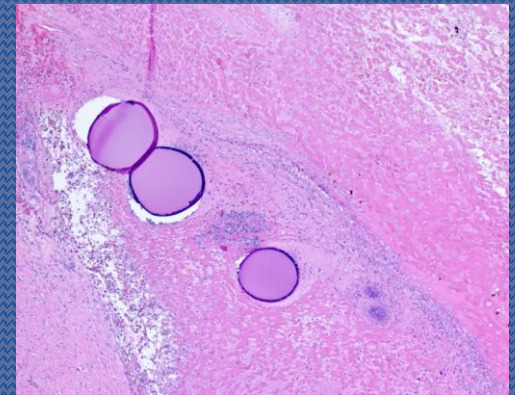
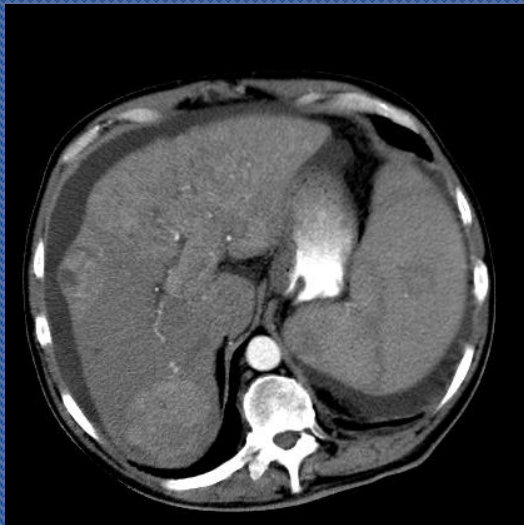
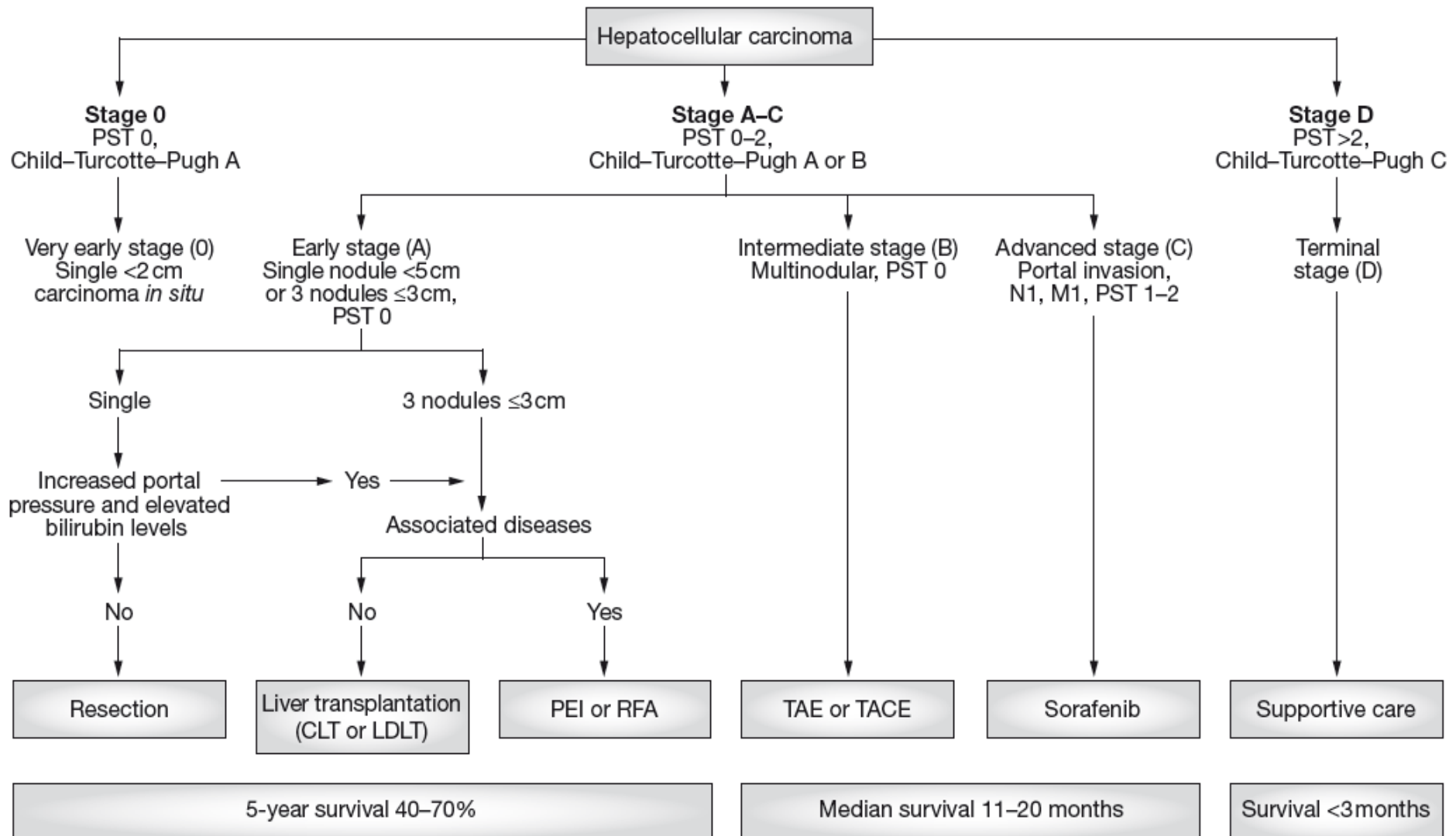


Resultater efter palliation for hepatocellulært carcinom

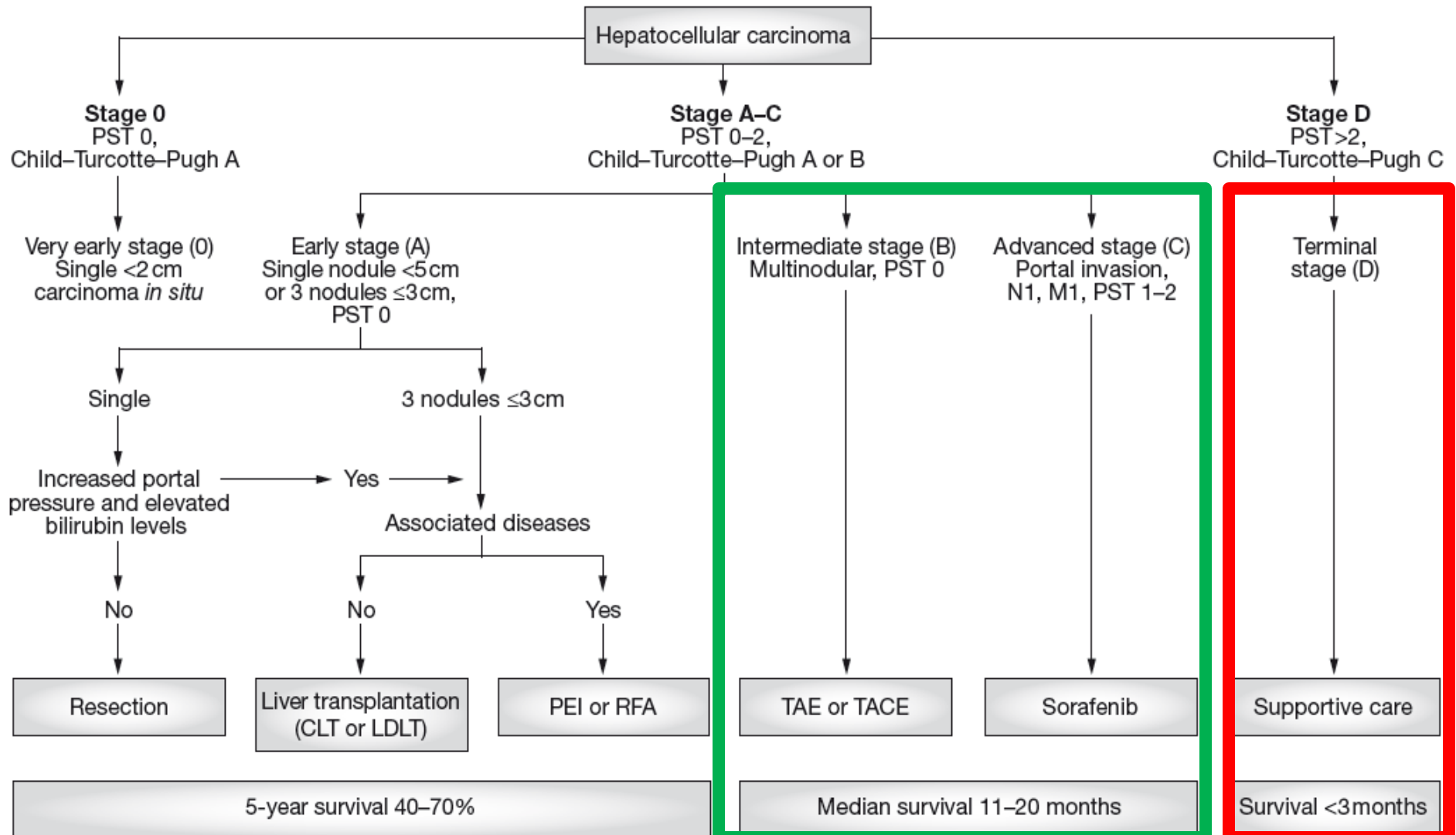
Henning Grønbæk, overlæge ph.d.
Medicinsk Afdeling V
Aarhus Universitetshospital



Multidisciplinær = MDT



Ikke palliation- livsforlængende



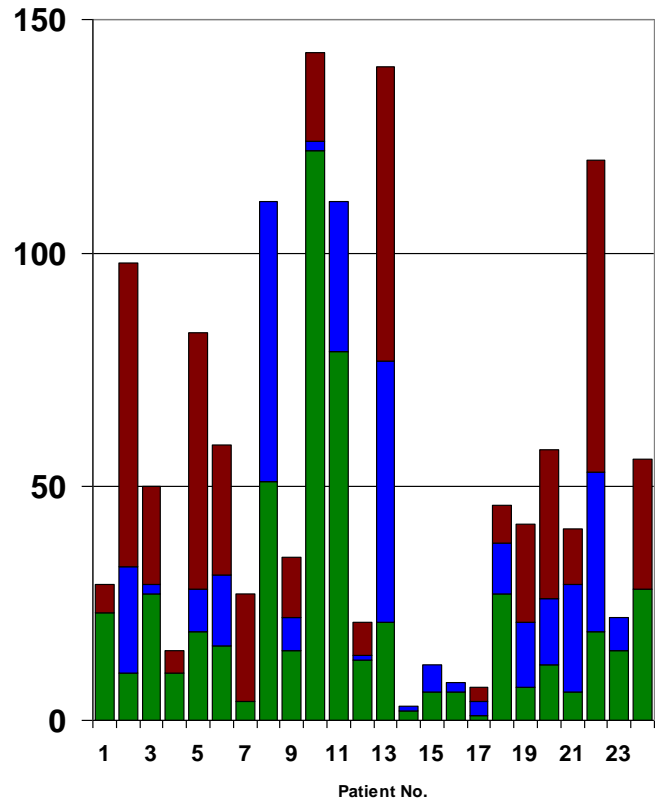
Aarhus MDT =

Multidisciplinære lever team

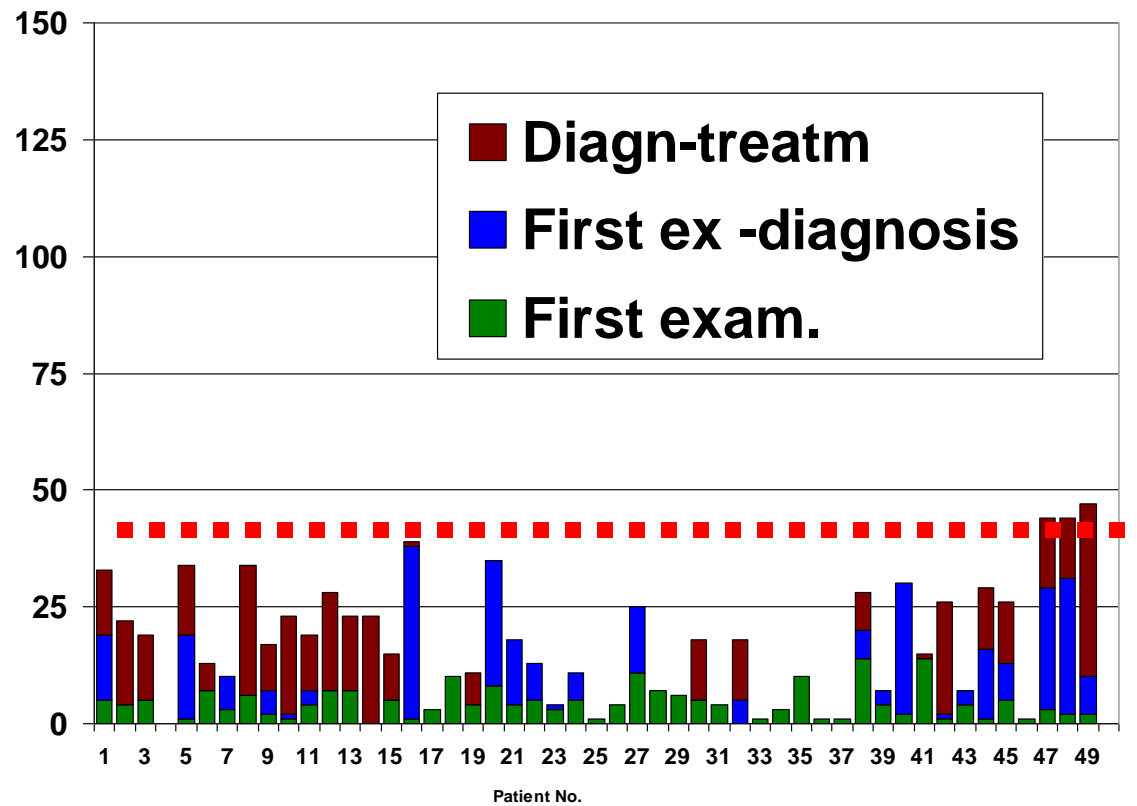
- Etableret 2002, 2 konferencer/uge
- **Hepatologer**
- Lever kirurger
- Radiologer (CT/MRI/US, biopsi, RFA, kemo-embolisering)
- PET Center (FDG^{Gal}-PET, FDG-PET)
- Onkologer (Sorafenib, stereotaktisk strålebehandling)
- Patologer

EFFEKT AF CANCER PAKKEFORLØB

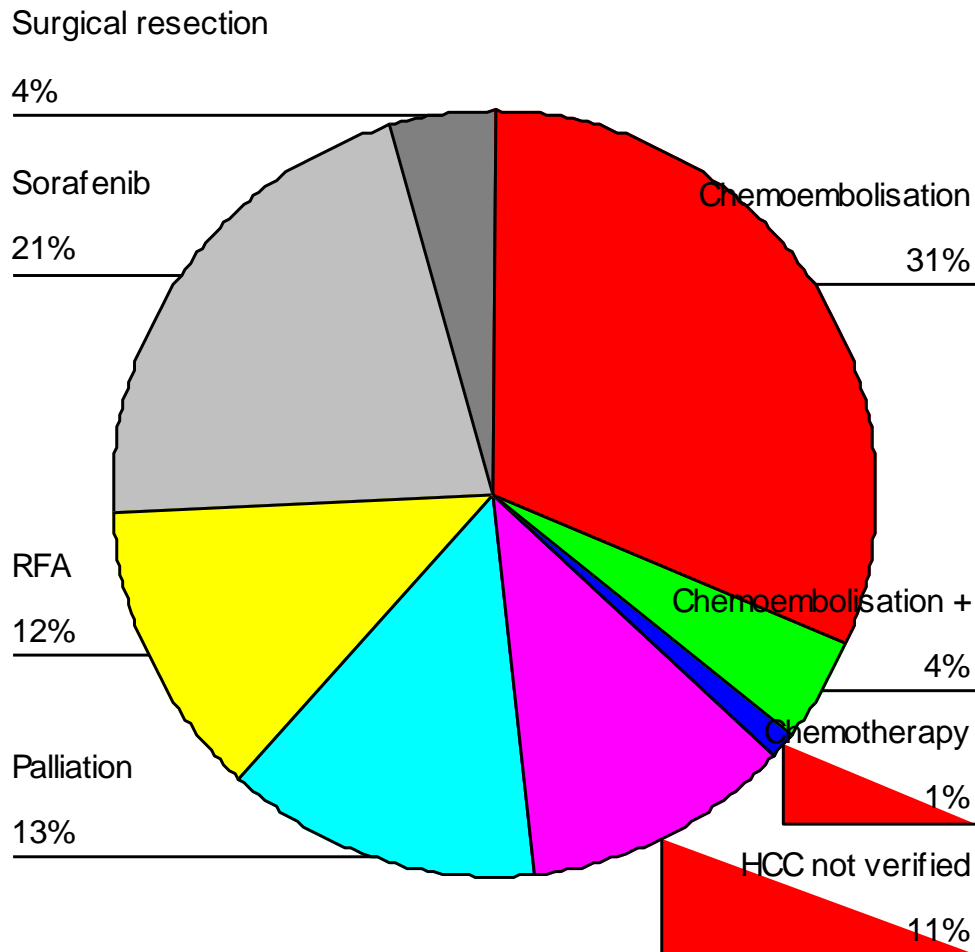
2006



2009



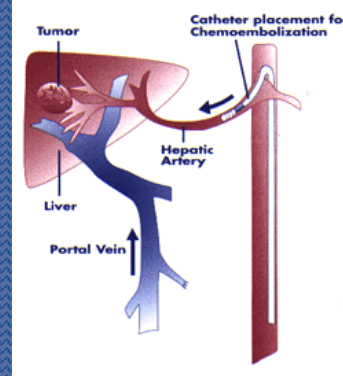
89 patienter 1. MDT beslutning



Median tid MDT - behandling

- RFA 19 (6-45) days
- Kemoembolisering 20 (7-90)
- Kirurgi 56 (20-60)
- Sorafenib 2-3 weeks

TACE og HCC



- Positiv effekt på overlevelse

Patient

PS, CP-A

Alfa-foetoprotein

Leverfunktion

Mænd

Høj albumin

Sygdomspræsentation Behandling

TNM

Unifokal

Hypervaskulær

Ingen PVT

Selektiv TACE

> 1 TACE

I dag: DC Beads Tidligere: Lipiodol + Doxyrubicin

TACE kontraindikationer

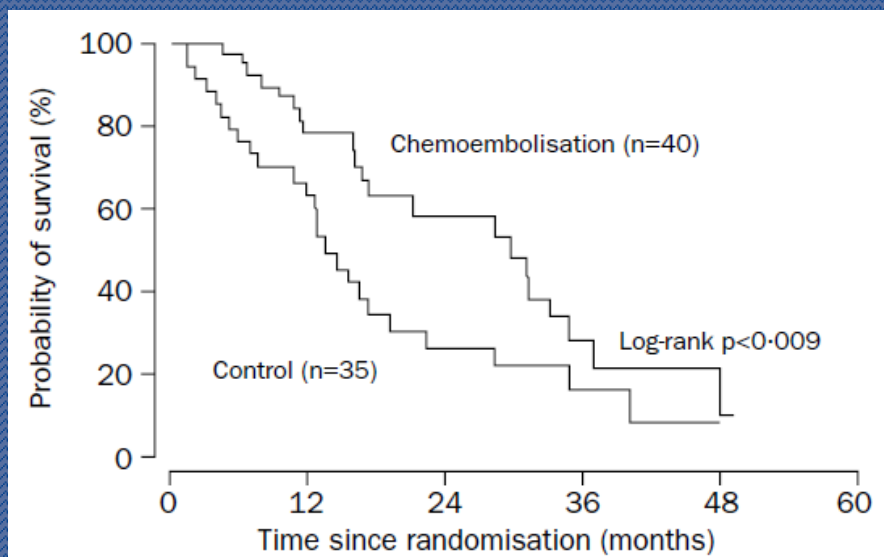
Absolutte

- CP > 8
- Ikterus,
- HE
- Refraktær ascites, HRS
- Bilobær ekstensiv tumor
- PVT

Relative

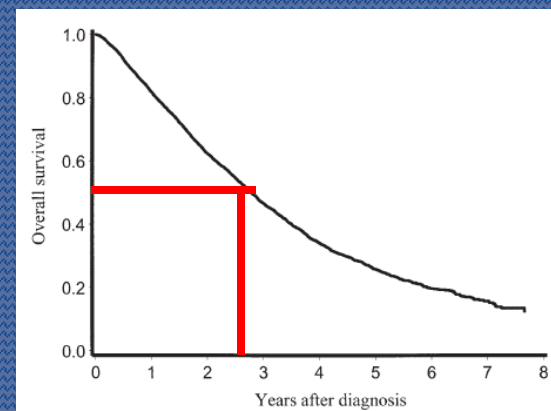
- Tumor > 10 cm
- Komorbiditet
 - CVD
 - COLD
- Grad 3 varicer
- Galdegangsookklusion

TACE - resultater

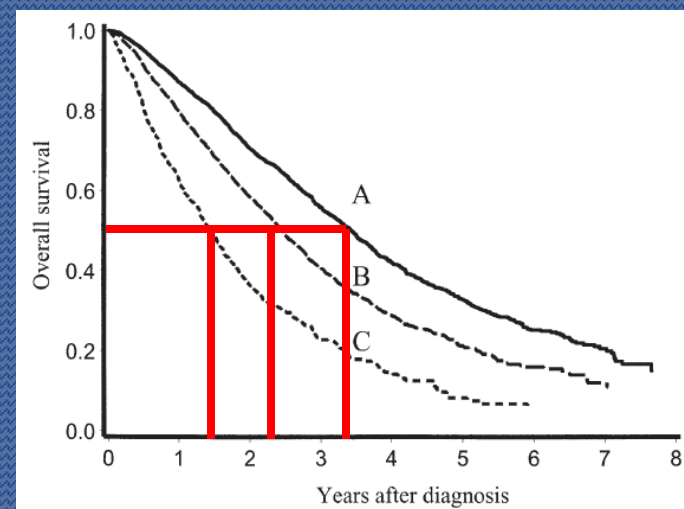


Llovet et al. Lancet 2002; 359: 1734–39

- 44/8510 døde pga. TACE
- 18 leversvigt,
 - 8 cancer relateret, 7 intraperitoneal HCC ruptur,
 - 5 varice blødning, 6 andre årsager
- Relateret til CP score: A: 7 patienter, B: 14, C 17, ukendt 6



Median overlevelse 34 mdr.
1-, 3-, 5-, års overlevelse 82, 47, 26%



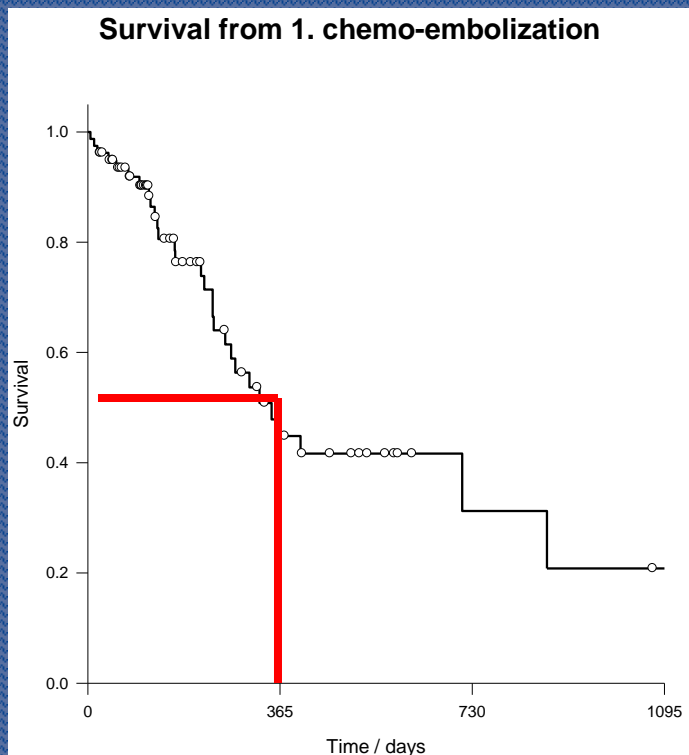
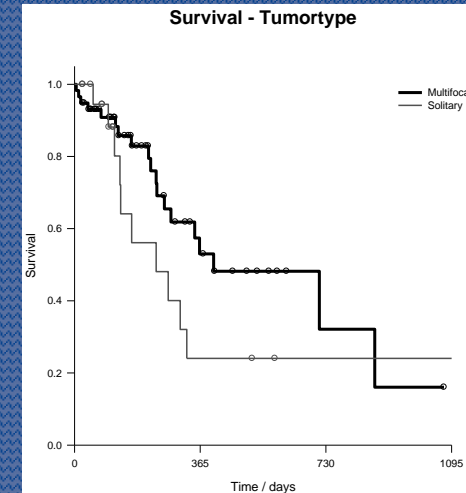
Takayasu et al. GASTROENTEROLOGY 2006;131:461–469

TACE overlevelse - AARHUS

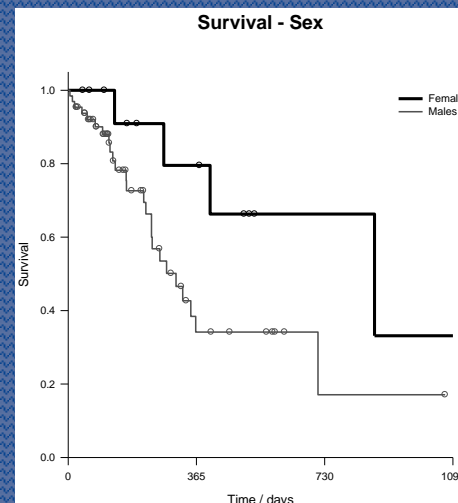
80 patienter,
66 Mænd 14 kvinder
Median alder 67 (25 - 85)
Solitær 21, multifokal 59

Cirrose				Non Cirrose
Hep B.	Hep C.	Alkohol	Anden årsag	
2	11	28	9	35

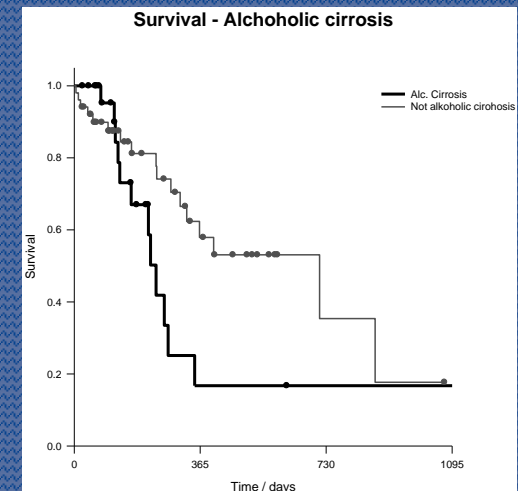
TUMOR TYPE



KØN



CIRROSE



TACE komplikationer

- Infarkt
- Abscess
- Blødning
- Lever svigt

- Feber, smerter, CRP↑ er forventet

- Oftest infarkt feber og ikke infektion – de fleste sendes hjem dag 2-3
- Ingen antibiotika.

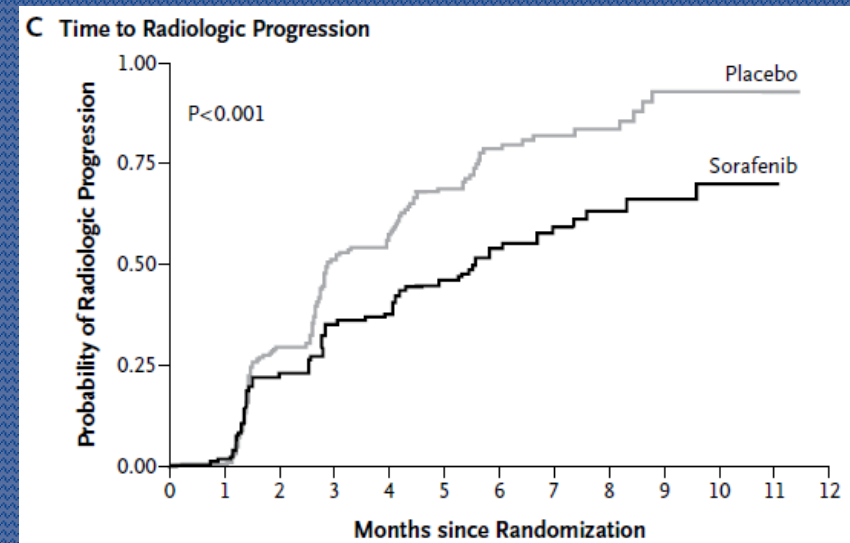
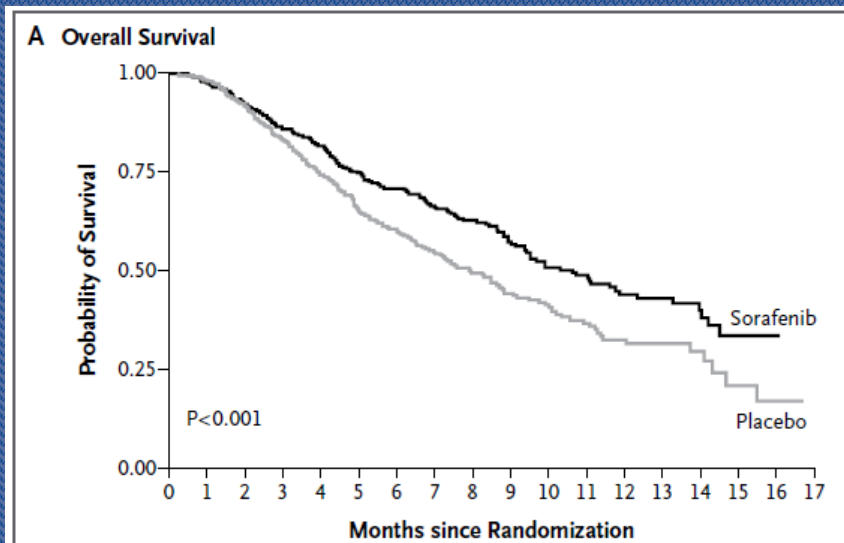
Sorafenib og HCC



HCC er en hypervaskulær tumor

RAF/MEK/ERF og VEGF af betydning for HCC tumorgenese og angiogenese

Sorafenib er en multikanase hæmmer af bl.a. RAF, VEGF og andre kinaser



Median overall survival 10.7 vs. 7.9, HR: 0.69; 95%CI: 0.55- 0.87; P < 0.001).

Ingen forskel for symptomatisk progression (4.1 months vs. 4.9 months, P = 0.77).

Median time to radiologic progression 5.5 months vs. 2.8 months (P < 0.001).

Randomized Controlled Trials of Sorafenib in Advanced Hepatocellular Carcinoma

Study characteristics	SHARP Study ¹	Asia Study ²
Median age	65 yrs	51 yrs
BCLC-B stage	18%	4%
Previous treatments	67%	na
HBV etiology of cirrhosis	19%	71%
<u>TTP (control)</u>	5.5 mo. (2.8 mo.)	2.8 mo. (1.4 mo.)
<u>Median survival (control)</u>	10.7 mo. (7.9 mo.)	6.5 mo. (4.2 mo.)
Grade 3/4 toxicity	30%	24%

¹ Llovet JM et al *NEJM* 2008;359:378-390; ² Cheng A et al *Lancet Oncol* 2009;10:25-34

Sorafenib behandling Aarhus Universitetshospital

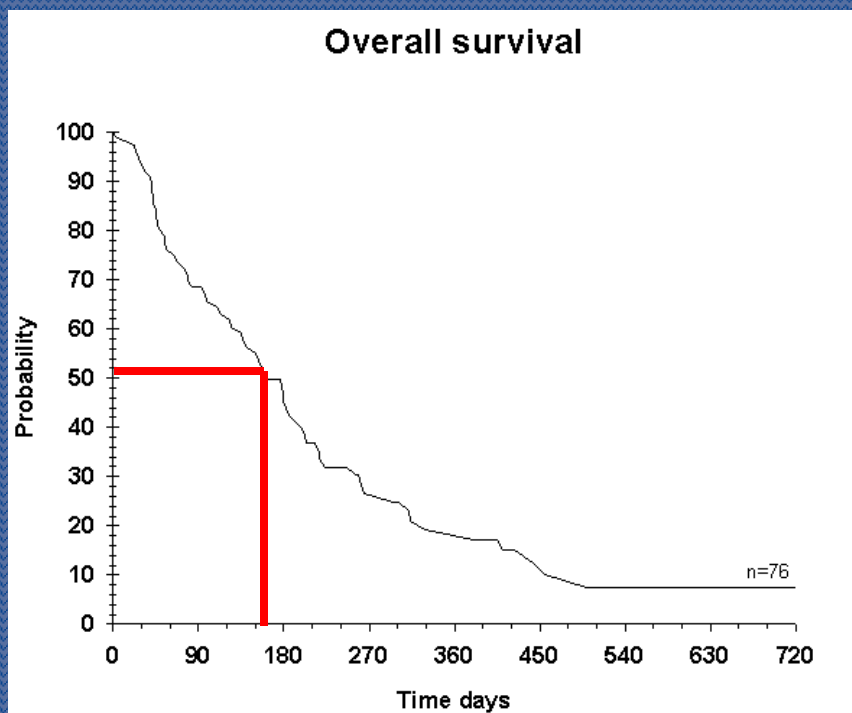
76 patienter

59 mænd, 17 kvinder

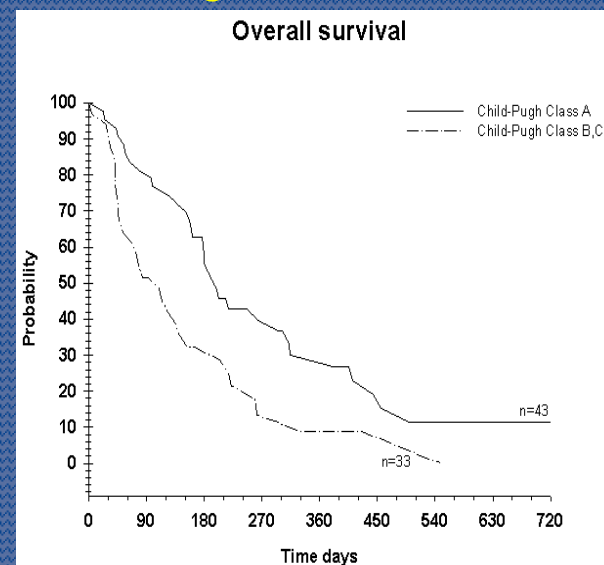
Median alder 63 (25-85)

CP-A: 43, CP-B: 29, CP-C: 4

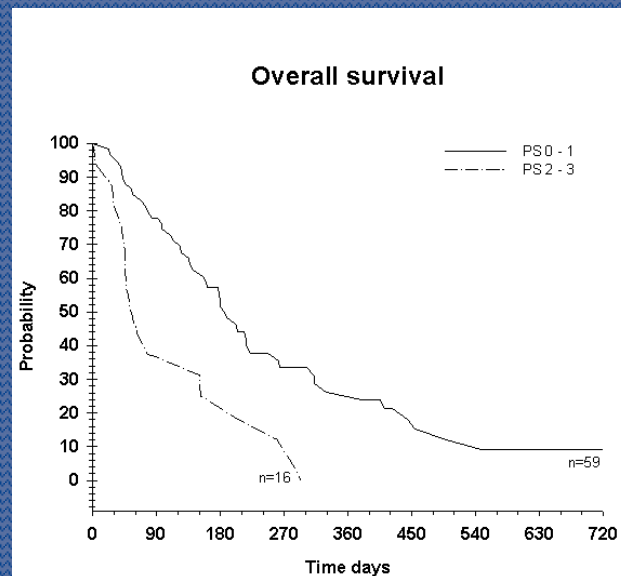
Alle patienter: Child-Pugh A-C, PS 0-3)



Child-Pugh score



WHO Performance status



Sorafenib toxicitet



Bivirkninger	Alle grader	Grad 3-4
Træthed	68%	12%
Anorexi	47%	3%
Diarre	42%	10%
Hududslæt (Rash)	33%	4%
Kvalme	32%	3%
”Hand-foot” syndrom	26%	12%
Hypertension	18%	3%
Opkast	16%	3%
Trombocytopeni	5%	3%
Knoglemarvs suppression	4%	--

Behandling af HCC i fremtiden

- MDT – centraliseret, cirrose patienter, hepatologer, selektion af patienter
- Screening – mangler dokumentation i DK
- Incidens – uændret? NASH?
- Levertransplantation – donor antal?
- Kombinationsbehandlinger (RFA + TACE + medicin)
- Targeteret behandling (sunitinib? Everolimus? Bevacizumab?)

