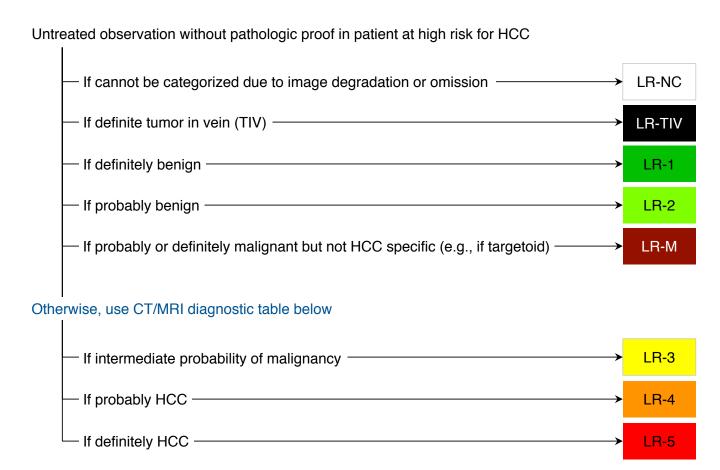


# CT/MRI LI-RADS® v2017 ESSENTIALS



### **CT/MRI Diagnostic Table**

Arterial phase hyperenhancement (APHE)		No APHE		APHE (not rim)		
Observation size (mm)		< 20	≥ 20	< 10	10-19	≥ 20
Count major features:  • "Washout" (not peripheral)  • Enhancing "capsule"  • Threshold growth	None	LR-3	LR-3	LR-3	LR-3	LR-4
	One	LR-3	LR-4	LR-4	LR-4 LR-5	LR-5
	≥Two	LR-4	LR-4	LR-4	LR-5	LR-5



Observations in this cell are categorized LR-4, except:

- LR-5g, if ≥ 50% diameter increase in < 6 months (equivalent to OPTN 5A-g)
- LR-5us, if "washout" and visibility at screening ultrasound (per AASLD HCC criteria)



### What's New in LI-RADS® v2017?

### New algorithms:

- US Screening and Surveillance
- CEUS Diagnosis
- CT/MRI Treatment Response Assessment

### New or revised categories for CT/MRI LI-RADS:

- LR-NC (new)
- LR-TIV (previously LR-5V)

### Threshold growth definition modified

New explicit criteria for LR-M

Updated algorithmic display for CT/MRI LI-RADS

New list-view displays to supplement algorithmic displays

Ancillary features are now optional and their use is clarified

New ancillary feature favoring malignancy: ultrasound visibility

Name change for ancillary feature: distinctive rim → nonenhancing capsule

Improved schematic diagrams, new time-intensity curves

#### **New FAQs**

#### Clarifies:

- Distinction between non-rim arterial phase hyperenhancement (major feature of HCC) vs. rim arterial phase hyperenhancement (feature of LR-M)
- Distinction between nonperipheral "washout" (major feature of HCC) vs. peripheral "washout" (feature of LR-M)
- Distinction between enhancing "capsule" (major feature of HCC) vs. nonenhancing "capsule" (ancillary feature favoring HCC)
- That ancillary features favoring malignancy include some favoring malignancy in general and others favoring HCC in particular
- · That CT/MRI LI-RADS can be used in liver transplant candidates with HCC
- · Categorization of tumor in vein and malignancy with infiltrative appearance

## Why is This Update Needed?

As new evidence emerges and based on feedback from users, LI-RADS evolves to better meet clinical, educational, and research needs. LI-RADS v2017 is the next step in this evolution.



### CT/MRI LI-RADS® v2017

### Apply in patients at high risk for HCC, namely those with:



- Cirrhosis OR
- Chronic hepatitis B viral infection OR
- Current or prior HCC

Including adult liver transplant candidates and recipients posttransplant

### Do not apply in patients:



- · Without the above risk factors
- < 18 years old</li>
- With cirrhosis due to congenital hepatic fibrosis
- With cirrhosis due to a vascular disorder such as hereditary hemorrhagic telangiectasia, Budd-Chiari syndrome, chronic portal vein occlusion, cardiac congestion, or diffuse nodular regenerative hyperplasia



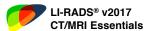
### Apply for multiphase exams performed with:

- CT or MRI with extracellular contrast agents (ECA) OR
- MRI with hepatobiliary contrast agents (HBA)

### Do not assign LI-RADS categories for observations:

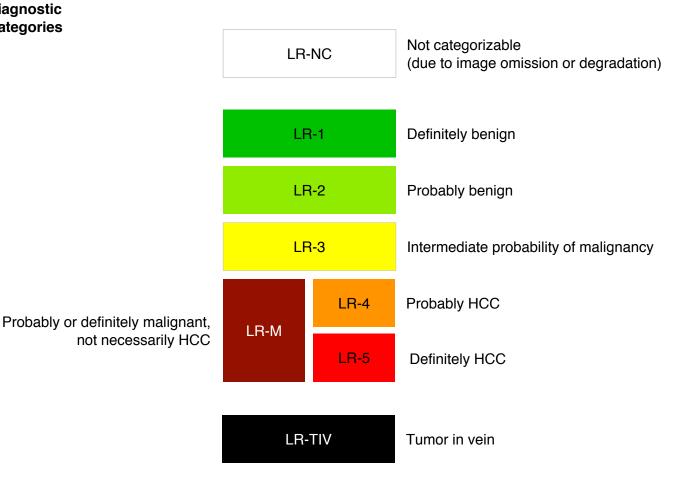


- · That are path-proven malignancies OR
- That are path-proven benign lesions of non-hepatocellular origin such as hemangiomas

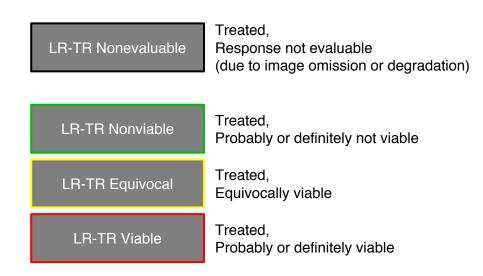


# CT/MRI LI-RADS® v2017 Categories

### Diagnostic **Categories**



### **Treatment Response** Categories



LR-4

LR-5



# Step 1. Apply CT/MRI LI-RADS® Diagnostic Algorithm

Untreated observation without pathologic proof in patient at high risk for HCC

If cannot be categorized due to image degradation or omission 

LR-NC

If definite tumor in vein (TIV)

If definitely benign

LR-1

If probably benign

LR-2

If probably or definitely malignant but not HCC specific (e.g., if targetoid)

LR-M

Otherwise, use CT/MRI diagnostic table below

If intermediate probability of malignancy

# **CT/MRI Diagnostic Table**

If probably HCC -

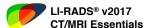
If definitely HCC —

Arterial phase hyperenhancement (APHE)		No APHE		APHE (not rim)		
Observation size (mm)		< 20	≥ 20	< 10	10-19	≥ 20
Count major features:  • "Washout" (not peripheral)  • Enhancing "capsule"  • Threshold growth	None	LR-3	LR-3	LR-3	LR-3	LR-4
	One	LR-3	LR-4	LR-4	LR-4 LR-5	LR-5
	≥Two	LR-4	LR-4	LR-4	LR-5	LR-5



Observations in this cell are categorized LR-4, except:

- LR-5g, if ≥ 50% diameter increase in < 6 months (equivalent to OPTN 5A-g)</li>
- LR-5us, if "washout" and visibility at screening ultrasound (per AASLD HCC criteria)

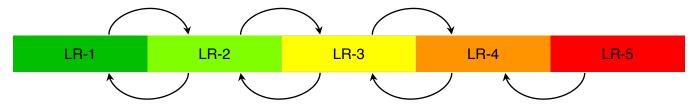


# Step 2. Optional: Apply Ancillary Features (AFs)

Ancillary features may be used **at radiologist discretion** for: Improved detection, increased confidence, or category adjustment

For category adjustment (upgrade or downgrade), apply ancillary features as follows:

One or more ancillary features favoring malignancy: upgrade by 1 category up to LR-4 (Absence of these ancillary features should not be used to downgrade)



One or more ancillary features favoring benignity: downgrade by 1 category (Absence of these ancillary features should not be used to upgrade)

If there are conflicting AFs (i.e., one or more favoring malignancy <u>and</u> one or more favoring benignity):

Do not adjust category

Ancillary features cannot be be used to upgrade to LR-5

### **Ancillary features favoring malignancy**

# Favoring malignancy in general, not HCC in particular

- US visibility as discrete nodule
- Subthreshold growth
- · Restricted diffusion
- Mild-moderate T2 hyperintensity
- Corona enhancement
- Fat sparing in solid mass
- Iron sparing in solid mass
- Transitional phase hypointensity
- Hepatobiliary phase hypointensity

#### Favoring HCC in particular

- Nonenhancing "capsule"
- Nodule-in-nodule
- Mosaic architecture
- Blood products in mass
- Fat in mass, more than adjacent liver

### **Ancillary features favoring benignity**

- Size stability > 2 yrs
- Size reduction
- · Parallels blood pool
- Undistorted vessels
- Iron in mass, more than liver
- Marked T2 hyperintensity
- Hepatobiliary phase isointensity

If unsure about presence of any ancillary feature: characterize that feature as absent

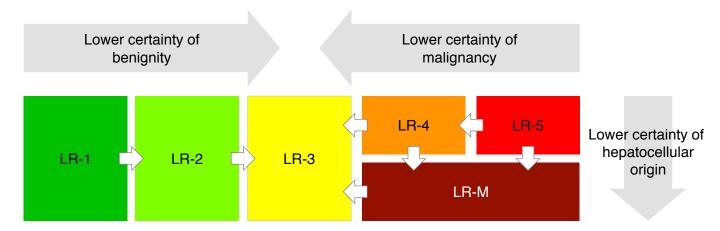


# Step 3. Apply Tiebreaking Rules if Needed

If unsure about presence of TIV, do not categorize as LR-TIV



If unsure between two categories, choose the one reflecting lower certainty



# **Step 4. Final Check**

After Steps 1, 2, and 3 -

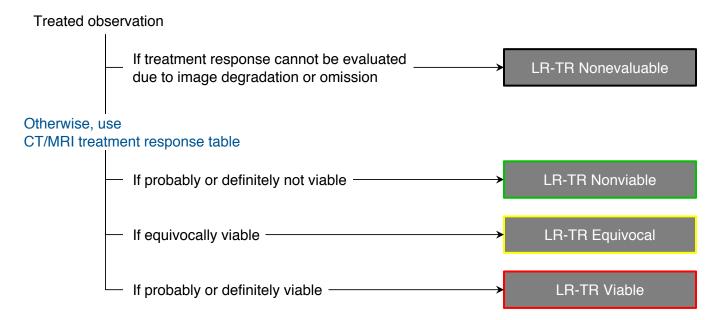
Ask yourself if the assigned category seems reasonable and appropriate

If YES: You are done, move on the next observation (if any).

**If NO:** Assigned LI-RADS category may be inappropriate, so reevaluate.



# Step 1. Apply LI-RADS® CT/MRI Treatment Response Algorithm

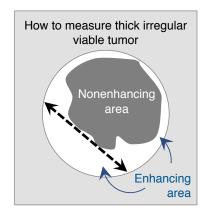


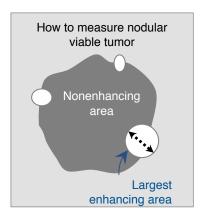
### **CT/MRI Treatment Response Table**

Response Category	Criteria
LR-TR Nonviable	<ul> <li>No lesional enhancement <b>OR</b></li> <li>Treatment-specific expected enhancement pattern</li> </ul>
LR-TR Equivocal	Enhancement atypical for treatment-specific expected enhancement pattern and not meeting criteria for probably or definitely viable
LR-TR Viable	Nodular, masslike, or thick irregular tissue in or along the treated lesion with any of the following:  • Arterial phase hyperenhancement <b>OR</b> • Washout appearance <b>OR</b> • Enhancement similar to pretreatment



# **Step 2. Measure Viable Tumor Size**





# Size of equivocally, probably, or definitely viable tumor

Longest dimension through enhancing area of treated lesion, not traversing nonenhancing area

# Step 3. Apply Tiebreaking Rule if Needed

If unsure between two categories, choose the one reflecting lower certainty as illustrated below

Lower certainty of nonviability

Lower certainty of viability

LR-TR Nonviable



LR-TR Equivocal



LR-TR Viable

# Step 4. Final Check

After Steps 1, 2, and 3 -

Ask yourself if the assigned response category seems reasonable and appropriate

**If YES:** You are done, move on the next observation (if any).

**If NO:** Assigned LI-RADS category may be inappropriate, so reevaluate.