

Hepatocellular Carcinoma Management Guidelines

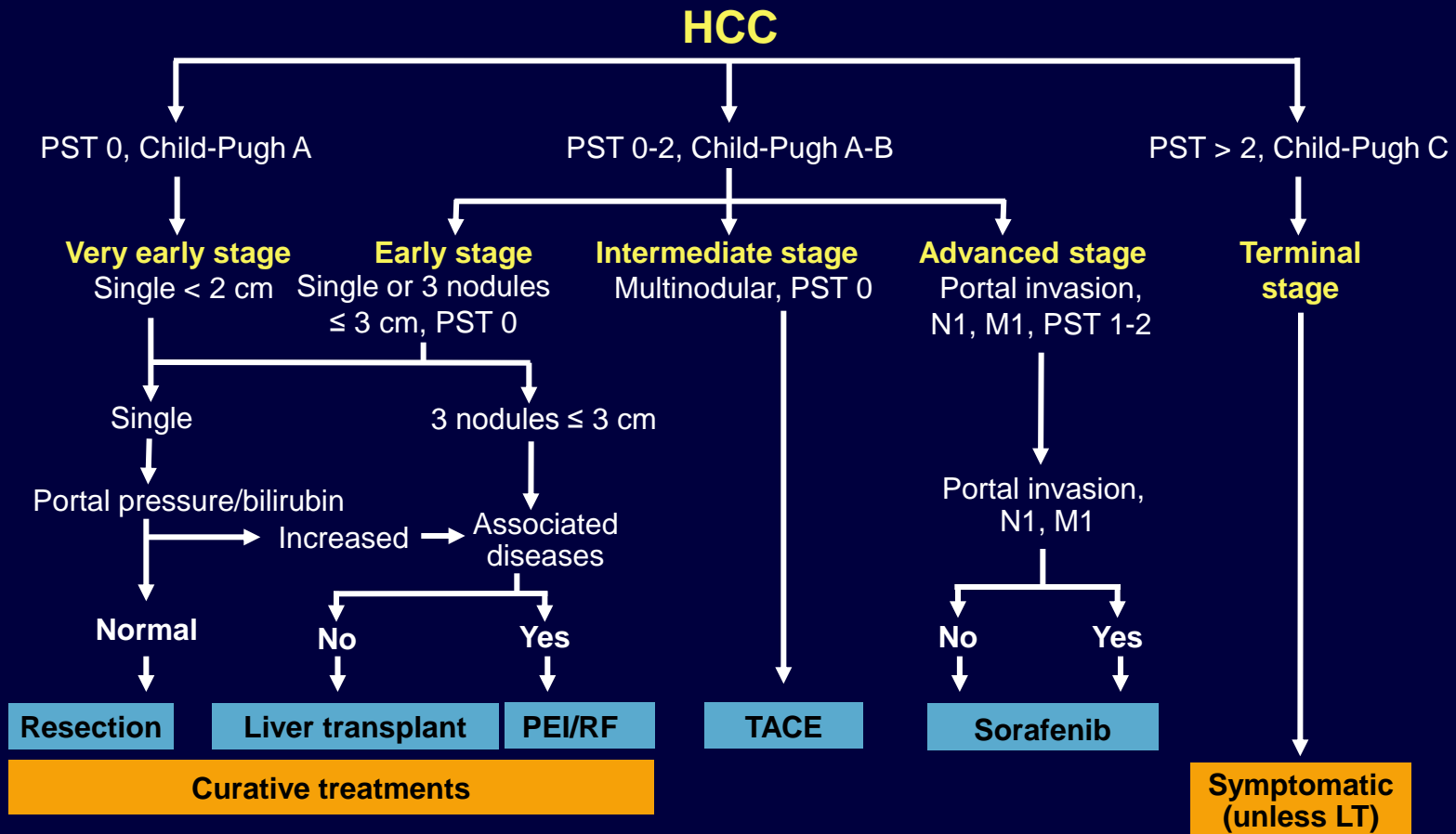
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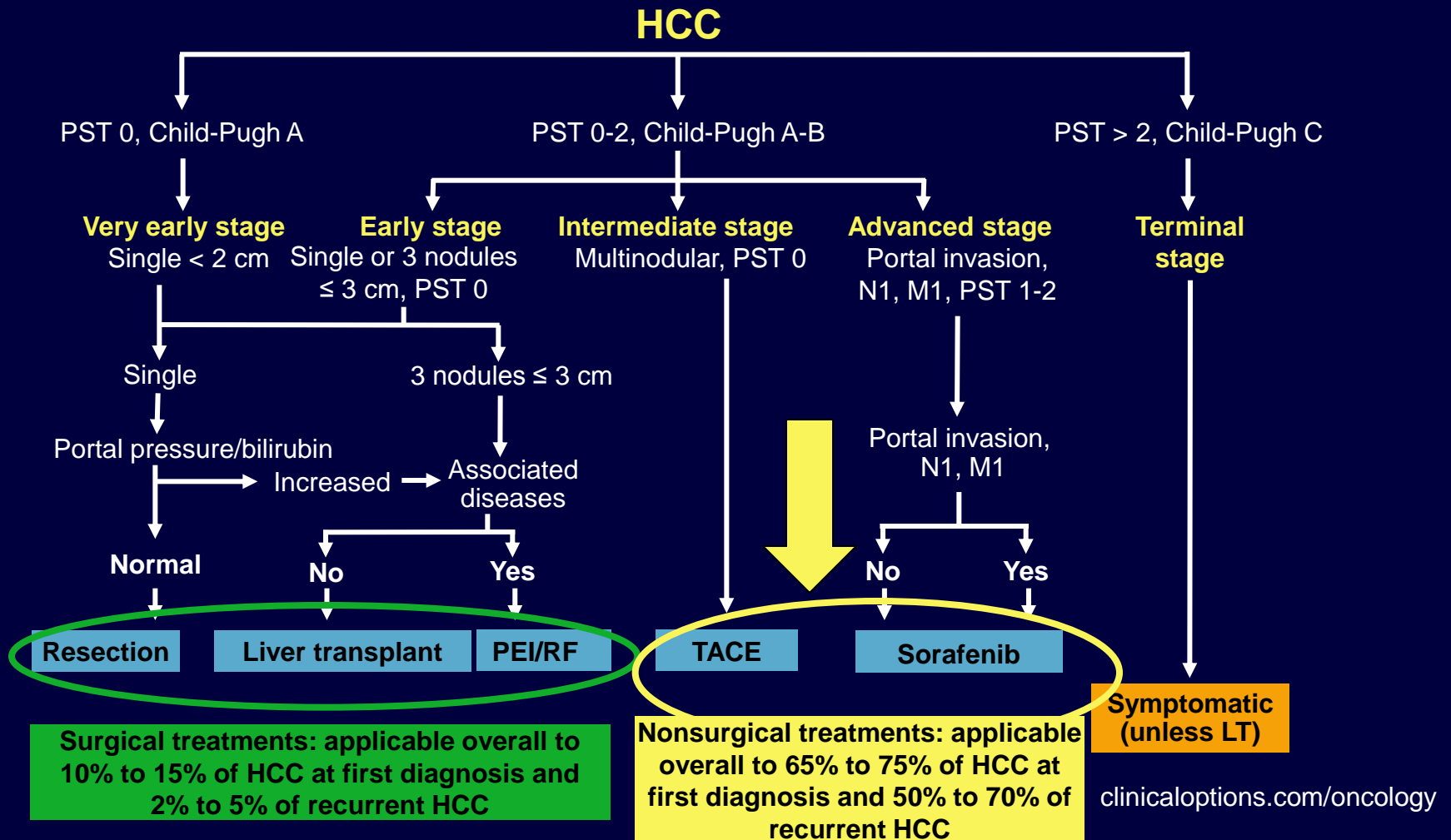
Cairo University

Staging Strategy and Treatment for Patients With HCC



Llovet JM, et al. J Natl Cancer Inst. 2008;100:698-711.
Bruix J, et al. Hepatology. 2005;42:1208-1236.

Staging Strategy and Treatment for Patients With HCC



Approved Curative Treatments for Unresectable HCC: Percutaneous Ablation

- Local ablation: safe and effective therapy for patients who cannot undergo resection or as a bridge to transplantation (level II)
- Alcohol injection and radiofrequency are equally effective for tumors < 2 cm
 - However, necrotic effect of radiofrequency is more predictable in all tumor sizes
 - In addition, efficacy is clearly superior to that of alcohol injection in larger tumors (level I)

Approved & Investigational Noncurative Agents for Unresectable HCC

- AASLD 2005 recommendations
 - **Chemoembolization (TACE)** (with doxorubicin, cisplatin, or mitomycin) **is recommended** as first-line, noncurative therapy for nonsurgical patients with large/multifocal HCC who do not have vascular invasion or extrahepatic spread (and are not eligible for percutaneous ablation) (level I)
 - Tamoxifen, octreotide, antiandrogens, and hepatic artery ligation/embolization **are not recommended** (level I); other options such as drug-eluting beads, radiolabelled yttrium glass beads, radiolabelled lipiodol, or immunotherapy **cannot be recommended** as standard therapy for advanced HCC outside clinical trials

Treatment of Advanced HCC (BCLC Stage C)

- AASLD 2005 recommendation: no standard therapy; patients should enroll in a randomized clinical trial^[1]
- 2008 recommendation: sorafenib has become the standard of care for advanced HCC^[2]
 - Prolongs OS by 3 months^[3]
 - 1-year survival: 44%^[4]

1. Bruix J, et al. Hepatology. 2005;42:1208-1236.

2. Llovet JM, et al. J Hepatol. 2008;48:S20-S37.

3. Llovet J, et al. ASCO 2007. Abstract LBA 1.

4. Llovet J, et al. N Engl J Med. 2008;359:378-390.

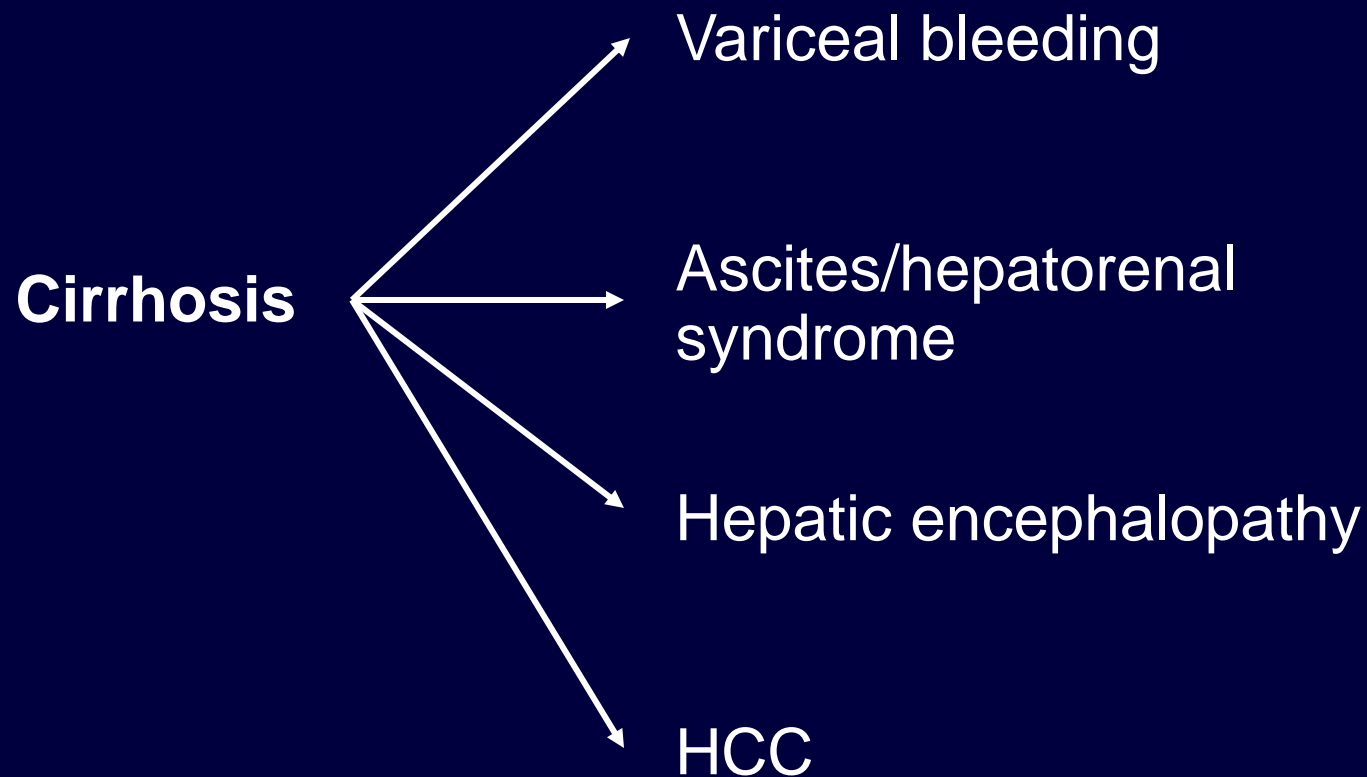
Intermediate/Advanced HCC: Future Directions

- 499 trials registered at clinicaltrials.gov for HCC as of August 21, 2008, including
 - Improving efficacy of RF and TACE (drug-eluting beads)
 - Exploring alternative treatments for intermediate HCC (yttrium-90)
 - Molecularly targeted agents in combination regimens (advanced HCC)
 - Molecularly targeted agents in combination with current modalities (early/intermediate HCC)
 - Improving tumor targeting of chemotherapeutic agents
 - New molecular targets and new molecularly targeted agents

Treatment of Liver Disease

- Hepatitis C: IFN + RBV
- Hepatitis B: IFN, lamivudine, adefovir, entecavir
- Alcohol: Abstinence
- Primary biliary cirrhosis: Ursodeoxycholic acid
- Hemochromatosis: Phlebotomy
- Alpha-1 ATD: None
- Nonalcoholic fatty liver: Diet and exercise
- Wilson's disease: Zinc, trientine
- Sclerosing cholangitis: Ursodeoxycholic acid, biliary stents
- Autoimmune hepatitis: Immunosuppression

Complications of Cirrhosis



Management of HCC

- Liver transplantation
- Resection
- Tumor ablation
 - Radiofrequency thermal ablation
 - Alcohol injection
 - Chemoembolization
- Targeted molecular therapy
- Chemotherapy
 - Regional/systemic



**Potentially
curative**

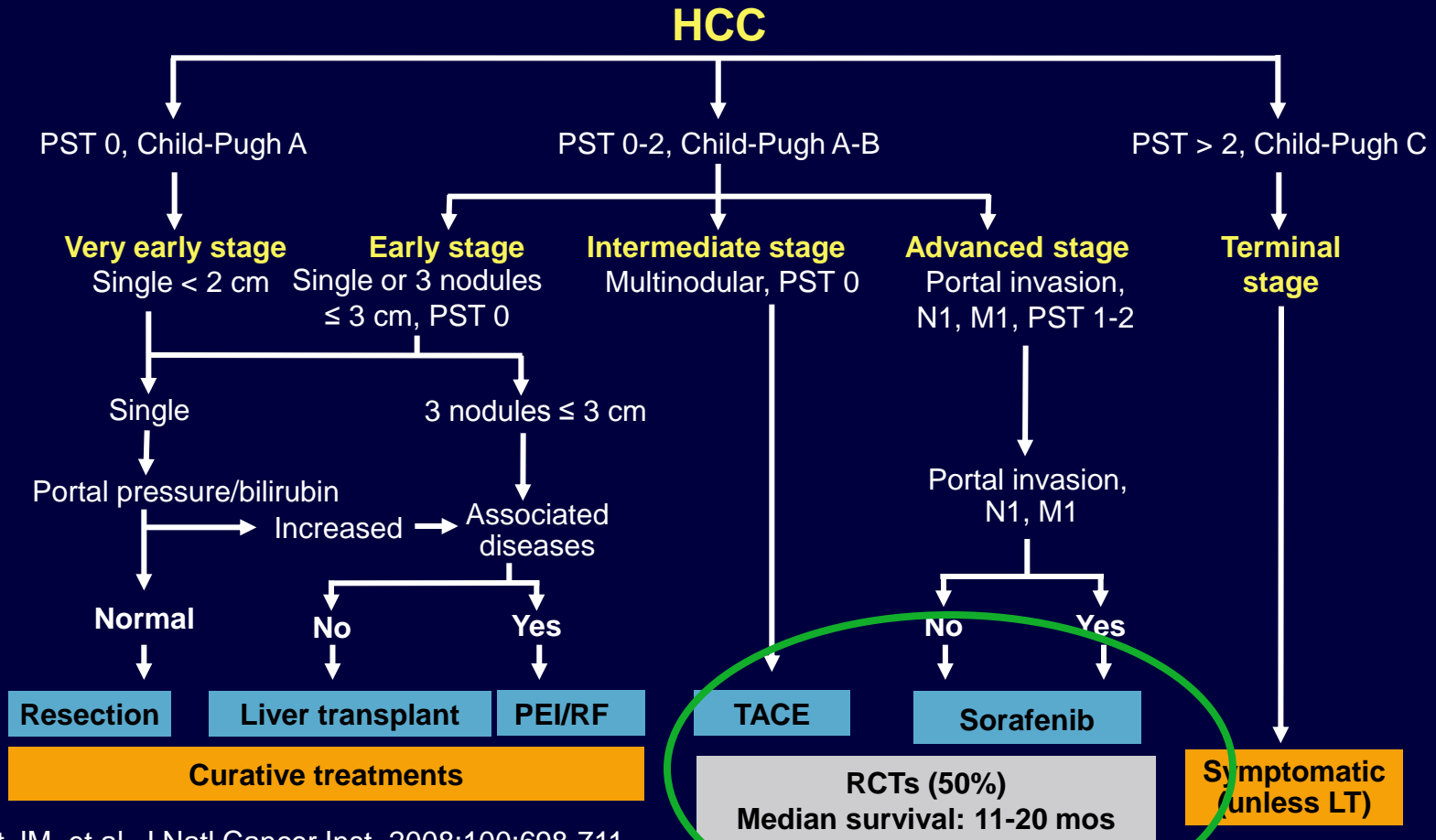
Evidence of Benefit in Treatment of HCC

Treatment	Benefit	Evidence
Surgical treatments		
Resection	Increased survival	Case series
▪ Adjuvant therapies	Uncertain	Randomized trial, meta-analysis, nonblinded
Liver transplantation	Increased survival	Case series
▪ Neoadjuvant therapies	Treatment response	Nonrandomized trials
Locoregional treatment		
Percutaneous treatment	Increased survival	Case series
RFA vs PEI	Better local control	Randomized trial, meta-analysis, nonblinded
Chemoembolization	Increased survival	Randomized trial, meta-analysis, nonblinded
Arterial chemotherapy	Treatment response	Case series
Internal radiation	Treatment response	Case series

Evidence of Benefit in Treatment of HCC (cont'd)

Treatment	Benefit	Evidence
Systemic therapies		
Sorafenib	Increased survival	Randomized trial, meta-analysis, double blinded
Tamoxifen	No benefit	Randomized trial, meta-analysis, double blinded
Chemotherapy	No benefit	Randomized trial, meta-analysis, nonblinded
IFN	No benefit	Randomized trial, meta-analysis, nonblinded

Staging Strategy and Treatment for Patients With HCC

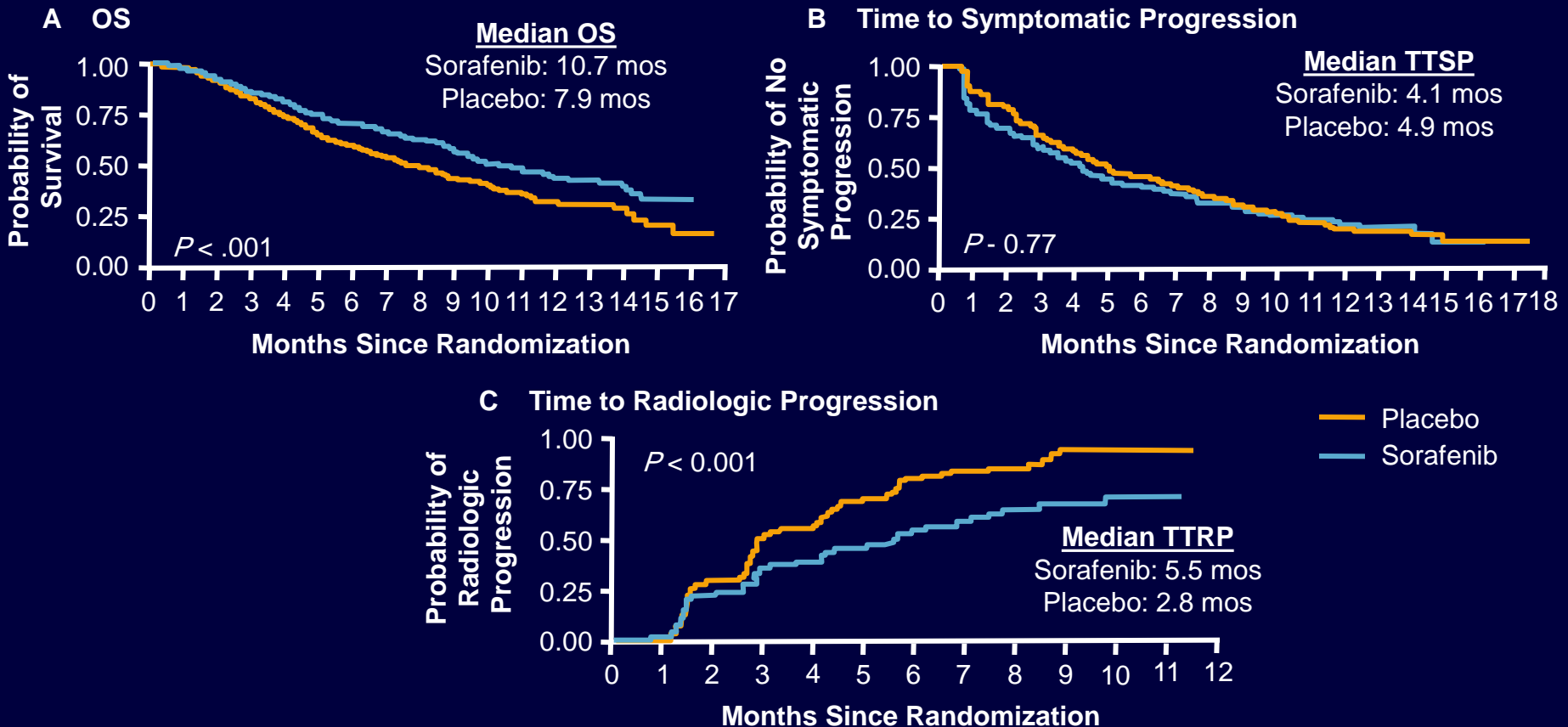


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Bruix J, et al. Hepatology. 2005;42:1208-1236.

Sorafenib in Advanced HCC: The SHARP Trial

- Entry criteria
 - Advanced HCC
 - Not eligible for or failed surgical or locoregional therapies
 - Child-Pugh class A disease
 - At least 1 untreated target lesion
 - Exclusions
 - Previous chemotherapy
 - Previous molecularly targeted therapy

The SHARP Trial: OS and Time to Progression



Strategies for Managing AEs

- Hand-foot syndrome
 - Creams and lotions
 - Avoid tight footwear
 - May require dose reduction
- Diarrhea
 - Antidiarrheal agents if severe
- Fatigue
 - Consider modafinil or methylphenidate if severe
- Hypertension
 - Start or adjust antihypertensives

Intra-arterial Locoregional Therapy

- Established
 - TACE
 - Radioembolization: yttrium-90 radioactive microspheres
- Undergoing clinical trials
 - Drug-eluting beads

Chemoembolization: Randomized Trials (Nearly Identical Techniques)

Lo et al^[1]: N = 80 with newly diagnosed unresectable HCC, 80% HBV positive, 7-cm tumors (60% multifocal)

Technique	Survival, %		
	Year 1	Year 2	Year 3
TACE	57	31	26
Supportive care	32	11	3

Llovet et al^[2]: N = 112 with unresectable HCC, 80% to 90% HCV positive, 5-cm tumors (~ 70% multifocal)

Technique	Survival, %	
	Year 1	Year 2
TACE	82	63
Supportive care	63	27

1. Lo CM, et al. Hepatology. 2002;35:1164-1171.
2. Llovet JM, et al. Lancet. 2002;359:1734-1739.

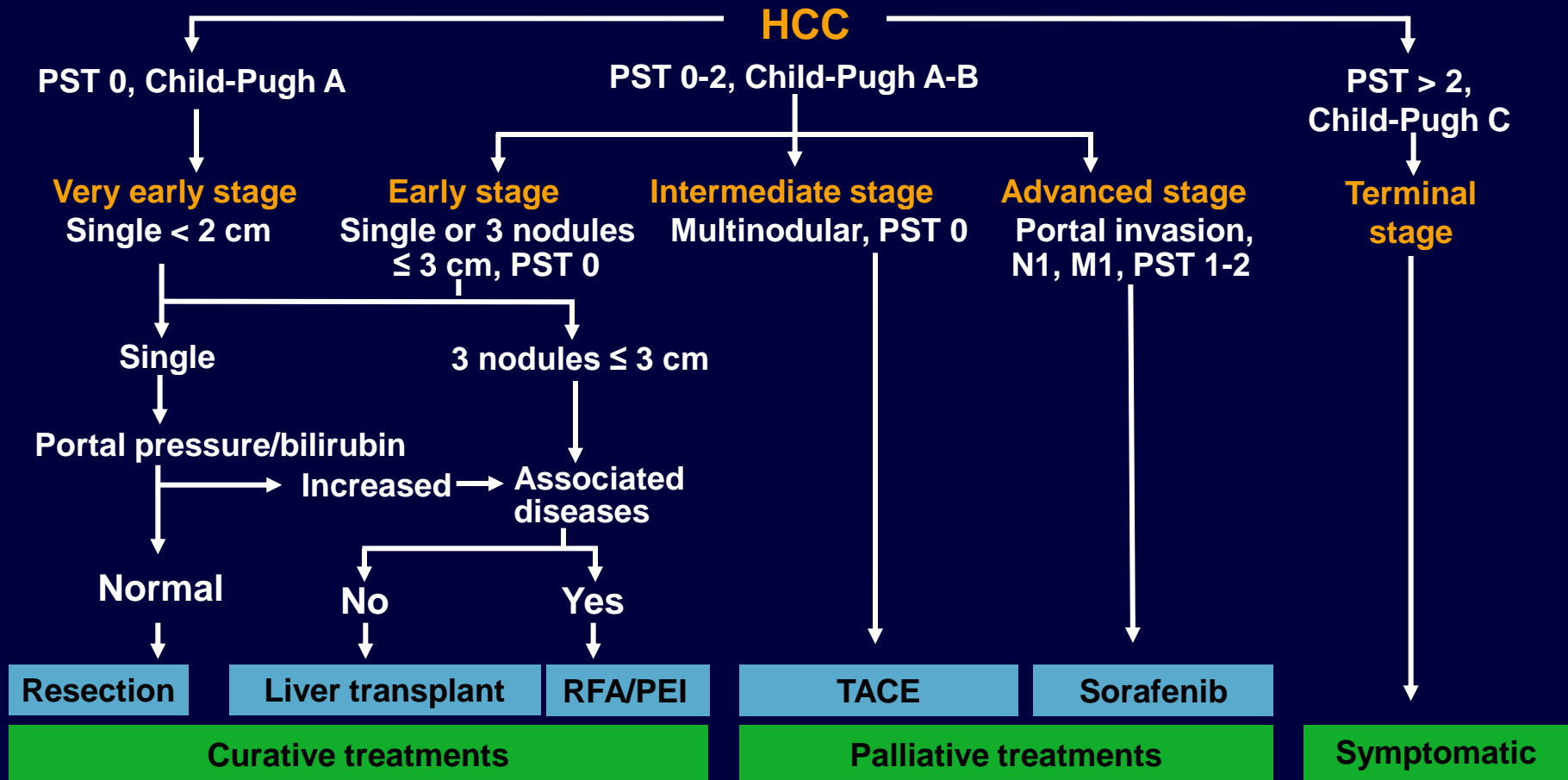
Chemoembolization: Ineligibility Criteria

- Absolute contraindications
 - Child-Pugh class C disease
 - Poor performance status (ECOG PS > 2)
- Relative contraindication
 - Extrahepatic disease (benefit unclear)
- Former contraindication
 - PVT
 - Minimize embolization and be more selective

Conclusions

- TACE accepted as treatment of choice for unresectable (nonablatable?) HCC
- Prolonged survival established through randomized trials and prospective studies
- Best vs good performance status, Child-Pugh class A-B
- Role for yttrium-90 microspheres
- Growing role for doxorubicin-loaded beads, pending outcome of clinical trials

AASLD Guidelines: Staging Strategy and Treatment for Patients With HCC

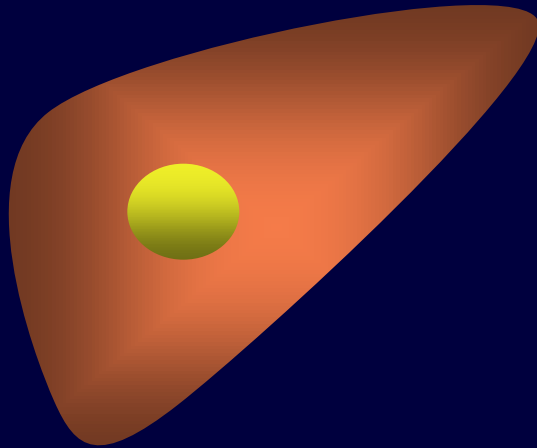


- WHY LT for HCC?
- LT is attractive because both the tumour as well as cirrhosis presents in 50-90% which is the fertile soil for the development of new lesions are removed by this procedure

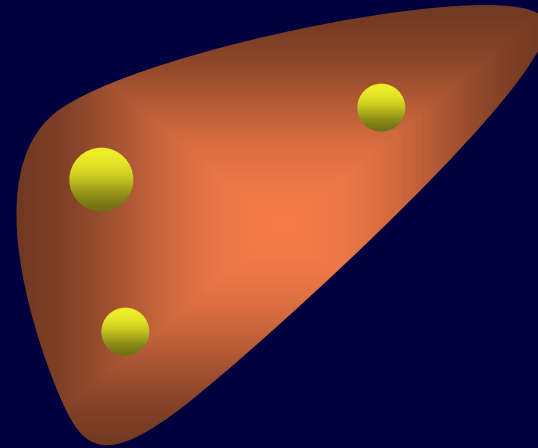
- What should be a suitable criteria for liver transplantation for HCC?

Liver Transplantation for HCC: Milan Criteria (Stage 1 and 2)

Single tumor, not > 5 cm



Up to 3 tumors, none > 3 cm



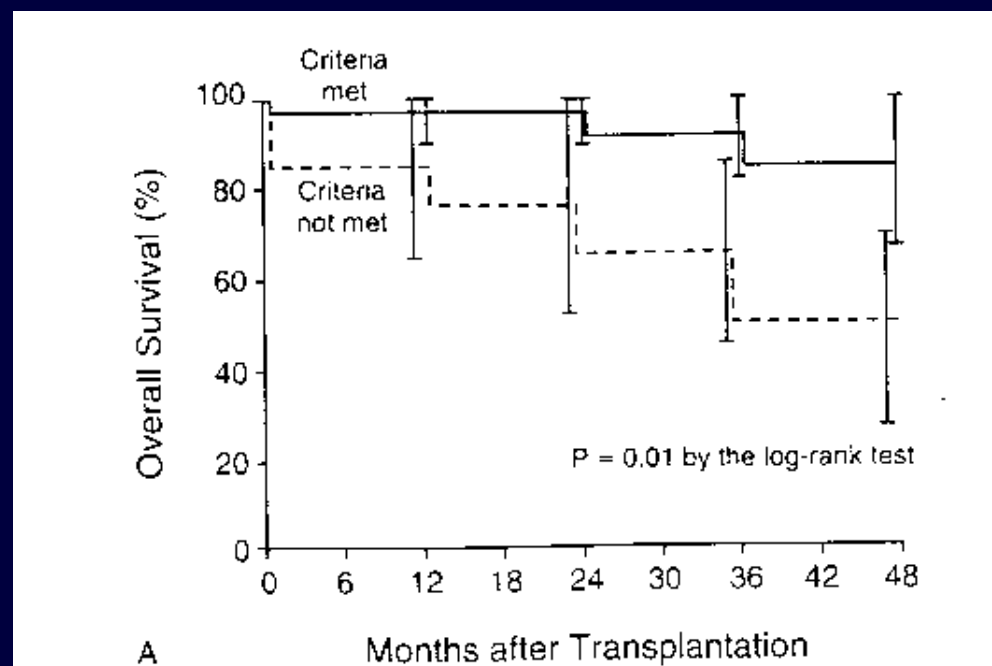
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**Absence of macroscopic vascular invasion,
absence of extrahepatic spread**

Current indications

Mazzafero et al. N Engl J Med 1996

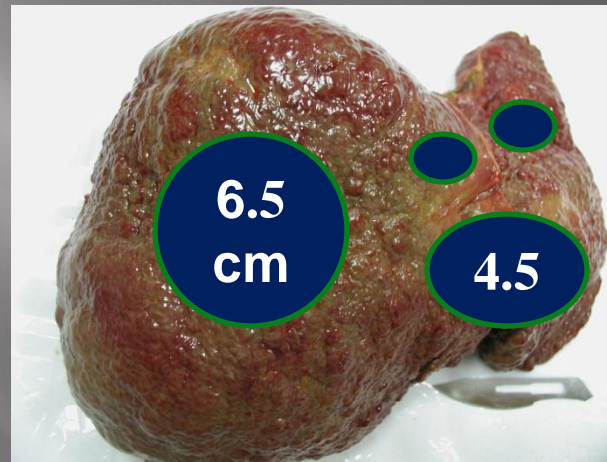
- Single tumour < 5 cm
or ≤ 3 nodules < 3 cm
- No portal thrombosis
- Overall survival
 - 1-year : 90 %
 - 4-year : 75 %
- Recurrence : 8 %



Survival according the Milan's Criteria
on the explanted liver

- Can we expand the Milan criteria for hepatocellular carcinoma in liver transplantation?

UCSF Criteria



Total \leq 8 cm

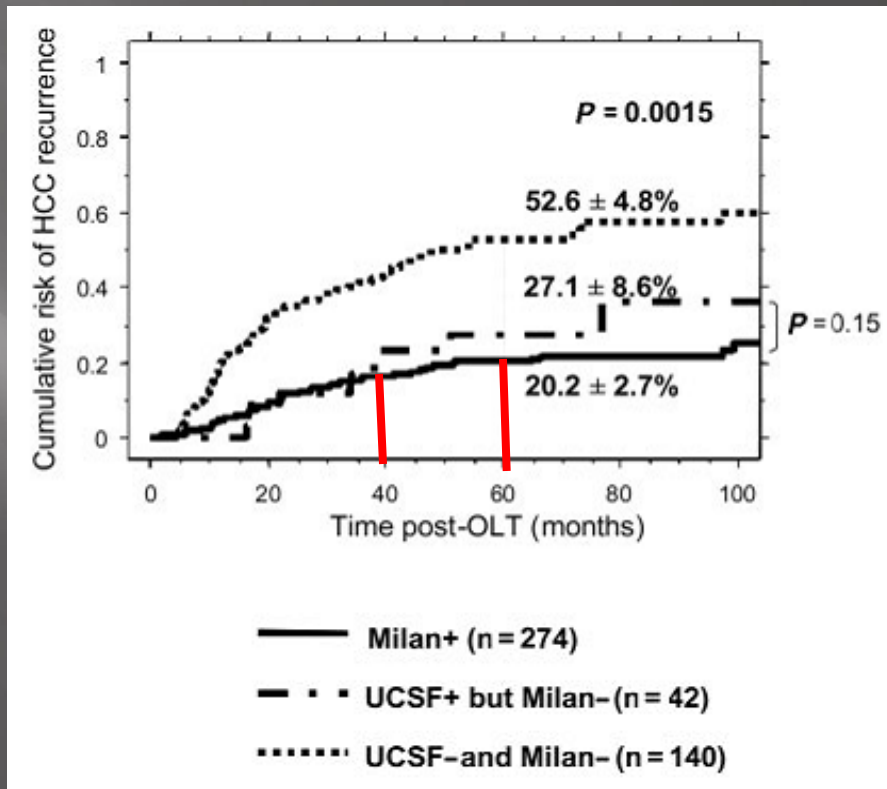
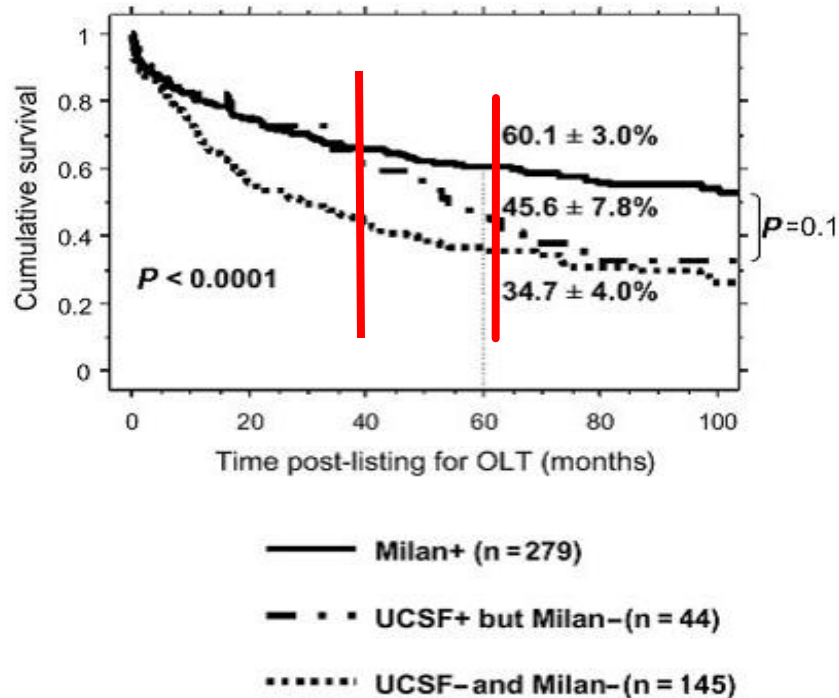
- * Explant pathology: *Criticised*
- * Clinical applicability

UCSF Criteria

Results:

- Tumors within UCSF criteria
 - 1 yr survival 90%
 - 5 yr survival 75%
- Tumors outside UCSF criteria
 - 1 yr survival 50%
 - 5 yr survival < 30%

Milan & UCSF Criteria Radiologic Staging



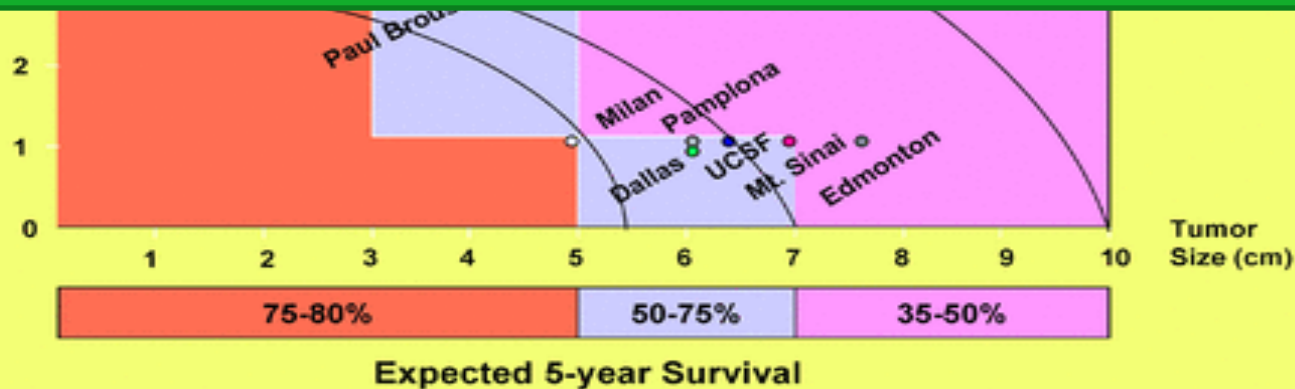
Beyond Milan Criteria–HCC “Metro Ticket”

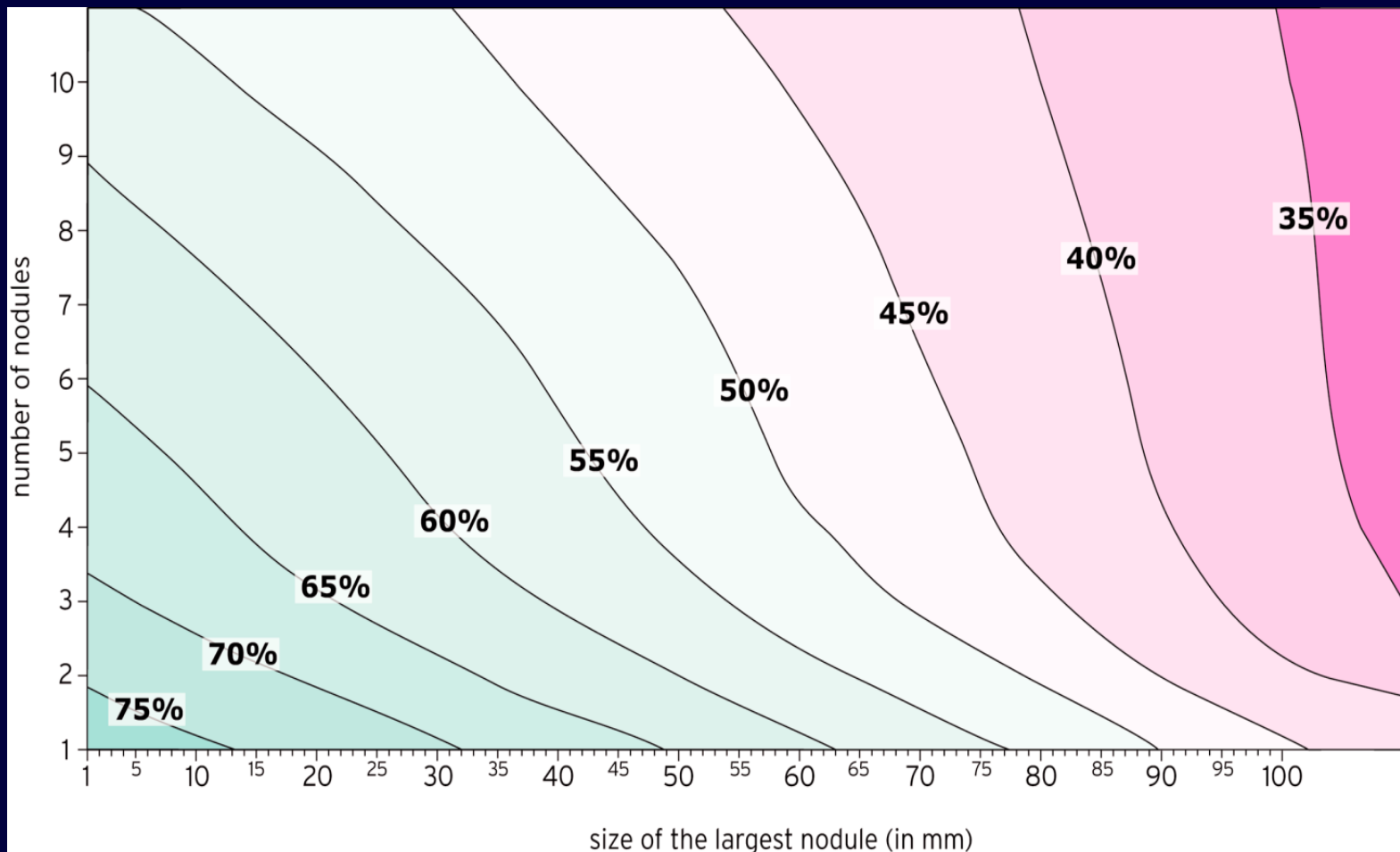
HCC “Metro Ticket” - The further the distance, the higher the price

Number of nodules

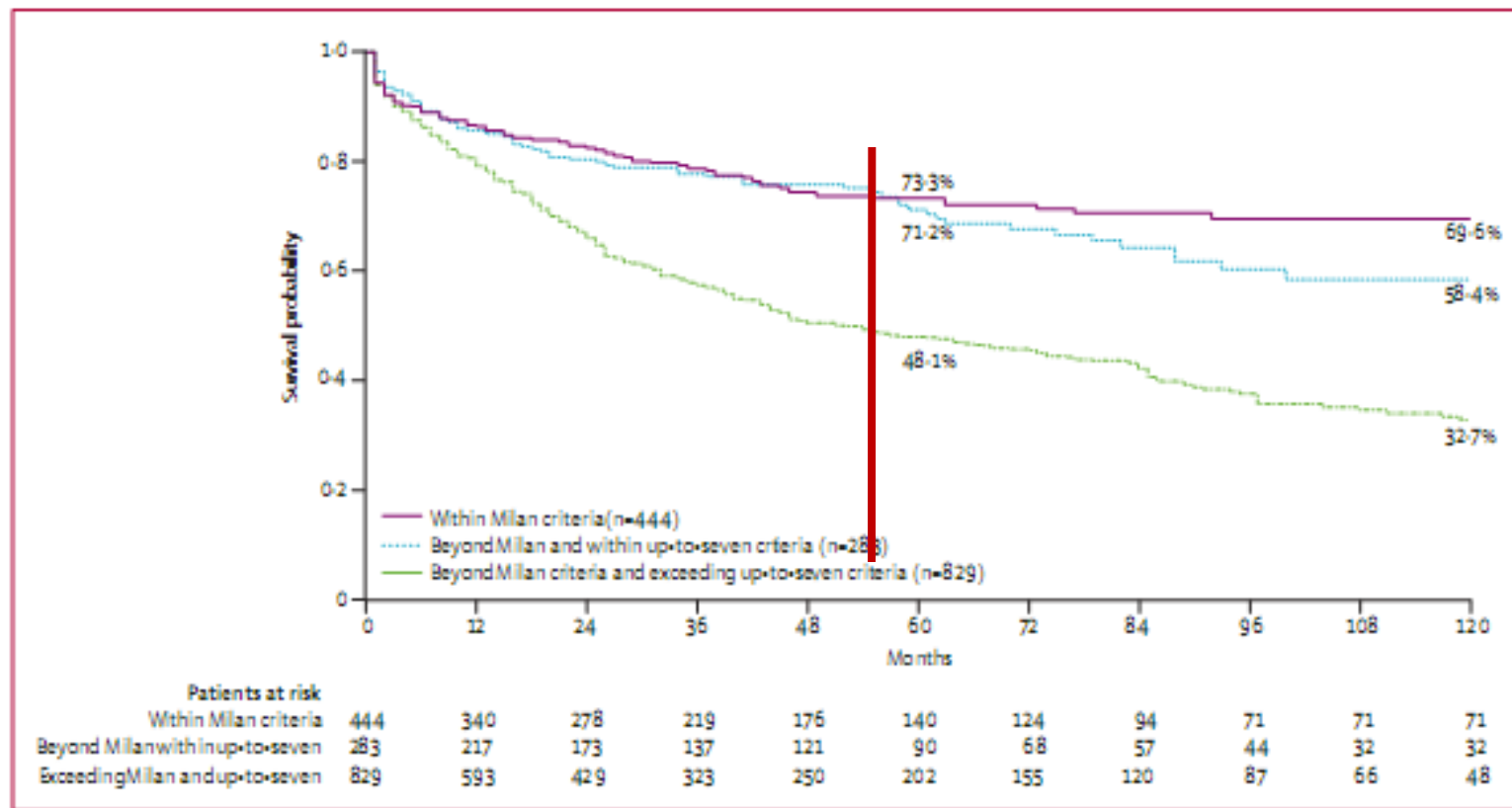
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Metroticket: “Up To Seven” Criteria
Largest tumor + tumor number ≤ 7



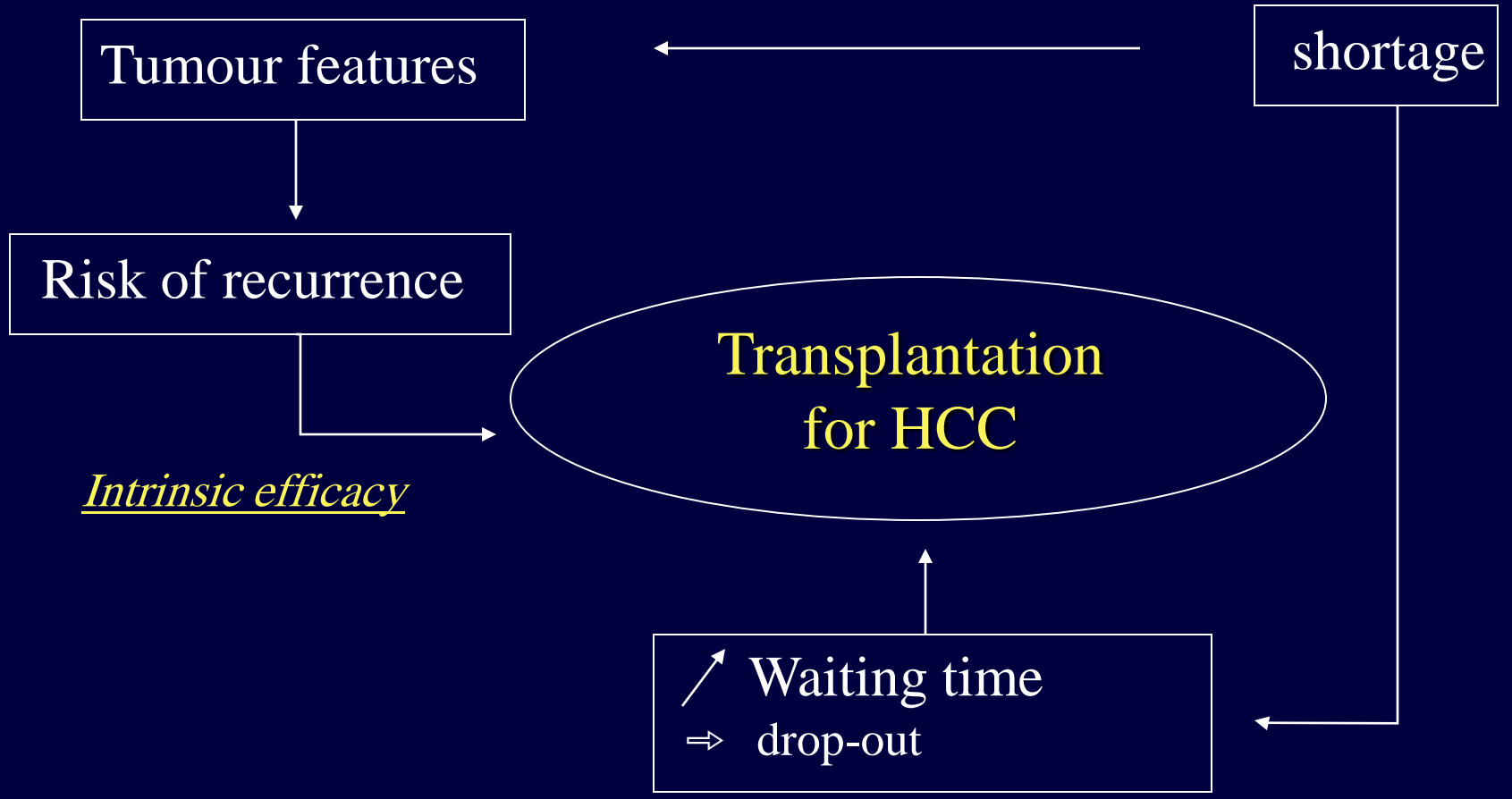


Milan vs “Up to 7”



N indications

N grafts



Intrinsic efficacy

Intent-to-transplant efficacy

Definitions

- Neoadjuvant treatment.
- Bridging
- Downstaging

- Downstaging

lowering the stage to allow for transplannation for patients
who when firstly seen don't qualify fro LTx

- Bridging

Strategy to allow patient to wait for a longer time without progression

TACE RFA

- Neoadjuvant treatment.

Treatment before a procedure to improve outcome

- TACE

-RFA

Excellent outcome following down-staging of HCC prior to liver transplantation: an intention-to-treat analysis

Criteria for downstaging

- 1 lesion > 5 cm and up to 8 cm
- 2–3 lesions with 1 or more lesions > 3 cm and not > 5 cm, with total tumor diameter up to 8 cm
- 4–5 lesions with none > 3 cm, with total tumor diameter up to 8 cm

Excellent outcome following down-staging of HCC prior to liver transplantation: an intention-to-treat analysis

Table 4. Down-Staging Treatments Received by the 61 Patients

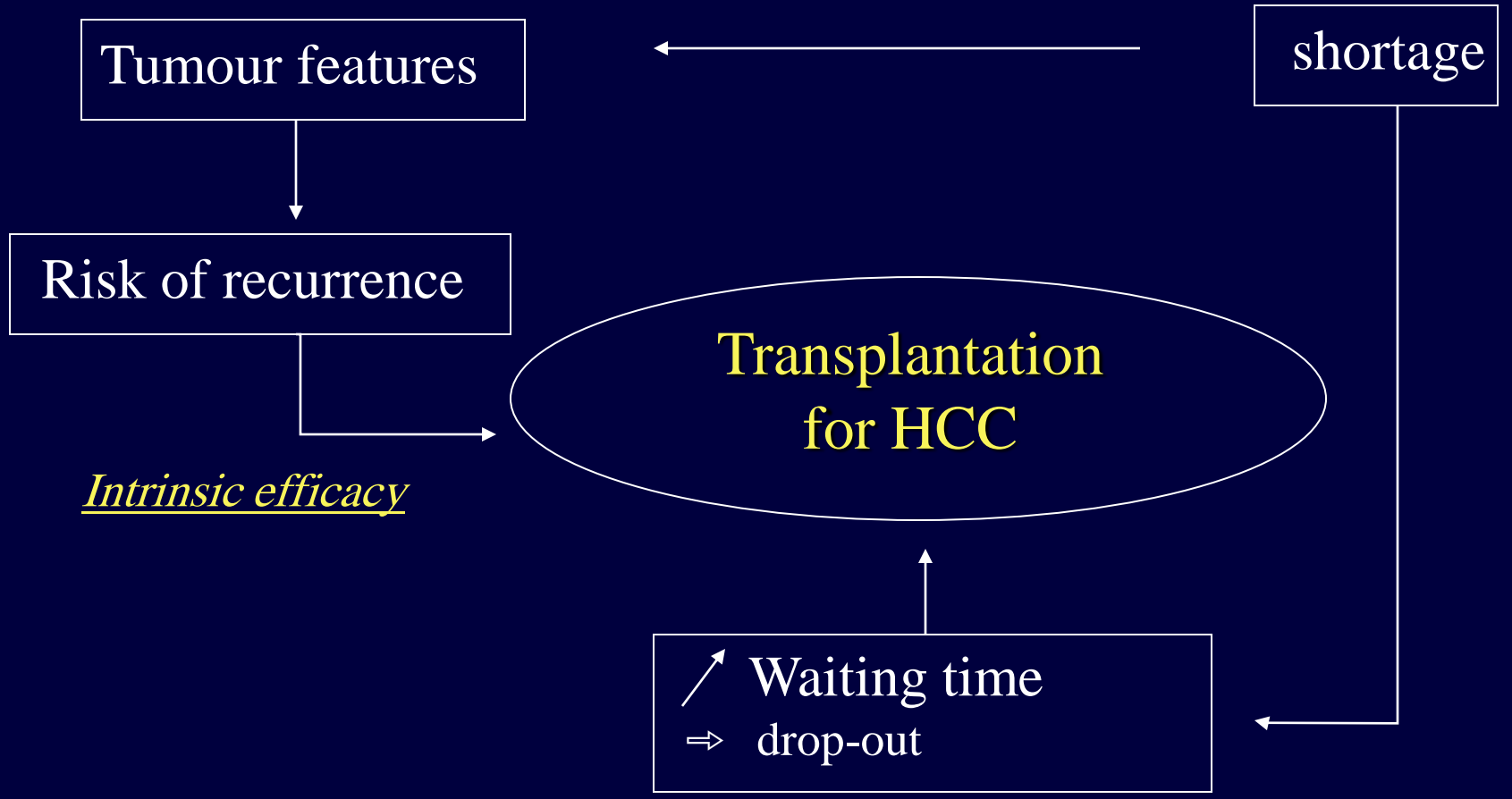
Treatment	No. of Patients (No. of Treatments)
Laparoscopic/open RFA only*	11 (11)
TACE only	15 (34)
TACE + percutaneous ablation	15 (54)
TACE + percutaneous ethanol ablation	6 (27)
TACE + percutaneous RFA	9 (27)
Laparoscopic RFA + TACE	14 (34)
Resection†	6 (6)

*Two received open RFA, nine received laparoscopic RFA.

†One of these patients underwent resection despite a high preoperative CTP score of 11. This patient had a 5.3-cm lesion very close to the liver surface at risk for rupture. The other five patients had a CTP score of ≤ 7 before resection.

N indications

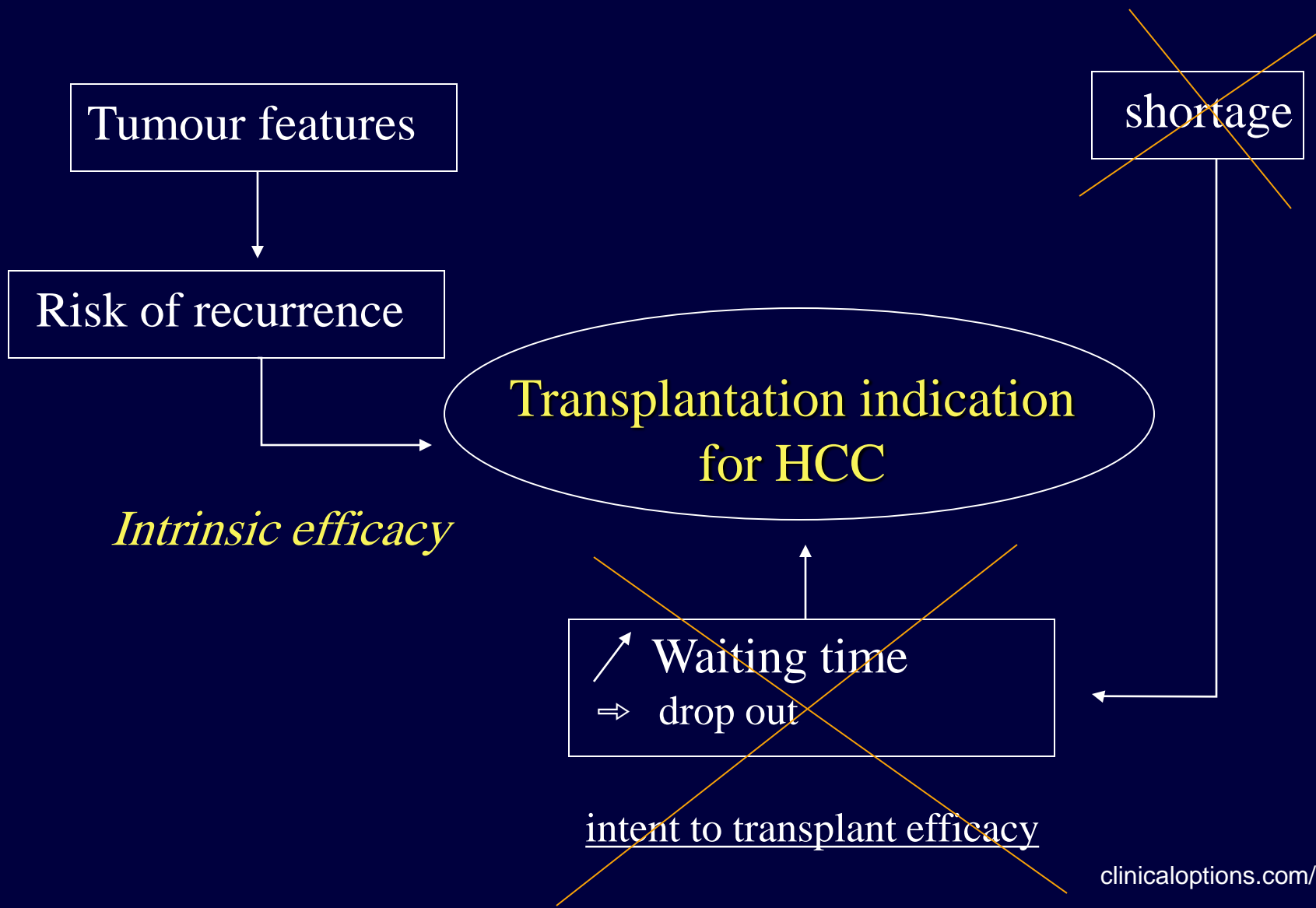
N grafts



Intrinsic efficacy

Intent-to-transplant efficacy

Living donor transplantation and HCC



Is LDLT for HCC as efficacious
as DDLT ?

Is LDLT for HCC as efficacious as DDLT ?

PRO

Thuluvath PJ, Yoo HY. Graft and patient survival after adult live donor liver transplantation compared to a matched cohort who received a deceased donor transplantation. *Liver Transpl* 2004;10:1263-8.

Lo CM, Fan ST, Liu CL, et al. The role and limitation of living donor liver transplantation for hepatocellular carcinoma. *Liver Transpl* 2004;10:440-7.

CON

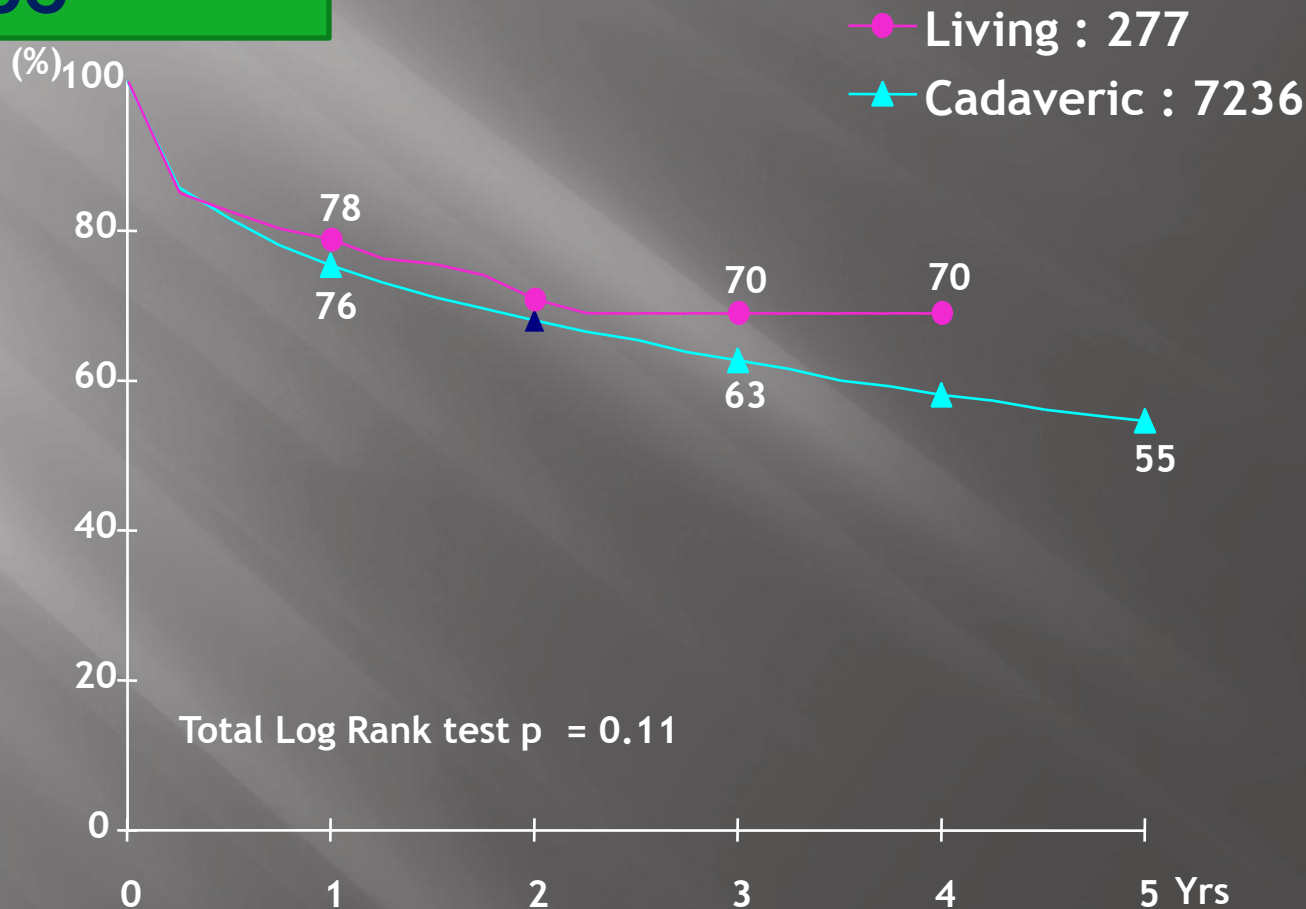
Fisher RA, Kulik LM, Freise CE, et al. Hepatocellular carcinoma recurrence and death following living and deceased donor liver transplantation. *Am J Transplant* 2007;7:1601-8.

Lo CM, Fan ST, Liu CL, et al. Living donor versus deceased donor liver transplantation for early irresectable hepatocellular carcinoma. *Br J Surg* 2007; 94:78-86.

Kulik L, Abecassis M. Living donor liver transplantation for hepatocellular carcinoma. *Gastroenterology* 2004;127(5 Suppl 1):S277-82.

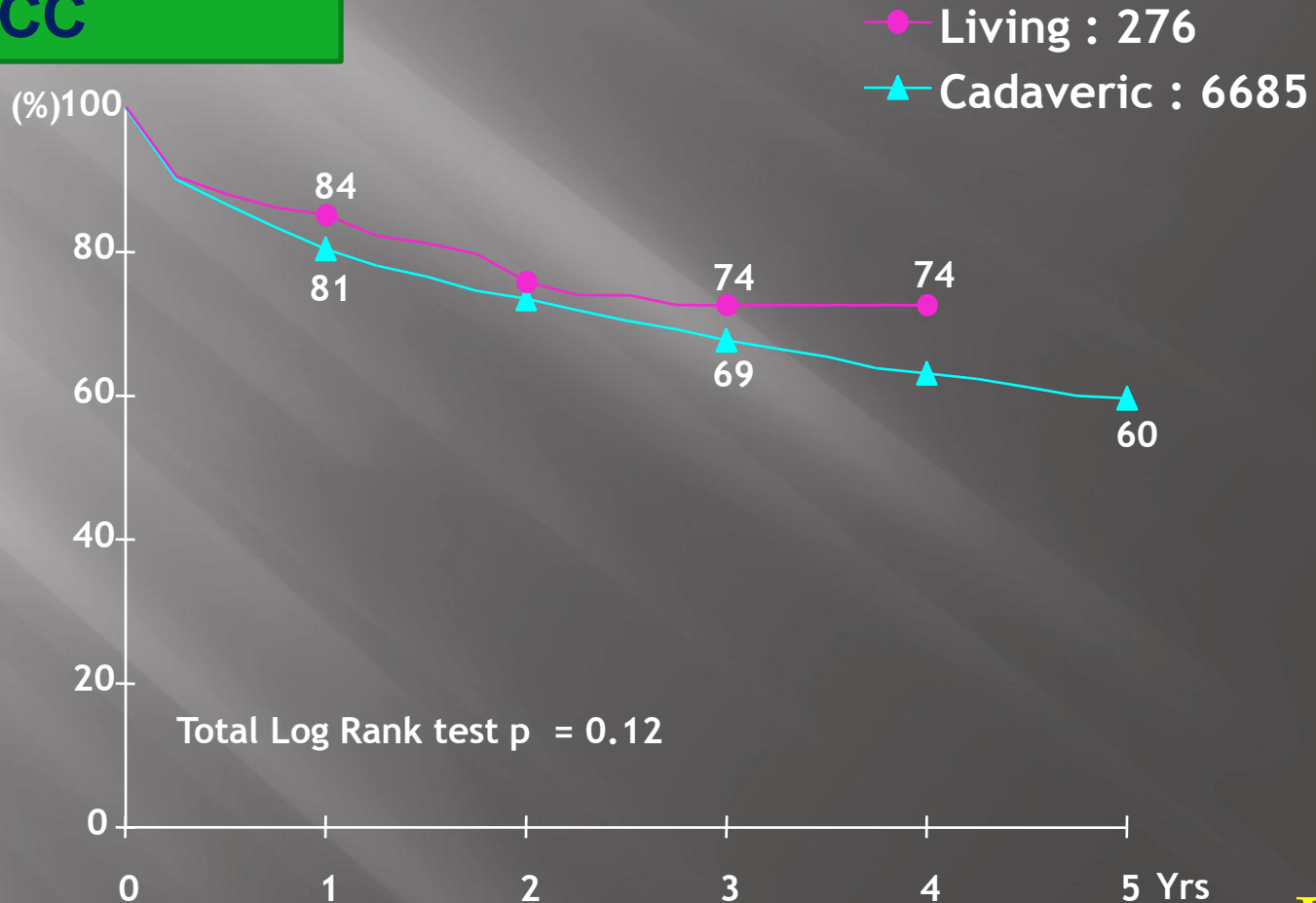
Is LDLT for HCC as efficacious as DDLT ?

Graftsurvival in HCC



Is LDLT for HCC as efficacious as DDLT ?

Patients survival in HCC



Predictors of Recurrence after LT

- 1.L.N involvement
- 2.gross vascular invasion (angio / CT)
- 3.microscopic invasion (in the specimen)
- 4.> 5cm
- 5.multiple lesions
- 6.infiltrating rather than circumscribed lesion
7. More than one lobe
- 8.pTNM staging

How to Minimize Risk of Recurrence ?

- **HCC biology**
- **Refinement of immunosuppression: “ mTOR” ?**
- **Radiologic identification of VI**
- **Prospective multicenter RCT: is the key.**

Conclusion

- HCC patients exceeding Milan criteria can still be cured ; nodules 5-7cm and with no gross vascular invasion >> good survival & higher recurrence rate.
- HCC is a prime indication for LDLT ...>.lower dropout ,extending the acceptance of HCC for LT without waiting for cadaveric LT.
- However is no consensus on the use of LDLT for HCC due to lack of adequate data

Thank
You

