Preoperative Visualization of the Artery of Adamkiewicz by Intra-Arterial CT Angiography


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BACKGROUND AND PURPOSE: CT and MR angiographies have been reported to visualize the artery of Adamkiewicz (AKA) noninvasively to prevent spinal cord ischemia in surgery of thoracic descending aortic aneurysms. The purpose of this work was to compare the usefulness of CT angiography (CTA) with intra-arterial contrast injection (IACTA) with that of conventional CTA with intravenous contrast injection (IVCTA).

MATERIALS AND METHODS: We enrolled 32 consecutive patients with thoracic or thoracoabdominal aortic aneurysms who were scheduled for surgical repair or endovascular stent-graft treatment. All of the CTA images were obtained using a 16-detector row CT scanner and 100 mL of contrast material (370 mg/mL) injected at a rate of 5 mL/s. Contrast was injected via the antecubital veins of 15 patients and via a pig-tail catheter placed at the proximal portion of the descending aorta in 17 patients who underwent IVCTA and IACTA, respectively. Two datasets were reconstructed from 2 consecutive scans. The AKA was identified as a characteristic hairpin curved vessel in the anterior midsagittal surface of the spine and by the absence of further enhancement in the second rather than in the first phase. Continuity between the AKA and aorta was confirmed when the vessel could be traced continuously by panning the oblique coronal multiplanar reconstruction or original axial images.

RESULTS: Intra-arterial contrast injection was significantly more sensitive in identifying the AKA than IVCTA: 16 (94.1%) of 17 versus 9 (60.0%) of 15 (P = .033). Continuity between the AKA and aorta through intercostal or lumbar artery was confirmed in 14 (87.5%) of 16 and 5 (55.6%) of 9 of the IACTA and IVCTA groups, respectively.

CONCLUSION: Intra-arterial contrast injection detected the AKA at a high rate and verified continuity from the aorta to the AKA.

The AKA is a small vessel with a diameter of 0.5–1.5 mm that is surrounded by osseous structures. In addition, intercostal or lumbar arteries and dorsal branches run very close to the osseous structures (Fig 1). Due to these anatomic features, the contrast-to-noise ratio (CNR) in the spinal canal is decreased, and the AKA and its continuity with the aorta can be obscured. Robust aortic contrast enhancement is necessary to detect small vessels, but CTA with intravenous contrast injection (IVCTA) has limitations with respect to elevating aortic enhancement, because contrast material is diluted in the circulation of the right side of the heart.

Methods

Patients

Between March 2005 and September 2006, 32 consecutive patients with TAA or TAAA who were scheduled for surgical repair or an endovascular stent-graft underwent CTA to visualize the AKA. The patient group was composed of 22 men and 10 women aged 22–87 years (mean, 68.1 years). Twenty patients had TAAA and 12 had TAA. Of these, 11 had dissecting aneurysms. One patient had a background of Takayasu disease, and another had Maffan syndrome.

The patients were assigned to 2 groups according to whether the injection method was IVCTA (n = 15) or IACTA (n = 17; Table 1). Thirteen patients underwent IVCTA before July 25, 2005, when IACTA was initiated. From then until September 2006, 17 patients underwent IACTA. After July 25, 2005, 2 patients with insufficient renal function (serum creatinine >1.5 mg/dL) underwent IVCTA.
Fig 1. Anatomic course of the AKA. Right anteriosuperior view of a 3D volume-rendered CT image of IACTA with semitransparent skeletal system. Intercostal and lumbar arteries (1) originate from the aorta, and divide into posterior (2) and anterior (3) branches. Anterior branches run through the intercostal groove. Posterior branches subdivide into the radiculomedullary artery (4) and muscular branch (5). Radiculomedullary artery courses to the spine and enters the vertebral foramen. The AKA (6) is the largest anterior radiculomedullary artery and joins the anterior spinal artery (7) in a characteristic hairpin curve.

| Table 1: Patient data and results of CT angiography |
|-----------------|-----------------|-----|
| Variable | IVCTA | IACTA | P |
| No. of patients | 15 | 17 | |
| Age (range), years | 70.6 (54–87) | 65.9 (22–83) | .168 |
| Male/female | 9/6 | 13/4 | .450 |
| TAA:TAAA | 3:12 | 9:8 | .125 |
| Dissecting:true | 7.8 | 4:13 | .266 |
| AKA detection, n/N (%) | 9/15 (60.0) | 16/17 (94.1) | .033 |
| Confirmation of continuity, n/N (%) | 5/9 (55.6) | 14/16 (87.5) | .142 |
| CT value of aorta, mean ± SD, HU | 352.2 ± 46.0 | 965.4 ± 634.1 | .002 |

Note: — IVCTA indicates intravenous CT angiography; IACTA, intra-arterial CT angiography; TAA, thoracic aortic aneurysm; TAAA, thoracoabdominal aortic aneurysm; HU, Hounsfield unit; AKA, artery of Adamkiewicz.

or more days after coronary angiography to avoid a large amount of contrast medium in a single day. Thus, 15 patients underwent IVCTA.

This study was concordant with our institutional ethics guidelines, and all of the patients provided written informed consent to participate in the study.

Scanning Protocol
We obtained CT images from the levels of the seventh thoracic and the third lumbar vertebrae in the craniocaudal direction using a 16-detector row multissection CT (Brilliance16; Philips Medical Systems, Best, the Netherlands). Scanning was consecutively repeated to obtain early and late-phase images. Scanning variables were composed of 0.75 seconds per rotation, 0.75-mm collimation, 0.666–0.875 pitch, 120 kV, and 300–400 mA (according to individual physique).

Injection Protocol and Scan Timing
IVCTA. The contrast material, iopamidol (Iopamiron; Schering, Berlin, Germany), was injected (100 mL; 370 mg/mL; 5 mL/s) via a 20-gauge plastic intravenous catheter placed in the antecubital vein and followed by a 30-mL saline flush. The start time of the scan was determined by using conventional real-time monitoring (Bolus Tracking, Phillips Medical Systems). Regions of interest were placed on the descending aorta at the level of the seventh thoracic vertebra. Low-dose dynamic monitoring scanning (120 kV; 100 mA) started 15 seconds after the intravenous contrast material injection started, and 1 image per second was obtained. The trigger threshold was set at 100 Hounsfield units (HU) above the baseline (absolute value, 140–160 HU). Scanning automatically started 4 seconds after reaching the threshold.

IACTA. Patients underwent IACTA after coronary angiography as a routine preoperative evaluation of coronary stenosis. After coronary angiography (radial or brachial arterial approach), a 5F Pigtail Catheter (Terumo, Tokyo, Japan) was advanced to the proximal portion of the descending aorta under fluoroscopy, and visualization of intercostal arteries was confirmed by manual injection of contrast material. Patients were then transferred to the CT room, and 100 mL of the same contrast material was injected via the catheter at a rate of 5 mL/s. The CT scan started 4 seconds after starting the injection.

Image Procession and Diagnosis
Two datasets were reconstructed from the 2 consecutive scans with a section width of 0.8 mm, a reconstruction interval of 0.4 mm, and 200-mm FOV. Image datasets were transferred to a workstation (Zio M900; Ziosoft, Tokyo, Japan). From the first phase image, an oblique coronal multiplanar reconstruction (MPR; 0.8-mm thickness, 0.8-mm intervals) fitting the curvature of the AKA was obtained. Exactly the same MPR sections were obtained from the second phase to distinguish the AKA from the radiculomedullary vein, which can be similarly shaped. Curved planar reformation (CPR) and 3D volume rendering were applied to display continuity between the AKA and the aorta.

The AKA was identified by the consensus of 2 experienced radiologists (K.U. and A.K.) who specialize in cardiac and vascular imaging. The analytical criteria of the AKA comprised detection of the AKA and confirmation of continuity between the AKA and aorta. The detection criteria of the AKA were the presence of a characteristic hairpin curved vessel ascending from the intercostal foramen to the anterior midsagittal surface of the spine and the absence of further enhancement in the second phase compared with the first phase (Figs 2 and 3). Continuity was confirmed when the vessel could be continuously traced via the intercostal or lumbar artery by paging oblique coronal MPR or original axial images. To compare the contrast bolus between the 2 groups, the CT value of the aorta was measured in sections of the midlevel of the scan range (approximately the 10th or 11th thoracic vertebra). A region of interest was placed in the dorsal portion of the aorta from which the intercostal or lumbar artery arose. To assess differences in the features of the contrast bolus between the 2 groups, we visually evaluated sagittal partial maximum intensity projection (MIP) images to determine whether contrast material was dorsally concentrated or homogeneously distributed.

Statistical Analysis
Data were processed using StatMate (ATMS, Tokyo, Japan). Variables were compared between the IVCTA and IACTA groups using the 2-sided Fisher exact test for proportions and a 2-tailed t test for the means of CT values and patient ages.

Results
No complications were associated with the angiographic procedure or CT imaging. Table 1 summarizes the results. The
AKA was detected more sensitively by IACTA than IVCTA: 16 (94.1%) of 17 versus 9 (60.0%) of 15 (P = .033). Continuity between the AKA and aorta through the intercostal artery (arrow) was confirmed in 14 (87.5%) of 16 and in 5 (55.6%) of 9 of the IACTA and IVCTA groups, respectively. The CT value of the aorta was significantly higher in IACTA than in IVCTA (965.4 ± 634.1 versus 352.2 ± 46.0 HU; P = .002). Contrast material concentrated in the dorsal portion of the aorta in IACTA but homogeneously enhanced the aorta in IVCTA. The intercostal and lumbar arteries were displayed more clearly by IACTA than by IVCTA (Fig 4). The AKA originated from a left intercostal or lumbar artery in 72% of the patients and at the level of the ninth to 12th intercostal arteries in 84% (Fig 5). We found hairpin curved vessels that were visible in the first phase images but were enhanced stronger in the second phase images in 10 patients of the IACTA group and 9 of the IVCTA group. They were identified as radiculomedullary veins that may mimic the AKA in the first phase image.
Discussion

The detection rates of the AKA and the confirmation rates of continuity were higher with IACTA compared with those of previous reports using IVCTA (Table 2). The AKA and its anatomic course from the intercostal or lumbar arteries to the anterior spinal artery are very thin, and the artery runs close to osseous structures, which frequently reduce the CNR of the spinal vasculature and complicate vessel tracking. We applied a slower rotation speed and smaller helical pitch than usual to obtain higher mAs and, thus, elevate the CNR. Furthermore, contrast material is diluted in aortic aneurysms, which reduces the CNR. To obtain sufficient arterial opacification, rapid injection (4.0–4.5 mL/s) and a relatively high dose (130–150 mL) of contrast material are considered necessary to visualize the AKA by using IVCTA. Aortic visualization is enhanced by IACTA because contrast material is directly injected without dilution in the circulation of the right heart. Similar to the report of Nojiri et al., we injected 100 mL of contrast material at a rate of 5.0 mL/s through a Pigtail catheter placed in the proximal portion of the descending aorta and started CT scanning with a 4-second delay after the injection start. The dorsal portion of the aorta and intercostal or lumbar arteries tended to have extremely high attenuation values in IACTA (Fig 4) because of the high specific gravity of the contrast material (1.405 at 370 mg/mL of Iopamidol; 37°C). High z-axis (craniocaudal) resolution is also important to visualize thin arteries. Collimation of 1 mm improved the spatial resolution and arterial tracking compared with 2-mm collimation. When IACTA was applied using 0.75-mm collimation and intra-arterial contrast injection, the detection rate of the AKA was 94%, and continuity was confirmed at a rate of 88%. We could not detect the AKA by using IACTA in 1 patient despite sufficient enhancement of the aorta and intercostal arteries. The large anterior radiculomedullary artery might not have been located within the scan range of this patient. One of 2 patients in whom the AKA was detected but continuity was not confirmed underwent selective intercostal angiography in the course of stent-graft deployment, and the AKA originating from the 11th intercostal artery visualized by using IACTA was confirmed by selective spinal arterial angiography (Fig 6).

A few points should be considered when performing IACTA. The catheter tip should be advanced to a suitable position to visualize the intercostal and lumbar arteries when a dissecting ascending aortic aneurysm is associated with a patent false lumen. We placed the catheter tip immediately proximal to the entry to enhance both true and false lumens in patients with dissection and confirmed visualization of intercostal and lumbar arteries by test injections of contrast material. None of the patients studied here had dissection of the ascending aorta or aortic arch. Patients with Stanford type A dissection had undergone total arch replacement before IACTA. Collateral circulation from branches of the subclavian artery, such as the internal thoracic artery, might not have been visualized when contrast material was injected from the proximal descending aorta.

Radiculomedullary veins can have a similar shape to the AKA, and they can run close to the radiculomedullary arteries at the level of the vertebral foramen. Thus, they can be enhanced in the first phase image because of a scan duration of 20–25 seconds, whereas the estimated interval between the arterial and venous phases is 10–15 seconds. Some vessels with a hairpin curve configuration in the first phase image that were more enhanced in the second were regarded as radiculomedullary veins (Fig 3). Without multiphase dynamic scanning, such vessels might be misdiagnosed as arteries if continuity with intercostal or lumbar arteries is not confirmed. Thus, confirmation of continuity between the aorta and the AKA by multiphase scanning is critical to avoid misdiagnosis.

Selective spinal arterial angiography is the “gold standard” for confirmation of the AKA. Selective spinal arterial angiography is the “gold standard” for confirmation of the AKA.
for diagnosing the AKA. However, previous diagnoses of the AKA by using CTA and MRA did not have a “gold standard” probably because of the risk of selective angiography. Only one report has proved that MRA can be diagnostic in a subgroup of 3 patients who underwent selective spinal angiography.13 Thus, the true detection rates of the AKA by MRA and CTA are unknown. The detection rate of the AKA by using IVCTA in the present study was not as high as in previous reports (Table 2). One possible cause of this variance is that diagnostic criteria were not identical among the studies. Multiphase dynamic imaging was performed by using MR imaging,14–16 whereas single-phase imaging was performed by using CT.19–21,24 We believe that multiphase imaging can improve the precision of AKA diagnosis. The present study showed that the detection rates of IACTA were better than those of IVCTA when using the same scan variables, contrast dose, injection rate, and diagnostic criteria.

Arterial catheterization causes IACTA to be more invasive than IVCTA and MRA but is nevertheless safer and less time consuming than selective spinal angiography, which is associated with complications such as retroperitoneal bleeding, cerebral ischemia, renal failure, and spinal cord ischemia in 1.2%–4.6% of patients.10–11 We have not yet experienced any complications related to IACTA, which proceeded in addition to routine preoperative coronary angiography, with an elevated total dose of contrast material. For patients with insufficient renal function, we performed IVCTA on a different day from coronary angiography.

Although the contribution of information about the AKA to the outcome of thoracoabdominal aortic surgery remains controversial,9,11,12 some reports have suggested that preoperative AKA localization is useful in reducing the risk of postoperative ischemic spinal complications14,15 and in shortening surgical duration.16 Preoperative AKA localization aids in planning surgical strategies, and protection for the spinal cord can be more reliable.7 We believe that IACTA will improve diagnostic accuracy and would help to reduce postoperative spinal complications. In this study, 14 and 15 patients from the IVCTA and IACTA groups, respectively, underwent surgical repair or stent grafting, and 1 patient who underwent IVCTA developed hemiaplasia despite preoperative AKA demonstration. However, the study cohort was small, and further study is needed to confirm the efficacy of IACTA in the clinical setting.

Conclusions
The rate of detecting the AKA and the confirmation rate of continuity from the aorta to the AKA were high using IACTA. A high concentration of contrast material delivered via intraarterial injection can improve the accuracy of visualizing the AKA. The AKA should be correctly identified so that radiologists can provide surgeons with the critical information needed for reconstruction of the intercostal and lumbar arteries. We believe that IACTA is a suitable preoperative means of investigating patients with TAA and TAAA.

References