



grupo brasileiro
de tumores
gastrointestinais

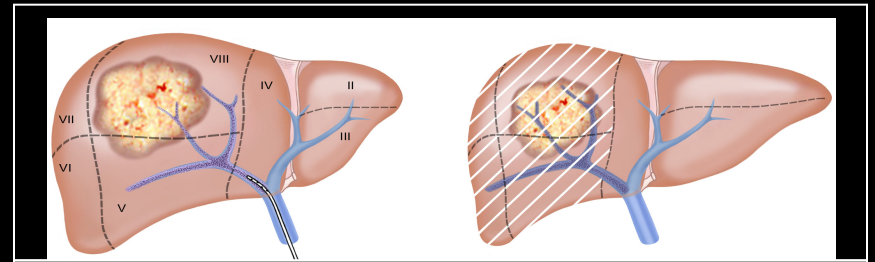
Associating Liver Partition and Portal Vein Ligation for Staged hepatectomy

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Indução de Hipertrofia

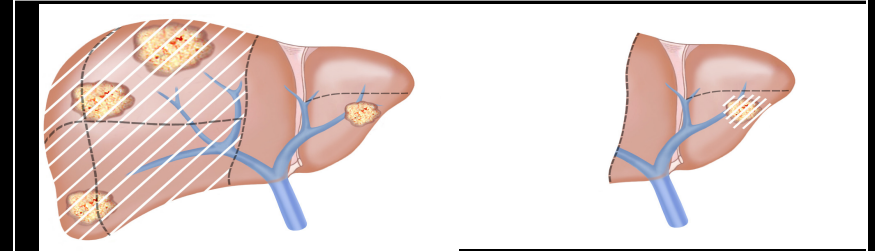
Embolização Portal (EP)

Makuuchi M, et al. Surgery 1990



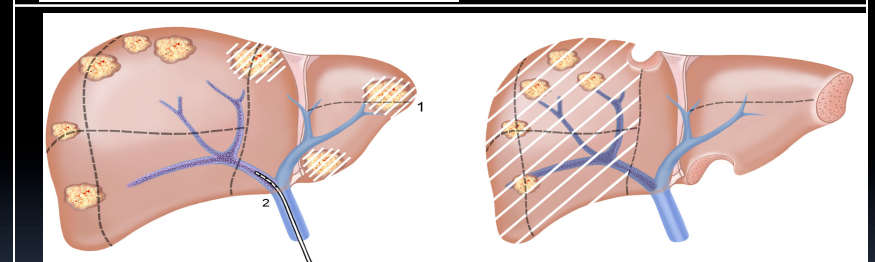
Hepatectomia em 2 tempos

Adam R, et al. Ann Surg. 2000



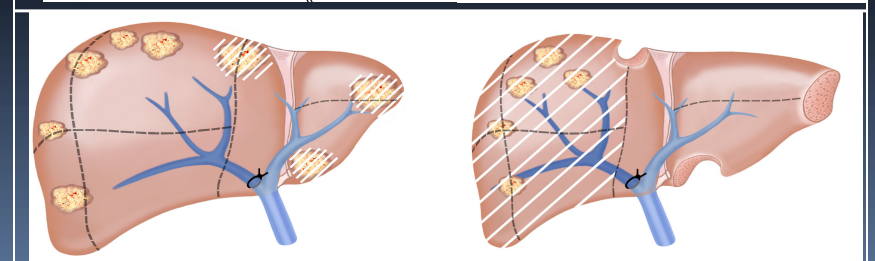
Hepatectomia 2 tempos + EP

Jaeck D, et al. Ann Surg. 2004



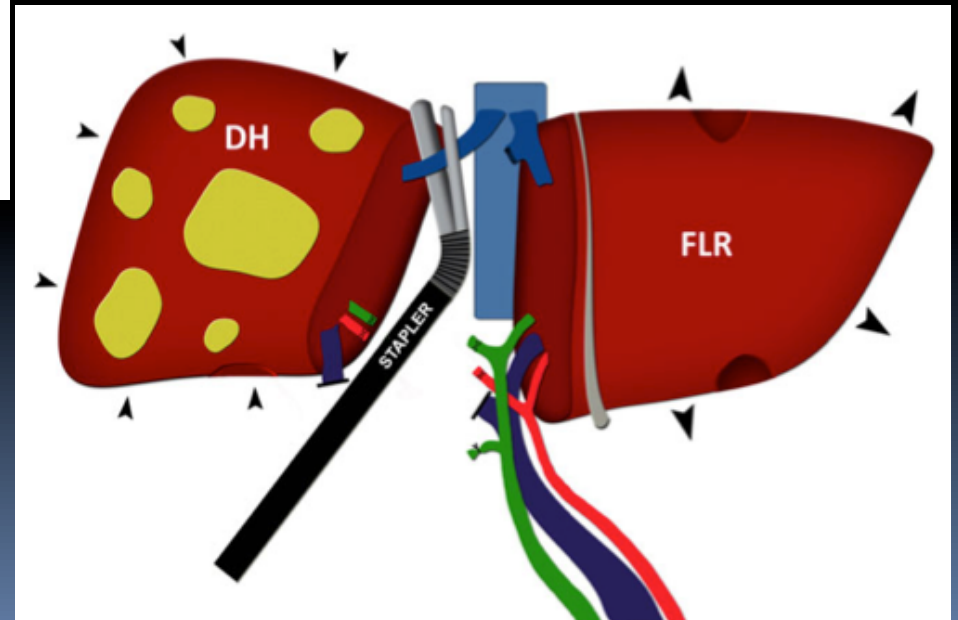
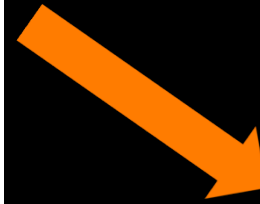
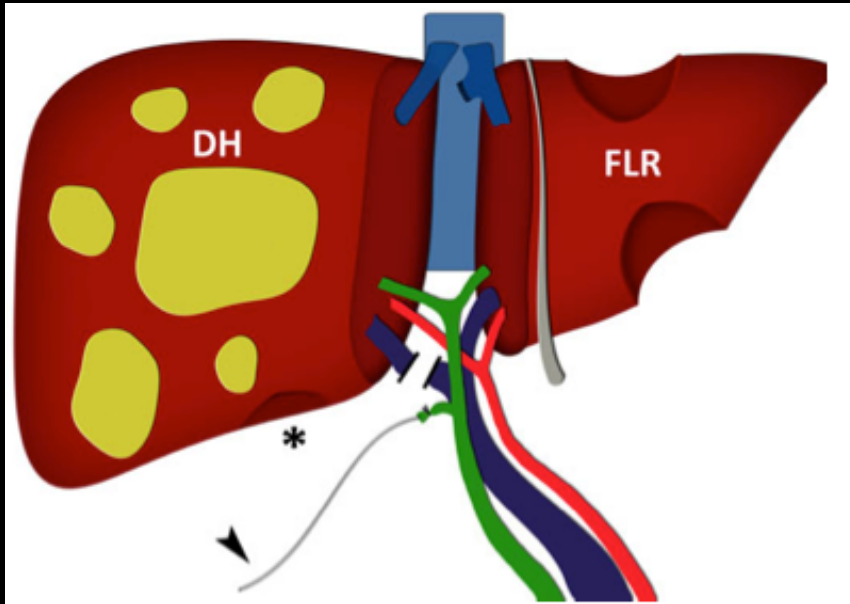
Hepatectomia 2 tempos + LP

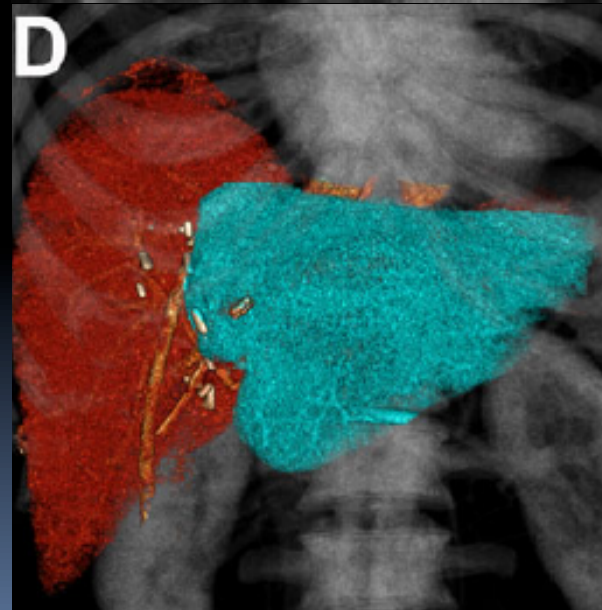
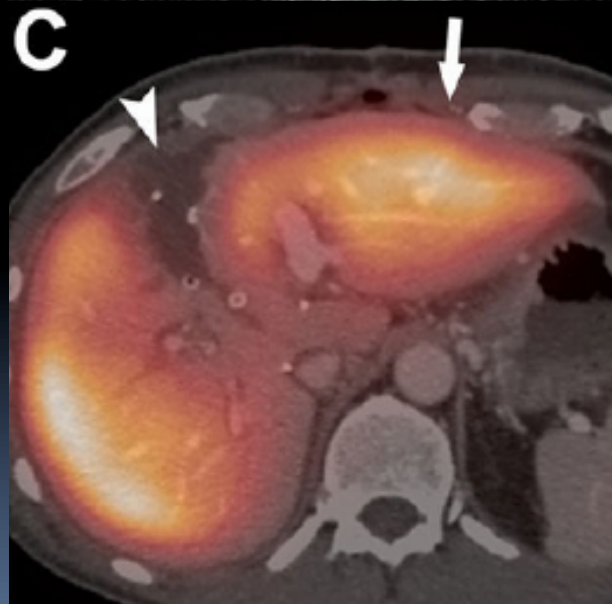
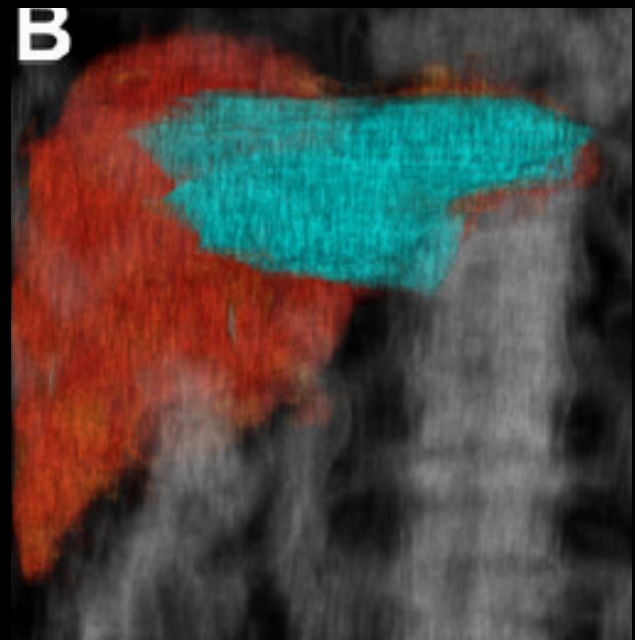
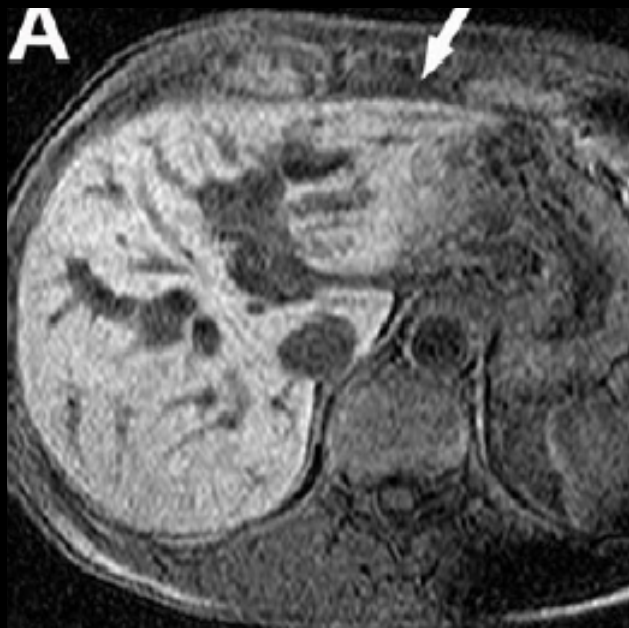
J Belghiti, Clavien AP, et al. Hepatology 2008



Aspectos técnicos

ALPPS





ALPPS

major advances to surgically induce fast liver hypertrophy. The rapid worldwide adoption of ALPPS, after being described in Germany, has resulted in preliminary single center and cooperative experiences showing a high morbidity and mortality of this emerging method. In the large multicenter experiences reported, the Brazilian¹⁶ with 39 patients and the German⁹ with 25 patients, a morbidity of 59% to 68% and a mortality of 12% to 12.8% were reported. Proponents of PVE argue that the ALPPS has excessively high morbidity and mortality rates. However, the recently reported data from 87 patients who underwent a major hepatectomy after chemotherapy and PVE or PVL at the Beaujon Hospital in France for initially unresectable

Cortesía Dr. Eduardo de Santibanes (Buenos Aires - Argentina)

Factibilidade (Completar resecção R0)

ALPPS vs Cirurgia em dois tempos

| Author | Year | Nº patients | Completed 2nd Stage | Resectability |
|--|------|-------------|---------------------|---------------|
| Lamb <i>et al</i> (Systematic review) ¹ | 2013 | 459 | 352 patients | 76% |
| Abbot <i>et al</i> (MD Anderson) ² | 2013 | 82 | 56 patients | 68% |
| Cardona <i>et al</i> (MSKCC) ³ | 2013 | 40 | 35 patients | 88% |
| Tsai <i>et al</i> (J Hopkins) ⁴ | 2010 | 45 | 35 patients | 78% |
| Belghiti J (Beaujon) ⁵ | 2008 | 35 | 26 patients | 74% |
| Adam <i>et al</i> (Paul Brousse) ⁶ | 2008 | 59 | 41 patients | 69% |
| Torres <i>et al</i> (Brazil) ⁷ | 2013 | 39 | 37 patients | 94,9% |
| ALPPS Registry ⁸ | 2013 | 202 | 197 patients | 98% |

¹ Lamb *et al*. HPB 2013

³ Cardona *et al*. Ann Surg Oncol. 2013

⁵ Belghiti J *et al*. J Gastrointest Surg. 2008

⁷ Torres *et al*. ABCD 2013

² Abbot *et al*. J Surg Oncol. 2013

⁴ Tsai *et al*. HPB 2010

⁶ Adam *et al*. Ann Surg. 2008

⁸ Schdde *et al*. Annals of surgery 2014

SEGURANÇA

| Author | Year | N° patients | Morbidity (major) | Liver Failure | Mortality |
|--|-------------|-------------|-------------------|---------------|-----------|
| Lamb <i>et al</i> (Systematic review) ¹ | 2013 | 459 | 40% | - | 3% |
| Cardona <i>et al</i> (MSKCC) ² | 2013 | 40 | 45% | - | 0% |
| Brouquet <i>et al</i> (MD Anderson) ³ | 2011 | 65 | 29% | 5% | 6.4% |
| Tsai <i>et al</i> (J Hopkins) ⁴ | 2010 | 45 | 28% | 8.6% | 8.8% |
| Belghiti J (Beaujon) ⁵ | 2008 | 35 | 46% | 15% | 0% |
| Adam <i>et al</i> (Paul Brousse) ⁶ | 2008 | 59 | 59% | 22% | 7% |
| ALPPS Registry (CRLM)^{7*} | 2013 | 202 | 38% | 6% | 8% |

¹ Lamb *et al*. HPB 2013

³ Brouquet *et al*. J Clin Oncol 2011

⁵ Belghiti J *et al*. J Gastrointest Surg. 2008

⁷ www.alpps.net

² Cardona *et al*. Ann Surg Oncol. 2013

⁴ Tsai *et al*. HPB 2010

⁶ Adam *et al*. Ann Surg. 2008

Resultados oncológicos a curto prazo

| Author | Year | N° patients | OS | | DFS | |
|--|------|-------------|------|-----|-----|-----|
| | | | 1 yr | 2yr | 1yr | 2yr |
| Lamb <i>et al</i> (Systematic review) ¹ | 2013 | 459 | - | - | - | - |
| Tsai <i>et al</i> (J Hopkins) ² | 2010 | 45 | 88% | 74% | 85% | 68% |
| Brouquet <i>et al</i> (MD Anderson) ³ | 2011 | 65 | 89% | 80% | 39% | - |
| Adam <i>et al</i> (Paul Brousse) ⁴ | 2008 | 41 | 87% | 76% | 60% | 40% |
| ALPPS Registry (CRLM) ⁵ | 2013 | 141 | 88% | 74% | 59% | 41% |

¹ Lamb *et al* HPB. 2013

² Tsai *et al*. HPB 2010

³ Brouquet *et al*. J Clin Oncol 2011

⁴ Adam *et al*. Ann Surg. 2008

⁵ Schadde *et al*. Annals of Surgery 2014

O procedimento ALPPS é factível, seguro e com resultados oncológicos semelhantes aos procedimentos semelhantes com a mesma finalidade.

ORIGINAL ARTICLE

Associating Liver Partition and Portal Vein Ligation for Staged Hepatectomy Offers High Oncological Feasibility With Adequate Patient Safety

A Prospective Study at a Single Center

*Fernando A. Alvarez, MD, Victoria Ardiles, MD, Martin de Santibañes, MD, Juan Pekolj, MD, PhD,
and Eduardo de Santibañes, MD, PhD*

Cortesía Dr. Eduardo de Santibanes (Buenos Aires – Argentina)

TABLE 1. Baseline characteristics of the study population

| Variable | Patients (n= 30) |
|--|-------------------------|
| Age/ yrs, median (range) | 58.6 (35-81) |
| Sex/ male, n° (%) | 19 (63) |
| BMI (kg/m ²), median (range) | 25.2 (17.7-31.7) |
| ASA operative risk, n° (%) | |
| < 3 | 22 (73) |
| ≥ 3 | 8 (27) |
| Charlson Index, median (range) | 7 (2-10) |
| Tumor type, n° (%) | |
| Colorectal liver metastases | 19 (63.3) |
| Neuroendocrine metastases | 3 (10) |
| Hepatocellular carcinoma | 3 (10) |
| Hilar cholangiocarcinoma | 1 (3.3) |
| Lobar cholangiocarcinoma | 1 (3.3) |
| Other* | 3 (10) |
| Bilateral disease, n° (%) | 19 (63) |
| Colorectal liver metastases | 15 |
| Neuroendocrine metastases | 3 |
| NCRNNE | 1 |
| Chemotherapy, n° (%) | 18 (60) |
| ≥ 6 cycles | 13 |
| ≥ 2 lines | 6 |
| Targeted therapy | 12 |
| Diseased parenchyma, n° (%) | 14 (46.6) |
| Chemotherapy-induced | 11 |
| Non chemotherapy-induced | 3 |
| Previous abdominal intervention, n° (%) | 19 (63) |
| Synchronous primary tumor, n° (%) | 9 (30) |
| % FLR/TLV, mean (s.d.) | 22.1 (6.1) |
| % FLR/BW (cc/kg), mean (s.d.) | 0.47 (0.14) |

ASA, American Society of Anesthesiologists; BMI, body mass index; NCRNNE, non-colorectal non-neuroendocrine metastases; FLR, future liver remnant; TLV, total liver volume; BW, body weight.

* Metastatic breast cancer (n= 1), metastatic leiomyosarcoma (n= 1), metastatic esophageal cancer (n= 1).

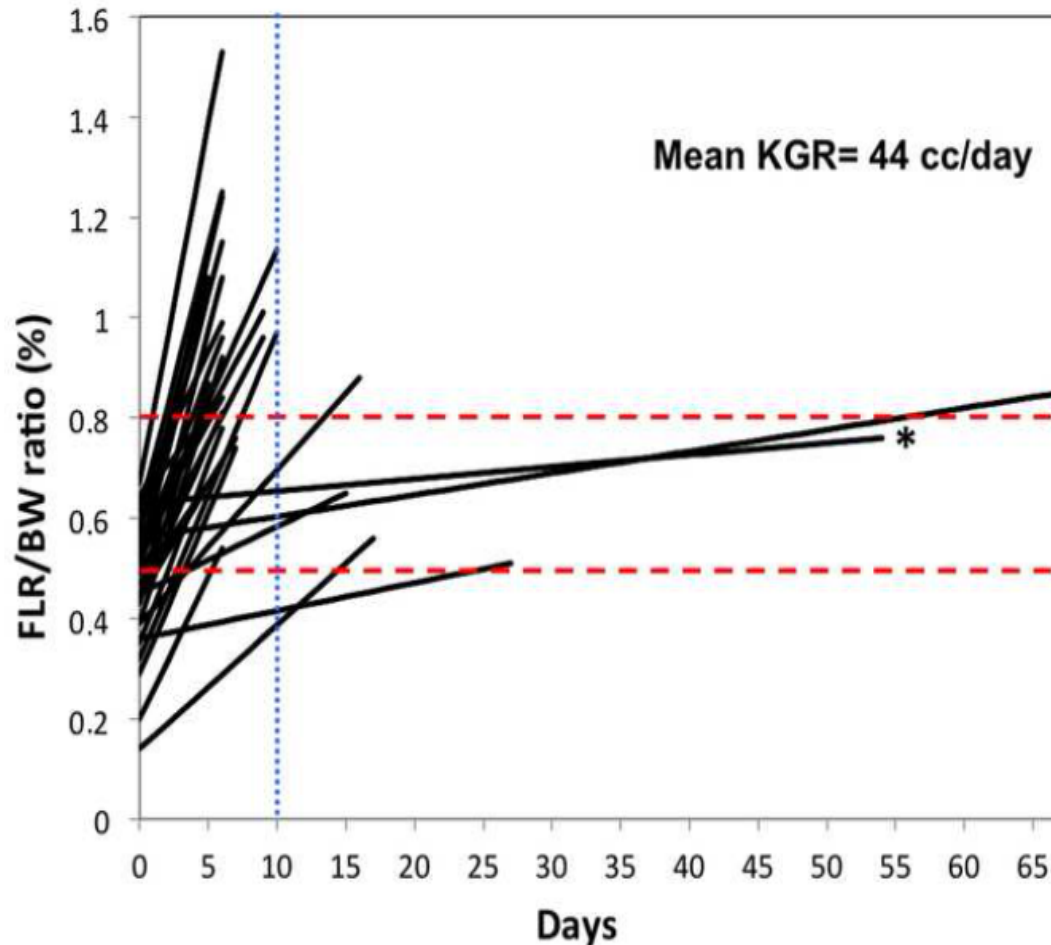
TABLE 2. Intraoperative data in 30 patients treated.

| Characteristic | 1 st Stage | 2 nd Stage |
|---------------------------------------|-----------------------|-----------------------|
| Type of major liver resection, n° (%) | | |
| Right hepatectomy | - | 8 (27) |
| Right trisectionectomy | - | 21 (70) |
| Left trisectionectomy | - | 1 (3) |
| FLR clean-up, n° (%) | 19 (63) | 0 |
| Lesions resected, median (range) | 2 (1-20) | |
| Extrahepatic procedures, n° (%) | 11 (37) | 0 |
| Colorectal resection | 4 | - |
| Colorectal resection + RYH | 1 | - |
| RYH | 1 | - |
| Small bowel resection | 1 | - |
| Distal spleno-pancreatectomy | 1 | - |
| Distal spleno-pancreatectomy + LRN | 1 | - |
| Colostomy | 1 | - |
| Diaphragmatic resection | 1 | - |
| Operative time (min), median (range) | 315 (160-480) | 120 (60-300) |
| Intermittent Pringle maneuver, n° (%) | 9 (30) | 0 |
| Duration (min), median (range) | 29 (7-50) | - |
| Plastic bag used, n° (%) | 15 (50) | |
| RBC transfusion *, n° (%) | 8 (26.6) | 13 (43.3) |
| Packs per patient, median (range) | 2 (1-4) | 2 (1-4) |
| Portal pressure (mmHg), n° (%) | 9 (30) | 3 (10) |
| Baseline, median (range) | 12 (7-18) | 15 (13-17) |
| After PV ligation, median (range) | 15 (12-21) | - |
| After RHA clamping, median (range) | 18 (12-23) | 15 (13-17) |
| R0 resection †, n° (%) | 19 (100) | 27 (93.1) |

FLR, future liver remnant, RHY, roux-en-Y hepaticojejunostomy; LRN, left radical nephrectomy; RBC, red blood cells; PV, portal; vein; RHA, Right hepatic artery.

* Three patients received blood transfusion during both stages.

† 1st stage= % in patients with FLR clean-up. 2nd stage= % in patients who completed the strategy.



24 pacientes alcançaram o volume hepático necessário em 10 dias

A media de hipertrofia foi de 89% em 6 dias

FIGURE 2. Volumetric increase in terms of FLR to BW ratio in all patients treated. As observed, only 6 patients did not reach a sufficient hypertrophy within 10 days of the interval period (blue dashed line). The red dashed lines indicate the critical FLR/BW ratios for safe liver surgery of 0.5% and 0.8% respectively. *In this patient the FLR/total liver volume ratio was 35% and the sectorial FLR function by scintigraphy was 41% of the overall liver function before reoperation.

TABLE 4. Postoperative morbidity according to severity grade*

| Stage | Grade ²⁴ | Number | Event (n) |
|-----------------|---------------------|--------|--------------------------------|
| 1 st | I | 2 | Acute renal insufficiency (2) |
| | II | 3 | Pancreatic fistula (2) |
| | | | Catheter-related infection (1) |
| | IIIa | 2 | Intra-abdominal collection (2) |
| | IIIb | 4 | Abdominal pain (2) |
| | | | Reduced portal flow (1) |
| | | | Portal vein thrombosis (1) |
| | IVa | 1 | Acute renal insufficiency (1) |
| | IVb | 1 | Intra-abdominal hemorrhage (1) |
| | V | 1 | Multi-organ failure (1) |
| 2 nd | I | 5 | PHLF/ Grade A (3) |
| | | | Acute renal insufficiency (2) |
| | II | 1 | Intra-abdominal hemorrhage (1) |
| | IIIa | 3 | Pleural effusion (3) |
| | IIIb | 4 | Biliary fistula (4) |
| | IVa | 1 | PHLF/ Grade B (1) |
| | IVb | 1 | Pneumonia (1) |
| | V | 1 | Multi-organ failure (1) |

PHLF, posthepatectomy liver failure.

* A total of 30 events were recorded in 16 patients, 10 of which occurred in 2 patients who died due to multi-organ failure.

Morbidade: 53%

Mortalidade: 6,6%

39% de
complicações
>IIIb

TABLE 3. Risk factors associated with a reduced kinetic growth of the future liver remnant.

| Variables | KGR<35cc/day | Univariate | | Multivariate | |
|-------------------------------------|--------------|----------------|-------|------------------|------|
| | Yes, n° (%) | OR (95% CI) | P | OR (95% CI) | P |
| Age, yr | | | | | |
| ≥ 60 | 6 (50) | 5 (0.9-26.8) | 0.06 | 2.7 (0.1-65.3) | 0.55 |
| < 60 | 3 (16.7) | | | | |
| Sex | | | | | |
| Male | 4 (21) | 3.1 (0.6-15.8) | 0.16 | | |
| Female | 5 (45.5) | | | | |
| ASA operative risk | | | | | |
| ≥ 3 | 6 (75) | 19 (2.5-141) | 0.004 | 10.3 (0.8-136.8) | 0.08 |
| < 3 | 3 (13.6) | | | | |
| Charlson Index | | | | | |
| ≥ 7 | 8 (47) | 10.6 (1.1-101) | 0.039 | 0.9 (0.04 – 21) | 0.93 |
| < 7 | 1 (7.7) | | | | |
| BMI, kg/m ² | | | | | |
| ≥ 25 | 4 (25) | 0.6 (0.1-2.9) | 0.52 | | |
| < 25 | 5 (35.7) | | | | |
| Chemotherapy | | | | | |
| Yes | 4 (22.2) | 0.4 (0.1-1.9) | 0.26 | | |
| No | 5 (41.6) | | | | |
| FLR/TLV, % | | | | | |
| < 20 | 4 (44.4) | 0.4 (0.1-2) | 0.26 | | |
| ≥ 20 | 5 (23.8) | | | | |
| Transfusion, 1 st stage | | | | | |
| Yes | 4 (50) | 3.4 (0.6-18.7) | 0.16 | | |
| No | 5 (22.7) | | | | |
| Extrahepatic procedure | | | | | |
| Yes | 3 (27.3) | 0.8 (0.1-4.1) | 0.8 | | |
| No | 6 (31.6) | | | | |
| Pringle use | | | | | |
| Yes | 2 (22.2) | 0.6 (0.1-3.5) | 0.54 | | |
| No | 7 (33.3) | | | | |
| Liver partition | | | | | |
| Total | 5 (55.6) | 5.3 (0.1-29.2) | 0.055 | 5.4 (0.3-106.3) | 0.26 |
| Partial | 4 (19.1) | | | | |
| Complication, 1 st stage | | | | | |
| Yes | 6 (60) | 8.5 (1.4-49.5) | 0.017 | 2.9 (0.2-37.1) | 0.42 |
| No | 1 (5.6) | | | | |

OR, odds ratio; CI, confidence interval, ASA, American Society of Anesthesiologists; BMI, body mass index; FLR, future liver remnant; TLV, total liver volume.

Early Survival and Safety of ALPPS

First Report of the International ALPPS Registry

Erik Schadde, MD, FACS, Victoria Ardiles, MD,† Ricardo Robles-Campos, MD,‡ Massimo Malago, MD, FACS,§
Marcel Machado, MD,¶ Roberto Hernandez-Alejandro, MD,|| Olivier Soubrane, MD,**
Andreas A. Schnitzbauer, MD,†† Dimitri Raptis, MD,* Christoph Tschuor, MD,* Henrik Petrowsky, MD, FACS,*
Eduardo De Santibanes, MD, PhD, FACS,† and Pierre-Alain Clavien, MD, PhD, FACS*§§; On behalf of the ALPPS
Registry Group*

ALPPS Registry

TABLE 1. Main Characteristics of 202 Patients in the ALPPS Registry

| Variable of Interest | All Patients (n = 202) |
|---|------------------------|
| Age, median (IQR), yr | 60 (53–68) |
| Sex, male/female, number (%) | 121/81(60%/40%) |
| Ethnic origin | |
| White, n (%) | 188 (93) |
| Asian, n (%) | 10 (5) |
| Other*, n (%) | 4 (2) |
| Tumor type | |
| CRLM, n (%) | 141 (70) |
| HCC, n (%) | 17 (8) |
| PHCC, n (%) | 11 (5) |
| IHCC, n (%) | 8 (4) |
| NET, n (%) | 8 (4) |
| Gallbladder cancer, n (%) | 6 (3) |
| Others, n (%) | 11 (5) |
| Charlson Index (1–14)†, median (IQR) | 8 (6–9) |
| Histological abnormalities, data available (100%) | n = 150 (100%) |
| Abnormal liver histology (fibrosis/steatosis/chemotherapy-related changes), n (%) | 79 (52) |

ALPPS Registry

| | |
|--|----------|
| Location of ALPPS patients | |
| Total centers (no. centers registered) | 75 |
| Total (no. patients/no. centers) | 202/41 |
| Europe (no. patients/no. centers) | 136/27 |
| South America (no. patients/no. centers) | 43/4 |
| North America (no. patients/no. centers) | 13/4 |
| Asia (no. patients/no. centers) | 9/5 |
| Middle East (no. patients/no. centers) | 1/1 |
| Year in which ALPPS was performed | |
| 2011, n (%) | 28 (14) |
| 2012, n (%) | 112 (55) |
| 2013, n (%) | 62 (30) |
| Low- and high-volume centers | |
| <8 procedures, no. patients/no. centers | 75/31 |
| ≥8 procedures, no. patients/no. centers | 127/10 |

“Data available” refers to the number of patients in the registry with complete information about the respective variable.

*Other include 3 African patients and 1 Indian patient.

†Charlson Index is a validated method to quantify comorbidities.

CRLM indicates colorectal liver metastasis; HCC, hepatocellular carcinoma; IHCC, intrahepatic cholangiocarcinoma; IQR, interquartile range; NET, neuroendocrine tumor; PHCC, perihilar cholangiocarcinoma.

ALPPS Registry

TABLE 2. Main Operative Characteristics of 202 Patients in the ALPPS Registry

| Variable | All Patients (n = 202) |
|--|------------------------|
| Laparoscopic/robotic ALPPS, n (%) | 5 (3) |
| Type of ALPPS* | |
| <i>Right hepatectomy ALPPS, n (%)</i> | 106 (52) |
| <i>Right trisectionectomy ALPPS + Sg 1, n (%)</i> | 69 (34) |
| <i>Right trisectionectomy ALPPS – Sg 1, n (%)</i> | 17 (8) |
| Other types†, n (%) | 10 (5) |
| Mean operative time ALPPS stage I, minutes, mean (SD) | 327 (±119) |
| Mean operative time ALPPS stage II, minutes, mean (SD) | 156 (±75) |
| Pringle maneuver, data available | n = 134 (100%) |
| Performed in n (%) of cases | 65 (49) |
| Cumulative time performed, minutes, median (IQR) | 30 (16–45) |
| CVP, data available | n = 68 (100%) |
| mm Hg, median (IQR) | 5 (3–6) |

ALPPS Registry

| | |
|--|----------------|
| Blood loss ALPPS stage I, data available | n = 159 (100%) |
| <100 mL, n (%) | 23 (14) |
| 101–600 mL, n (%) | 77 (48) |
| 601–1000 mL, n (%) | 35 (22) |
| >1000 mL, n (%) | 24 (15) |
| Blood loss ALPPS stage II, data available | n = 145 (100%) |
| <100 mL, n (%) | 60 (41) |
| 101–600 mL, n (%) | 67 (46) |
| 601–1000 mL, n (%) | 10 (7) |
| >1000 mL, n (%) | 8 (6) |
| RBC transfusion ALPPS stage I | n = 189 (100%) |
| Patients transfused, n (%) | 53 (28) |
| Units of RBC, median (IQR) | 3 (2–4) |
| RBC transfusion ALPPS Stage II, data available | n = 184 (100%) |
| Patients transfused, n (%) | 44 (24) |
| Units of RBC, median (IQR) | 2 (2–3) |

ALPPS Registry

TABLE 3. Main Postoperative Outcomes of 202 Patients in the ALPPS Registry

| Variable | All Patients (n = 202) |
|--|------------------------|
| Failure to reach stage II, n (%) | 5 (2) |
| 30-d mortality, n (%) | 5 (2) |
| In-hospital mortality, n (%) | 18 (9) |
| 90-d mortality | |
| In all patients n (%) | 19 (9) |
| In CRLM, n (%) (no. total CRLM) | 11 (8%) (n = 141) |
| In HCC, n (%) (no. total HCC) | 2 (12%) (n = 17) |
| In PHCC, n (%) (no. total PHCC) | 3 (27%) (n = 11) |
| In IHCC, n (%) (no. total IHCC) | 1 (13%) (n = 8) |
| In NET, n (%) (no. total NET) | 0 (0%) (n = 8) |
| In gallbladder cancer (%) (no. total gallbladder cancer) | 2 (33%) (n = 6) |
| In subgroup ≤ 60 yr + CRLM, n (%) (no. total) | 4 (5.1%) (n = 78) |
| Highest complication \geq grade IIIa in both stages | |
| All patients, n (%) (no. total) | 80 (40%) (n = 202) |
| In CRLM group, n (%) (no. total) | 51 (36%) (n = 141) |
| In subgroup ≤ 60 yr + CRLM, n (%) (no. total) | 23 (29%) (n = 78) |

ALPPS Registry

| | |
|---|--------------------|
| Highest complication \geq grade IIIb in both stages | |
| All tumor types, n (%) | 56 (28%) (n = 202) |
| In CRLM group, n (%), (no. total) | 30 (21%) (n = 141) |
| In subgroup \leq 60 yr + CRLM, n (%) (no. total) | 12 (16%) (n = 78) |

ALPPS Registry

TABLE 4. Survival and Disease-Free Survival in 202 Patients in the ALPPS Registry

| Patients According to Tumor Type | All Patients (n = 202) |
|---|------------------------|
| All patients | 202 |
| R-status available | n = 185 (100%) |
| Incomplete resection (R1/R2), n (%) | 16 (9) |
| Median follow-up, mo (IQR) | 9 (6–13) |
| Median survival, mo | 25 |
| Survival at 1 yr (patients at risk) | 73% (52) |
| Survival at 2 yr (patients at risk) | 59% (5) |
| Median disease-free survival, mo | 14 |
| Disease-free survival at 1 yr (patients at risk) | 60% (27) |
| Disease-free survival at 2 yr (patients at risk) | 42% (1) |
| CRLM, no. patients | 141 |
| R-status available | n = 130 (100%) |
| Incomplete resection (R1/R2), n (%) | 12 (9) |
| Survival at 1 yr* (patients at risk) | 76% (41) |
| Survival at 2 yr* (patients at risk) | 62% (6) |
| Disease-free survival at 1 yr* (patients at risk) | 59% (28) |
| Disease-free survival at 2 yr* (patients at risk) | 41% (9) |
| Subgroup <60 yr + CRLM only, number of patients | 78 |
| R-status available | n = 73 (100%) |
| Incomplete resection (R1/R2), n (%) | 6 (8) |
| Disease-free survival at 1 yr* (patients at risk) | 55% (17) |

ALPPS Registry

| | |
|---|---------------|
| Subgroup <60 yr + CRLM only, number of patients | 78 |
| R-status available | n = 73 (100%) |
| Incomplete resection (R1/R2), n (%) | 6 (8) |
| Disease-free survival at 1 yr* (patients at risk) | 55% (17) |
| Disease-free survival at 2 yr* (patients at risk) | 36% (7) |
| Survival at 1 yr* (patients at risk) | 88% (33) |
| Survival at 2 yr* (patients at risk) | 74% (10) |
| HCC, no. patients | 17 |
| R-status available | n = 15 (100%) |
| Incomplete resection (R1/R2), n (%) | 0 |
| Disease-free survival at 1 yr* (patients at risk) | 87% (1) |
| Survival at 1 yr* (patients at risk) | 61% (1) |
| PHCC, no. patients | 11 |
| R-status available | n = 9 (100%) |
| Incomplete resection (R1/R2), n (%) | 2 (22) |
| Disease-free survival at 1 yr* | NA† |
| Survival at 1 yr* | NA† |
| IHCC, no. patients | 8 |
| R-status available | n = 7 (100%) |
| Incomplete resection (R1/R2) in % | 1 (14%) |
| Disease-free survival at 1 yr* (patients at risk) | 31% (1) |
| Survival at 1 yr* (patients at risk) | 73% (1) |

ALPPS Registry

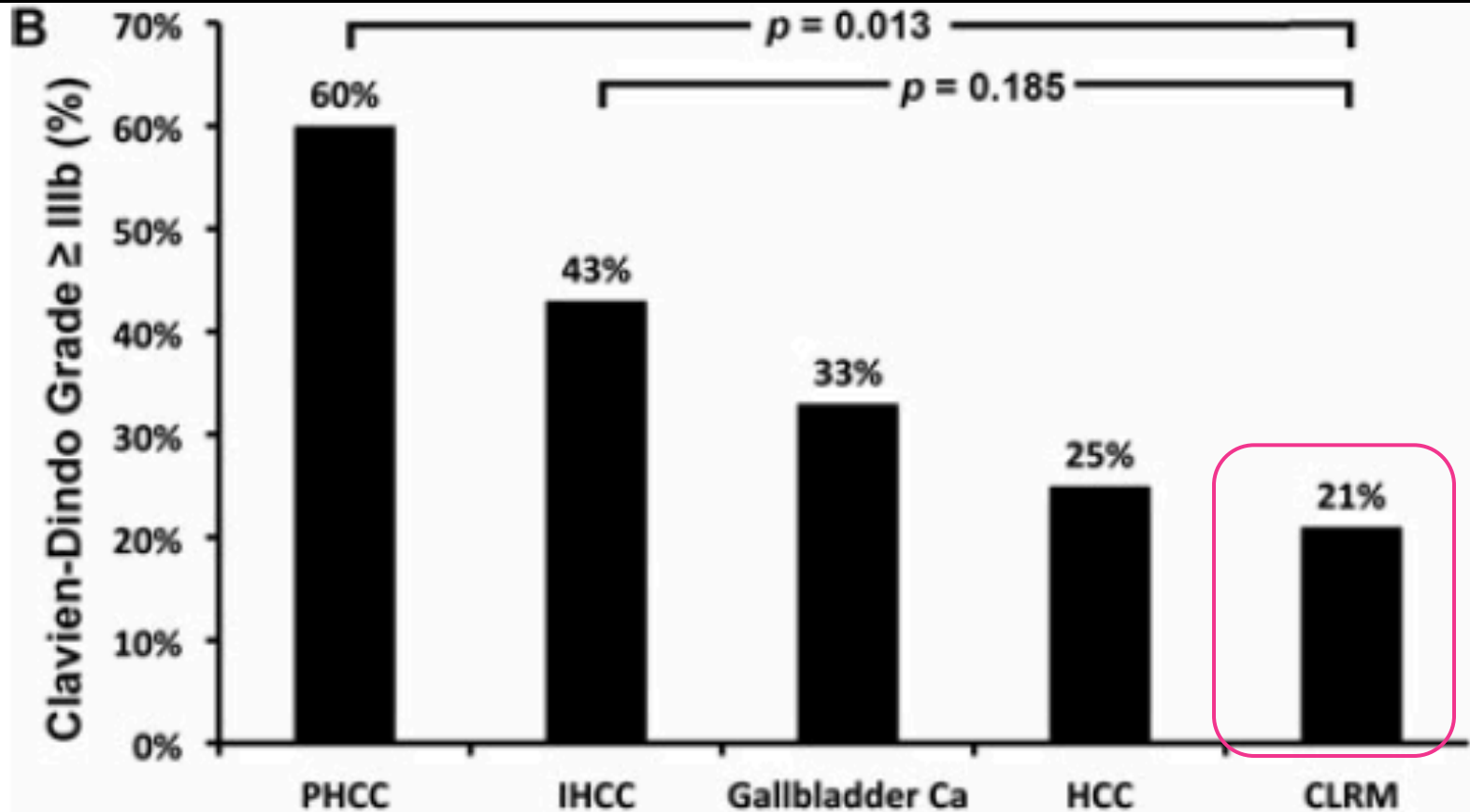
| | |
|---|--------------|
| IHCC, no. patients | 8 |
| R-status available | n = 7 (100%) |
| Incomplete resection (R1/R2) in % | 1 (14%) |
| Disease-free survival at 1 yr* (patients at risk) | 31% (1) |
| Survival at 1 yr* (patients at risk) | 73% (1) |
| NET, no. patients | 8 |
| R-status available | n = 8 (100%) |
| Incomplete resection (R1/R2), n (%) | 1 (13) |
| Disease-free survival at 1 yr* (patients at risk) | 83% (5) |
| Survival at 1 yr* (patients at risk) | 73% (1) |
| Gallbladder cancer, no. patients | 6 |
| R-status available | n = 6 (100%) |
| Incomplete resection (R1/R2), n (%) | 0 |
| Disease-free survival at 1 yr* | NA† |
| Survival at 1 yr* | NA† |

*Cumulative proportion surviving at the time according to Kaplan-Meier estimates.

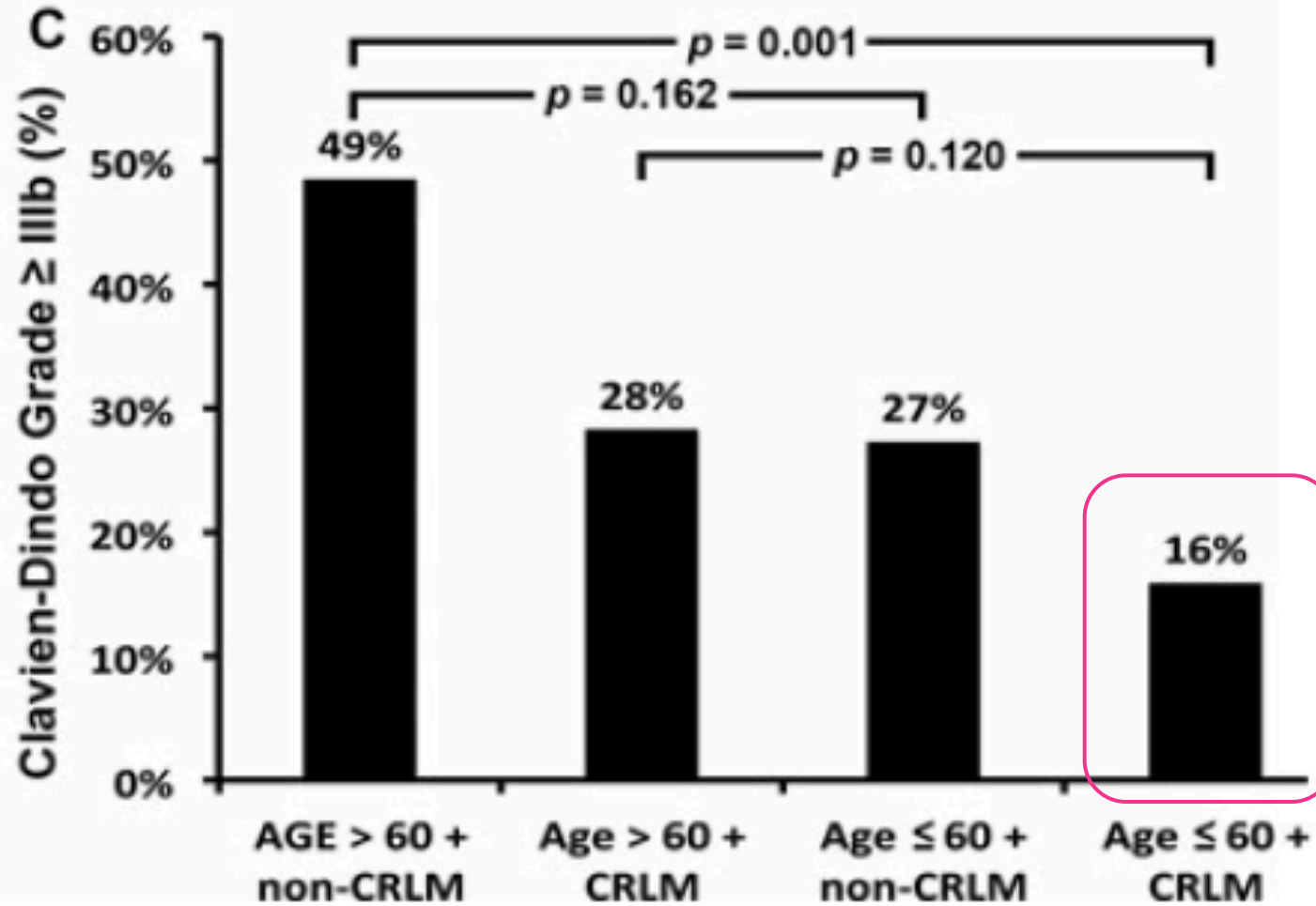
†Not available: follow-up not long enough to assess.

CRLM indicates colorectal liver metastases; HCC, hepatocellular carcinoma; IHCC, intrahepatic cholangiocarcinoma; IQR, interquartile range; NET, neuroendocrine tumor; PHCC, perihilar cholangiocarcinoma.

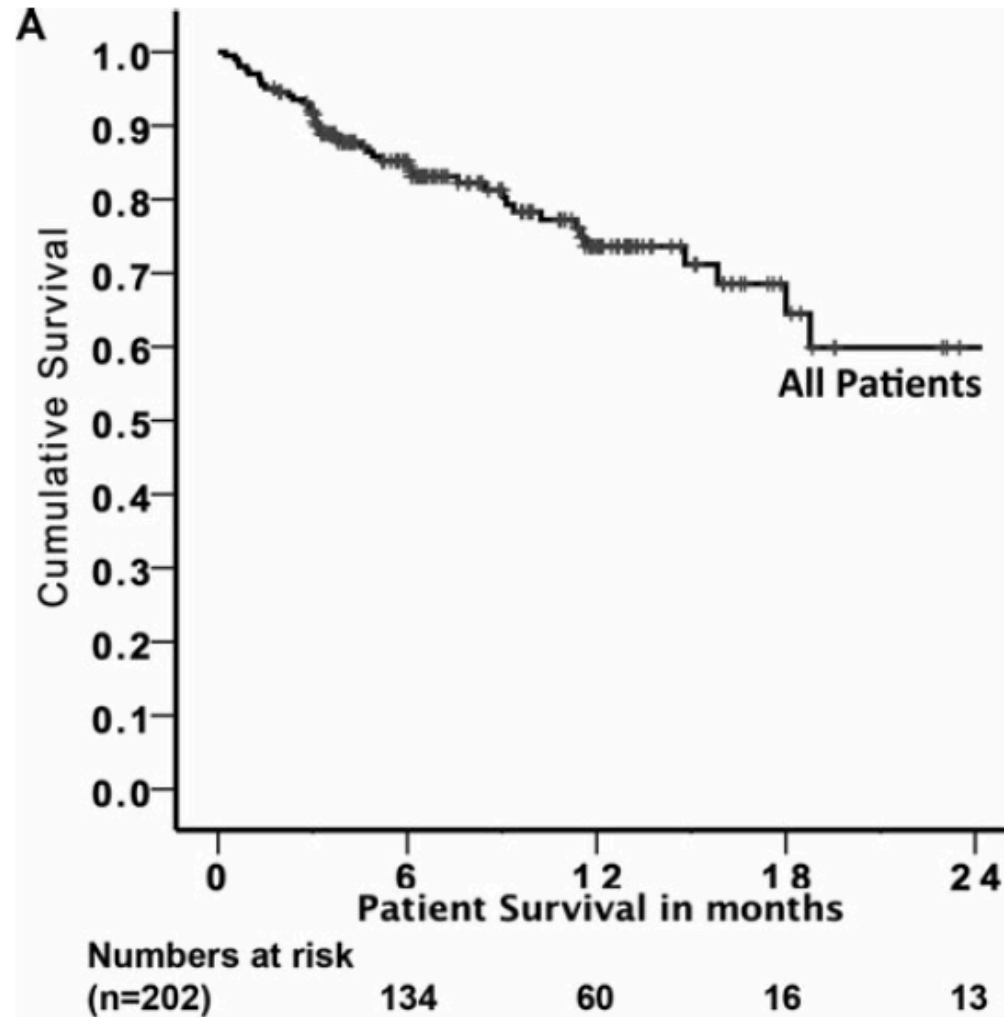
ALPPS Registry



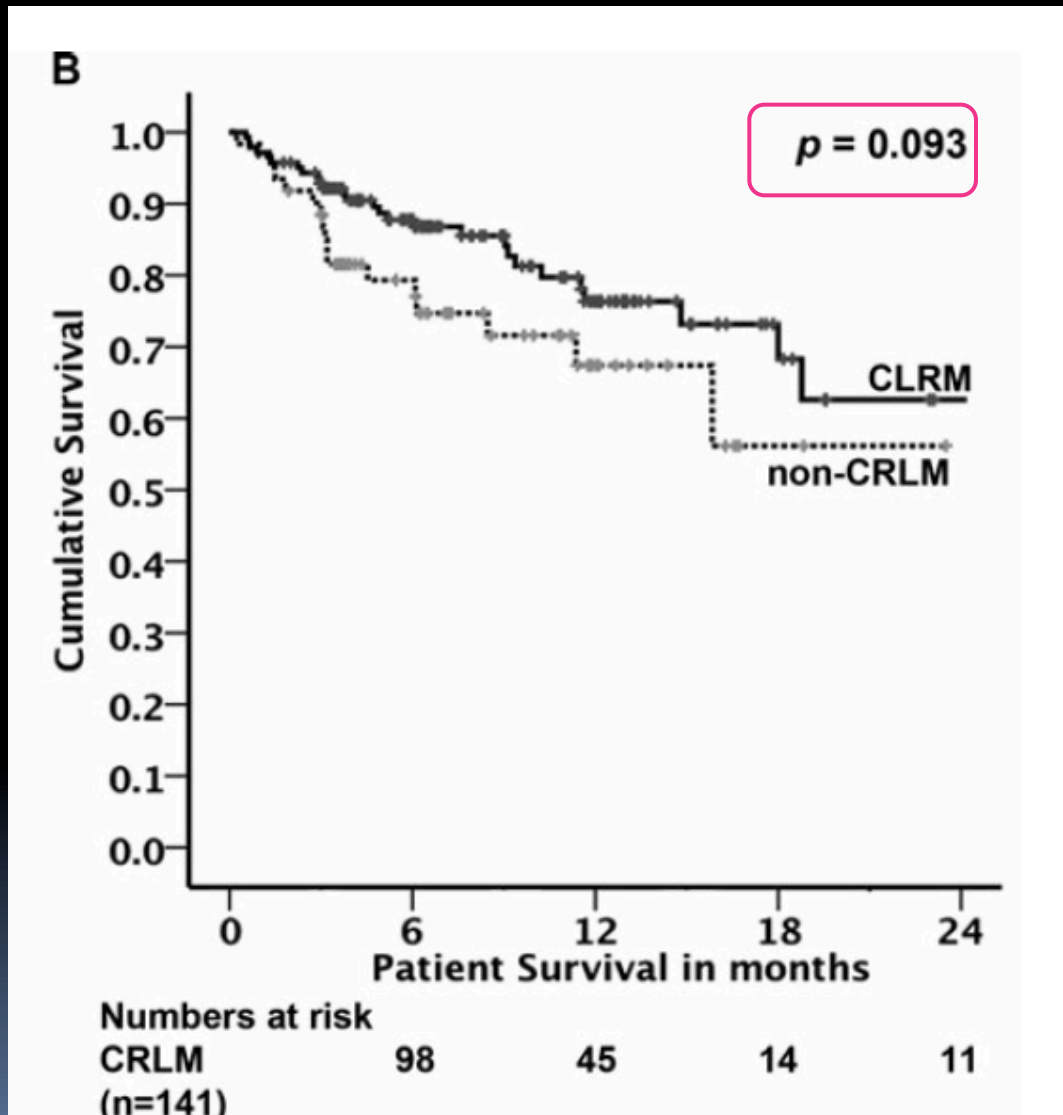
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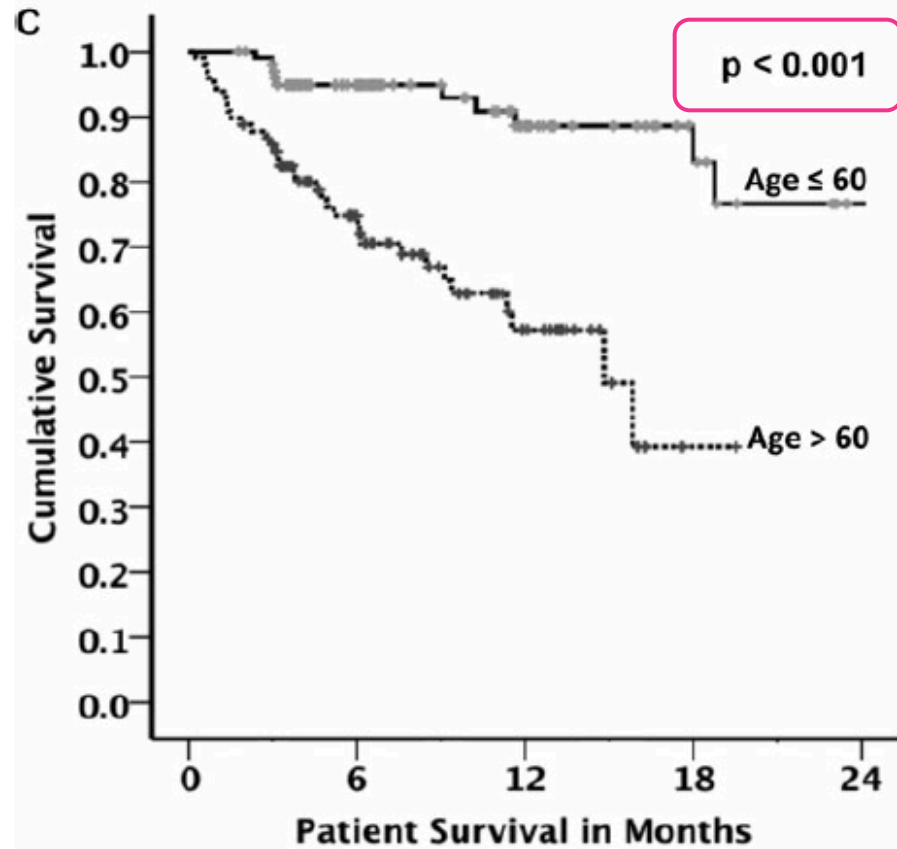
ALPPS Registry



ALPPS Registry



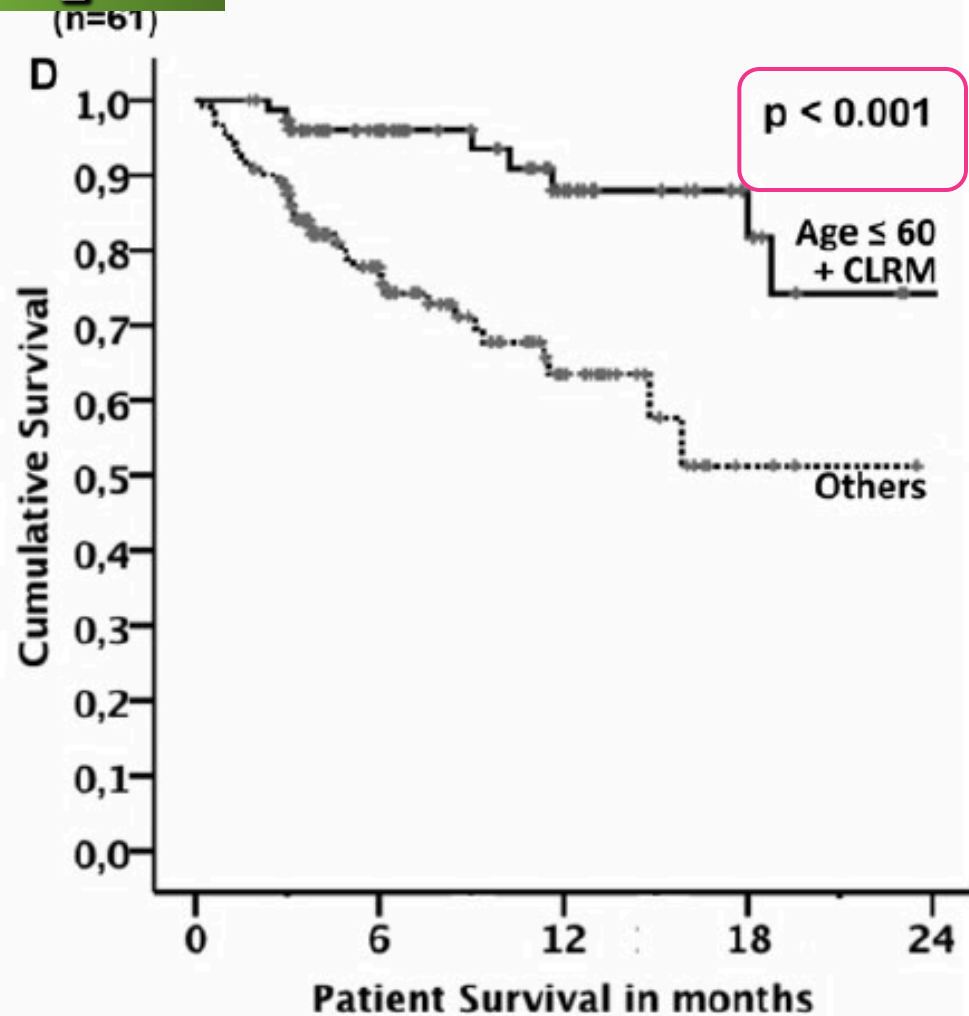
ALPPS Registry



Numbers at risk

| | | | | |
|---------------------|----|----|----|----|
| Age ≤ 60 (n=101) | 90 | 39 | 15 | 12 |
| Age > 60 (n=98) | 57 | 20 | 4 | 0 |

ALPPS Registry



Numbers at risk

| | | | | |
|---------------------------|----|----|----|----|
| Age ≤ 60 + CLRM (n=78) | 71 | 30 | 13 | 10 |
| Others (n=121) | 72 | 29 | 8 | 0 |

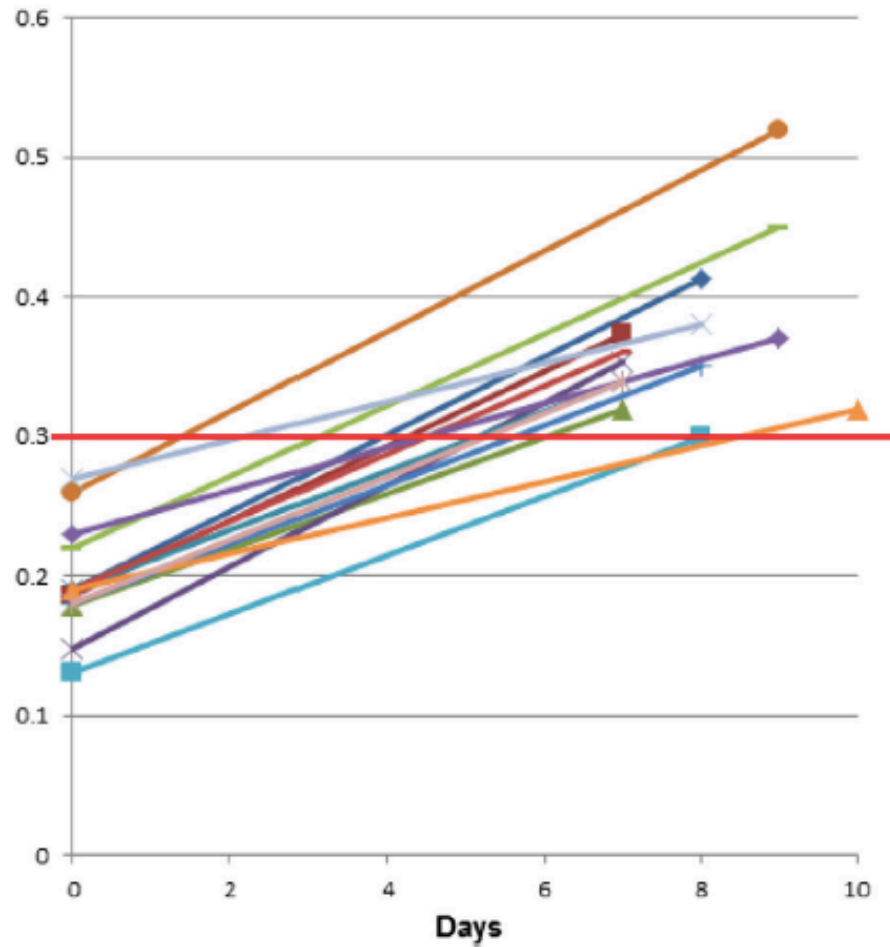
Deve ser indicado no remanescente < 30% ou < 0,5% do peso corporal para fígado sadio (< 40% ou 0,8% em fígados doentes), doença bilobar, fracasso na EVP ou LVP, extensão do tumor inesperada durante a operação e necessidade de hipertrofia > 65%.

Can we improve the morbidity and mortality associated with the associating liver partition with portal vein ligation for staged hepatectomy (ALPPS) procedure in the management of colorectal liver metastases?

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**Cortesía Dr. Roberto Hernandez-Alejandro
(London-Ontario - Canada)**

Standardized Future Liver Remnant (FLR)



| Patient | sFLR0 | sFLR1 |
|---------|-------|-------|
| 1 | 19% | 41% |
| 2 | 19% | 37% |
| 3 | 18% | 32% |
| 4 | 15% | 35% |
| 5 | 19% | 34% |
| 6 | 26% | 52% |
| 7 | 18% | 35% |
| 8 | 19% | 36% |
| 9 | 23% | 40% |
| 10 | 25% | 41% |
| 11 | 13% | 30% |
| 12 | 19% | 32% |
| 13 | 27% | 38% |
| 14 | 17% | 34% |

Fig. sFLR in ALPPS.

Table I. Summary table of ALPPS patients

| | <i>ALPPS (N = 14)</i> |
|---|-----------------------|
| Age, y | 57 (31–66) |
| Sex, male | 9 (64%) |
| Number of lesions | 9 (4–15) |
| Simultaneous resection | 4 (29%) |
| Reversal approach | 2 (14%) |
| OR time, min, stage 1 | 385 ± 59 |
| Blood loss, mL, stage 1 | 725 ± 85 |
| Units of RBC transfused stage 1 | 0.5 ± 0.8 |
| OR time, min, stage 2 | 144 ± 41 |
| Blood loss, mL, stage 2 | 178 ± 80 |
| Units of RBC transfused stage 2 | 0 ± 0 |
| Any complication | 5 (36%) |
| Severe complications (Clavien-Dindo ≥ IIIB) | 2 (14%) |
| Duration of stay between stages, mean ± SD (median) days | 8 ± 1 (8) |
| Total duration of stay, mean ± SD (median) days | 23 ± 12 (18) |

ALPPS, Associating liver partition with portal vein ligation for staged hepatectomy; *OR*, operating room; *RBC*, red blood cell.

Table II. Complications

| <i>Patient</i> | <i>Clavien-Dindo</i> | <i>Description</i> | <i>50/50 criteria</i> | <i>Ascites</i> | <i>Prolonged cholestasis</i> | <i>R resection</i> |
|----------------|----------------------|-------------------------------------|-----------------------|----------------|------------------------------|--------------------|
| 1 | IVA | Liver dysfunction | Yes | Yes | Yes | R0 |
| 2a | I | Wound infection | No | No | No | R0 |
| 3b | 0 | N/A | No | No | No | R0 |
| 4 | 0 | N/A | No | No | No | R0 |
| 5c | IVA | Liver dysfunction, wound dehiscence | Yes | Yes | Yes | R1 |
| 6d | IIIA | Abscess requiring drain | No | No | No | R0 |
| 7d,e | 0 | N/A | No | No | No | R1* |
| 8 | 0 | N/A | No | No | No | R0 |
| 9 | II | Ileus, TPN | Yes | Yes | No | R0 |
| 10d | 0 | N/A | No | No | No | R0 |
| 11d | 0 | N/A | No | No | No | R0 |
| 12 | 0 | N/A | No | No | No | R0 |
| 13b | 0 | N/A | Yes | No | No | R0 |
| 14 | 0 | N/A | No | No | No | R0 |

*R0 resection achieved with resection of IVC at second stage.

a = failed previous PVE; b = reversal approach; c = segment 4 ALPPS; d = simultaneous approach; e = vena cava resection during stage 2 of ALPPS involving the region of previous caudate R1 resection.

IVC, Inferior vena cava; N/A, not applicable; TPN, total parenteral nutrition.

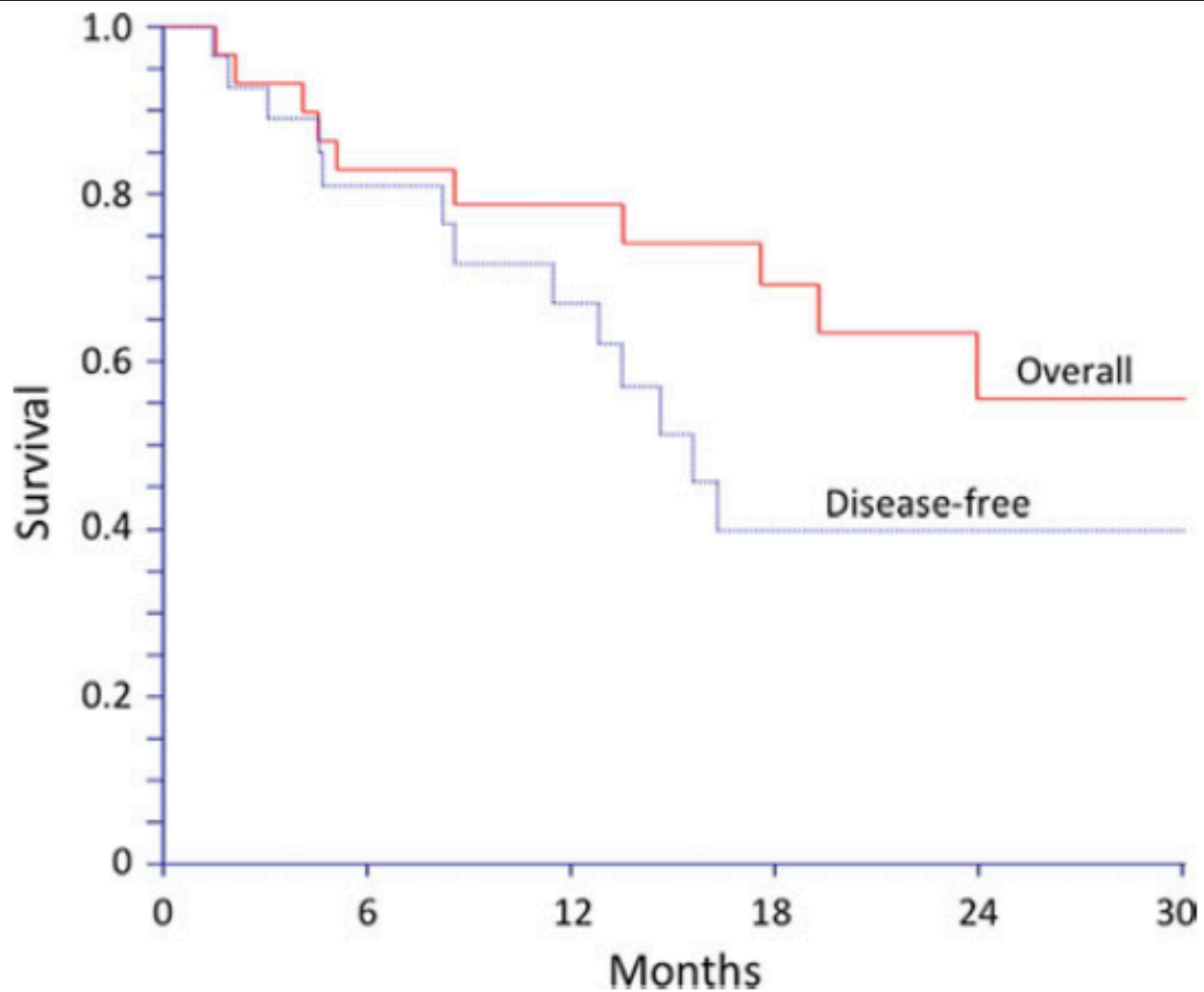


FIGURE 4. DFS and OS of the study population.

Conclusões

O procedimento ALPPS é factível, seguro e com resultados oncológicos semelhantes aos procedimentos semelhantes com a mesma finalidade.

Deve ser indicado no remanescente $< 30\%$ ou $< 0,5\%$ do peso corporal para fígado sadio ($< 40\%$ ou $0,8\%$ em fígados doentes), doença bilobar, fracasso na EVP ou LVP, extensão do tumor inesperada durante a operação e necessidade de hipertrofia $> 65\%$.

O procedimento ALPPS deve ser realizado em pacientes com metástase hepática de origem colo-retal, naqueles com idade igual ou inferior a 60 anos.

Evidência 3
Recomendação C