Resectability of bilobar liver tumours after simultaneous portal and hepatic vein embolization versus portal vein embolization alone: meta-analysis

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Abstract

Background: Many patients with bi-lobar liver tumours are not eligible for liver resection due to an insufficient future liver remnant (FLR). To reduce the risk of posthepatectomy liver failure and the primary cause of death, regenerative procedures intend to increase the FLR before surgery. The aim of this systematic review is to provide an overview of the available literature and outcomes on the effectiveness of simultaneous portal and hepatic vein embolization (PVE/HVE) versus portal vein embolization (PVE) alone.

Methods: A systematic literature search was conducted in PubMed, Web of Science, and Embase up to September 2022. The primary outcome was resectability and the secondary outcome was the FLR volume increase.

Results: Eight studies comparing PVE/HVE with PVE and six retrospective PVE/HVE case series were included. Pooled resectability within the comparative studies was 75 per cent in the PVE group (n = 252) versus 87 per cent in the PVE/HVE group (n = 166, OR 1.92 (95% c.i., 1.13–3.25)) favouring PVE/HVE (P = 0.015). After PVE, FLR hypertrophy between 12 per cent and 48 per cent (after a median of 21–30 days) was observed, whereas growth between 36 per cent and 67 per cent was reported after PVE/HVE (after a median of 17–31 days). In the comparative studies, 90-day primary cause of death was similar between groups (2.5 per cent after PVE versus 2.2 per cent after PVE/HVE), but a higher 90-day primary cause of death was reported in single-arm PVE/HVE cohort studies (6.9 per cent, 12 of 175 patients).

Conclusion: Based on moderate/weak evidence, PVE/HVE seems to increase resectability of bi-lobar liver tumours with a comparable safety profile. Additionally, PVE/HVE resulted in faster and more pronounced hypertrophy compared with PVE alone.

Introduction

Primary liver cancers and colorectal liver metastases (CRLM) are among the leading causes of cancer-related deaths1,2. Resection or ablation of the tumours in the affected liver segments are the only potentially curative treatment3. In patients undergoing liver resection for extensive bi-lobar disease, post-hepatectomy liver failure (PHLF) continues to be the primary cause of the primary cause of death4–6. In metastatic disease, only up to 20 per cent of patients are eligible for liver resection at the time of diagnosis, due to the extent of disease or an insufficient future liver remnant (FLR) volume7–10.

Generally, an FLR of approximately 30–40 per cent is considered sufficient, depending on the patient’s co-morbidities, underlying liver disease, and history of chemotherapy11–13. In patients with an estimated insufficient FLR at high risk of PHLF, FLR-hypertrophy-inducing procedures are a possibility to improve resectability14. Liver growth after portal vein embolization (PVE) was first reported in the 1980s15. In PVE, portal blood flow to the affected liver lobe later to be resected is occluded, which causes the de-portalized lobe to shrink, whereas growth is induced in the unaffected contralateral FLR16; however, 20–30 per cent of patients still do not qualify for surgery after PVE. Irresectability is mainly due to tumour progression during waiting time until sufficient hypertrophy has been achieved (typically 4–8 weeks) to allow resection or overall insufficient liver growth after PVE15,17.
More recently, several groups have developed techniques that aim for a faster and more pronounced liver hypertrophy, allowing a shorter timeframe between embolization and resection, and increase the number of patients eligible for surgery. Some of these hypertrophy-inducing techniques, however, are based on invasive surgical procedures such as associated liver partition and portal vein ligation for staged hepatectomy (ALPPS). In 2016, Guiu et al. introduced a new technique called liver venous deprivation (LVD). In LVD, the portal and hepatic vein of the diseased side of the liver are simultaneously occluded by using a combination of vascular plugs and glue to also occlude small collaterals. Depending on the research group applying the method, different variants have been described for the same intervention. In the literature, bi-embolization, double vein embolization, radiological simultaneous portohepatic vein embolization (RASPE), and combined PVE and hepatic vein embolization (HVE) have been reported. The latter consists of a combined PVE/HVE technique in which the portal vein is embolized (using glue and particles/plugs) and the hepatic vein is occluded by using vascular plugs simultaneously.

In 2019, Esposito et al. presented a systematic review evaluating the effectiveness of simultaneous performed PVE/HVE and its technical variants. They demonstrated that PVE/HVE is a safe and effective intervention to increase postoperative FLR volume, allowing 85% per cent of patients to undergo surgery, however, studies comparing PVE/HVE with PVE alone could not be included as they were not available at that time. In the meantime, multiple studies comparing these two procedures have been published. The aim of this systematic review, therefore, is to provide an update of the effectiveness of PVE/HVE compared with PVE.

Methods

This systematic review was written according to the PRISMA guidelines (Appendix S1). Randomized clinical trials (RCTs), retrospective single-arm cohort studies examining PVE/HVE, and cohort studies comparing PVE with PVE/HVE were included. Conference abstracts were excluded from this systematic review. Articles with patients aged 18 years and older diagnosed with primary or secondary liver cancer undergoing PVE or simultaneous PVE/HVE were considered eligible for inclusion. All technical variations of PVE and PVE/HVE were included, irrespective of embolization techniques and/or materials used. No other exclusion criteria with regard to patient characteristics were applied. Sequential portal and hepatic vein embolization, defined as a staged procedure with more than 48 h in between the PVE and HVE procedure, was excluded because hypertrophy of FLR induced by staged procedures is limited compared with simultaneous embolization.

The primary endpoint of this systematic review was resectability, which was defined as a surgically successful liver resection procedure, irrespective of 90-day primary cause of death. Secondary outcomes of interests were FLR volume increase (absolute and per cent), degree of hypertrophy and/or kinetic growth rate (KGR), time interval between intervention and liver volumetry, time interval between embolization and surgery, reasons for non-resectability, and 90-day primary cause of death. From each included study, the indication for PVE/HVE, embolization technique and materials used were extracted.

A systematic literature search was performed in September 2022 in PubMed, Web of Science, and Embase with no restriction on date of publication. Only studies written in English meeting the selection criteria were reviewed. Keywords and/or Medical Subject Heading (MeSH) terms were formulated and adapted to individual search engines, equivalent free-text terms were used. Search strategies specific for each search engine and the MeSH and/or free-text terms used are in Appendix S2.

Titles and abstracts were independently screened by two reviewers (J.v.Z. and R.K.) using Rayyan software. Articles were excluded if both excluded the record at the title/abstract stage. Subsequently, the same reviewers independently performed full-text screening. Disagreements regarding inclusions or exclusions of studies were resolved by discussion between the reviewers and a third reviewer (B.O.). The minimal requirement for inclusion of an article was the presentation of our primary outcome measure, resectability. Secondary outcomes that were not presented in the used articles could not be recalculated because absolute liver volumes or formula’s used were lacking. Only full-text articles on human participants in small or large cohorts were evaluated. Case reports, non-human studies, and studies in which the PVE/HVE procedure was staged (more than 48 h) were excluded. Single-arm PVE/HVE cohort studies were included only to present a comprehensive overview. The selection process is presented in a PRISMA flow chart.

A standard data collection form designed in Microsoft Excel version 16.57 (Microsoft, Redmond, WA, USA) was used. One of the authors (J.v.Z.) extracted data from each included study. Data on the following variables were collected: sample size, patient characteristics (age, BMI, underlying liver disease status, and patient exclusion criteria), and primary and secondary outcomes. The primary outcome was expressed as a percentage.

Study quality was assessed using the Effective Public Health Practice Project (EPHPP) tool by one of the authors (J.v.Z.). Included studies were assessed on quality based on criteria in six fields: selection bias, study design, confounders, blinding, data collection methods, and withdrawal and dropouts. Quality in each field could be classified as ‘strong’, ‘moderate’, or ‘weak’. An overall rating of ‘strong’ was given by the authors according to EPHPP to studies that did not score ‘weak’ on any of the fields, ‘moderate’ to studies that had only one ‘weak’ assessment, and ‘weak’ when two or more fields were rated as ‘weak’.

Review Manager version 5.4.1 (RevMan [Computer program]. Version 5.4, The Cochrane Collaboration, 2020.) was used to pool data and calculate ORs. The Mantel–Haenszel (M–H) statistical method in combination with a fixed-effects model was used to compare the pooled resectability between groups. Results were presented in a forest plot.

Results

A total of 1858 articles were identified by the search strategy. After screening of titles and abstracts, 47 articles were eligible for full-text screening. Four reports written in German could not be retrieved, and an additional nine duplicates were removed. Out of the resulting 34 articles, 22 were excluded after full-text screening for the following reasons: because of a sequential embolization procedure (n = 2), reporting of different outcome measures (n = 2), conference abstracts (n = 7), lack of intervention (n = 1), full text not available in English or Dutch (n = 1), case studies (n = 2), animal studies (n = 2), or summaries or commentaries (n = 5). Two articles were added to the review through screening of the reference lists of the articles obtained during the full-text screening. The complete process of study identification and selection is shown in the flow chart (Fig. 2). Overall, this process resulted in a total of 14 studies, eight...
comparative studies and six case series. The included studies were published between 2016 and 2022. Two comparative studies, FVE/HVE versus ALPS, were included in this systematic review and handled as two case series on FVE/HVE.

The characteristics of the patients are presented in Table 1. Sample size of the included studies ranged from 6 to 160 for FVE and 6 to 39 participants for FVE/HVE. The mean age ranged from 55 to 67 years across the intervention groups. Three studies included patients with underlying liver disease, such as liver steatosis, fibrosis, and cirrhosis. More patients with underlying liver disease were included in the FVE/HVE groups. The retrospective DRAGON Trial Collaborative analysis was the only study that reported a significant difference in underlying liver disease between the two groups (p = 0.021), but multivariable analysis showed that this difference did not impact the resectability rate. In the other four studies, underlying liver disease status was not documented in two studies and was an explicit exclusion criterion in four studies. A total of 334 patients in the FVE group and 190 patients in the FVE/HVE group were included by the comparative studies. Data from 175 FVE/HVE patients from an additional six cohort studies were included in this systematic review.

Information on selection of baseline per cent FLR, definition of PHLF, additional segment four embolization, additional middle hepatic vein occlusion, and formulae used for baseline FLR are presented in Table S1.

Resectability rates are presented in Table 2. In most studies, resectability did not differ significantly between treatments and ranged from 64 to 94 per cent and from 67 to 100 per cent for FVE and FVE/HVE respectively. Summary analysis showed that resection was possible in 252 (75 per cent) patients after FVE and 166 (87 per cent) patients after FVE/HVE in the comparative studies. In the single-arm studies, 175 patients underwent FVE/HVE, of whom 146 patients underwent successful liver resection (83 per cent). Overall, reasons for non-resectability were insufficient post-embolization FLR volume in 22 (6.6 per cent) versus 5 (1.4 per cent) patients, peritoneal carcinomatosis in 11 (0.3 per cent) versus 5 (1.4 per cent) patients, or disease progression in 47 (14.1 per cent) versus 38 (10.4 per cent) patients, for FVE and FVE/HVE respectively. In the retrospective DRAGON analysis, a significant difference in resectability was found between groups (p = 0.007). Kobayashi et al. did not present a p value for resectability, so it is uncertain whether the difference was statistically significant in this article. Other studies reported non-significant differences or did not report significance. For each comparative study, ORs (with 95 per cent confidence intervals) for resectability were calculated and are graphically presented in Fig. 2. After pooling of results, resectability was more frequently possible after FVE/HVE (OR 1.92 (1.13 to 3.25),
P = 0.015) with a 9 per cent heterogeneity (I²) across the included studies (Fig. 2). Eleven out of 14 articles reported on 90-day primary cause of death. In the comparative studies, 90-day primary cause of death was similar between the groups (2.53 per cent after PVE and 2.23 per cent after PVE/HVE, respectively), whereas a higher 90-day primary cause of death was reported in the single-arm PVE/HVE cohort studies (6.9 per cent after PVE/HVE).

Median days between embolization and resection ranged from 35–44 days for PVE and 23–49 days for PVE/HVE with considerable variation.

Liver volume before and after embolization, time interval between embolization and liver volumetry, and mean or median KGR are presented in Table 3. At baseline pre-embolization, FLR volume was expressed in ml or cc, and either per cent FLR or per cent standardized FLR (sFLR) was reported. A comparison of the studies on FLR volume was difficult due to the usage of different volume metrics. All studies reported an increase in FLR volume and percentage from baseline after both PVE and PVE/HVE. Baseline FLR volume ranged from 294 to 542 ml for PVE and from 281 to 547 ml for PVE/HVE. A significant difference was reported for per cent FLR at baseline between the groups only in the study by Laurent et al. (P < 0.001).

Contrary to baseline FLR, post-embolization FLR volume was expressed in ml or cc by five studies, whereas two studies only reported on the degree of hypertrophy. Different formulae were used to calculate liver hypertrophy; 

| Author | Year | n | Tumour type(s) | Age (years) | Underlying liver disease | BMI | Exclusion criteria |
|--------|------|---|----------------|-------------|-------------------------|-----|------------------|
| Hoque et al. | 2018 | PVE 6, PVE/HVE 6 | pHCC 6 | FLR | PVE | NA |
| Panaro et al. | 2019 | PVE 16, PVE/HVE 13 | CRLM 5, HCC 9 | FLR | PVE | NA |
| Le Roy et al. | 2020 | PVE 41, PVE/HVE 31 | CRLM 26, HCC 2, pHCC 2 | FLR | PVE | NA |
| Kobayashi et al. | 2020 | PVE 39, PVE/HVE 21 | CRLM 20, pHCC 8 | FLR | PVE | NA |
| Laurent et al. | 2020 | PVE 36, PVE/HVE 37 | CRLM 20, pHCC 4, NET 3 | FLR | PVE | NA |
| Guis et al. | 2020 | PVE 22, PVE/HVE 29 | CRLM 17, HCC 2, iHCC 1 | FLR | PVE | NA |
| Heil et al. | 2021 | PVE 160, PVE/HVE 39 | CRLM 85, HCC 11, iHCC 22, pHCC 25, GBC 9 | FLR | PVE | NA |
| Boning et al. | 2022 | PVE 14, PVE/HVE 14 | CRLM 4, pHCC 10 | FLR | PVE | NA |

Case series

Guis et al. | 2016 | PVE/HVE 7 | NA | FLR | PVE | NA |
| Le Roy et al. | 2017 | PVE/HVE 10 | NA | FLR | PVE | NA |
| Chebaro et al. | 2021 | PVE/HVE 124 | NA | FLR | PVE | NA |
| Ghosh et al. | 2021 | PVE/HVE 12 | NA | FLR | PVE | NA |
| Cassese et al. | 2022 | PVE/HVE 15 | NA | FLR | PVE | NA |

All values are median (range or i.q.r.) as extracted from the included studies. PVE, portal vein embolization; PVE/HVE, simultaneous PVE and hepatic vein embolization; CRLM, colorectal liver metastasis; HCC, hepatocellular carcinoma; iHCC, intrahepatic carcinoma; pHCC, perihilar carcinoma; GBC, gallbladder carcinoma; NET, neuroendocrine tumour; NR, not reported; NA, not applicable; NASH, non-alcoholic steatohepatitis; i.q.r., interquartile range.
Table 2 Waiting time, resectability rates, and 90-day primary cause of death

| Author                  | Time between embolization and surgery (days) | Resectability (%) | Reasons for exclusion from resection | 90-day primary cause of death |
|-------------------------|---------------------------------------------|-------------------|-------------------------------------|------------------------------|
|                         | PVE | PVE/HVE | PVE | PVE/HVE | PVE | PVE/HVE | PVE | PVE/HVE | PVE | PVE/HVE |
| **Comparative studies** |     |         |     |         |     |         |     |         |     |         |
| Hocquelet et al.        | NR  | 21      | 83  | 67      |      |          |     |         |     |         |
| Panaro et al.           | 37  | 38      | 94  | 100     |      |          |     |         |     |         |
| Le Roy et al.           | NR  | NR      | 76  | 81      |      |          |     |         |     |         |
| Kobayashi et al.        | 35  | (20–181)| 77  | 95      |      |          |     |         |     |         |
| Laurent et al.          | 44  | (21–78) | 89  | 86      |      |          |     |         |     |         |
| Guiu et al.             | 36  | (22–55) | 91  | 97      |      |          |     |         |     |         |
| Heil et al.             | 41  | (28–61)| 68  | 90      |      |          |     |         |     |         |
| Boning et al.           |     |         | 71  | 64      |      |          |     |         |     |         |
| **Case series**         |     |         |     |         |     |          |     |         |     |         |
| Guiu et al.             | NA  | 23      | NA  | 85.6    |      |          |     |         |     |         |
| Le Roy et al.           | NA  | 31      | NA  | 90      |      |          |     |         |     |         |
| Chebaro et al.          | NA  | 37      | NA  | 80.6    |      |          |     |         |     |         |
| Ghosn et al.            | NA  | 39      | NA  | 83.3    |      |          |     |         |     |         |
| Cassese et al.          | NA  | 39      | NA  | 100     |      |          |     |         |     |         |

Values are median (range or i.q.r.), or mean(s.d.) as extracted from the included studies. PVE, portal vein embolization; PVE/HVE, simultaneous portal and hepatic vein embolization; FLR-V, future remnant liver volume; FLR-F, future remnant function; NR, not reported; NA, not applicable. *Not clear whether this is the 30-day or 90-day primary cause of death.

Table 4 An overview of the study quality assessment of the included studies

| Study or subgroup | PVE/HVE | PVE | Odds ratio M–H, fixed, 95% c.i. | Odds ratio M–H, fixed, 95% c.i. |
|-------------------|---------|-----|--------------------------------|--------------------------------|
| Panaro et al.     | 28      | 13  | 2.61 (0.10, 69.63)              |                                 |
| Le Roy et al.     | 21      | 25  | 1.34 (0.43, 4.21)               |                                 |
| Laurent et al.    | 20      | 32  | 0.80 (0.20, 3.25)               |                                 |
| Kobayashi et al.  | 32      | 20  | 0.80 (0.20, 3.25)               |                                 |
| Hocquelet et al.  | 38      | 4   | 0.40 (0.03, 6.18)               |                                 |
| Heil et al.       | 21      | 35  | 0.40 (0.13, 12.14)              |                                 |
| Guiu et al.       | 18      | 28  | 0.80 (0.24, 33.04)              |                                 |
| Boning et al.     | 29      | 9   | 0.72 (0.15, 3.54)               |                                 |
| Total (95% c.i.)  |         | 100 | 1.92 (1.13, 3.25)               |                                 |
| Total events      | 166     | 252 |                               |                                 |

Heterogeneity: $\chi^2 = 7.66, 7$ d.f., $P = 0.36; I^2 = 9%$
Test for overall effect: $Z = 2.42, P = 0.02$

Three studies reported a significant difference on KGR between PVE and PVE/HVE (all $P < 0.001$)21,29,32. Le Roy et al. (2020) presented liver growth as degree of hypertrophy: 8 per cent versus 19 per cent, for PVE and PVE/HVE respectively ($P = 0.026$)31. These rates were comparable to the degree of hypertrophy of 13 per cent versus 21 per cent for PVE and PVE/HVE reported by Heil et al. ($P < 0.001$)21. An overview of the study quality assessment of the included studies is presented in Table 4. According to EPHP, two articles were rated as ‘weak’28,38, and the others as ‘moderate’. No RCTs

Figure 2. Forest plot of calculated ORs (95% c.i.) for resectability in comparative studies

PVE, portal vein embolization; PVE/HVE, simultaneous PVE and hepatic vein embolization.

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four studies finding significant differences between the two groups21,29,31,32. Two studies did not report on per cent hypertrophy18,28.

Mean or median time (days) between embolization and liver volumetry was 21–30 and 17–31 days for PVE and PVE/HVE respectively. Heil et al. (2021) reported a significant difference on interval between embolization and liver volumetry, favouring the PVE/HVE group (24 versus 17 days, $P = 0.009$). Laurent et al. (2020) showed a non-significant difference ($P = 0.95$20, whereas the remaining studies did not report on significance.
Discussion
This systematic review and meta-analysis shows that PVE/HVE, as a regenerative procedure to enhance FLR hypertrophy in patients scheduled for extended liver resection, results in a significantly higher resection rate (87% versus 75% per cent) compared with PVE only. The KGR tends to be higher in the PVE/HVE group but could not be statistically analysed due to the different ways of reporting this variable by different studies. The 90-day primary cause of death was comparable in the two embolization groups. These results suggest superiority of PVE/HVE over PVE in

Table 3 Volumetry and growth data

| Author                | Baseline FLR volume (ml, cc, %) | Time between embolization and liver volumetry (days) | Hypertrophy (%) | KGR |
|-----------------------|---------------------------------|-----------------------------------------------|-----------------|-----|
|                       | PVE                             | PVE/HVE                                        |                 |     |
| Comparative studies   |                                 |                                               |                 |     |
| Hocquelet et al.38    | 429 (391–560)                   | 517 (310–828)                                  | 23.5* (15–29)   | 67  |
|                       | 31 FLR (24–33)                  | 35.3 (23–35.3)                                 | 23.5* (15–29)   |     |
|                       | 31.2 (6.5) FLR                  |                                               | 31.3 (10.57–43.27) |     |
|                       | 394 (262–478)                   |                                               |                 |     |
| Panaro et al.28       | NR (266–547)                    | 21* (27)                                       | 21* (27)        |     |
|                       | 523 (420–659)                   | 31.2 (6.5) FLR                                 | 31.9 (34)       |     |
|                       | 24 sFLR (20–30)                 |                                               | 40.8 (7.9) FLR |     |
|                       | 547 (453–656)                   |                                               | 51.2 (41.7)     |     |
|                       | 25 sFLR (32–31)                 |                                               | 4.8 (4) cc/day  |     |
|                       | 387 (203–263)                   |                                               | 8 (13)%/week    |     |
|                       | 22.9 FLR (16–23)                |                                               | 19 (18)%/week   |     |
|                       | 30 (25–43)                      |                                               |                 |     |
|                       | 31 (21–40)                      |                                               | 29              |     |
|                       | 61.2 NR NR                      |                                               |                 |     |
| Laurent et al.30      | 468 (23–845)                    |                                               | 22.5*           |     |
|                       | 31.03 FLR (18.33–38.95)         |                                               |                 |     |
|                       | 387 (16.55–32.15)               |                                               | 18.6 (−10.7–102.2) |     |
|                       | 22.91 FLR (16.6–37.7)           |                                               | 52.6 (1.0–175.6) |     |
|                       | 24 (19–37)                      |                                               | NR NR           |     |
|                       | 17 (13–32)                      |                                               |                 |     |
|                       | 48 (24–69)                      |                                               |                 |     |
|                       | 59 (45–79)                      |                                               |                 |     |
|                       | 30 (25–43)                      |                                               |                 |     |
|                       | 31 (21–40)                      |                                               |                 |     |
|                       | 49.4 (28.9)                     |                                               |                 |     |
|                       | 48.2 (22.2)                     |                                               |                 |     |
|                       | 30* (25–43)                     |                                               |                 |     |
|                       | 44.9 (28.9)                     |                                               |                 |     |
|                       | NR NR                           |                                               |                 |     |
| Case series           |                                 |                                               |                 |     |
| Guiu et al.29         | 542 (236–1119)                  |                                               | 21              |     |
|                       | 27 FLR (13.7–47.7)              |                                               | 18.6 (−10.7–102.2) |     |
|                       | 548 (233–805)                   |                                               | 52.6 (1.0–175.6) |     |
|                       | 22.6 FLR (16.6–37.7)            |                                               | NR NR           |     |
|                       | 281 (234–592.1)                 |                                               |                 |     |
|                       | 18 sFLR (16–23)                 |                                               |                 |     |
|                       | 30 (25–43)                      |                                               |                 |     |
|                       | 31 (21–40)                      |                                               |                 |     |
|                       | 30* (25–43)                     |                                               |                 |     |
|                       | 44.9 (28.9)                     |                                               |                 |     |
|                       | 48.2 (22.2)                     |                                               |                 |     |
|                       | 30* (25–43)                     |                                               |                 |     |
|                       | 44.9 (28.9)                     |                                               |                 |     |
|                       | NR NR                           |                                               |                 |     |
| Guiu et al.18         | NA 409 (345–601)                | NA 23 (13–30)                                  | NA NR           |     |
|                       | 28.2% FLR (22.4–33.3)           | NA 21                                           | 63.3            |     |
|                       | 28 (4–52)                       |                                               | 63              |     |
|                       | 28(7.6)                         |                                               | 45 (19–28)      |     |
|                       | 28 (7.9)                        |                                               | 2.5%/week       |     |
|                       | 2%/day                          |                                               |                 |     |
|                       | 3.5%/week(2.3)                  |                                               |                 |     |
|                       | NA 29.3(6.8)                    |                                               |                 |     |
|                       | 29(7.9)                         |                                               |                 |     |
|                       | 29(7.9)                         |                                               |                 |     |
|                       | 29(7.9)                         |                                               |                 |     |
|                       | 29(29)                          |                                               |                 |     |
|                       | 49(29)                          |                                               |                 |     |
|                       | 49(29)                          |                                               |                 |     |
|                       | 0.2%/day(2.0)                   |                                               |                 |     |

Values are median (range or i.q.r.), or mean(s.d.) as extracted from the included studies. FLR, future liver remnant; KGR, kinetic growth rate; PVE, portal vein embolization; PVE/HVE, simultaneous portal and hepatic vein embolization; NR, not reported. *Authors did not provide specific numbers for each group, only a mean was presented. †No information was given on FLR hypertrophy, instead post-embolization FLR percentage was presented.

Table 4 The Effective Public Health Practice Project quality assessment for included studies

| Author                | Selection bias | Study design | Confounders | Blinding | Data collection methods | Withdrawals and dropouts | Global rating |
|-----------------------|----------------|--------------|-------------|----------|-------------------------|--------------------------|---------------|
| Hocquelet et al.38    | −              | ±            | −           | ±        | −                       | −                        | Weak          |
| Panaro et al.28       | −              | ±            | −           | ±        | +                       | +                        | Weak          |
| Le Roy et al.31       | −              | ±            | −           | ±        | +                       | +                        | Moderate       |
| Kobayashi et al.32    | −              | ±            | −           | ±        | +                       | ±                        | Moderate       |
| Laurent et al.30      | −              | ±            | ±           | −        | +                       | ±                        | Moderate       |
| Guiu et al.29         | −              | ±            | +           | ±        | −                       | ±                        | Moderate       |
| Heil et al.21         | −              | ±            | ±           | ±        | +                       | +                        | Moderate       |
| Boning et al.39       | −              | ±            | −           | ±        | +                       | −                        | Weak          |
| Guiu et al.18         | −              | ±            | −           | ±        | +                       | +                        | Weak          |
| Guiu et al.33         | −              | ±            | −           | ±        | +                       | +                        | Weak          |
| Le Roy et al.19       | −              | ±            | −           | ±        | +                       | −                        | Weak          |
| Chebaro et al.35      | −              | ±            | −           | ±        | +                       | −                        | Weak          |
| Ghosn et al.40        | −              | ±            | −           | ±        | +                       | +                        | Weak          |
| Cassese et al.37      | −              | ±            | −           | ±        | +                       | +                        | Weak          |

Weak quality score (−), Moderate quality score (+), strong quality score (+).
regenerative effectiveness; however, this systematic review included only retrospective studies of moderate or weak quality at best. In addition, only a small number of participants was included in most of the studies.

The relatively large multicentre study of Heil et al. was the only study able to demonstrate a significant difference in liver resectability, which was primarily responsible for the results of the pooled primary outcome of this meta-analysis.

The lack of difference in resectability presented by most of the studies individually may be explained by relatively large FLR volumes before the embolization, long intervals between embolization and resection, small sample sizes, patient selection, use of interval chemotherapy, or differing practice regarding chemotherapy-free intervals. Patients with CRLM who are kept off chemotherapy for long intervals before complete resection may have a propensity of progression in the process of waiting.

In another review by Heil et al. a resection success rate of 87 per cent after PVE/HVE was reported. An 85 per cent resection rate was reported after PVE/HVE by Esposito et al. Both their reviews included studies with patients with multiple tumour types. In the present systematic review, resection rates after PVE/HVE ranged from 67 to 100 per cent. This demonstrates that the rates vary widely depending on the type of study and the number of patients who were included and are influenced by selection and treatment biases in each centre.

We recommend researchers to use one standardized growth outcome measure in future publication to be able to compare future studies more easily. We propose to use KGR as this is a standardized outcome measure. KGR reflects the regenerative capacity of the FLR over time and therefore can be compared more reliably among different studies. The KGR also has a predictive value for the potential risk for future PHLF; however, KGR as an outcome measure carries the risk of bias as the majority of growth is achieved in the first week after embolization, until a plateau phase is reached after approximately 21 days, as shown by two studies in the present meta-analysis.

A consensus among included studies seems to exist to obtain volumetric data 21 days or less after embolization. A longer waiting interval until volumetry consequently leads to underestimation of the KGR when compared with a KGR with a shorter time interval. Therefore, standardization of time intervals after both PVE and PVE/HVE is required for adequate comparison. We recommend researchers to present the KGR over the first 3 weeks after embolization, until a plateau phase is reached after approximately 21 days, as shown by two studies in the present meta-analysis.

A consensus among included studies seems to exist to obtain volumetric data 21 days or less after embolization. A longer waiting interval until volumetry consequently leads to underestimation of the KGR when compared with a KGR with a shorter time interval. Therefore, standardization of time intervals after both PVE and PVE/HVE is required for adequate comparison. We recommend researchers to present the KGR over the first 3 weeks after embolization, until a plateau phase is reached after approximately 21 days, as shown by two studies in the present meta-analysis.

The potential that surgery may be performed earlier after PVE/HVE may be a further advantage of this combined procedure, which may translate into long-term oncological survival benefits. Although, the included studies probably did not focus on decreasing the time from intervention to surgery, as shown by the long interval, the interval was reported to be significantly shorter by Heil et al. The time interval to obtain volumetric data in the included studies suggests that performing the first postembolization volumetry at week 1 can shorten the interval time between embolization and surgery. In cases where the FLR is not sufficient at 1 week after embolization volumetry, an estimation of the eventually expected FLR hypertrophy can be made based on the KGR. Consequently, the smallest time window to resection can be achieved to further reduce the risk of tumour progression and increase the feasibility of resection.

There are some limitations regarding the included studies and the systematic review itself. First, in contrast to a previous systematic review and a scoping review, it was decided to present outcome measures in several metrics, so that the included studies could be compared more easily. This showed that research groups opt for various formulae to calculate outcome measures. In particular, the formula to calculate the outcome measure ‘hypertrophy’ was not consistent across the included studies, therefore readers of the present systematic review should be cautious about the interpretation of this outcome. This is a major concern and hinders an accurate comparison between reported results. For reliable comparison of future studies, the formula sFLR1 − sFLR0 for the degree of hypertrophy is recommended by the authors of this systematic review as well as the formula for KGR:

\[ \text{KGR} = \frac{sFLR1 - sFLR0}{sFLR0} \]

Another limitation of this review is the small sample size in the different studies. Third, included articles were best qualified as ‘moderate’ according to the EPHP. The risk of selection bias therefore is very high due to the retrospective design, and it is often difficult or even impossible to determine how the authors dealt with this issue.

Although patients are selected for embolization based on guidelines obtained by CT volumetry, several studies included patients in the PVE group with high baseline FLR percentages. This suggests that patients with higher baseline FLRs (25–30 per cent) are ideal candidates, but further research needs to clarify cutoffs in different types of liver quality and/or function. Laurent et al. reported that one-third of the included patients had no initial indication for embolization to induce liver hypertrophy before liver resection. This raises questions about the selection of patients as appropriate controls in these studies. Therefore, it is recommended that both KGR and baseline FLR volumes are reported. Moreover, choosing an appropriate matched control group seems to be important to allow for valid comparison, as two studies had relatively large differences between the control and intervention group.

Furthermore, an era bias was observed in some studies, as PVE was completely replaced by PVE/HVE. This observation is supported by the fact that the same selection criteria were used for PVE and PVE/HVE. Considering the assumption that PVE/HVE induces faster and more growth, some might have expected lower baseline per cent FLR in the PVE/HVE group; however, baseline per cent FLR did not statistically differ in most of the included studies.

RCTs with larger sample sizes are needed to make reliable statements on the actual effect of PVE/HVE compared with PVE. Two prospective trials are currently in progress, the HYPER-LIV01 (registration number: NCT03841305; http://www.clinicaltrials.gov) and DRAGON 1 trial (registration number: NCT04272931; http://www.clinicaltrials.gov) was just ethically approved and will commence later this year.

PVE/HVE seems to be more effective than PVE with regard to resection rate and seems to offer increased KGR. Therefore, we suggest that, to induce hypertrophy before major liver resection, PVE/HVE could be considered in patients with a small FLR. Propective trials to examine the exact role of PVE/HVE are currently underway.
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**Funding**
The DRAGON Trials Collaborative received unrestricted financial support from the Dutch Cancer Foundation, National Institute for Health and Care Research (UK), Abbott Laboratories, Maastricht University Medical Center+, and Guerbet.

**Acknowledgements**
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**Disclosure**
The authors declare no conflict of interest.

**Supplementary material**
Supplementary material is available at BJS Open online.

**Data availability**
All data generated or analysed during this study are included in this published article and its supplementary information files.

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