

## MRI PATIENT HISTORY AND CONTRAST CONSENT FORM

Patient Information

History/Symptoms:						
Injury:  Yes  No	Туре:		Date:			
Surgical History:						
Head/Brain:						
Neck/Chest:						
Spine:						
Bone/Joint:						
Abdomen/Pelvis:						
Have you ever been di	agnosed with cancer?	☐ Yes ☐ No Type:	Date:			
Radiation Therapy:	☐ Yes ☐ No Anaton	ny:				
Chemo Treatment	☐ Yes ☐ No					
Have you been diagn	osed with:					
☐ Asthma	☐ Alzheimer's	☐ Atherosclerosis	☐ Coronary Artery Disease			
□ COPD	□ CVA	☐ Congestive Heart Failure	☐ Chronic Kidney Disease			
☐ Diabetes	☐ Dialysis Treatments	□ DVT □ HTN	☐ Migraine Headaches			
☐ Osteoarthritis	□ PVD	☐ Osteoporosis	☐ Rheumatoid Arthritis			
Signature of Patient/	Responsible Party(relation	onship) Date	e and Time			
* For unresponsive pat			e and Time medical images will be assessed by a & medical history:			
☐ Exam is ordered	without IV contrast					
☐ Exam is ordered Continue to back page	with IV contrast					



## **MRI PATIENT HISTORY AND CONTRAST CONSENT FORM**

Creatinine results are required for patients 50 years of age and older, diabetic history, renal disease, multiple myeloma, or family history of renal failure prior to any contrast injection.  MR Exam:	Patient Information			
MR Exam:	Are you currently taking any home medications (presc If YES, please complete home medication record.	ription and/or non-prescription) ☐ Ye	s 🗆 No	
Creatinine:			disease, multiple	
Have you ever had an allergic reaction to IV contrast:	MR Exam:			
Type of reaction:  I UNDERSTAND THAT THIS CONTRAST AGENT (LIKE MANY NEW DRUGS) MAY CAUSE AN ALLERGIC REACTION.  Understand that my physician may have requested the use of an intravenous contrast media that will assist the radiologist in better distinguishing certain anatomy or abnormalities that would otherwise be difficult or impossible to see.  I understand that the procedure to be performed on me involves the use of a high strength magnetic field, and possibly insertion needles and gadolinium containing solution, which may enhance the diagnostic accuracy of the procedure.  I understand that I may be receiving an intravenous contrast media and/or oral contrast media to enhance the visibility of certain tissues. Possible side effects may include, but are not limited to, pain or swelling at the site of injection, nausea, vomiting, and a warm, flushed sensation. Also potential allergic reactions including, but not limited to, hives, wheezing, difficulty breathing, and in rare instances, anaphylactic shock (with severe allergic reactions).  The purpose, benefits, and complications of the contrast procedure will be explained prior to any injection that may take place, hereby consent to any measure necessary to correct complications, which may occur. I am aware that the practice of medicine is not an exact science and I acknowledge that no guarantee has been made to me concerning the results of the examination.  Confirm that the information I have provided is complete and accurate to the best of my knowledge.  IUNDERSTAND THE RISKS, BENEFITS, AND ALTERNATIVES INVOLVED IN THE PROCEDURE. I HAVE HAD THE DPPORTUNITY TO ASK ANY QUESTIONS AND ALL OF MY QUESTIONS HAVE BEEN ANSWERED TO MY SATISFACTION.  Signature of Patient or Responsible Party (relationship)  Date and Time  V Contrast Media/Amount:  V/Injection Site:  Treatment:  IV/Injection Site after injection and after Needle Removal:	Creatinine: Result Date:	GFR:		
I UNDERSTAND THAT THIS CONTRAST AGENT (LIKE MANY NEW DRUGS) MAY CAUSE AN ALLERGIC REACTION.  I understand that my physician may have requested the use of an intravenous contrast media that will assist the radiologist in better distinguishing certain anatomy or abnormalities that would otherwise be difficult or impossible to see.  I understand that the procedure to be performed on me involves the use of a high strength magnetic field, and possibly insertion needles and gadolinium containing solution, which may enhance the diagnostic accuracy of the procedure.  I understand that I may be receiving an intravenous contrast media and/or oral contrast media to enhance the visibility of certain tissues. Possible side effects may include, but are not limited to, pain or swelling at the site of injection, nausea, vomiting, and a warm, flushed sensation. Also potential allergic reactions including, but not limited to, hives, wheezing, difficulty breathing, and in rare instances, anaphylactic shock (with severe allergic reactions).  The purpose, benefits, and complications of the contrast procedure will be explained prior to any injection that may take place. I hereby consent to any measure necessary to correct complications, which may occur. I am aware that the practice of medicine is not an exact science and I acknowledge that no guarantee has been made to me concerning the results of the examination.  I confirm that the information I have provided is complete and accurate to the best of my knowledge.  UNDERSTAND THE RISKS, BENEFITS, AND ALTERNATIVES INVOLVED IN THE PROCEDURE. I HAVE HAD THE DPPORTUNITY TO ASK ANY QUESTIONS AND ALL OF MY QUESTIONS HAVE BEEN ANSWERED TO MY SATISFACTION.  Signature of Patient or Responsible Party (relationship)  Date and Time  V Contrast  Vedia/Amount:  Treatment:  IV/Injection Site after injection and after Needle Removal:	Have you ever had an allergic reaction to IV contrast:	Yes ☐ No Date of Reaction:		
understand that my physician may have requested the use of an intravenous contrast media that will assist the radiologist in better distinguishing certain anatomy or abnormalties that would otherwise be difficult or impossible to see.  I understand that the procedure to be performed on me involves the use of a high strength magnetic field, and possibly insertion needles and gadolinium containing solution, which may enhance the diagnostic accuracy of the procedure.  I understand that I may be receiving an intravenous contrast media and/or oral contrast media to enhance the visibility of certain tissues. Possible side effects may include, but are not limited to, pain or swelling at the site of injection, nausea, vomiting, and a warm, flushed sensation. Also potential allergic reactions including, but not limited to, hives, wheezing, difficulty breathing, and in rare instances, anaphylactic shock (with severe allergic reactions).  The purpose, benefits, and complications of the contrast procedure will be explained prior to any injection that may take place. I hereby consent to any measure neccessary to correct complications, which may occur. I am aware that the practice of medicine is not an exact science and I acknowledge that no guarantee has been made to me concerning the results of the examination.  I confirm that the information I have provided is complete and accurate to the best of my knowledge.  I UNDERSTAND THE RISKS, BENEFITS, AND ALTERNATIVES INVOLVED IN THE PROCEDURE. I HAVE HAD THE DPPORTUNITY TO ASK ANY QUESTIONS AND ALL OF MY QUESTIONS HAVE BEEN ANSWERED TO MY SATISFACTION.  Signature of Patient or Responsible Party (relationship)  Date and Time  V Contrast  Wedia/Amount:  Treatment:  IV/Injection Site after injection and after Needle Removal:	Type of reaction:			
understand that the procedure to be performed on me involves the use of a high strength magnetic field, and possibly insertion needles and gadolinium containing solution, which may enhance the diagnostic accuracy of the procedure.  I understand that I may be receiving an intravenous contrast media and/or oral contrast media to enhance the visibility of certain tissues. Possible side effects may include, but are not limited to, pain or swelling at the site of injection, nausea, vomiting, and a warm, flushed sensation. Also potential allergic reactions including, but not limited to, hives, wheezing, difficulty breathing, and in rare instances, anaphylactic shock (with severe allergic reactions).  The purpose, benefits, and complications of the contrast procedure will be explained prior to any injection that may take place. I hereby consent to any measure neccesary to correct complications, which may occur. I am aware that the practice of medicine is not an exact science and I acknowledge that no guarantee has been made to me concerning the results of the examination.  confirm that the information I have provided is complete and accurate to the best of my knowledge.  IUNDERSTAND THE RISKS, BENEFITS, AND ALTERNATIVES INVOLVED IN THE PROCEDURE. I HAVE HAD THE OPPORTUNITY TO ASK ANY QUESTIONS AND ALL OF MY QUESTIONS HAVE BEEN ANSWERED TO MY SATISFACTION.  Signature of Patient or Responsible Party (relationship)  Date and Time  Witness/Technologist  Patient Tolerated Procedure Well:  Witness/Technologist  V Contrast  Wedia/Amount:  Note that the procedure Well:  V/Injection Site after injection and after Needle Removal:	I UNDERSTAND THAT THIS CONTRAST AGENT (LIKE N	IANY NEW DRUGS) MAY CAUSE AN ALLE	RGIC REACTION.	
and gadolinium containing solution, which may enhance the diagnostic accuracy of the procedure.  Junderstand that I may be receiving an intravenous contrast media and/or oral contrast media to enhance the visibility of certain tissues. Possible side effects may include, but are not limited to, pain or swelling at the site of injection, nausea, vomiting, and a warm, flushed sensation. Also potential allergic reactions including, but not limited to, hives, wheezing, difficulty breathing, and in rare instances, anaphylactic shock (with severe allergic reactions).  The purpose, benefits, and complications of the contrast procedure will be explained prior to any injection that may take place. The purpose, benefits, and complications of the contrast procedure will be explained prior to any injection that may take place. The purpose, benefits, and complications of the contrast procedure will be explained prior to any injection that may take place. The purpose, benefits, and complications of the contrast procedure will be explained prior to any injection that may take place. The purpose, benefits, and complications, which may occur. I am aware that the practice of medicine is not an exact science and I acknowledge that no guarantee has been made to me concerning the results of the examination.  It understand that the information I have provided is complete and accurate to the best of my knowledge.  INDURERSTAND THE RISKS, BENEFITS, AND ALTERNATIVES INVOLVED IN THE PROCEDURE. I HAVE HAD THE DEPORTUNITY TO ASK ANY QUESTIONS AND ALL OF MY QUESTIONS HAVE BEEN ANSWERED TO MY SATISFACTION.  Signature of Patient or Responsible Party (relationship)  Date and Time  Witness/Technologist  Patient Tolerated Procedure Well:  Patient Tolerated Procedure Well:  Treatment:  IV/Injection Site after injection and after Needle Removal:			radiologist in better	
Possible side effects may include, but are not limited to, pain or swelling at the site of injection, nausea, vomiting, and a warm, flushed sensation. Also potential allergic reactions including, but not limited to, hives, wheezing, difficulty breathing, and in rare instances, anaphylactic shock (with severe allergic reactions).  The purpose, benefits, and complications of the contrast procedure will be explained prior to any injection that may take place. Increby consent to any measure neccesary to correct complications, which may occur. I am aware that the practice of medicine is not an exact science and I acknowledge that no guarantee has been made to me concerning the results of the examination.  Confirm that the information I have provided is complete and accurate to the best of my knowledge.  UNDERSTAND THE RISKS, BENEFITS, AND ALTERNATIVES INVOLVED IN THE PROCEDURE. I HAVE HAD THE OPPORTUNITY TO ASK ANY QUESTIONS AND ALL OF MY QUESTIONS HAVE BEEN ANSWERED TO MY SATISFACTION.  Signature of Patient or Responsible Party (relationship)  Date and Time  Witness/Technologist  Date and Time  Patient Tolerated Procedure Well:Yes No Reaction: Type: Time: Treatment: Injection Site after injection and after Needle Removal:			ossibly insertion needles	
I hereby consent to any measure neccesary to correct complications, which may occur. I am aware that the practice of medicine is not an exact science and I acknowledge that no guarantee has been made to me concerning the results of the examination.  I confirm that the information I have provided is complete and accurate to the best of my knowledge.  I UNDERSTAND THE RISKS, BENEFITS, AND ALTERNATIVES INVOLVED IN THE PROCEDURE. I HAVE HAD THE OPPORTUNITY TO ASK ANY QUESTIONS AND ALL OF MY QUESTIONS HAVE BEEN ANSWERED TO MY SATISFACTION.  Signature of Patient or Responsible Party (relationship)  Date and Time  Witness/Technologist  Date and Time  Patient Tolerated Procedure Well: Yes No Reaction: Type: Time: Treatment: Injection Time: IV/Injection Site after injection and after Needle Removal:	Possible side effects may include, but are not limited to, pain or sw	elling at the site of injection, nausea, vomiting,	and a warm, flushed	
UNDERSTAND THE RISKS, BENEFITS, AND ALTERNATIVES INVOLVED IN THE PROCEDURE. I HAVE HAD THE OPPORTUNITY TO ASK ANY QUESTIONS AND ALL OF MY QUESTIONS HAVE BEEN ANSWERED TO MY SATISFACTION.    Signature of Patient or Responsible Party (relationship)   Date and Time	I hereby consent to any measure neccesary to correct complication	ns, which may occur. I am aware that the prac	tice of medicine is not an	
Signature of Patient or Responsible Party (relationship)  Date and Time  Witness/Technologist  Date and Time  V Contrast Media/Amount:  V/Injection Site:  Injection Time:  Date and Time  Patient Tolerated Procedure Well:  Reaction: Type:  Treatment:  IV/Injection Site after injection and after Needle Removal:	I confirm that the information I have provided is complete and accu	rate to the best of my knowledge.		
Witness/Technologist  Date and Time  V Contrast Media/Amount:  Patient Tolerated Procedure Well: Reaction: Type: Time:  Treatment:  Njection Time:  IV/Injection Site after injection and after Needle Removal:				
V Contrast Media/Amount:	Signature of Patient or Responsible Party (relationship)	Date and Time	<del></del>	
Media/Amount: Reaction: Type: Time:    Treatment:  Njection Time:  IV/Injection Site after injection and after Needle Removal:	Witness/Technologist	Date and Time	<del></del>	
V/Injection Site:     Treatment:       njection Time:     IV/Injection Site after injection and after Needle Removal:	IV Contrast	Patient Tolerated Procedure Well:	☐ Yes ☐ No	
njection Time: IV/Injection Site after injection and after Needle Removal:	Media/Amount:	Reaction: Type:	Time:	
	IV/Injection Site:	Treatment:		
□ No Redness □ No Swelling □ Catheter Intact	Injection Time: IV/Injection Site after injection and after		r Needle Removal:	
	□ No Redness □ No Swelling □ Catheter In			

