

AmeriHealth Caritas Louisiana

National Imaging Associates, Inc.	
Clinical guidelines	Original Date: September 1997
CHEST MRA/MRV	
CPT Codes: 71555	Last Revised Date: March 2022 April 2023
Guideline Number: NIA_CG_022-2	Implementation Date: January 20234

GENERAL INFORMATION

- It is an expectation that all patients receive care/services from a licensed clinician. All appropriate supporting documentation, including recent pertinent office visit notes, laboratory data, and results of any special testing must be provided. If applicable: All prior relevant imaging results and the reason that alternative imaging cannot be performed must be included in the documentation submitted.
- Where a specific clinical indication is not directly addressed in this guideline, medical necessity determination will be made based on widely accepted standard of care criteria. These criteria are supported by evidence-based or peer-reviewed sources such as medical literature, societal guidelines and state/national recommendations.

INDICATIONS FOR CHEST MRA

Chest Magnetic Resonance Angiography (MRA) is ordered for evaluation of the intrathoracic blood vessels. Chest MRI and Chest MRA should not be approved at the same time.

Vascular Disease

- Superior vena cava (SVC) syndrome¹Superior vena cava (SVC) syndrome¹
- Subclavian Steal Syndrome after positive or inconclusive ultrasound^{2, 3}
- Thoracic Outlet Syndrome⁴⁻⁶⁴⁻⁶
- Takayasu's arteritis⁷
- Clinical concern for acute aortic dissection^{8, 9}Clinical concern for acute aortic dissection^{8, 9}
 - o Sudden painful ripping sensation in the chest or back and may include
 - New diastolic murmur
 - Cardiac tamponade

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- Distant heart sounds
- Hypotension or shock
- For MRPA (MR Pulmonary Angiography) in patients with intermediate pretest probability with a positive D-dimer or high pretest probability (but only at centers that routinely perform it well and only for patients for whom standard tests are contraindicated)
 - Risk can be determined by the parameters detailed in Background section

Initial/Screening for Thoracic Aortic Disease¹⁰⁻¹²10-12

- Echocardiogram or chest x-ray show aneurysm
- Screening of first-degree relatives of individuals with a thoracic aortic aneurysm (defined as <u>></u> 50% above normal) or dissection
- KnownEvaluation in patients with known or suspected connective tissue disease or genetic

conditions that predisposes to aortic aneurysm or dissection (e.g., Marfan syndrome, such as Marfan's, Ehlers-Danlos-or, get a one-time study or for Loeys-Dietz syndromessyndrome- allow imaging at diagnosis and then every two years, or more frequently if abnormalities are found (Imaging may include head, neck, chest, abdomen and pelvis)(MRA preferred due to cumulative radiation risk)

- Screening of the thoracic aorta after a diagnosis of a bicuspid aortic valve (dilation of the ascending aorta may not be seen on echocardiogram)^{13, 1413, 14}
 - If normal, reimage every three to five years
- Screening of first-degree relatives of patients with a bicuspid aortic valve
- Turner's syndrome Screen for coarctation or aneurysm of the thoracic aorta
 - o If normal results, screen every 5-10 years
 - o If abnormal, screen annually
- Suspected vascular cause of dysphagia or expiratory wheezing with other imaging is suggestive or inconclusive

Follow-up after established Thoracic Aneurysm^{10 12_14-16}

- Six months follow-up after initial finding of a dilated thoracic aorta, for assessment of rate of change
 - Aortic Root or Ascending Aorta (in cm)
 - 3.5 to 4.4 <u>–</u>annual
 - 4.5 to 5.5 or growth rate >≥ 0.5 cm5cm/year -- every 6 months
 - Genetically mediated (Marfans syndrome, Aortic Root or Ascending Aorta) (in cm)
 - 3.5 to 4.4 <u>–</u> annual
 - 4.5 to 5.05 or growth rate >≥ 0.5 cm5cm/year every 6 months
 - Surgery generally recommended over 5.0 cm
 - Descending Aorta (Braverman, 2011)

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<u>o</u> Descending Aorta (in cm)¹⁷

- 4.0 to 5.0 <u>–</u> annual
- 5.0 to 6.0 <u>–</u>every 6 months
- Follow-up post medical treatment of aortic dissection:
 - Acute dissection: 1 month, 6 months, then annually
 - Chronic dissection: annually
- Follow-up <u>TEVAR surveillance at 1 month, then 1 year</u> post either root repair or AVR plus ascending aortic root/arch repair: baseline post-op if stable, then annually
- Follow-up open repair if no residual aortopathy within first post op year, then every 5 years (if have residual aortopathy or abnormal findings on surveillance, annual followup if needed)
- Re-evaluation of known ascending aortic dilation or history of aortic dissection with a change in clinical status or cardiac exam or when findings may alter management

Congenital Malformations

- Thoracic malformation on other imaging (chest x-ray, echocardiogram, gastrointestinal study, or inconclusive CT)¹⁵⁻¹⁸¹⁵⁻¹⁸
- Congenital heart disease with pulmonary hypertension¹⁹ or vascular anomalies
- Pulmonary Sequestration²⁰
 - Pulmonary Sequestration²⁰

Pulmonary Hypertension based on other testing^{21, 22}21, 22

- Echocardiogram
- Right heart catheterization

Atrial fibrillation with ablation planned²³

Atrial fibrillation with ablation planned²³

Pre-operative/procedural evaluation

- Pre-operative evaluation for a planned surgery or procedure
- Pre-transplant CT or CTA/MRA chest approvable for surgical planning (to evaluate for vascular anatomy, mediastinal pathology, malignancy screening etc.)

Post-operative/procedural evaluation

- Post-operative complications^{24, 25}
- Routine post-operative^{26, 27}

■ <u>1 month</u>

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- More frequent follow-up/possible intervention if complication detected
- If stable, annual for 5 years
- See above indications for TAA follow up

Chest MRA and Abdomen MRA or Abdomen/Pelvis MRA

- Acute aortic dissection⁸
- Takayasu's arteritis⁷
 - -Transcatheter Aortic Valve Replacement (TAVR)
 - Acute aortic dissection
 - Takayasu's arteritis
 - Post-operative complications
 - To evaluate for an embolic source of lower extremity vascular disease (may also approved as a combination chest MRA, Abdominal MRA and a single LE MRA when LE runoff disease needs to be evaluated as well). Echocardiography is also needed, since the heart is the most commonly reported source of lower extremity emboli i, accounting for 55 to 87 percent of events.

Other Indications

Further evaluation of indeterminate findings on prior imaging (unless follow up is otherwise specified within the guideline):

- For initial evaluation of an inconclusive finding on a prior imaging report that requires further clarification
- One follow-up exam of a prior indeterminate MR/CT finding to ensure no suspicious interval change has occurred. (No further surveillance unless specified as highly suspicious or change was found on last follow-up exam.)

BACKGROUND

Magnetic resonance angiography (MRA) is a noninvasive technique used to provide crosssectional and projection images of the thoracic vasculature, including large- and medium-sized vessels, e.g., the thoracic aorta. MRA provides images of both normal and diseased blood vessels, and it quantifies blood flow through these vessels. Successful vascular depiction relies on the proper imaging pulse sequences. MRA may use a contrast agent, gadolinium, which is non-iodine-based, for better visualization. It can be used in patients who have history of contrast allergy and who are at high risk of kidney failure.

OVERVIEW

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MRA and Coarctation of the Aorta – One of the most common congenital vascular anomalies is coarctation of the aorta, characterized by obstruction of the juxtaductal aorta. Clinical symptoms, e.g., murmur, systemic hypertension, difference in blood pressure in upper and lower extremities, absent femoral or pedal pulses, may be present. Gadolinium-enhanced 3D MRA may assist in preoperative planning as it provides angiographic viewing of the aorta, the arch vessels, and collateral vessels. It may also assist in the identification of postoperative complications.

- MRA and Pulmonary Embolism (PE) Note: D-Dimer blood test in patients at low risk* for DVT is indicated prior to MRA imaging. Negative D-Dimer suggests alternative diagnosis in these patients.
- *Low risk is not approved. Low risk is defined as **NO** to **ALL** of the following questions with intermediate and high risk defined based on the number of positive responses²⁸:
- Evidence of current or prior DVT;
- HR > 100;
- Cancer diagnosis;
- Recent surgery or prolonged immobilization;
- Hemoptysis;
- History of PE;
- Oral hormone use;
- Another diagnosis beside PE is less likely

<u>Pulmonary Embolism (PE)</u> –Studies show mixed results regarding the value of MRA versus CTA in detecting pulmonary embolism. A systematic review and meta-analysis found MRA to be inferior to CTA in detecting PE. Therefore, MRA should be used only if CTA is not available or contraindicated in a specific patient.²⁹²⁶

MRA and Thoracic Aortic Aneurysm — One of the most common indications for thoracic MRA is thoracic aortic aneurysm, most often caused by atherosclerosis. These aneurysms may also be due to aortic valvular disease. Aneurysms are defined by their enlargement, and patients with rapidly expanding aortas, or with aortic diameters greater than five or six centimeters, are at high risk of rupture and may require surgery.

MRA and Thoracic Aortic Dissection – The most common clinical symptom of aortic dissection is tearing chest pain, and the most common risk factor is hypertension. An intimal tear is the hallmark for aortic dissection and intramural hematoma may also be detected. Unfortunately, patients with aortic dissection may be unstable and not good candidates for routine MR evaluation; MRA may be indicated as a secondary study. 3D MRA is also useful in postoperative evaluation of patients with repaired aortic dissections.

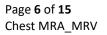
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MRA and Central Venous Thrombosis – <u>CTA/</u>MRA is useful in the identification of venous thrombi. Venous thrombosis can be evaluated by gadolinium-enhanced 3D MRA as an alternative to CTA, which may not be clinically feasible due to allergy to iodine contrast media or renal insufficiency.

MRI and Patent Ductus Arteriosus – Patent ductus arteriosus (PDA) is a congenital heart problem in which the ductus arteriosus does not close after birth. It remains patent allowing oxygen-rich blood from the aorta to mix with oxygen-poor blood from the pulmonary artery. MRI can depict the precise anatomy of a PDA to aid in clinical decisions. It allows imaging in multiple planes without a need for contrast administration. Patients are not exposed to ionizing radiation.

Other MRA Indications – MRA is useful in the assessment for postoperative complications of pulmonary venous stenosis.



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POLICY HISTORY

Date	Summary
<u>April 2023</u>	Simplified PE indications and removed other details from
	background)
	Clarified and updated follow up after repair of TAA
	General Information moved to beginning of guideline with added
	statement on clinical indications not addressed in this guideline
	Added statement regarding further evaluation of indeterminate
	findings on prior imaging
March 2022	No significant changes
April 2021	 Follow-up recommendations for bicuspid aortic valve
	 Added suspected vascular cause of dysphagia or expiratory
	wheezing
	 Combined follow-up surveillance recommendations for
	endovascular and open ascending aorta repair as per
	literature review
	 Added indications for combination studies and for ordering
	combination studies
	 Added Pulmonary Embolism criteria to Overview
	 Clarified pre-operative evaluation for a planned surgery or
	procedure
May 2020	Thoracic Aortic Disease
	 Initial/Screening
	Follow-up of known aneurysm/vascular
	pathology
	 Removed: 'Annual follow up of
	enlarged thoracic aorta that is above
	top normal for age, gender, and body
	surface area'
May 2019	 Removed pulmonary embolism indication
	 Added indications specifying criteria for follow-up of
	thoracic aneurysm
	 Added statement: "For MRPA (MR Pulmonary Angiography
	in patients with intermediate pretest probability with a
	positive D-dimer or high pretest probability (but only at
	centers that routinely perform it well and only for patients
	for whom standard tests are contraindicated)"
	Expanded criteria for congenital malformations
	Updated thoracic aortic disease section for consistency with
	, cardiac guidelines



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Reviewed / Approved by NIA Clinical Guideline Committee

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GENERAL INFORMATION

It is an expectation that all patients receive care/services from a licensed clinician. All appropriate supporting documentation, including recent pertinent office visit notes, laboratory data, and results of any special testing must be provided. If applicable: All prior relevant imaging results and the reason that alternative imaging cannot be performed must be included in the documentation submitted.

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