

National Imaging Associates, Inc. [*]		
Clinical guidelines	Original Date: September 1997	
MUGA (Multiple Gated Acquisition) Scan		
CPT Codes: 78472, 78473, 78494, +78496	Last Revised Date: March 2021	
Guideline Number: NIA_CG_027	Implementation Date: January 2022	

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. All prior relevant imaging results, and the reason that alternative imaging cannot be performed must be included in the documentation submitted.

Indications for Multiple Gated Acquisition (MUGA) Scan

(Doherty, 2019)

- To evaluate left ventricular function in a patient with coronary artery disease, valvular heart disease, myocardial disease, or congenital heart disease, in any of the following scenarios:
 - When ventricular function is required for management, and transthoracic echocardiography (TTE) or other imaging has proven inadequate (Patel, 2013; Yancy, 2013)
 - When there are conflicting results between other testing (i.e. Myocardial Perfusion Imaging and TTE) in the measurement of ejection fraction (EF), and the results of the MUGA will help in the management of the patient
 - Prior TTE has demonstrated systolic dysfunction (EF < 50%) and management will change based on the results of the MUGA scan
- In the course of cardiotoxic chemotherapy when TTE images are inadequate to evaluate left ventricular systolic function (Patel, 2013; Plana, 2014; Yancy, 2013; Zamorano, 2016):
 - <u>Previous low LV ejection fraction was < 50% and receiving potentially cardiotoxic</u> <u>chemotherapy</u>
 - Prior to cardiotoxic chemotherapy, and subsequently for monitoring and follow up. The frequency of testing should be left to the discretion of the ordering physician, but generally no more often than at baseline and every 6 weeks thereafter

^{*} National Imaging Associates, Inc. (NIA) is a subsidiary of Magellan Healthcare, Inc.

BACKGROUND

(Friedman, 2006; Mitra, 2012; Patel, 2013; Ritchie, 1995)

Multiple-gated acquisition (MUGA) scanning uses radio-labelled red blood cells to scan right and left ventricular images in a cine loop format that is synchronized with the electrocardiogram.

A prior MUGA scan is not an indication for repeat MUGA (if another modality would be suitable, i.e., TTE).

Abbreviations

EF	Ejection Fraction
MUGA	Multiple Gated Acquisition (nuclear scan of ventricular function)
TTE	Transthoracic echocardiography

POLICY HISTORY

Date	Summary	
March 2021	Added the following statement: Previous low LV ejection fraction was < 50% and receiving potentially cardiotoxic chemotherapy	
<u>March 2020</u>	 Added general information section as Introduction which outlines requirements for documentation of pertinent office notes by a licensed clinician, and inclusion of laboratory testing and relevant imaging results for case review Added statement to Background that a prior MUGA scan is not an indication for repeat MUGA (if another modality would be suitable. i.e. TTE) Removed statements from Background that CMR is recommended when TTE is inadequate and/or candidacy for cardiotoxic chemotherapy based upon LVEF is questionable and that MUGA can also be considered when CMR is not available. 	
<u>July 23, 2019</u>	 Removed chart on individual dosing for specific chemotherapeutic agents Added indication for when there are conflicting results between other testing (i.e. MPI and TTE) in the measurement of ejection 	

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fraction, and the	results of the MUGA will help in the management		
of the patient			
Removed section	n on Radionuclide Angiography, Combination of		
Other Studies wi	th MUGA, section on TTE and strain		
Removed CAD in	Removed CAD indication		
Added indication	n for cardiotoxicity as follows:		
○ In the	 In the course of cardiotoxic chemotherapy when TTE 		
image	es are inadequate to evaluate left ventricular		
systol	systolic function (Patel 2013, Plana 2014, Yancy 2013,		
Zamorano 2016):			
<u>o</u>	Prior to cardiotoxic chemotherapy, and		
	subsequently for monitoring and follow up. The		
	frequency of testing should be left to the		
	discretion of the ordering physician, but		
	generally no more often than at baseline and		
	every 6 weeks thereafter		
<u>o</u>	In patients with EF < 50% on TTE receiving		
	potentially cardiotoxic chemotherapy, more		
	frequent monitoring (every 4 weeks) may be		
	<u>appropriate</u>		
<u>o</u>	Removed section on Radionuclide Angiography,		
	Combination of Other Studies with MUGA,		
	section on TTE and strain		

July 23, 2019

- Removed chart on individual dosing for specific chemotherapeutic agents
- Added indication for when there are conflicting results between other testing (i.e. MPI and TTE) in the measurement of ejection fraction, and the results of the MUGA will help in the management of the patient
- Removed section on Radionuclide Angiography, Combination of Other Studies with MUGA, section on TTE and strain
- Removed CAD indication
- Added indication for cardiotoxicity as follows:
 - In the course of cardiotoxic chemotherapy when TTE images are inadequate to evaluate left ventricular systolic function (Patel 2013, Plana 2014, Yancy 2013, Zamorano 2016):
 - Prior to cardiotoxic chemotherapy, and subsequently for monitoring and follow up. The frequency of testing should be left to the discretion of the ordering physician, but generally no more often than at baseline and every 6 weeks thereafter

- In patients with EF < 50% on TTE receiving potentially cardiotoxic chemotherapy, more frequent monitoring (every 4 weeks) may be appropriate
- Removed section on Radionuclide Angiography, Combination of Other Studies with MUGA, section on TTE and strain

March 2020

- Added general information section as Introduction which outlines requirements for documentation of pertinent office notes by a licensed clinician, and inclusion of laboratory testing and relevant imaging results for case review
- Added statement to Background that a prior MUGA scan is not an indication for repeat MUGA (if another modality would be suitable. i.e. TTE)
- Removed statements from Background that CMR is recommended when TTE is inadequate and/or candidacy for cardiotoxic chemotherapy based upon LVEF is questionable and that MUGA can also be considered when CMR is not available.

March 2021:

- <u>Added the following statement:</u>
- ____

<u>Previous low LV ejection fraction was < 50% and receiving potentially cardiotoxie</u> <u>chemotherapy</u>

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Reviewed / Approved by NIA Clinical Guideline Committee

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