

# Acquired renal arteriovenous malformation: the diagnostic value of three-dimensional multidetector-row computed tomography\*

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## Abstract

**Objective** To evaluate the diagnostic value of three-dimensional multidetector-row computed tomography (MDCT) in detecting acquired renal arteriovenous malformation (RAVM) and to compare its performance with that of ultrasonography and digital subtraction angiography (DSA).

**Methods** The institutional review board approved this retrospective study and written informed consent was obtained from all patients before examination. All 14 patients with acquired RAVM underwent MDCT, including cortical and medullary phase enhancement angiography and three-dimensional (3D) reconstruction. Five and nine patients were further examined and their diagnoses confirmed by DSA and surgery, respectively. The MDCT images, including 3D reconstructions, were analyzed for RAVM independently and in consensus by two observers using a workstation.

**Results** Among the 14 patients with acquired RAVM, 12 with maximum lesion diameter  $\geq 10$  mm, and one with a maximum lesion diameter between 5 and 10 mm, were correctly diagnosed with MDCT angiography. Among these patients, four diagnoses were confirmed by DSA. One patient with a lesion 5–10 mm in diameter was misdiagnosed with a renal aneurysm by MDCT angiography. The other one with the maximum diameter of the lesion between 5 mm and 10 mm was misdiagnosed as renal aneurysm with MDCT angiography, which was diagnosed as renal arteriovenous malformation with DSA. Among 14 lesions in 14 patients, eight and six originated in the left and right kidney, respectively.

**Conclusion** MDCT angiography can accurately diagnose RAVM and improve our understanding of the disease, which will allow clinicians to provide better care.

**Key words:** arteriovenous malformation; kidney; multidetector row computed tomography; digital subtraction angiography

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With the development of renal biopsy and improved treatment modalities for renal trauma, the incidence of acquired renal arteriovenous malformation (RAVM) has gradually increased in recent years [1–4]. Most cases are unilateral and characterized by direct communications between renal arteries and veins via enlarged, tortuous vascular spaces. The symptoms of RAVM usually include hematuria, vascular masses, and intractable hypertension [5–8]. Although digital subtraction angiography (DSA) has long been considered the reference standard for diagnosis of RAVM [9], it is an invasive technique less useful as a

conventional means to screen patients suspected of having RAVM.

With advances in multidetector-row computed tomography (MDCT) technology, MDCT angiography offers improved spatial resolution and has replaced traditional imaging work up, including DSA, for most visceral arteries [10–11]. Compared to traditional axial CT, maximum intensity projection (MIP), volume rendering (VR), multiplanar volume reconstruction (MPVR), and three-dimensional (3D) reconstruction MDCT imaging can offer more detailed visualization of vascular disorders.

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Thus, this study retrospectively analyzed and compared MDCT angiography imaging data to that of DSA for diagnosis of acquired RAVM.

## Materials and methods

### Patients

This study was approved by the review board of Tongji Hospital at Tongji Medical College of the Huazhong University of Science and Technology, and written informed consent was obtained from all patients before examination. From September 2006 to July 2012, double phase contrast enhanced MDCT was routinely performed according to a local protocol on patients with suspected RAVM based on ultrasound findings and clinical features. Among these, 14 patients (10 women and 4 men; age range, 13–54 years; mean age,  $34.71 \pm 13.57$  years), each with an RAVM lesion at least 5 mm in diameter (mean diameter,  $34.79 \pm 19.99$  mm; range, 6–81 mm), were retrospectively included in our study group. All patients had a history of percutaneous renal biopsy or penetrating trauma, and were clinically normal before the examinations. None of these subjects had concomitant renal cancer or other renal diseases. Five patients were further examined by DSA and all cases were confirmed as RAVM by DSA or surgery at Tongji Hospital.

### MDCT evaluation

MDCT scans of all patients were performed with a 64-detector MDCT (LightSpeed VCT, GE Healthcare Technologies, USA). The intravenous contrast protocol was 80–100 mL of nonionic contrast medium (Iopamiro 370 mg/mL, Brocco Sine, Shanghai, China) and physiological saline injected into the antecubital vein with an 18 gauge intravenous cannula using a double tube high-pressure injector (Stellant, Medred Co., USA) at a rate of 4 mL/s.

Patients were in a supine position. The MDCT scanning parameters were as follows: tube voltage 120 kV; tube current 150–440 mA; rotation time 0.5 s; detector pitch 1.375 : 1; detector rows 64 mm  $\times$  0.625 mm; matrix 512  $\times$  512; table speed 39.37 mm/rotation; and slice thickness/interval 0.625 mm.

Renal cortical phase scanning was initiated by bolus tracking (Smartprep, GE Healthcare Technologies) when a threshold enhancement of 120 Hu was reached in the abdominal aorta [12–13]. Medullar phase imaging was initiated 15 s after completion of arterial phase scanning.

### DSA evaluation

Using the Seldinger technique, a 5F Yahsiro polyethylene catheter was placed in the main renal artery to obtain a baseline selective angiography with cut film or digital subtraction techniques (Toshiba-2000 DSA system, Toshiba, Japan). The DSA parameters were as fol-

lows: tube voltage 66 kV; tube current 400 mA; exposure frequency 15 p/s. The intra-arterial contrast protocol included 35–45 mL of Iohexol (Omnipaque 350 mg/mL, GE Healthcare Technologies) injected using a high-pressure injector (Stellant, Medred Co., USA) at a rate of 3–4 mL/s.

### Image analysis

All MDCT image data were processed on a workstation (Advantage Windows 4.3; GE Medical Systems). Double phase MDCT images were independently analyzed by two radiologists (DY.H. and Z.L, with 27 and 12 years of experience, respectively, of reporting abdominal CT findings) at a GE Advantage Windows 4.3 workstation.

The source axial enhancement images were reconstructed using the ADW4.3 protocol. We reconstructed the kidney for three-dimensional (3D) analysis by volume rendering (VR), multiplanar volume reconstruction (MPVR), maximum intensity projection (MIP), and (3D) reconstruction technique. The combined source images were assessed for signs of RAVM and the findings compared to retrospective MDCT angiography and DSA findings.

### Statistical analysis

MDCT angiography and DSA images were retrospectively analyzed and data compared using Fisher's exact test (SPSS Statistics for Windows, Version 17.0., SPSS Inc., Chicago, IL). *P* values less than 0.05 were considered statistically significant.

## Results

Among 14 lesions from 14 patients, eight and six lesions originated from the left and the right kidney, respectively. Eight lesions were located in the junction between the renal cortex and medulla; four were giant, and some abnormal vessels were located beside the kidney. Two lesions were located in the inferior renal pole, and were complicated with angiomyolipoma (Table 1 and Fig. 1).

Of the 14 patients, 12 with lesions of maximum diameter greater than 10 mm, and one with a 5–10 mm maximum lesion diameter, were correctly diagnosed with RAVM by MDCT angiography. Diagnoses in four of these patients were confirmed by DSA. One additional patient, with a 5–10 mm diameter lesion, was misdiagnosed with renal aneurysm by MDCT angiography (Fig. 2), a finding confirmed by DSA (Table 1).

Both MDCT angiography and DSA showed thickened, circuitous, and well-defined arterial and venous elements as well as cavernous channels. In one case, the thickened and circuitous vein filled with contrast medium in the arterial phase – also visible by DSA – was incorrectly con-

Table 1 Diagnosis and therapy of 14 patients with acquired RAVM

| No. | Sex/age | Diagnosis   | Therapy | Origin | Cause | Size of AVM (mm) |     |         |
|-----|---------|-------------|---------|--------|-------|------------------|-----|---------|
|     |         |             |         |        |       | CTA              | DSA | Surgery |
| 1   | M/19    | MDCTA       | Surgery | LK     | PRB   | 13               | –   | 17      |
| 2   | F/25    | MDCTA       | Surgery | RK     | PRB   | 35               | –   | 32      |
| 3   | M/52    | MDCTA       | Surgery | RK     | PT    | 58               | –   | 55      |
| 4   | F/49    | MDCTA & DSA | DSA     | LK     | PRB   | 19               | 18  | –       |
| 5   | F/38    | MDCTA       | Surgery | RK     | PT    | 34               | –   | 37      |
| 6   | F/45    | MDCTA       | Surgery | LK     | PRB   | 32               | –   | 30      |
| 7   | M/24    | MDCTA & DSA | DSA     | LK     | PT    | 79               | 81  | –       |
| 8   | F/53    | MDCTA & DSA | DSA     | RK     | PT    | 43               | 44  | –       |
| 9   | F/31    | MDCTA       | Surgery | LK     | PT    | 51               | –   | 50      |
| 10  | F/29    | MDCTA       | Surgery | LK     | PRB   | 38               | –   | 36      |
| 11  | M/37    | MDCTA & DSA | DSA     | RK     | PT    | –                | 6   | –       |
| 12  | F/51    | MDCTA       | Surgery | LK     | PRB   | 18               | –   | 20      |
| 13  | F/18    | MDCTA       | Surgery | LK     | PT    | 45               | –   | 48      |
| 14  | F/15    | MDCTA & DSA | DSA     | RK     | PRB   | 9                | 9   | –       |

AVM, arteriovenous malformation; MDCTA, 64-detector MDCT angiography; DSA, digital subtraction angiography; LK, left kidney; RK, right kidney; PRB, percutaneous renal biopsy; PT, penetrating trauma

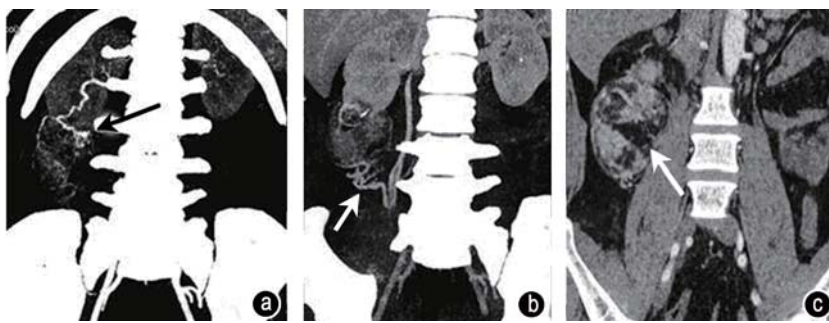


Fig. 1 MIP-3D of cortical phase (a) and intravenous phase (b) images showing AVM of the right kidney, complicated with angiomyolipoma, inferior lobe artery feeding (short arrow in A) the AVM (long arrow in A), and circuitous vein (white arrow in B) draining to right renal vein. Coronal MPVR image shows the lesion fat element (white arrow in C). These findings were confirmed by subsequent surgery. MIP-3D, maximum intensity projection (3 dimensional); AVM, arteriovenous malformation; MPVR, multiplanar volume reconstruction

considered evidence of renal aneurysm by MDCT angiography.

DSA can measure the speed and pressure of blood flow in AVM. MDCT angiography, on the other hand, can show the whole shape and surrounding structures.

Among the 14 patients included in this study, nine had continuous hypertension and macroscopic hematuria, three had continuous hypertension and microscopic hematuria, and two had microscopic hematuria. In addition, a renal vascular bruit was present in one patient. Hematuria disappeared after surgical and DSA-assisted repair. Hypertension resolved in eight patients after surgical and DSA-assisted repair (Table 2).

## Discussion

RAVM is usually classified as aneurysmal or cirroid [1]. Aneurysmal RAVM is characterized by solitary arteriovenous communications, with single, scrotiform channels, and well-defined arterial and venous elements. In contrast, cirroid types are considered the true congenital form and are characterized by multiple vascular communications, typically near the collecting system [2–3]. The

latter, also called acquired RAVM are the most common and occur as a result of penetrating trauma, percutaneous renal biopsy, and nephrectomy [2, 14–17].

The main symptoms include continuous or intractable hypertension and hematuria (microscopic or macroscopic); the hypertension severity appears to be correlated with the size of the arteriovenous malformation [3]. Vascular mass may be the only finding during health examinations.

Renal vascular bruits are rare signs, found only in relatively large RAVM. One patient had mega-malformation of the left kidney and showed this rare sign of the disorder (Fig. 3). After surgical or DSA-assisted repair, signs typically disappear or diminish. At four-month follow-up examinations after treatment, hematuria, hypertension, and renal vascular bruit had resolved in all patients, eight patients, and one patient, respectively (Table 2). DSA is the accepted reference standard for diagnosis of RAVM; it can clearly show lesion number, size, and location, especially for small malformations. In addition to its use as a diagnostic technique, DSA is also a good treatment option for RAVM [9, 15–19] (Fig. 4). The features of RAVM in conventional CT include masses of vascular density located

Table 2 Comparison of multidetector-row computed tomography angiography and DSA findings

| Item                       | CTA | DSA |
|----------------------------|-----|-----|
| No. of case                | 14  | 5   |
| Signs                      |     |     |
| Fed artery                 | -   | +   |
| Varicose configured vessel | +   | +   |
| Draining vein              | +   | +   |
| Early venous drainage      | -   | +   |
| Surrounding structures     | +   | -   |
| Complication               | +   | -   |
| Diagnosis*                 |     |     |
| No. of positive cases      | 13  | 5   |
| No. of negative cases      | 1   | 0   |
| Reference standard         |     |     |
| DSA                        | 5   | 5   |
| Surgery                    | 9   | 0   |

\* Fisher's exact test,  $P=0.304$ ; +, full view; -, lack of full view; DSA, digital subtraction angiography; CTA, multidetector-row computed tomography angiography

in the renal sinus and surrounding the pelvicalyceal system. Renal and left gonadal vein dilation are also some-

times observed [19]. However, during conventional CT, enhancement techniques such as the amount of contrast medium and speed of injection can significantly affect diagnosis and differential diagnosis [20-21].

MDCT is considered a dependable and noninvasive tool for assessment of almost all types of vascular disorders [10, 21]. This technique helps to identify anomalous kidney vessels and their courses. However, because RAVM is relatively rare, only a few case reports describe cases of RAVM. In our study, MDCT detected 13 of 14 RAVM lesions, visualizing irregular and varicose vessels and aneurismal dilatation, as well as feeding arteries and draining veins. Furthermore, more anatomic details and surrounding renal structures can be displayed on reformatted CT images [13]. For those patients who choose surgery instead of DSA therapy, MDCT angiography can show more anatomic details and pathological information before surgery. Only one patient with RAVM was misdiagnosed with renal aneurysm by MDCT angiography. The maximum lesion diameter was 6 mm; while imaging showed an abnormal vascular lesion, it did not clearly show nonvascular complications such as angiomyolipoma or renal hematoma. This finding suggests that MDCT may

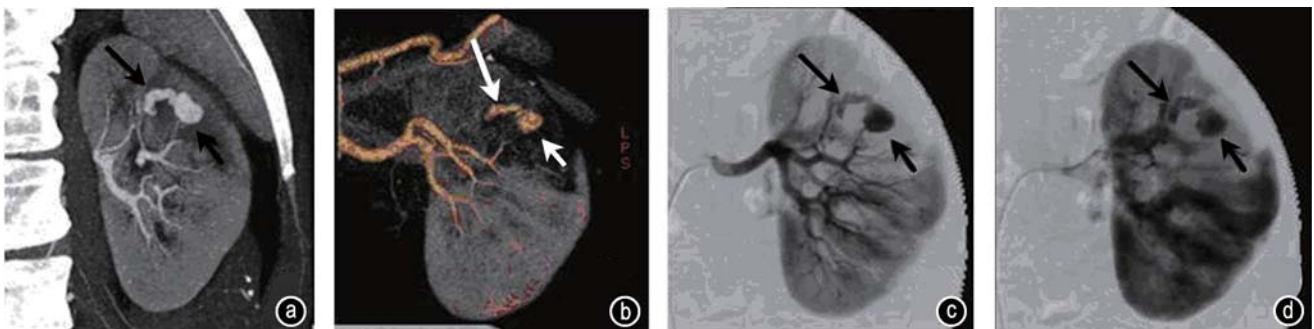


Fig. 2 MIP-3D (A) and VR-3D (b) images misleadingly suggest the lesion to be an aneurysm (short arrow in a, b), and feeding artery (long arrow). Subsequent selective angiography (c, d) images confirm that the feeding arterial actually is a draining vein (long arrow) associated with RAVM; these images also show aneurismal dilatation (short arrow). MIP-3D, maximum intensity projection (3 dimensional); VR-3D, volume rendering (3 dimensional); RAVM, renal arteriovenous malformation

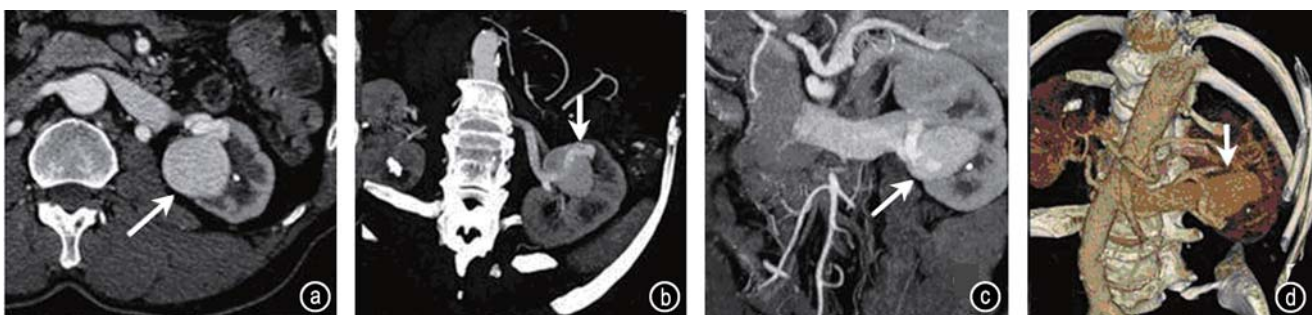


Fig. 3 Left renal AVM (a-d). Gross hematuria without pain had emerged two years prior. Vascular malformation (long arrow) and feeding artery were observed in the cortical phase axial CT images (a). The lesion is visible as a round, bulky vascular malformation. Cortical phase enhancement of maximum intensity projection images (b and c) show AVM and feeding artery (arrow). 3D volume rendering images (d) show the feeding arterial and draining vein associated with RAVM, as well as the aneurismal dilatation (arrow). AVM, renal arteriovenous malformation; RAVM, renal arteriovenous malformation



Fig. 4 Left renal AVM in a 49-year-old woman (a–e). Axial (a) and coronal (b, c) maximum intensity projection CT images show AVM in cortical phase enhancement (short arrow and long arrows). Subsequent selective angiography images (d, e) confirm that the feeding arterial actually is a draining vein associated with RAVM; the images also show aneurismal dilatation (long arrow). AVM, arteriovenous malformation; RAVM, renal arteriovenous malformation

not be as reliable as DSA for diagnosis of small RAVMs (maximum diameter < 10 mm). The reason for this lack of sensitivity may be that the lesion was too small to be displayed well by MDCT. Another reason may be that the double-phase enhancement scans of MDCT are independent processes rather than continuous images as is the case for DSA. DSA can more easily display the continuous movement of blood flow in small RAVMs. However, one case does not provide conclusive evidence, and advanced research is necessary to confirm these observations.

Compared to DSA, MDCT axial source images can be reconstructed as 2D or 3D images, which permits visualization of the number, size, and location of malformations at any projection without repetitive radiation exposure or extra contrast agent burden<sup>[12]</sup>. MIP-3D imaging can clearly display renal blood vessel circuitry, dilation, and stenosis, as well as the relationship between the lesion and surrounding structures, similar to DSA. VR-3D imaging is a common imaging reconstruction technique that can show large RAVM lesions, including the origin, circuitry, and dilation of anomalous vessels and surrounding structures. MPVR is a 2D imaging technique that can supplement VR to show RAVM shape and size, as well as the wall and lumen structures of abnormal vessels. MDCT angiography is a multidirectional imaging technique better for visualizing normally distributed kidney sub-arteries and anomalous channels between the artery and vein. Data from MDCT angiography are beneficial for future treatment by DSA or surgery.

This study has several limitations. First, because the disease is relatively rare, the study population was small. Second, there was only one small lesion (less than 10 mm) in the group, and this case was misdiagnosed by MDCT. This sample size and observation is too small to definitively conclude that either method is better. Finally, DSA was performed on only five patients, further emphasizing the need for more advanced research.

In conclusion, MDCT achieved results similar to those of DSA for diagnosis of RAVMs in this study, suggesting that 2D and 3D reconstruction imaging can provide

additional information for diagnosis. Compared to DSA, MDCT reconstruction imaging is more easily understood by clinical doctors, and is more acceptable in clinical practice as a non-invasive tool. Further study with a larger sample size is required to better determine the role of MDCT in diagnosis of RAVM.

#### Conflicts of interest

The authors indicated no potential conflicts of interest.

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