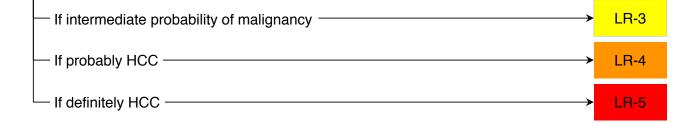
# CT/MRI LI-RADS<sup>®</sup> v2017 ESSENTIALS

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)	→ LR-TIV
If definitely benign	→ LR-1
— If probably benign ———————————————————————————————————	→ LR-2
	→ LR-M

#### Otherwise, use CT/MRI diagnostic table below



### **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:	None	LR-3	LR-3	LR-3	LR-3	LR-4
<ul><li> "Washout" (not peripheral)</li><li> Enhancing "capsule"</li><li> Threshold growth</li></ul>	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  $\geq$  50% diameter increase in < 6 months (equivalent to OPTN 5A-g)

• LR-5us, if "washout" and visibility at screening ultrasound (per AASLD HCC criteria)

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



### 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  $\rightarrow$  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<sup>®</sup> 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
- 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:

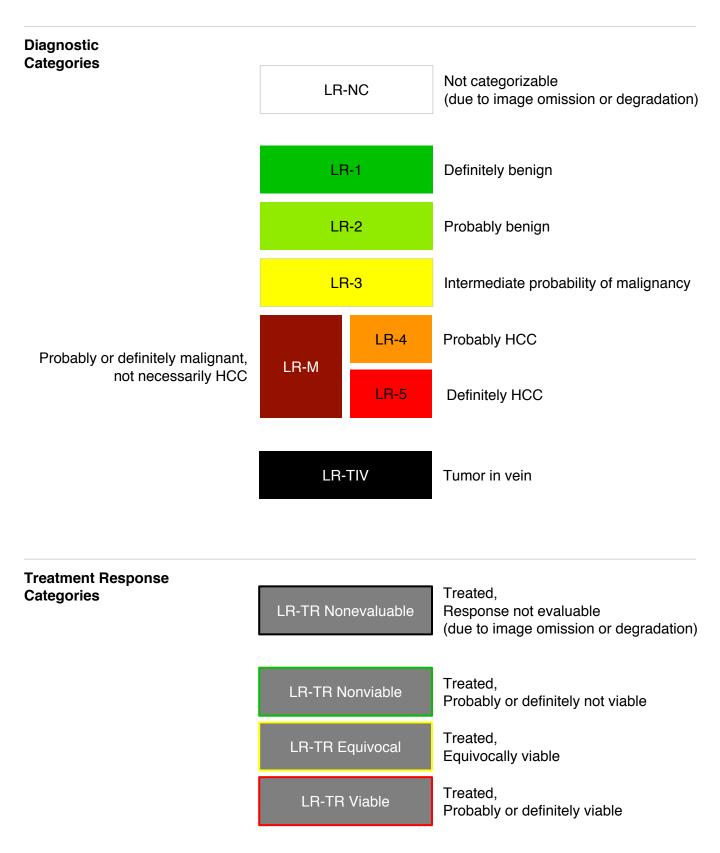


- That are path-proven malignancies OR
- That are path-proven benign lesions of non-hepatocellular origin such as hemangiomas





# CT/MRI LI-RADS® v2017 Categories





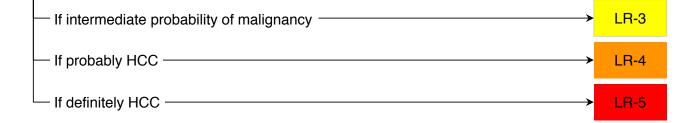


# Step 1. Apply CT/MRI LI-RADS® Diagnostic Algorithm

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

If cannot be categorized due to image degradation or omission	$\rightarrow$	LR-NC
If definite tumor in vein (TIV)	$\rightarrow$	LR-TIV
If definitely benign	$\rightarrow$	LR-1
— If probably benign ———————————————————————————————————	<b>→</b>	LR-2
- If probably or definitely malignant but not HCC specific (e.g., if targetoid)	$\rightarrow$	LR-M

#### Otherwise, use CT/MRI diagnostic table below



### **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:	None	LR-3	LR-3	LR-3	LR-3	LR-4
<ul><li> "Washout" (not peripheral)</li><li> Enhancing "capsule"</li><li> Threshold growth</li></ul>	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)

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



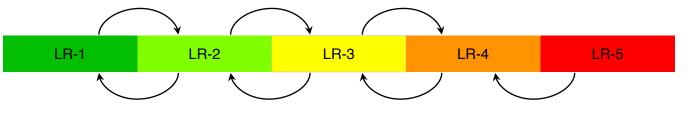
# Step 2. Optional: Apply Ancillary Features (AFs)

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	Ancillary features favoring benignity
<ul> <li>Favoring malignancy in general, not HCC in particular</li> <li>US visibility as discrete nodule</li> <li>Subthreshold growth</li> <li>Restricted diffusion</li> <li>Mild-moderate T2 hyperintensity</li> <li>Corona enhancement</li> <li>Fat sparing in solid mass</li> <li>Iron sparing in solid mass</li> <li>Transitional phase hypointensity</li> <li>Hepatobiliary phase hypointensity</li> </ul>	<ul> <li>Size stability &gt; 2 yrs</li> <li>Size reduction</li> <li>Parallels blood pool</li> <li>Undistorted vessels</li> <li>Iron in mass, more than liver</li> <li>Marked T2 hyperintensity</li> <li>Hepatobiliary phase isointensity</li> </ul>
<ul> <li>Favoring HCC in particular</li> <li>Nonenhancing "capsule"</li> <li>Nodule-in-nodule</li> <li>Mosaic architecture</li> <li>Blood products in mass</li> <li>Fat in mass, more than adjacent liver</li> </ul>	

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

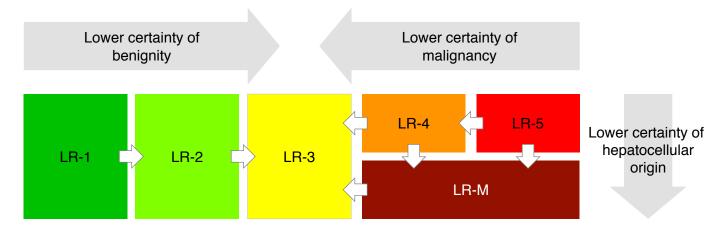


# Step 3. Apply Tiebreaking Rules if Needed

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 -

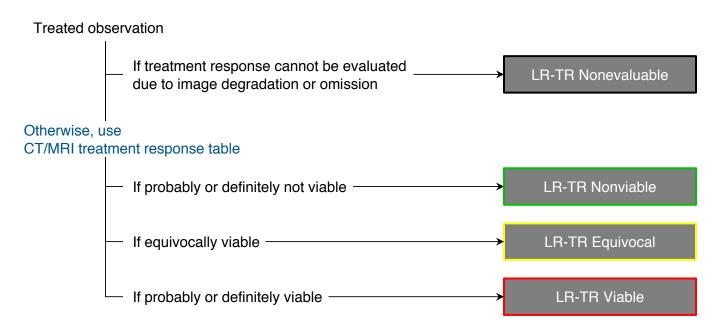
Ask yourself if the assigned category seems reasonable and appropriate

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<sup>®</sup> CT/MRI Treatment Response Algorithm

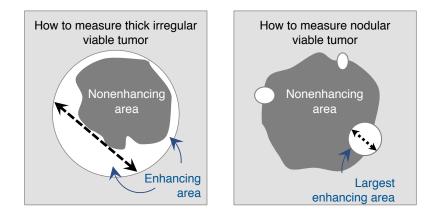


#### **CT/MRI Treatment Response Table**

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	<ul> <li>Nodular, masslike, or thick irregular tissue in or along the treated lesion with any of the following:</li> <li>Arterial phase hyperenhancement OR</li> <li>Washout appearance OR</li> <li>Enhancement similar to pretreatment</li> </ul>



## Step 2. Measure Viable Tumor Size

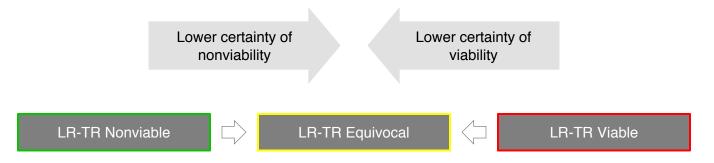


# Size of equivocally, probably, or definitely viable tumor

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



### 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.