformations of sensitivity and specificity from the individual studies. These bivariate models can be analysed using linear mixed model techniques that are now widely available in statistical packages, such as STATA

gllamm.^{14 15} A hierarchical summary receiver operating characteristic curve analysis was performed, yielding point estimates for each trial and pooled characteristics, including the 95% prediction region and the 95% confidence region.

Sources of statistical heterogeneity were explored by subgroup analyses, sensitivity analysis and meta-regression analysis,¹⁶ which involved the I^2 statistic; the following interpretation was applied for I^2 : <50%=low heterogeneity, 50% to 75%=moderate heterogeneity and >75%=high heterogeneity.

Several studies demonstrated that MRI has poorer resolution and slower acquisition than CT, with a risk of respiratory artefacts and that MRI is inferior to adrenal CT in PA subtype evaluation.^{17–20} Contrast materials can improve the visibility of adrenal structures imaged by CT and MRI scans and might have a positive effect on diagnosis accuracy.²¹ Thus, imaging methods and contrast materials were thought to be confounders for subgroup analyses. Moreover, a large sample size may represent experienced interventional radiologists and support the credibility of the included studies. Thus, a small sample size was thought of as another confounder for subgroup analyses. The different diagnostic criteria for PA, the AVS procedure (with or without ACTH stimulation), different cut-offs for the LI criteria and methodological quality might also affect the results for diagnosis accuracy.²⁰ Therefore, we also performed subgroup analyses stratified by these parameters. Thus, subgroup analyses were performed by the following factors: imaging methodology (CT or CT/MRI), contrast use, AVS procedure (with or without ACTH stimulation), cut-off value for the LI (2 or 4), diagnostic criteria for PA, sample size (divided by 100 subjects) and methodological quality (high quality, low quality and unclear quality).

Potential publication bias was examined using the Deeks test.²² The Cohen κ test was employed to assess the inter-rater reliability between two observers for quality assessment. If there was not agreement, a third reviewer was involved to resolve disagreements, and final decisions were determined by consensus. Statistical analyses were performed using Stata V.13.0 (StataCorp) and Review Manager V.5.3.

RESULTS

Study selection

After removal of 548 duplicates, the systematic review generated 1022 references that were screened according to titles and abstracts for possible inclusion. Among them, 962 studies were excluded for the following reasons: 489 studies were not relevant; 280 studies were reviews or practice guidelines; 92 studies did not include humans and 101 studies were case reports/letters. After screening, 60 studies were identified as being potentially eligible, and their full texts were retrieved for detailed evaluation. A total of 35 studies were excluded for the following reasons: data to compute diagnostic accuracy were not provided

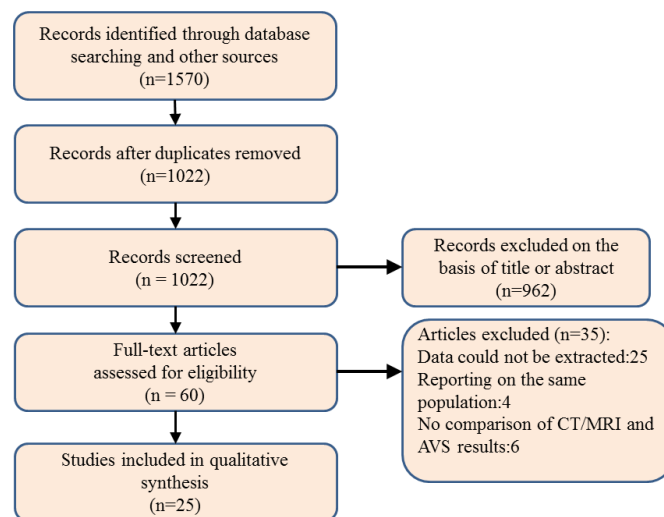


Figure 1 Flow diagram of the review process. AVS, adrenal vein sampling.

or could not be derived (25 papers), reporting on the same population (4 papers) and no comparison of CT/MRI and AVS results in individual patients (6 papers). Finally, 25 articles were deemed eligible and analysed in our meta-analysis^{17–20 23–43} (figure 1).

Study characteristics

Overall, a total of 4669 patients (mean age of 51 years; 54% male) from 25 articles were included. The sample sizes of the identified studies ranged from 35 to 1591, with the largest study recruiting over 1000 participants.⁴¹ Five studies including 724 participants underwent cross-sectional imaging either CT or MRI, and the remaining 20 studies including 3945 patients only used CT scans (eight studies administered contrast material). Seventeen studies performed AVS with ACTH stimulation, seven studies performed it without ACTH stimulation, and the remaining one study provided the above two methods. The aldosterone-to-renin ratio (ARR) was used as a screening tool for PA in 21 of the included articles, and an ARR >20 was commonly used as the threshold for a positive PA screening. The remaining four studies did not use the ARR as a screening test for PA. In 12 articles, a salt-loading test was performed to confirm the diagnosis of PA. Eight studies used additional options, including the fludrocortisone suppression test, captopril challenge test, upright-furosemide loading test and postural stimulation test, as a confirmatory test for PA. The diagnosis of PA was not confirmed in the remaining five studies by one of the confirmatory tests. The 2016 Endocrine Society Guideline recommends more strict criteria for the LI (2.0 or greater under unstimulated conditions and/or 4 for ACTH stimulation) and SI (2.0 or greater under unstimulated conditions and/or 3 for ACTH stimulation).⁸ In the meta-analysis, one included study used less permissive criteria for the LI,⁴² and six included studies used less permissive criteria for the

SI.^{20 27–30 42} The threshold of the SI was not accessible in four studies,^{18 23 33 34} and it was not accessible for the LI in two studies.^{1 26} Further details about the eligible and analysed studies are shown in [table 1](#) and online supplemental table S2.

Quality assessment

Overall, the identified studies were of excellent quality in terms of applicability and risk of bias. Out of 175 QUADAS-2 items (25 articles×7 items), the 172 (98%) were agreed on by the two reviewers, with an inter-rater agreement of $\kappa=0.9$. [Figure 2](#) summarises the QUADAS-2 assessment, and online supplemental table S3 displays each of the 25 individual QUADAS-2 evaluations.

The risk of bias from the reference standard was high in three studies,^{28 38 42} and it was unclear in five studies^{18 20 25 26 30} because it was not clear whether the reference standard was interpreted without knowledge of the adrenal imaging results or whether the cut-off values of the SI and LI correctly classified the target condition. The risk of bias regarding flow and timing was unclear in six studies^{20 24 25 30 39 41} because the time interval between the index test and the reference standard was unclear; it was high in one study²³ because not all patients had the same reference standard ([figure 2](#)). Finally, 13 studies were considered to be high-quality studies,^{17 19 27 29 31–38 43} 7 were considered low-quality studies^{20 23 25 28 30 38 42} and 5 were unclear-quality studies.^{18 24 26 39 41}

Overall analysis

Using the bivariate model, statistical heterogeneity was found for sensitivity ($I^2=86.9\%$; $p=0.001$), specificity ($I^2=86.9\%$; $p=0.00$), the positive LR ($I^2=76.3\%$; $p=0.00$), the negative LR ($I^2=79.2\%$; $p=0.00$) and DOR ($I^2=100.0\%$; $p=0.00$), indicating high between-study heterogeneity for all pooled measures, which might compromise the credibility of the study.

In the overall analysis, the pooled sensitivity, specificity, positive LR, negative LR and DOR for adrenal imaging were 68% (95% CI 61% to 74%), 57% (95% CI 50% to 65%), 1.6 (95% CI 1.4 to 1.9), 0.56 (95% CI 0.47 to 0.68) and 3 (95% CI 2 to 4), respectively ([figures 3 and 4](#)).

Subgroup analyses

Subgroup analysis, stratified by the imaging methodology, found more favourable specificity (60%) for CT than CT/MRI (45%). Notably, subgroup analysis showed an increase in sensitivity when contrast material was administered during the CT scan, compared with the traditional CT group (77% vs 58%). There was low heterogeneity detected on sensitivity in the CT/MRI group ($I^2=30\%$). However, the heterogeneity was high in all other groups, regardless of sensitivity or specificity ($I^2>75\%$).

Subgroup analysis based on AVS procedure (with or without ACTH stimulation) revealed a slight decrease in sensitivity when ACTH was administered during the AVS procedure (66% vs 70%). Sensitivity and specificity were

higher when LI was ≥ 4 vs LI was ≥ 2 . However, a large degree of heterogeneity was observed in all groups (I^2 all $>75\%$).

Subgroup analysis stratified by screening test showed that the sensitivity of the ARR group was higher than that of the non-ARR group (78% vs 66%). The heterogeneity was high ($I^2=87.7\%$) in the ARR subgroup, whereas it disappeared (0%) in the non-ARR subgroup. Regarding specificity, the heterogeneity was high for both groups (86.2% vs 89.1%). Subgroup analysis stratified by the confirmatory test for PA demonstrated an increase in sensitivity (71% vs 57%) and a slight decrease in specificity (60% vs 66%) for the salt-loading test group compared with additional options group, with moderate to high heterogeneity observed in all the above groups (I^2 all $>50\%$).

Subgroup analysis based on methodological quality (high quality, low quality and unclear quality) revealed that there was low heterogeneity for sensitivity in all the above groups (I^2 all $<50\%$). The diagnostic pooled sensitivity for the high-quality group was the highest, followed by the unclear-quality group and the low-quality group (78% vs 62% vs 48%). The unclear-quality group had the highest specificity, followed by the high-quality group and the low-quality group (69% vs 62% vs 51%). Regarding specificity, heterogeneity was decreased but still high in all the groups.

There were four studies that reported diagnostic accuracy in PA patients with an age of 40 years or younger. Using the bivariate model, the pooled sensitivity and specificity were 71% (95% CI 54% to 84%) and 79% (95% CI 37% to 96%), respectively, with moderate heterogeneity (53.1% vs 70.1%) (online supplemental figure S1). Summary estimates for pooled measures of diagnostic accuracy are shown in [table 2](#).

Meta-regression analysis

Results of meta-regression analysis showed that the sample size was the only covariate with a negative effect on sensitivity. Additionally, there was a significant interaction between lower age, as well as high methodological quality, and higher specificity of CT/MRI for the detection of unilateral forms of PA (online supplemental figure S2).

Sensitivity analysis

Goodness-of-fit and bivariate normality analyses (online supplemental figure S3A,B) showed that the bivariate model was moderately robust. Influence analysis and outlier detection identified four outliers (online supplemental figure S3C,D). After we excluded these outliers, the overall results did not change significantly, which suggested that the results of this study were statistically reliable ([table 2](#)).

Publication bias

Neither Deeks' funnel plot nor Deeks test ($t=0.46$, $p=0.65$) showed evidence of publication bias (online supplemental figure S4).

Table 1 Study characteristics

Study, year	Location	Male (%)	Age (years)	Sample	Imaging methodology	Contrast used	AVS characteristics	Screening test for PA (ng/ml)	Confirmatory test for PA
Li <i>et al</i> 2019 ⁴³	China	61 (50)	48.5	122	CT	yes	without ACTH, SI ≥ 2	ARR	Captopril test or salt-loading test
Campbell <i>et al</i> 2019 ¹⁹	USA	45 (61)	55.6	74	CT or MRI	no	with ACTH, SI ≥ 5 , LI ≥ 4	not ARR	no
Aono <i>et al</i> 2019 ⁴²	Japan	NA	56	317	CT	no	with ACTH, SI ≥ 2 , LI ≥ 2	ARR > 20	Captopril test, furosemide upright test or salt-loading test
Sam <i>et al</i> 2019 ²⁰	Canada	201 (59)	52.1	342	CT or MRI	yes	with ACTH, SI ≥ 2 , LI ≥ 4	ARR	no
Umakoshi <i>et al</i> 2018 ⁴¹	Japan	762 (47.9)	53	1591	CT	yes	with ACTH, SI ≥ 5 , LI ≥ 4	ARR	Captopril test, furosemide upright test or salt-loading test
Nanba <i>et al</i> 2017 ⁴⁰	USA	87 (59)	54	147	CT	no	with ACTH, SI ≥ 5 , LI ≥ 4	ARR	Salt-loading test
Kamemura <i>et al</i> 2017 ³⁹	Japan	177 (45)	54	393	CT	no	with ACTH, SI ≥ 5 , LI ≥ 4	ARR	Captopril test, furosemide upright test or salt-loading test
Zhu <i>et al</i> 2016 ³⁸	China	NA	46	394	CT	no	without ACTH, SI ≥ 3 , LI ≥ 2	ARR	Fludrocortisone test or salt-loading test
Pedersen <i>et al</i> 2016 ¹⁸	Denmark	24 (54)	51	45	CT or MRI	no	without ACTH, SI NA	ARR	Fludrocortisone test or postural test
Kocjan <i>et al</i> 2016 ³⁷	Slovenia	46 (69)	56	67	CT	no	with ACTH, SI ≥ 5 , LI > 4	ARR	Salt-loading test
Asmar <i>et al</i> 2015 ¹⁷	USA	148 (63)	55	235	CT or MRI	no	with ACTH, SI ≥ 5 , LI ≥ 4	ARR	no
Riester <i>et al</i> 2014 ³⁵	Germany	NA	35	28	CT or MRI	no	without ACTH, SI ≥ 2 , LI ≥ 4	ARR	Salt-loading test
Sze <i>et al</i> 2014 ³⁶	UK	42 (56)	50.5	75	CT	yes	with ACTH, SI ≥ 5 , LI ≥ 4	not ARR	salt-loading test
Küipers <i>et al</i> 2012 ³⁰	France	53 (61)	46	87	CT	no	with ACTH, SI ≥ 2 , LI ≥ 4	ARR	Salt-loading test
Lau <i>et al</i> 2012 ³²	UK	24 (64)	51.8	39	CT	no	with ACTH, SI ≥ 5 , LI ≥ 4	not ARR	salt-loading test
Burton <i>et al</i> 2012 ³⁴	UK	NA	50.9	40	CT	Yes	without ACTH, LI ≥ 4	not ARR	no
Salem <i>et al</i> 2012 ³³	UK	16 (44)	44.7	38	CT	no	without ACTH, LI ≥ 4	ARR	Salt-loading test
Oh <i>et al</i> 2012 ³¹	Korea	45 (52)	50.7	86	CT	yes	with ACTH, SI ≥ 3 , LI > 4	ARR	Salt-loading test

Continued

Table 1 Continued

Study, year	Location	Male (%)	Age (years)	Sample	Imaging methodology	Contrast used	AVS characteristics	Screening test for PA (ng/ml)	Confirmatory test for PA
Sarlon-Bartoli <i>et al</i> 2011 ²⁹	France	NA	52	58	CT	yes	without ACTH, SI \geq 1, LI $>$ 2	ARR	no
Mathur <i>et al</i> 2010 ²⁸	USA	63 (55)	50.6	114	CT	no	with ACTH, SI $>$ 2, LI \geq 4	ARR	Captopril test, posture test or salt-loading test
Mulatero <i>et al</i> 2008 ²⁷	Italy	NA	52.4	70	CT	yes	65 with ACTH; five without ACTH, SI $>$ 2, LI \geq 4	ARR	Salt-loading test
Minami <i>et al</i> 2008 ²⁶	Japan	12 (34)	54	35	CT	no	with ACTH SI / LI NA	ARR	Salt-loading test
Nwariaku <i>et al</i> 2006 ²⁵	USA	27 (67)	51	40	CT	yes	with ACTH, SI \geq 3, LI $>$ 4	ARR	Captopril test or posture test
Young <i>et al</i> 2004 ²⁴	USA	163 (84)	53	194	CT	no	with ACTH, SI $>$ 5, LI $>$ 4	ARR	Salt-loading test
Magill <i>et al</i> 2001 ²³	USA	27 (71)	51	38	CT	no	with ACTH, SI \geq 3, LI \geq 4	ARR	Salt-loading test

ACTH, adrenocorticotrophic hormone; ARR, aldosterone-to-renin ratio; AVS, adrenal vein sampling; LI, lateralisation index; NA, not available; PA, primary aldosteronism; SI, selectivity index.

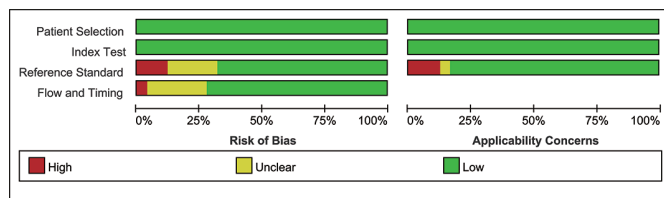


Figure 2 Assessment of methodological quality of included studies using the QUADAS-2 Criteria. Stacked bars represent the proportion of studies with a high (red), or unclear (yellow) or low (green) risk of bias and applicability concerns. QUADAS-2, Quality Assessment of Diagnostic Accuracy Studies-2 criteria.

DISCUSSION

Main findings

The accurate differentiation of unilateral and bilateral PA is critical for optimal clinical management. Although AVS is the 'gold-standard' test for subtype diagnosis,⁸ numerous studies have investigated the underlying diagnostic value of CT/MRI for subtype diagnosis due to several insurmountable shortcomings of AVS. The present meta-analysis, involving 4669 individuals from 25 studies, demonstrated that CT/MRI has poor sensitivity (68%) and specificity (57%) in subtype classification when AVS was used as the reference standard.

In the subtype diagnosis of PA, AVS was initially used in the 1960s. Subsequently, CT was adopted as the primary method for distinguishing the unilateral and bilateral forms of PA. Owing to its less invasive nature, lower cost and wide availability, many physicians prefer to perform CT/MRI as the first, and sometimes only, investigation of PA subtype. However, its sensitivity and specificity vary widely. The reported sensitivities ranged from 29%³ to 94%,³¹ and the reported specificities ranged from 18%¹⁹ to 87%.³⁰ Although the sensitivities reportedly exceeded 80% in five studies,^{19 27 29 31 37} relatively poor specificities were reported, with only one study showing a specificity of 72%.⁴² Similarly, three studies reported the sensitivities

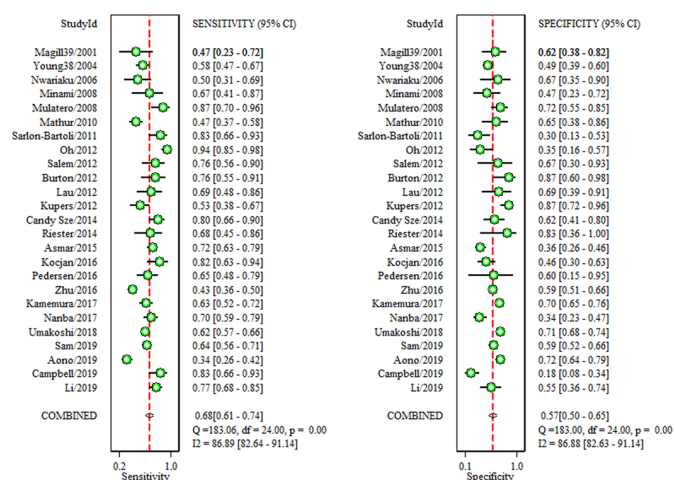


Figure 3 Forest plots of sensitivity and specificity of adrenal imaging compared with AVS. Horizontal lines are the 95% CIs. AVS, adrenal vein sampling.

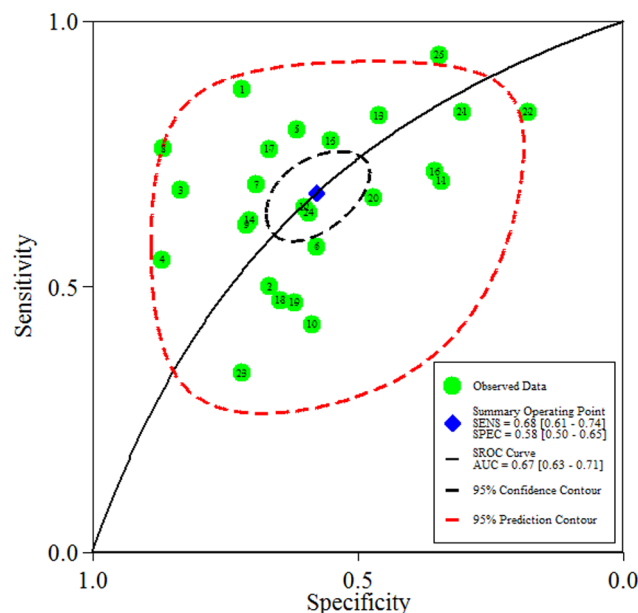


Figure 4 Hierarchical SROC plot showing average sensitivity and specificity estimate of the study results with 95% confidence region. The 95% prediction region represents the confidence region for a forecast of the true sensitivity (SENS) and specificity (SPEC) in a future study. AUC, area under the curve; SROC, summary receiver operating characteristic.

to be over 80%,^{30 34 35} but the specificities were reported to be lower than 76%.³⁴ The present meta-analysis showed that the pooled sensitivity was 68% and the specificity was 57%, which means that treatment decisions based on the presence of unilateral disease on CT/MRI alone could result in inappropriate unilateral adrenalectomy in 43% of patients. Basing this decision on CT/MRI alone would miss the possibility of a potentially curative procedure by surgery in 32% of patients. However, failure to make an early diagnosis and provide specific treatment for PA places these patients at higher risk of irreversible renal and cardiovascular damage. Our results suggest that CT/MRI does not have satisfactory diagnostic performance in classifying the subtypes of PA.

Stimulation with ACTH during the AVS procedure was introduced in 1979 and remains popular at many centres. Today, the AVS procedure, with or without ACTH stimulation, is still controversial.⁴⁴ The present meta-analysis revealed that there was no significant difference between the two AVS procedures (with or without ACTH stimulation) in terms of the diagnostic accuracy of CT/MRI to identify unilateral PA. In theory, the application of more stringent lateralisation criteria (a condition that is more likely to capture true cases of unilateral PA) would result in increased sensitivity and decreased specificity of CT/MRI to identify unilateral PA. However, our analysis demonstrated that stricter thresholds for determining lateralisation on AVS would result in higher sensitivity and specificity, which is not completely consistent with the theoretical situation.

Table 2 Pooled summary results by subgroups

Subgroups	No of studies	Sensitivity (95% CI)	Sensitivity with I ² , %	Specificity (95% CI)	Specificity with I ² , %
Total	25	68 (61 to 74)	86.9%	57 (50 to 65)	86.9
Age					
≤40 years	4	71 (54 to 84)	53.1%	79 (39 to 96)	70.1
>40 years	21	68 (60 to 75)	87.4%	57 (49 to 64)	87.4
AVS procedure					
With ACTH	17	66 (57 to 73)	86.5%	56 (46 to 65)	90.5
Without ACTH	7	70 (58 to 79)	90.3%	60 (45 to 74)	79.2
Cutoff values of LI					
LI≥2	4	61 (37 to 80)	95.5%	54 (38 to 68)	88.3
LI≥4	18	69 (62 to 75)	76.4%	59 (50 to 68)	89.0
Screening test for PA					
ARR	21	66 (58 to 72)	87.7%	58 (50 to 65)	86.2
Not ARR	4	78 (69 to 84)	0%	59 (29 to 84)	89.1
Confirmatory test for PA					
Salt-loading test	12	71 (62 to 80)	78.6%	60 (49 to 70)	74.9
Additional options	8	57 (46 to 67)	90.6%	66 (60 to 72)	66
No	5	72 (64 to 79)	55.8%	42 (24 to 63)	90.3
Imaging methodology					
CT	20	67 (59 to 74)	88.6%	60 (53 to 67)	82.5
Contrast CT	8	77 (66 to 85)	86.4%	60 (49 to 69)	82.2
Nocontrast CT	12	58 (49 to 66)	83.8%	60 (51 to 68)	81.8
CT/MRI	5	69 (62 to 76)	30%	45 (27 to 64)	87.9
Quality of studies					
High-quality studies	13	78 (73 to 83)	48.6%	51 (39 to 63)	78.6
Unclear-quality studies	6	62 (58 to 65)	0%	62 (54 to 70)	85.1
Low-quality studies	6	44 (38 to 50)	46.4%	69 (60 to 78)	69.2
Sample size					
≥100	10	59 (51 to 67)	90.6%	58 (49 to 66)	92.1
<100	15	74 (67 to 81)	71.3%	60 (47 to 71)	78.1
Outlier excluded	20	65 (60 to 70)	77.1%	59 (52 to 66)	85.2

ACTH, adrenocorticotrophic hormone; ARR, aldosterone-to-renin ratio; AVS, adrenal vein sampling; LI, lateralisation index; PA, primary aldosteronism.

Although the overall analysis suggested that CT/MRI does not have satisfactory diagnostic performance in classifying the subtypes of PA, the results should be interpreted with caution because of moderate to high heterogeneity due to several underlying confounders. First, the screening test and confirmatory test for PA may influence the results. In our meta-analysis, some patients did not undergo a screening test and confirmatory test for PA, which is the diagnostic reference standard test, and some of them might not have PA. Generally, inadvertently including patients without PA should increase the specificity of CT/MRI in identifying unilateral PA, as these subjects would not show lateralisation on AVS or a unilateral aldosteronoma on CT/MRI. However, although the

difference in the screening test was responsible for the heterogeneity of sensitivity and specificity to some extent, according to our analysis, there is no evidence to indicate that the confirmatory test influences the specificity of CT/MRI.

Second, meta-regression analysis showed that the heterogeneity of specificity may partly be due to age. Given that non-functioning adrenocortical adenomas ('incidentaloma') are relatively uncommon in young people (≤40 years), the 2009 guidelines for managing PA contended that younger patients with an unequivocal biochemical diagnosis of PA and a clear-cut unilateral adenoma on adrenal CT scan should proceed directly to surgery, whereas the AVS procedure may be

skipped.⁴⁵ Among studies included in the present meta-analysis, four reported the diagnostic accuracy of CT/MRI in identifying unilateral PA in patients ≤ 40 years. By combining these four studies, our results demonstrated that although the sensitivity (71%) and specificity (79%) were improved, the diagnostic performance was still unsatisfactory because 21% of patients would have undergone unnecessary adrenalectomy based on imaging results alone. In 2016, the updated clinical practice guidelines were published and suggested that the age cut-off for sparing AVS be 35 years.⁸ Regarding patients aged ≤ 35 years, several retrospective studies have evaluated the diagnostic value of CT. The reported rate of concordance between CT and AVS ranges from 59% to 90%.^{19 38 41} Based on these data, it still seems that CT cannot replace AVS in patients aged ≤ 35 years. However, due to the lack of numbers of false-positives and true-negatives, we did not perform a pooled analysis. Further studies are needed to clarify the diagnostic value of CT in patients aged ≤ 35 years.

As mentioned above, although adrenal imaging is not a reliable method to differentiate subtypes of PA, it does not mean that CT/MRI must be wrong and should not be used as a basis for clinical management. In centres without AVS facilities currently, what should a physician do? In the past few years, there has been rapidly growing interest in testing the utility of hybrid steroids, such as 18-oxocortisol/18-hydroxycortisol, for PA subtypes and the results demonstrated that levels of 18-oxocortisol/18-hydroxycortisol plus an adenoma on CT/MRI might be of more assistance in those centres without AVS facilities, especially in Japan and China, given their very high percentage of KCNJ5 mutations.^{46–48} It is hoped that perhaps the possibility of multi-steroid fingerprints in peripheral blood samples that distinguish unilateral from bilateral PA with a high degree of accuracy can substantially reduce or replace the use of lateralisation by AVS.

Limitations

The present meta-analysis has several limitations. First, there was great heterogeneity among the included studies, which might have compromised the credibility. The results of the subgroup analyses and meta-regression suggested that the screening test for PA, age, study quality, sample size and other unknown factors may also contribute to the aforementioned heterogeneity. However, the results from the subgroup analyses and sensitivity analysis all confirmed the robustness of our meta-analysis's results. Second, a minority of study participants underwent cross-sectional imaging with either CT or MRI, but absolute numbers were not provided or could not be derived based on the specific imaging methodology used, which limited our ability to identify which imaging methodology can provide more accurate diagnostic performance. In addition, the possibility of selection bias that is present in all meta-analysis cannot be overlooked.

CONCLUSION

Based on these analyses, we conclude that CT/MRI has poor sensitivity (68%) and specificity (57%) in the detection of unilateral PA when AVS is used as the reference standard. Even in young patients (≤ 40 years), 21% would have undergone unnecessary adrenalectomy based on imaging results alone. Given these findings, we recommend routinely referring all patients for AVS, regardless of age and imaging results, if the centre has access to AVS. However, due to moderate to high heterogeneity, our study should be interpreted with caution, and further high-quality studies with larger sample sizes are needed.

Contributors PW is the guarantor. All authors provided substantial contribution to conception and design of the project; drafted and revised the manuscript. YZ led the literature search, and completed the study selection, data extraction and critical appraisal with PW. LJ accepts responsibility for the integrity of the data analyses. FR led the drafting of all sections of the article in consultation with all of the coauthors. DW, SC and PZ provided substantial contributions to the background, critical appraisal of prior studies and interpretation of meta-analysis findings. PZ provided substantial contribution to the methods section.

Funding This systematic review and meta-analysis are funded by the National Natural Sciences Foundation of China (NSFC) (Grant NO. 81970262).

Competing interests None declared.

Patient and public involvement statement Patients and the public were not involved in the design or conduct of the study.

Patient consent for publication Not required.

Provenance and peer review Not commissioned; externally peer reviewed.

Data availability statement Data are available in a public, open access repository. Extra data can be accessed via the Dryad data repository at <http://datadryad.org/> with the doi: 10.5061/dryad.jm63xsj8t.

Supplemental material This content has been supplied by the author(s). It has not been vetted by BMJ Publishing Group Limited (BMJ) and may not have been peer-reviewed. Any opinions or recommendations discussed are solely those of the author(s) and are not endorsed by BMJ. BMJ disclaims all liability and responsibility arising from any reliance placed on the content. Where the content includes any translated material, BMJ does not warrant the accuracy and reliability of the translations (including but not limited to local regulations, clinical guidelines, terminology, drug names and drug dosages), and is not responsible for any error and/or omissions arising from translation and adaptation or otherwise.

Open access This is an open access article distributed in accordance with the Creative Commons Attribution Non Commercial (CC BY-NC 4.0) license, which permits others to distribute, remix, adapt, build upon this work non-commercially, and license their derivative works on different terms, provided the original work is properly cited, appropriate credit is given, any changes made indicated, and the use is non-commercial. See: <http://creativecommons.org/licenses/by-nc/4.0/>.

ORCID iD

Yaqiong Zhou <http://orcid.org/0000-0002-9441-0291>

REFERENCES

- Monticone S, Burrello J, Tizzani D, *et al.* Prevalence and Clinical Manifestations of Primary Aldosteronism Encountered in Primary Care Practice. *J Am Coll Cardiol* 2017;69:1811–20.
- Catena C, Colussi G, Nadalin E, *et al.* Cardiovascular outcomes in patients with primary aldosteronism after treatment. *Arch Intern Med* 2008;168:80–5.
- Pillai PR, Griffith M, Schwarcz MD, *et al.* Primary aldosteronism: cardiovascular risk, diagnosis, and management. *Cardiol Rev* 2020;28:84–91.
- Hundemer GL, Curhan GC, Yozamp N, *et al.* Cardiometabolic outcomes and mortality in medically treated primary aldosteronism: a retrospective cohort study. *Lancet Diabetes Endocrinol* 2018;6:51–9.
- Funder JW. Primary aldosteronism: where are we now? Where to from here? *Horm Metab Res* 2020;52:459–66.

- 6 Williams TA, Reincke M. Management of endocrine disease: diagnosis and management of primary aldosteronism: the endocrine Society guideline 2016 revisited. *Eur J Endocrinol* 2018;179:R19–29.
- 7 Byrd JB, Turcu AF, Auchus RJ. Primary aldosteronism: practical approach to diagnosis and management. *Circulation* 2018;138:823–35.
- 8 Funder JW, Carey RM, Mantero F, et al. The management of primary aldosteronism: case detection, diagnosis, and treatment: an endocrine Society clinical practice guideline. *J Clin Endocrinol Metab* 2016;101:1889–916.
- 9 Ruco A, Dossa F, Timmouth J, et al. Social media and mobile health technology for cancer screening: a systematic review and meta-analysis protocol. *BMJ Open* 2020;10:e035411.
- 10 Kempers MJE, Lenders JWM, van Outheusden L, et al. Systematic review: diagnostic procedures to differentiate unilateral from bilateral adrenal abnormality in primary aldosteronism. *Ann Intern Med* 2009;151:329–37.
- 11 Liberati A, Altman DG, Tetzlaff J, et al. The PRISMA statement for reporting systematic reviews and meta-analyses of studies that evaluate health care interventions: explanation and elaboration. *Ann Intern Med* 2009;151:W65–94.
- 12 Whiting PF, Rutjes AWS, Westwood ME, et al. QUADAS-2: a revised tool for the quality assessment of diagnostic accuracy studies. *Ann Intern Med* 2011;155:529–36.
- 13 Devillé WL, Buntinx F, Bouter LM, et al. Conducting systematic reviews of diagnostic studies: didactic guidelines. *BMC Med Res Methodol* 2002;2:9.
- 14 Reitsma JB, Glas AS, Rutjes AWS, et al. Bivariate analysis of sensitivity and specificity produces informative summary measures in diagnostic reviews. *J Clin Epidemiol* 2005;58:982–90.
- 15 Chu H, Guo H, Zhou Y. Bivariate random effects meta-analysis of diagnostic studies using generalized linear mixed models. *Med Decis Making* 2010;30:499–508.
- 16 Lijmer JG, Bossuyt PMM, Heisterkamp SH. Exploring sources of heterogeneity in systematic reviews of diagnostic tests. *Stat Med* 2002;21:1525–37.
- 17 Asmar M, Wachtel H, Yan Y, et al. Reversing the established order: should adrenal venous sampling precede cross-sectional imaging in the evaluation of primary aldosteronism? *J Surg Oncol* 2015;112:144–8.
- 18 Pedersen M, Karlsen MA, Ankjærgaard KL, et al. Primary hyperaldosteronism diagnosed with adrenal vein sampling. Characteristics and follow-up after adrenalectomy in a Danish study. *Scand J Clin Lab Invest* 2016;76:45–50.
- 19 Campbell RA, Young DS, Shaver CN, et al. Influence of adrenal venous sampling on management in patients with primary aldosteronism independent of lateralization on cross-sectional imaging. *J Am Coll Surg* 2019;229:116–24.
- 20 Sam D, Kline GA, So B, et al. Discordance between imaging and adrenal vein sampling in primary aldosteronism irrespective of interpretation criteria. *J Clin Endocrinol Metab* 2019;104:1900–6.
- 21 Nandra G, Duxbury O, Patel P, et al. Technical and interpretive pitfalls in adrenal imaging. *Radiographics* 2020;40:1041–60.
- 22 Deeks JJ, Macaskill P, Irwig L. The performance of tests of publication bias and other sample size effects in systematic reviews of diagnostic test accuracy was assessed. *J Clin Epidemiol* 2005;58:882–93.
- 23 Magill SB, Raff H, Shaker JL, et al. Comparison of adrenal vein sampling and computed tomography in the differentiation of primary aldosteronism. *J Clin Endocrinol Metab* 2001;86:1066–71.
- 24 Young WF, Stanson AW, Thompson GB, et al. Role for adrenal venous sampling in primary aldosteronism. *Surgery* 2004;136:1227–35.
- 25 Nwariaku FE, Miller BS, Auchus R, et al. Primary hyperaldosteronism: effect of adrenal vein sampling on surgical outcome. *Arch Surg* 2006;141:497–502.
- 26 Minami I, Yoshimoto T, Hirono Y, et al. Diagnostic accuracy of adrenal venous sampling in comparison with other parameters in primary aldosteronism. *Endocr J* 2008;55:839–46.
- 27 Mulatero P, Bertello C, Rossato D, et al. Roles of clinical criteria, computed tomography scan, and adrenal vein sampling in differential diagnosis of primary aldosteronism subtypes. *J Clin Endocrinol Metab* 2008;93:1366–71.
- 28 Mathur A, Kemp CD, Dutta U, et al. Consequences of adrenal venous sampling in primary hyperaldosteronism and predictors of unilateral adrenal disease. *J Am Coll Surg* 2010;211:384–90.
- 29 Sarlon-Bartoli G, Michel N, Taieb D, et al. Adrenal venous sampling is crucial before an adrenalectomy whatever the adrenal-nodule size on computed tomography. *J Hypertens* 2011;29:1196–202.
- 30 Küpers EM, Amar L, Raynaud A, et al. A clinical prediction score to diagnose unilateral primary aldosteronism. *J Clin Endocrinol Metab* 2012;97:3530–7.
- 31 Oh EM, Lee KE, Yoon K, et al. Value of adrenal venous sampling for lesion localization in primary aldosteronism. *World J Surg* 2012;36:2522–7.
- 32 Lau JHG, Sze WCC, Reznick RH, et al. A prospective evaluation of postural stimulation testing, computed tomography and adrenal vein sampling in the differential diagnosis of primary aldosteronism. *Clin Endocrinol* 2012;76:182–8.
- 33 Salem V, Hopkins TG, El-Gayar H, et al. Adrenal venous sampling as a diagnostic procedure for primary hyperaldosteronism: experience from a tertiary referral centre. *Hormones* 2012;11:151–9.
- 34 Burton TJ, Mackenzie IS, Balan K, et al. Evaluation of the sensitivity and specificity of (11)C-metomidate positron emission tomography (PET)-CT for lateralizing aldosterone secretion by Conn's adenomas. *J Clin Endocrinol Metab* 2012;97:100–9.
- 35 Riester A, Fischer E, Degenhart C, et al. Age below 40 or a recently proposed clinical prediction score cannot bypass adrenal venous sampling in primary aldosteronism. *J Clin Endocrinol Metab* 2014;99:E1035–9.
- 36 Sze WCC, Soh LM, Lau JH, et al. Diagnosing unilateral primary aldosteronism - comparison of a clinical prediction score, computed tomography and adrenal venous sampling. *Clin Endocrinol* 2014;81:25–30.
- 37 Kocjan T, Janez A, Stankovic M, et al. A new clinical prediction criterion accurately determines a subset of patients with bilateral primary aldosteronism before adrenal venous sampling. *Endocr Pract* 2016;22:587–94.
- 38 Zhu L, Zhang Y, Zhang H, et al. Comparison between adrenal venous sampling and computed tomography in the diagnosis of primary aldosteronism and in the guidance of adrenalectomy. *Medicine* 2016;95:e4986.
- 39 Kamemura K, Wada N, Ichijo T, et al. Significance of adrenal computed tomography in predicting laterality and indicating adrenal vein sampling in primary aldosteronism. *J Hum Hypertens* 2017;31:195–9.
- 40 Nanba AT, Nanba K, Byrd JB, et al. Discordance between imaging and immunohistochemistry in unilateral primary aldosteronism. *Clin Endocrinol* 2017;87:665–72.
- 41 Umakoshi H, Tsuiji M, Takeda Y, et al. Significance of computed tomography and serum potassium in predicting subtype diagnosis of primary aldosteronism. *J Clin Endocrinol Metab* 2018;103:900–8.
- 42 Aono D, Kometani M, Karashima S, et al. Primary aldosteronism subtype discordance between computed tomography and adrenal venous sampling. *Hypertens Res* 2019;42:1942–50.
- 43 Li S, Ren Y, Zhu Y, et al. The potential clinical application of a lower bilateral adrenal limb width ratio (L/RW) in patients with bilateral primary hyperaldosteronism. *Endocr Pract* 2019;25:830–5.
- 44 Laurent I, Astère M, Zheng F, et al. Adrenal venous sampling with or without adrenocorticotrophic hormone stimulation: a meta-analysis. *J Clin Endocrinol Metab* 2019;104:1060–8.
- 45 Zeiger MA, Thompson GB, Duh Q-Y, et al. The American association of clinical endocrinologists and American association of endocrine surgeons medical guidelines for the management of adrenal incidentalomas. *Endocr Pract* 2009;15(Suppl 1):1–20.
- 46 Taguchi R, Yamada M, Nakajima Y, et al. Expression and mutations of KCNJ5 mRNA in Japanese patients with aldosterone-producing adenomas. *J Clin Endocrinol Metab* 2012;97:1311–9.
- 47 Wang B, Li X, Zhang X, et al. Prevalence and characterization of somatic mutations in Chinese aldosterone-producing adenoma patients. *Medicine* 2015;94:e708.
- 48 Satoh F, Morimoto R, Ono Y, et al. Measurement of peripheral plasma 18-oxocortisol can discriminate unilateral adenoma from bilateral diseases in patients with primary aldosteronism. *Hypertension* 2015;65:1096–102.

Search strategies for PubMed

#1. ((aldosteronism) OR (primary hyperaldosteronism) OR (primary aldosteronism) OR (conn syndrome) OR (conn) OR (aldosterone-producing adenoma) OR (APA) OR (idiopathic hyperaldosteronism) OR (IHA) OR (primary adrenal hyperplasia) OR (PAH) OR (bilateral adrenal hyperplasia) OR (BAH))

#2. ((adrenal venous sampling) OR (AVS) OR (adrenal vein sampling) OR (adrenal vein) OR (venous sampling) OR (vein sampling) OR (adrenal venous))

#3. ("2000/01/01"[Date - Entry] : "2020/01/01"[Date - Entry])

#4. #1 AND #2 AND #3

Table S2: original data

Study, Year	Age (years)	TP	FP	FN	TN
Li et al. 2019	48.5	72	13	21	16
	<35	11	1	0	2
Campbell et al.2019	55.6	29	32	6	7
Aono et al. 2019	56	52	46	102	117
	<35	3	2	0	0
Sam et al.2019	52.1	101	75	57	109
Umakoshi et al.2018	53	297	322	185	787
	<35	27	3	0	0
Nanba et al.2017	54	58	42	25	22
Kamemura et al.2017	54	60	88	36	209
Zhu et al. 2016	46	87	79	116	112
	<40	30	35	1	0
Pedersen et al.2016	51	26	2	14	3
Kocjan et al. 2016	56	23	21	5	18
Asmar et al. 2015	55	96	65	38	36
	<40	9	5	6	3
Riester et al. 2014	35	15	1	7	5
Candy Sze et al. 2014	50.5	39	10	10	16
Küpers et al.2012	46	26	5	23	33
	<40	9	0	6	10
Lau et al. 2012	51.8	18	4	8	9
Burton et al. 2012	50.9	19	2	6	13
Salem et al.2012	44.7	22	3	7	6
Oh et al. 2012	50.7	59	15	4	8
Sarlon-Bartoli et al.2011	52	29	16	6	7
Mathur et al. 2010	50.6	46	6	51	11
Mulatero et al. 2008	52.4	27	11	4	28
Minami et al.2008	54	12	9	6	8
Nwariaku et al. 2006	51	14	4	14	8
Young et al. 2004	53	57	48	42	47
Magill et al.2001	51	8	8	9	13

TP: true-positive; TN: true-negative; FP: false-positive; FN: false-negative.

Table S3: The quality assessment for each study -QUADAS-2 results

Study	Risk of bias				Applicability Concerns		
	Patient Selection	Index Test	Reference Standard	Flow and Timing	Patient Selection	Index Test	Reference Standard
Li et al	☺	☺	☺	☺	☺	☺	☺
Campbell et al	☺	☺	☺	☺	☺	☺	☺
Aono et al	☺	☺	⊗	☺	☺	☺	⊗
Sam et al et al	☺	☺	?	?	☺	☺	☺
Umakoshi et al	☺	☺	☺	?	☺	☺	☺
Nanba et al	☺	☺	☺	☺	☺	☺	☺
Kamemura et al	☺	☺	☺	?	☺	☺	☺
Zhu et al	☺	☺	⊗	☺	☺	☺	⊗
Pedersen et al	☺	☺	?	☺	☺	☺	☺
Kocjan et al	☺	☺	☺	☺	☺	☺	☺
Asmar et al	☺	☺	☺	☺	☺	☺	☺
Riester et al	☺	☺	☺	☺	☺	☺	☺
Candy Sze et al	☺	☺	☺	☺	☺	☺	☺
Kúpers et al	☺	☺	?	?	☺	☺	☺
Lau et al	☺	☺	☺	☺	☺	☺	☺
Burton et al	☺	☺	☺	☺	☺	☺	☺
Salem et al	☺	☺	☺	☺	☺	☺	☺
Oh et al	☺	☺	☺	☺	☺	☺	☺
Sarlon-Bartoli et al	☺	☺	☺	☺	☺	☺	☺
Mathur et al	☺	☺	⊗	☺	☺	☺	☺
Mulatero et al	☺	☺	☺	☺	☺	☺	☺
Minami et al	☺	☺	?	☺	☺	☺	☺
Nwariaku et al	☺	☺	?	?	☺	☺	☺
Young et al	☺	☺	☺	?	☺	☺	☺
Magill et al	☺	☺	☺	⊗	☺	☺	☺

☺= low risk, ? = unclear risk, ⊗=high risk

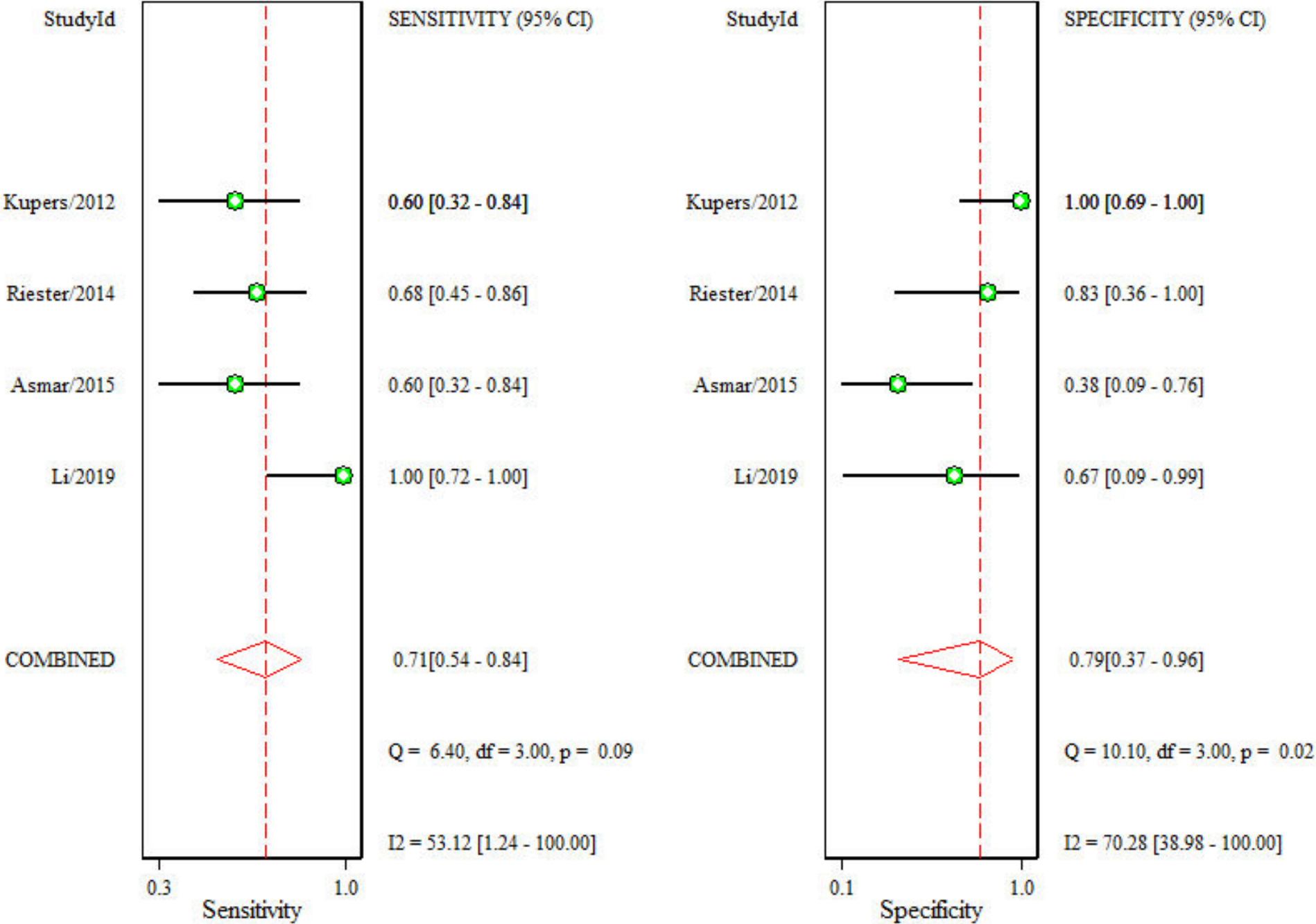
Supplementary Figure legends

Figure S1: Forest plots of sensitivity and specificity of adrenal imaging compared with AVS in young patients (≤ 40 years).

Figure S2: Graphical presentation of the generalised linear mixed model exploring the impact of selected variables on sensitivity and specificity of adrenal imaging.

Figure S3 Graphs for sensitivity analyses: a goodness of fit, b bivariate normality, c influence analysis, and d outlier detection.

Figure S4: Deeks' funnel plot for checking the publication bias. DOR: diagnostic OR.



Univariable Meta-regression & Subgroup Analyses

