

# Three-dimensional length from the center of the liver is a prognostic factor of colorectal cancer with liver metastasis: a retrospective analysis

**Jun Woo Bong**

Korea University College of Medicine and School of Medicine

**Yeonuk Ju**

Korea University College of Medicine and School of Medicine

**Jihyun Seo**

Korea University College of Medicine and School of Medicine

**Sang Hee Kang**

Korea University College of Medicine and School of Medicine

**Pyoung-Jae Park**

Korea University College of Medicine and School of Medicine

**Sae-Byeol Choi**

Korea University College of Medicine and School of Medicine

**Sun Il Lee** (✉ [silee@korea.ac.kr](mailto:silee@korea.ac.kr))

Korea University College of Medicine and School of Medicine

**Sang Cheul Oh**

Korea University College of Medicine and School of Medicine

**Byung Wook Min**

Korea University College of Medicine and School of Medicine

---

## Research

**Keywords:** colorectal cancer, liver metastasis, resectability

**Posted Date:** July 20th, 2020

**DOI:** <https://doi.org/10.21203/rs.3.rs-31845/v2>

**License:**   This work is licensed under a Creative Commons Attribution 4.0 International License.

[Read Full License](#)

---

# Abstract

**Background** Resectability of liver metastasis is important to establish a treatment strategy for colorectal cancer patients. We aimed to evaluate the effect of distance from metastasis to the center of the liver on the resectability and patient outcomes after hepatectomy.

**Methods** Clinical data of a total of 124 patients who underwent hepatectomy for colorectal cancer with liver metastasis were retrospectively reviewed. We measured the minimal length from metastasis to the bifurcation of the portal vein at the primary branch of the Glissonean tree and defined it as “Centrality”. Predictive effects on positive resection margin and overall survival of centrality were statistically analyzed.

**Results** The value as a predictive factor for the positive resection margin of centrality was analyzed by the receiver operating characteristic curve (area under the curve = 0.72,  $P < 0.001$ ) and centrality  $\leq 1.5$  cm was an independent risk factor the positive resection margin in multivariate analysis. Total number of metastases  $\geq 3$  and centrality  $\leq 1.5$  cm were significant risk factors of overall survival after Cox regression analysis. Patients with these two risk factors ( $n=21$ ) had worse 5-year overall survival (10.7%) than patients with one ( $n=35$ , 58.3%) or no risk factor ( $n=68$ , 69.2%).

**Conclusion** Centrality was related with the positive resection margin of deeply located liver metastasis. Centrality should be considered to establish the surgical strategy for patients with advanced colorectal cancer with liver metastasis.

## Introduction

Although curative resection of isolated hepatic metastasis improves long-term survival of patients with colorectal cancer, only 20–30% of colorectal liver metastasis (CRLM) is resectable at the time of diagnosis [1, 2]. Several trials assessing the expansion of liver resectability by staged resection or systemic chemotherapy to convert unresectable metastases into resectable metastases have been conducted [3, 4]. Generally, the tolerable liver remnant volumes are 20% in a normal liver, 30% in a liver with chemotherapy-induced injury, and 40% in cirrhotic liver [5]. Additionally, adequate resection means the removal of a negative margin in all viable tumors with adequate vascular inflow and outflow and biliary drainage [6]. Thus, a multidisciplinary setting is required to establish treatment strategies for CRLM to achieve favorable oncologic outcomes and preserve remnant liver function after curative resection.

Vascular proximity to the central vessels of the liver is an important factor in selecting a proper treatment strategy. With metastasis occasionally observed near the central vessels, a large-volume resection is required, and subsequently, complete resection is considered difficult. Thus, this study aimed to evaluate the effect of distance from metastasis to the center of the liver on the resectability of metastases and to demonstrate its association with patients' survival.

## Materials And Methods

### *Data collection*

We retrospectively reviewed the medical records of patients who underwent hepatectomy for CRLM in our institution from January 2012 to December 2017. Patients who were diagnosed with colorectal adenocarcinoma with liver-only metastasis and underwent resections with curative intent were included. Patients with other liver diseases, including liver cirrhosis or hepatocellular cell carcinoma, were excluded. The clinical characteristics of patients, such as age at diagnosis of the primary cancer, sex, primary cancer location, and TN categories, were investigated. Characteristics associated with liver metastasis, such as carcinoembryonic antigen (CEA) level at the diagnosis of liver metastasis, neoadjuvant chemotherapy (NAC), and the longest diameter, total number, and lobar involvement of liver metastases, were also investigated. Operative factors associated with hepatectomy, such as positive resection margin, length of the closest resection margin, intraoperative radiofrequency ablation (IORFA), and synchronous resection with primary tumor, were investigated. Positive resection margin was defined as the presence of malignant tumor cells on the border of the specimen's resection margin after hepatectomy.

### *Decision-making in a multidisciplinary team*

All the treatment strategies for these patients were established by a multidisciplinary team (MDT) in our institution. The MDT for CRLM comprised a group of experienced specialists from medical oncology, colorectal and hepatobiliary surgery, radiation oncology, pathology, and diagnostic and interventional radiology. The resectability of CRLM was assessed by two experienced hepatobiliary surgeons in MDT. IORFA was performed by radiologic interventionists in MDT only for the tumor located at the periphery of the liver and with no abutting to any vascular structure and less than 2cm in size.

### *Centrality*

“Centrality” was defined as the minimal length from metastasis to the center of the liver, the bifurcation of the portal vein at the primary branch of the Glissonean tree. To measure the centrality, a process of three-dimensional (3D) interpolation from two-dimensional (2D) computed tomography (CT) or magnetic resonance (MR) images was performed using a 3- or 5-mm slice thickness (a gap between two serial images) of CT or MR images (Figure 1). For 3D interpolation, three vertices of A, B, C were required (A = bifurcation of the portal vein, B = point where the liver lesion began to appear, C = projection of A on the axial image where B appears). First, the first axial image of A and the second axial image of B were identified. The length on the Y-axis was measured as the distance between the two images, a multiple (n) of slice thickness and the length between B and C ( $\omega$ ) can be measured on the same axial image. Thus, centrality was calculated using the Pythagorean theorem as follows:

$$\text{Centrality}^2 = (\text{slice thickness} \times n)^2 + \omega^2$$

In case of multiple metastatic lesions, centralities were calculated for all lesions close to the hilum, and the minimum value was selected and defined as a final centrality for that case.

### *Statistical analyses*

Data were analyzed using the Statistical Package for the Social Sciences (SPSS) software version 20.0 (International Business Machines Corporation SPSS, Chicago, IL, USA). Discrete values, such as sex, TN categories, and primary tumor locations, were compared using the Pearson's  $\chi^2$  test. Student's t-test was used to compare continuous values such as age, CEA level, and total number and longest diameter of liver metastases. To compare the predictive value of centrality for positive resection margin after hepatectomy, the receiver operating characteristic (ROC) curve analysis and multivariate analysis using the logistic regression model were used. Overall survival (OS), from diagnosis of liver metastasis to death or the last follow-up, was analyzed using the Kaplan-Meier method and log-rank test. Cox proportional hazards regression analysis was used to evaluate the risk factors for OS. A two-sided P value < 0.05 was considered statistically significant.

## **Results**

A total of 124 patients were included in this study (Table 1). The median age at diagnosis of primary tumor was 60 years (interquartile range [IQR], 54–70), and the present study predominantly comprised men (n=77, 62.1%). Left colon and rectal cancer were more frequently observed than right colon cancer (42.7%, 37.9%, and 19.4%, respectively). Eighty-one (65.3%) patients were simultaneously diagnosed with primary cancer and liver metastasis. The median length of liver metastasis was 2 cm (IQR, 1.2–4), and the median number of total liver metastasis was 2 (IQR, 1–4). The median centrality was 4.4 cm (IQR, 2.3–6.0), and 56 (45.2%) patients showed metastasis on both the right and left lobes. Neoadjuvant chemotherapy was performed in 42 (33.9%) patients, and among them, 83.3% received a combination of neoadjuvant chemotherapy with a target agent bevacizumab or cetuximab. Synchronous resections with primary tumor were performed in 73 (58.9%) patients, and 16 (12.9%) patients received IORFA. The median length of the closest resection margin of patients with negative resection margin was 0.4 cm (IQR, 0.1–0.8), and the proportion of patients with negative resection margin was 71.0% (n=88).

The distribution of centrality is shown in Figure 2(a), and a total of 25 (20.1%) patients had centrality  $\leq$  1.5 cm. The ability of centrality to predict positive resection margin was analyzed using the ROC curve (area under the curve=0.72,  $P<0.001$ , Figure 2(b)). Selecting cut-off value from ROC curve was not possible due to the optimal value with high sensitivity and specificity could not be obtained from ROC curve. We set the cut-off value of centrality as 1.5cm according to opinions of hepatobiliary surgeons in our institution. More major hepatectomies were performed in the group of patient with centrality  $\leq$  1.5cm than the group of patient with centrality > 1.5cm (9 (36.0%) vs 20 (20.2%)), but there was no statistical

difference ( $P=0.095$ , Supplement Table 1). In the multivariate analysis, centrality  $\leq 1.5$  cm and bilobar involvement were a significant risk factor for positive resection margin (Supplement Table 2). The characteristics of the two groups, patients with centrality  $> 1.5$  cm ( $n=99$ ) and patients with centrality  $\leq 1.5$  cm ( $n=25$ ), were compared, and the results are shown in Table 2. The length of the longest diameter, total number of metastasis, proportion of bilobar involvement, and NAC were higher in patients with centrality  $\leq 1.5$  cm than those in patients with centrality  $> 1.5$  cm. The proportion of positive resection margin after hepatectomy was also higher in patients with centrality  $\leq 1.5$  cm than that in patients with centrality  $> 1.5$  cm. Moreover, the resection margin in patients with centrality  $\leq 1.5$  cm was closer than that in patients with centrality  $> 1.5$  cm.

The median follow-up period of all patients was 39.2 months (min: 5.9, max: 145.8). According to a multivariate analysis, risk factors related with OS were the number of liver metastasis  $> 3$  and centrality  $\leq 1.5$  cm (Table 3). All patients were classified based on the number of the risk factors observed, and the distribution of patients is shown in Figure 3. Z0 (zone 0) represents patients with no risk factor (centrality  $> 1.5$  cm and total number of metastasis  $< 3$  [ $n=68$ ]). Z1 represents patients with single risk factor (centrality  $\leq 1.5$  cm or total number of metastasis  $\geq 3$  [ $n=35$ ]), and Z2 represents patients with both risk factors (centrality  $\leq 1.5$  cm and total number of metastasis  $\geq 3$  [ $n=21$ ]). Figure 4 presents the Kaplan-Meier curves of OS of Z0, Z1, and Z2 (5-year OS: Z0, 69.2%; Z1, 58.3%; Z2, 10.7%). Z2 showed worst OS among the three groups, and statistically significant differences in OS among the three groups were observed (Z0 and Z1,  $P=0.026$ ; Z1 and Z2,  $P=0.016$ ; Z0 and Z2,  $P<0.001$ ). Additionally, patients in Z0–Z2 were stratified based on NAC, and their differences in OS between subgroups were analyzed (Figure 5). After performing a subgroup analysis, patients receiving NAC in Z2 showed better OS than patients not receiving NAC in Z2 (3-year OS: 65.3% vs. 0%,  $P=0.017$ ). However, there was no statistically significant difference in OS between subgroups of Z0 and Z1 (Z0: 5-year OS, 70.2% vs. 72.9%,  $P=0.166$ ; Z1: 5-year OS, 65.0% vs. 51.5%,  $P=0.396$ ).

## Discussion

This study is considered significant considering that centrality, which is defined as the proximity with the liver hilum at the portal bifurcation of liver metastasis, was proven to be another morphologic factor in deciding the resectability of CRLM. During the resection of liver metastasis, how to optimally solve two conflicting issues of preserving maximum liver function and securing sufficient resection margin after hepatectomy should be significantly considered. Traditionally, volumetric parameters such as tumor size and number of CRLM have been shown to be significant factors associated with resectability and prognosis after resection [7, 8]. However, the concept of resectability of CRLM has been changed over the last decade with the introduction of several treatments. Surgical techniques, such as portal vein embolism (PVE) or two-stage hepatectomy (TSH) are proven to be safe and curative in selective cases of

advanced CRLM with inadequate future liver remnant (FLR) or underlying liver diseases [9, 10]. Combined resection with IORFA is beneficial in preserving FLR with favorable oncologic outcomes [11, 12]. Additionally, NAC with a target agent and rescue surgery for unresectable CRLM results in tumor downsizing with survival benefits [13]. Considering that the benefits of these treatments have been proven, active resections have been widely performed, and the criteria for resectability have focused on how to perform R0 resection with sufficient liver function [7]. Thus, factors associated with positive resection margin should be comprehensively studied to define the resectability of CRLM.

Although the standard width of the resection margin of CRLM remains unknown, traditionally, resection margin of 1 cm has been accepted as a minimal margin to obtain favorable oncologic outcomes [14]. Certainly, R1 resection, i.e., microscopic identification of malignant cells on the resection margin, is a negative factor for survival after resection based on previous reports [15]. However, other studies reported that the width of surgical margin, even < 1 cm, does not affect survival in patients with R0 resection [16]. Thus, hepatic resection tends to be actively performed even when the length of the resection margin is < 1 cm on imaging modality. However, all hepatectomy for CRLM should be performed with efforts to achieve sufficient resection margin length even for the centrally-located tumors.

Troisi et al. suggested several factors in affecting the difficulty of hepatectomy, such as previous hepatectomy, disease type (hepatocellular carcinoma, benign or metastatic lesion), liver function, and tumor size and number [17]. They also reported that performing hepatectomy is highly difficult when the tumor is located in deeper layers or proximal to the branch of the Glissonean tree. As the tumor is closer to the primary branch of the Glissonean tree, the hilar dissection or anatomical resection is required to completely resect the tumor. The centrality in this study was at the bifurcation of the primary Glissonean branch; thus, we could assume that the centrality was a factor indicating the difficulty of hepatectomy to secure sufficient resection margin near the center of the liver.

Subsequently, how should we approach CRLM with high centrality? Per et al. demonstrated that associating liver partition and portal vein ligation (ALPPS) made major resections possible with improved resection rates compared with conventional TSH [18]. They also reported the rate of severe complications and negative resection margins were comparable with TSH. Enhanced one-stage hepatectomy (E-OSH) is another technique that makes major hepatectomy possible especially for deep-located nodules [19, 20]. E-OSH utilizes the intraoperative ultrasound and vascular manipulations to achieve complete vascular control and detachment of liver metastasis from intrahepatic vascular structures. This approach was reported to be helpful to overcome the disease progression between the two hepatectomies which was a major drawback of TSH and also showed a comparable survival with ALPPS. These approaches, ALPPS and E-OSH, should be considered for advanced liver metastasis with high centrality. Additionally, recent studies reported a greater conversion rate of initially unresectable CRLM after NAC with regimens of oxaliplatin, irinotecan, and other target agents [21]. Thus, NAC should be considered in patients with high centrality, so that hepatectomy can be performed after securing sufficient resection margin at the central side. However, prolonged chemotherapy may also have a detrimental effect on the hepatic parenchyma due to oxaliplatin-induced sinusoidal obstruction or irinotecan-induced steatosis [22, 23]. Hence, liver

function and remnant liver volume should be carefully monitored during chemotherapy. Taken together, various treatment tools should be considered when establishing appropriate strategies for CRLM with high centrality, and treatments should be customized based on patient's condition.

This study has several limitations. First, the definition of the center of the liver, bifurcation of the portal vein at the first branch of the Glissonian tree, was subjective and did not include other important structures. For example, proximity to the inferior vena cava is an important factor in deciding adequate vascular outflow, but it was not considered in this study. Second, the factor of centrality  $\leq 1.5$  cm was selected empirically according to the opinion of hepatobiliary surgeons in our MDT team. According to the ROC curve analysis, the specificity for positive resection margin of centrality  $\leq 1.5$  cm was 91.9%. However, the sensitivity was only 30.6%, suggesting that factors other than centrality should be considered when deciding the positive resection margin of CRLM. Third, this study was retrospectively conducted based on previous pathologic reports, which contained only the shortest length of resection margin of specimens and did not indicate whether the resection margin was of the central side. Fourth, although the difference was not statistically significant, the difference of the long-term overall survival between of - NAC group in Z1 and that of + NAC group in Z1 was relatively significant. This might be caused by the small volume of subgroups of Z1 (number of NAC- in Z1 = 13 and number of NAC+ in Z1 = 22). Fifth, the reason why the centrality is related with a worse survival was not definite. More major hepatectomy were performed in the high centrality group and major hepatectomy is related with post-hepatic liver failure [24]. We carefully assumed that the liver function of the high centrality group might be deteriorated due to complications after major hepatectomy such as liver cirrhosis or biliary fistula in the mid or long term period of follow-up and treatments such as repeat resections for recurrent hepatic metastasis or compliance to the chemotherapy might be affected to survival of patients. However, the tumor burden of the high centrality group seemed to be higher than that of the low centrality group, thus the heterogeneity in tumor burden might affect the results of survival of patients with high centrality.

## Conclusion

Centrality, proximity to the central vessels of liver metastasis, had a potency in predicting the positive resection margin and a worse overall survival after hepatectomy for CRLM. Centrality might be beneficial in establishing treatment strategies especially for patients with advanced colorectal cancer liver metastasis.

## Abbreviations

CT: computed tomography; MR: magnetic resonance; CRLM: colorectal cancer with liver metastasis; CEA: carcinoembryonic antigen; NAC: neoadjuvant chemotherapy; IORFA: intraoperative radiofrequency ablation; MDT: multidisciplinary team; 3D: 3-dimensional; 2D: 2-dimensional; ROC: receiver operating characteristic; OS: overall survival; IQR: interquartile range ratio; Z1: zone1; Z2: zone2; Z3: zone3; PVE: portal vein embolization; TSH: two-stage hepatectomy; FLR: future liver remnant

## **Declarations**

### **Acknowledgements**

Not applicable.

### **Funding**

This work was supported by Institute for Information & Communications Technology Promotion (IITP) grant funded by the Korea Government (MSIT) (No. 2018-0-00861. Intelligent SW Technology Development for Medical Data Analysis)

### **Authors' contributions**

Jun Woo Bong: collection, analysis and interpretation of data, writing paper; Sun Il Lee: final approval of the manuscript version; Yeonuk Ju, Jihyun Seo and Sang Hee Kang: drafting of the work for intellectual content; Pyoung-Jae Park, Sae-Byeol Choi, Sang Cheul Oh and Byung Wook Min: design and conception of the work.

### **Ethics approval and consent to participate**

The Institutional Review Board of Korea University Guro Hospital approved this study with a waiver of informed consent (Approval Number: 2018GR0395).

### **Consent for publication**

Not applicable

### **Conflict of interest**

There is no conflict of interest in this study.

### **Data Availability Statement**

The data that support the findings of this study are available from the corresponding author upon reasonable request.

## **References**

1. Jönsson K, [Gröndahl G](#), Salö M, et al. Repeated liver resection for colorectal liver metastases: a comparison with primary liver resections concerning perioperative and long-term outcome. *Gastroenterol Res Pract*. 2012;2012:568214.
2. Simmonds PC, Primrose JN, Colquitt JL, et al. Surgical resection of hepatic metastases from colorectal cancer: a systematic review of published studies. *Br J Cancer*. 2006;94:982-999.
3. Matsuoka H, Morise Z, Tanaka C, et al. Repeat hepatectomy with systemic chemotherapy might improve survival of recurrent liver metastasis from colorectal cancer—a retrospective observational study. *World J Surg Oncol*. 2019;17:33.
4. Nanji S, Tsang ME, Wei X, Booth CM. Outcomes after repeat hepatic resection for recurrent metastatic colorectal cancer: A population-based study. *Am J Surg*. 2017;213:1053-1059.
5. Abdalla EK, Denys A, Chevalier P, et al. Total and segmental liver volume variations: implications for liver surgery. *Surgery*. 2004;135:404-410.
6. Charnsangavej C, Clary B, Fong Y, et al. Selection of patients for resection of hepatic colorectal metastases: expert consensus statement. *Ann Surg Oncol*. 2006;13:1261-1268.
7. Pawlik TM, Schulick RD, Choti MA. Expanding criteria for resectability of colorectal liver metastases. *Oncologist*. 2008;13:51-64.
8. Sasaki K, Morioka D, Conci S, et al. The Tumor Burden Score: a new “metro-ticket” prognostic tool for colorectal liver metastases based on tumor size and number of tumors. *Ann Surg*. 2018;267:132-141.
9. Farges O, [Belghiti J](#), [Kianmanesh R](#), et al. Portal vein embolization before right hepatectomy: prospective clinical trial. *Ann Surg*. 2003;237:208-217.
10. Jaeck D, [Oussoultzoglou E](#), [Rosso E](#), et al. A two-stage hepatectomy procedure combined with portal vein embolization to achieve curative resection for initially unresectable multiple and bilobar colorectal liver metastases. *Ann Surg*. 2004;240:1037-1049; discussion 1049-1051.
11. Leung EY, [Roxburgh CS](#), [Leen E](#), [Horgan PG](#). Combined resection and radiofrequency ablation for bilobar colorectal cancer liver metastases. *Hepatogastroenterology*. 2010;57:41-46.
12. Sasaki K, [Margonis GA](#), [Andreatos N](#), et al. Combined resection and RFA in colorectal liver metastases: stratification of long-term outcomes. *J Surg Res*. 2016;206:182-189.
13. Ye LC, [Liu TS](#), [Ren L](#), et al. Randomized controlled trial of cetuximab plus chemotherapy for patients with KRAS wild-type unresectable colorectal liver-limited metastases. *J Clin Oncol*. 2013;31:1931-1938.
14. Huiskens J, Bolhuis K, Engelbrecht MRW, et al. Outcomes of resectability assessment of the Dutch Colorectal Cancer Group Liver Metastases Expert Panel. *J Am Coll Surg*. 2019;229:523-532.e2.
15. Altendorf-Hofmann A, Scheele J. A critical review of the major indicators of prognosis after resection of hepatic metastases from colorectal carcinoma. *Surg Oncol Clin N Am*. 2003;12:165-192, xi.
16. Pawlik TM, Scoggins CR, Zorzi D, et al. Effect of surgical margin status on survival and site of recurrence after hepatic resection for colorectal metastases. *Ann Surg*. 2005;241:715-724.

17. Troisi RI, [Montalti R](#), [Van Limmen JG](#), et al. Risk factors and management of conversions to an open approach in laparoscopic liver resection: analysis of 265 consecutive cases. *HPB*. 2014;16:75-82.
18. Sandström, P, [Røsok BI](#), [Sparrelid E](#), et al., ALPPS Improves Resectability Compared With Conventional Two-stage Hepatectomy in Patients With Advanced Colorectal Liver Metastasis: Results From a Scandinavian Multicenter Randomized Controlled Trial (LIGRO Trial). *Annals of Surgery*, 2018. 267:833-840.
19. [Torzilli G](#), [Serenari M](#), [Viganò L](#), et al., Outcomes of enhanced one-stage ultrasound-guided hepatectomy for bilobar colorectal liver metastases compared to those of ALPPS: a multicenter case-match analysis. *HPB (Oxford)*, 2019. 21:1411-1418.
20. [Torzilli G](#), [Viganò L](#), [Cimino M](#), et al., Is Enhanced One-Stage Hepatectomy a Safe and Feasible Alternative to the Two-Stage Hepatectomy in the Setting of Multiple Bilobar Colorectal Liver Metastases? A Comparative Analysis between Two Pioneering Centers. *Dig Surg*, 2018. 35:323-332.
21. Nozawa H, [Ishihara S](#), [Kawai K](#), et al. Conversion to resection in patients receiving systemic chemotherapy for unresectable and/or metastatic colorectal cancer-predictive factors and prognosis. *Clin Colorectal Cancer*. 2018;17:e91-e97.
22. Pawlik TM, Olin K, Gleisner AL, et al. Preoperative chemotherapy for colorectal liver metastases: impact on hepatic histology and postoperative outcome. *J Gastrointest Surg*. 2007;11:860-868.
23. Rubbia-Brandt L, Audard V, Sartoretti P, et al. Severe hepatic sinusoidal obstruction associated with oxaliplatin-based chemotherapy in patients with metastatic colorectal cancer. *Ann Oncol*. 2004;15:460-466.
24. Kauffmann R, Fong Y, Post-hepatectomy liver failure. *Hepatobiliary Surgery and Nutrition*, 2014. 3: 238-246.

## Tables

Table 1. Clinicopathologic characteristics of colorectal cancer in patients with liver metastasis

Characteristics	Total N = 124
	No. (%)
<b>Patient characteristics</b>	
Age, years, median (IQR)	60 (54-70)
Sex, male	77 (62.1)
<b>Primary tumor characteristics</b>	
<b>Location</b>	
Right colon	24 (19.4)
Left colon	53 (42.7)
Rectum	47 (37.9)
<b>T categories</b>	
1	3 (2.4)
2	8 (6.5)
3	83 (66.9)
4	26 (21.0)
<b>N categories</b>	
0	36 (29.0)
1	72 (58.1)
2	12 (9.7)
<b>Liver metastasis</b>	
Synchronous	81 (65.3)
CEA, median (IQR)	6 (2.5-28.5)
Diameter, cm, median (IQR)	2 (1.25-4)
Number, median (IQR)	2 (1-4)
Bilobar involvement	56 (45.2)
NAC	42 (33.9)
Target agent	35 (83.3)
Centrality, cm, median (IQR)	4.4 (2.3-6.0)
<b>Operative factors</b>	
R0 resection	88 (71.0)
IORFA	16 (12.9)
Resection margin, cm, median (IQR)*	0.4 (0.1-0.8)
Synchronous resection	73 (58.9)
<b>Postoperative factors</b>	
Adjuvant chemotherapy	119 (96)

---

CEA: carcinoembryonic antigen, NAC: neoadjuvant chemotherapy before liver resection, IORFA: intraoperative radiofrequency ablation, \* : the median length of resection margin of patients with negative resection margin

Table 2. Comparison of characteristics after subgroup classification by centrality

Table 3. Univariate and multivariate analyses of overall survival

Characteristics	Centrality > 1.5 cm	Centrality ≤ 1.5 cm	P
	Total N = 99	Total N = 25	
	No. (%)	No. (%)	
<b>Patient characteristics</b>			
Age, years, mean±SD	60.9±11.2	61.5±9.8	0.813 <sup>a</sup>
Sex, male	61 (61.6)	16 (64.0)	0.826 <sup>b</sup>
<b>Primary tumor characteristics</b>			
<b>Location</b>			0.429 <sup>b</sup>
Right colon	19 (19.2)	5 (20.0)	
Left colon	45 (45.5)	8 (32.0)	
Rectum	35 (35.4)	12 (48.0)	
<b>T categories</b>			0.729 <sup>b</sup>
1	2 (2.1)	1 (4.0)	
2	7 (7.4)	1 (4.0)	
3	64 (67.4)	19 (76.0)	
4	22 (23.2)	4 (16.0)	
<b>N categories</b>			0.919 <sup>b</sup>
0	29 (30.5)	7 (28.0)	
1	57 (60.0)	15 (60.0)	
2	9 (9.5)	3 (12.0)	
<b>Liver metastasis</b>			
Synchronous	61 (61.6)	20 (80.0)	0.084 <sup>b</sup>
CEA, mean±SD	83.7±320.6	79.8±127.4	0.954 <sup>a</sup>
Diameter, cm, mean±SD	2.9±2.5	5.5±4.8	0.015 <sup>a</sup>
Number, mean±SD	2.8±3.4	7.3±6.9	0.004 <sup>a</sup>
Bilobar involvement	35 (35.4)	21 (84.0)	<0.001 <sup>b</sup>
NAC	26 (26.3)	16 (64.0)	<0.001 <sup>b</sup>
<b>Operative factors<sup>c</sup></b>			
Positive resection margin	22 (22.2)	13 (54.2)	0.002 <sup>b</sup>
Resection margin, cm, mean±SD	0.7±0.8	0.2±0.2	<0.001 <sup>a</sup>
IORFA	11 (11.1)	5 (20.0)	0.236 <sup>b</sup>
Synchronous resection	56 (56.6)	17 (68.0)	0.299 <sup>b</sup>
<b>Postoperative factors</b>			
Adjuvant chemotherapy	95 (96.0)	24 (96.0)	0.993 <sup>b</sup>

CEA: carcinoembryonic antigen, NAC: neoadjuvant chemotherapy before liver resection, IORFA:

intraoperative radiofrequency ablation, SD: standard deviation, a: analyzed by Student's t-test method, b: analyzed by Pearson's  $\chi^2$  test method, c: factors related with hepatectomy

Factors	Univariate analysis			Multivariate analysis		
	HR	95% CI	P	HR	95% CI	P
<b>Age, years</b>						
< 65	1					
≥ 65	1.092	0.570, 2.091	0.791			
<b>Sex</b>						
Female	1					
Male	1.808	0.851, 3.841	0.123			
<b>Location of primary tumor</b>						
Right colon	1		0.433			
Left colon	0.668	0.299, 1.493				
Rectum	0.559	0.227, 1.381				
<b>T categories</b>						
1,2	1		0.228			
3,4	1.48	0.782, 2.799				
<b>N categories</b>						
0	1		0.692			
1,2	1.154	0.569, 2.342				
<b>Synchronous metastasis</b>						
No	1					
Yes	1.235	0.627, 2.431	0.542			
<b>CEA</b>						
<5	1					
≥ 5	1.908	0.953, 3.818	0.068			
<b>Size</b>						
< 5 cm	1					
≥ 5 cm	1.667	0.829, 3.354	0.152			
<b>Number</b>						
< 3	1					
≥ 3	3.49	1.775, 6.865	<0.001	2.688	1.290, 5.602	0.008
<b>Lobar involvement</b>						
Unilobar	1					
Bilobar	2.979	1.524, 5.822	0.001	1.336	0.462, 3.864	0.593
<b>Centrality</b>						
> 1.5 cm	1					

≤ 1.5 cm	3.713	1.815, 7.595	<0.001	2.272	1.033, 4.999	0.041
<b>Resection margin</b>						
Negative						
Positive	2.243	1.060, 4.746	0.035	1.428	0.646, 3.154	0.379
<b>Combined resection of primary tumor</b>						
No	1					
Yes	1.433	0.736, 2.791	0.29			
<b>Adjuvant chemotherapy</b>						
No	1					
Yes	1.514	0.207, 11.071	0.683			

CEA: carcinoembryonic antigen, HR: hazard ratio, CI: confidence interval

Supplement Table 1. Surgical details of hepatectomy

	Centrality > 1.5cm	Centrality ≤ 1.5 cm
	n = 99 (%)	n = 25 (%)
Major hepatectomy*	20 (20.2)	9 (36.0)
Right hepatectomy	9 (9.1)	2 (8.0)
Right hepatectomy + WR	2 (2.0)	2 (8.0)
Right hepatectomy + PVE	1 (1.0)	
Right hepatectomy + WR + PVE	1 (1.0)	
Right trisectionectomy		1 (4.0)
Left hepatectomy	3 (3.0)	1 (4.0)
Central bisectionectomy	1 (1.0)	1 (4.0)
Central bisectionectomy + WR	1 (1.0)	1 (4.0)
ALPPS	1 (1.0)	
ALPPS + WR	1 (1.0)	1 (4.0)
Minor hepatectomy	79 (79.8)	16 (64.0)
Right anterior sectionectomy	1 (1.0)	
Right anterior sectionectomy + WR	1 (1.0)	
Right posterior sectionectomy	1 (1.0)	
Right posterior sectionectomy + WR	2 (2.0)	
Left lateral sectionectomy	4 (4.0)	
Left lateral sectionectomy + WR	4 (4.0)	2 (8.0)
Bisegmentectomy	4 (4.0)	
Bisegmentectomy + WR		1 (4.0)
Segementectomy	10 (10.1)	3 (12.0)
Segementectomy + WR	6 (6.1)	3 (12.0)
WR	46 (46.5)	7 (28.0)
Combined RFA	11 (11.1)	5 (20.0)

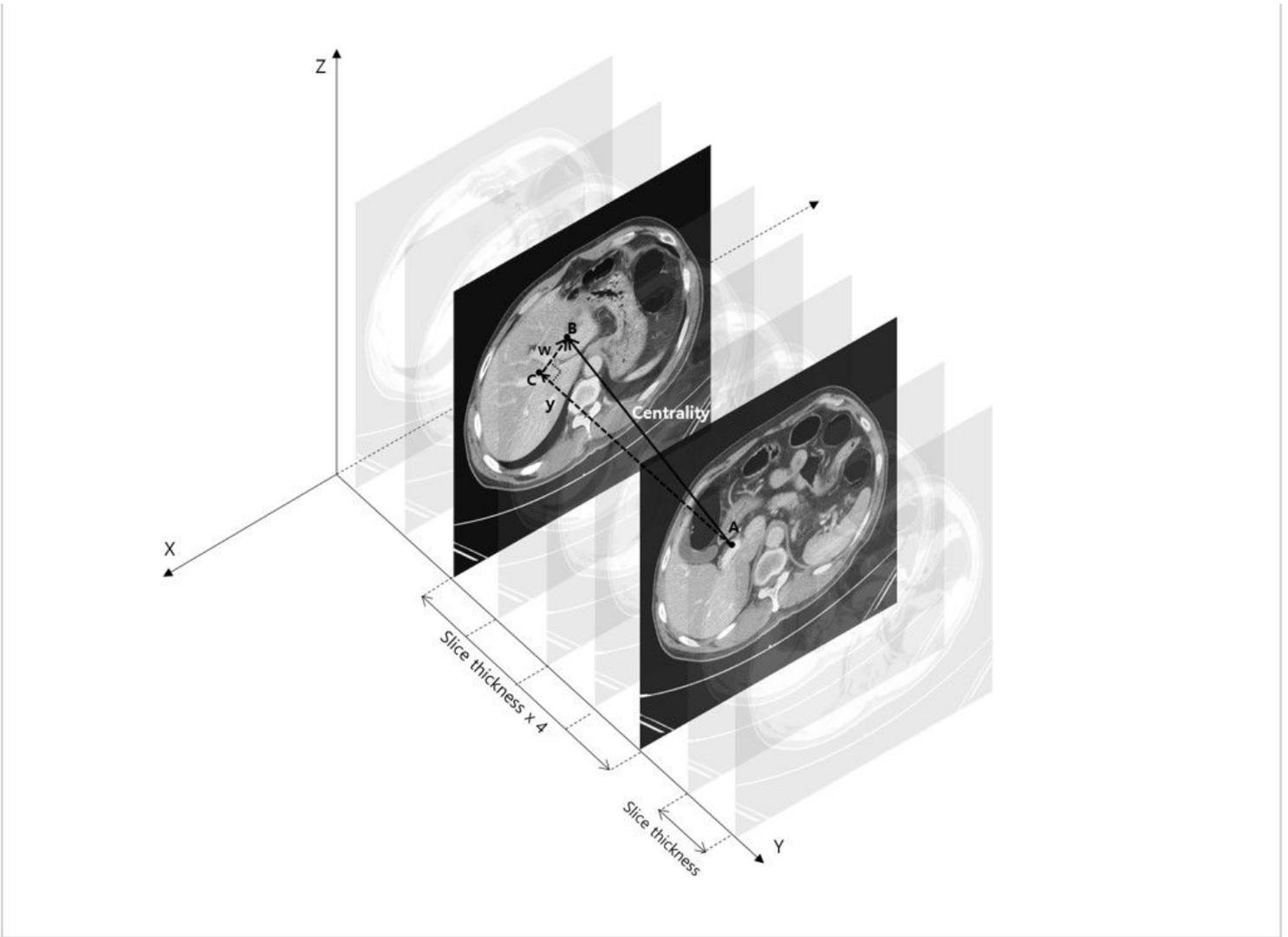
WR: wedge resection, PVE: portal vein embolization, \*: there was no statistical difference in the major hepatectomy rate between two groups (P=0.095).

Supplement Table 2. Univariate and multivariate analyses of positive resection margin

Factors	Univariate analysis			Multivariate analysis		
	OR	95% CI	P	OR	95% CI	P
<b>Age, years</b>						
< 65	1					
≥ 65	0.787	0.359, 1.725	0.550			
<b>Sex</b>						
Female	1					
Male	0.635	0.287, 1.406	0.263			
<b>Location of primary tumor</b>						
Right colon	1		0.731			
Left colon	0.743	0.251, 2.202				
Rectum	1.039	0.354, 3.052				
<b>T categories</b>						
1,2	1		0.925			
3,4	0.925	0.436, 2.012				
<b>N categories</b>						
0	1		0.683			
1,2	1.204	0.494, 2.931				
<b>Size</b>						
< 5 cm	1					
≥ 5 cm	1.392	0.554, 3.499	0.482			
<b>Number</b>						
< 3	1			1		
≥ 3	3.599	1.597, 8.111	0.002	1.735	0.499, 5.835	0.394
<b>Lobar involvement</b>						
Unilobar	1			1		
Bilobar	3.500	1.545, 7.929	0.003	2.487	1.026, 6.029	0.044
<b>Centrality</b>						
> 1.5 cm	1			1		
≤ 1.5 cm	4.455	1.774, 11.188	0.001	3.016	1.118, 8.138	0.029

CEA: carcinoembryonic antigen, OR: odds ratio, CI: confidence interval

## Figures



**Figure 1**

Measurement of centrality using three-dimensional interpolation from two-dimensional images of liver metastasis. Centrality was defined as the minimal length from liver metastasis to the center of the liver to the bifurcation of the portal vein at the primary branch of the Glissonian tree.  $Centrality^2 = y^2 + \omega^2$ ,  $y = \text{slice thickness} \times n$ ,  $\omega = \text{length between B and C}$ , A = bifurcation of the portal vein, B = point where the liver lesion began to appear, C = projection of A on the axial image where B appears

Figure 2 (a)

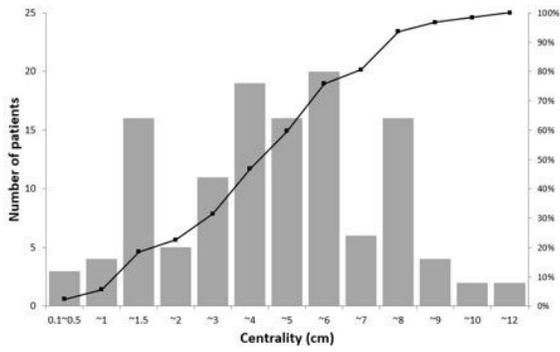
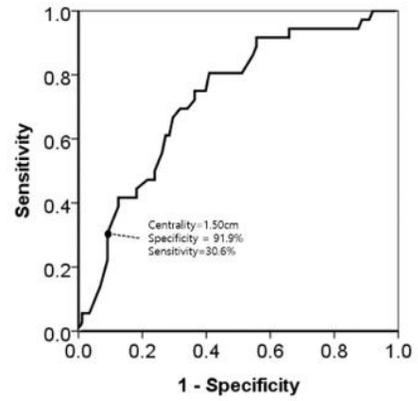
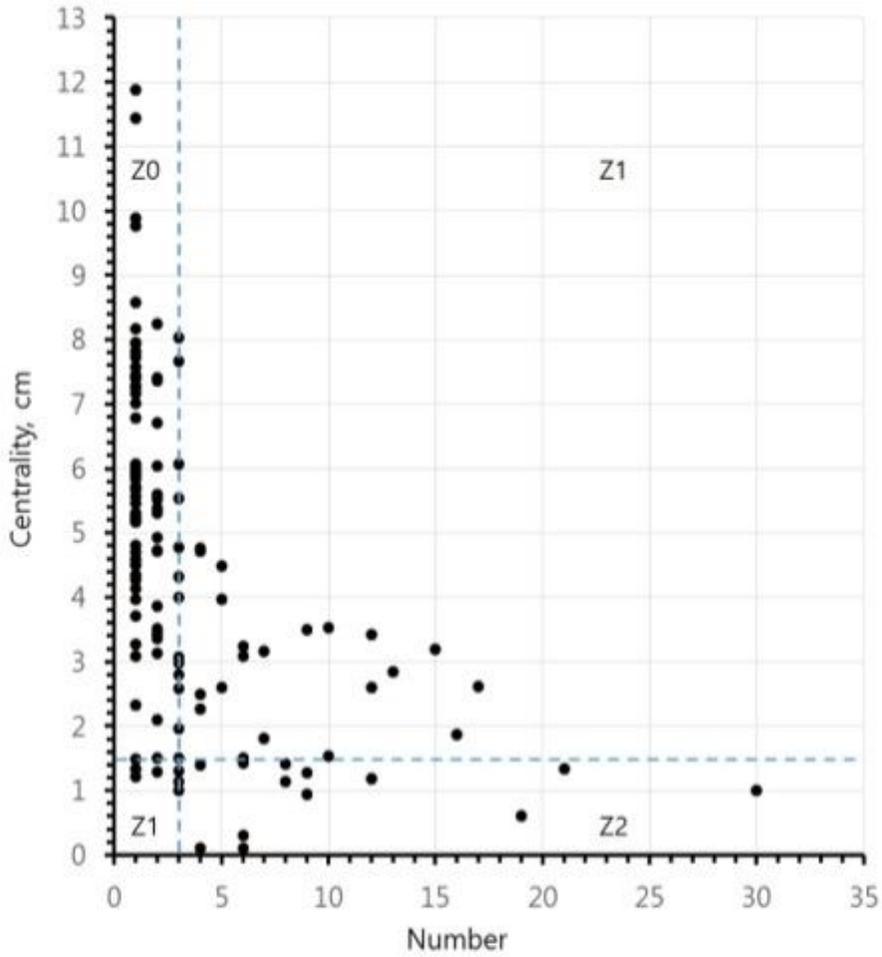


Figure 2 (b)



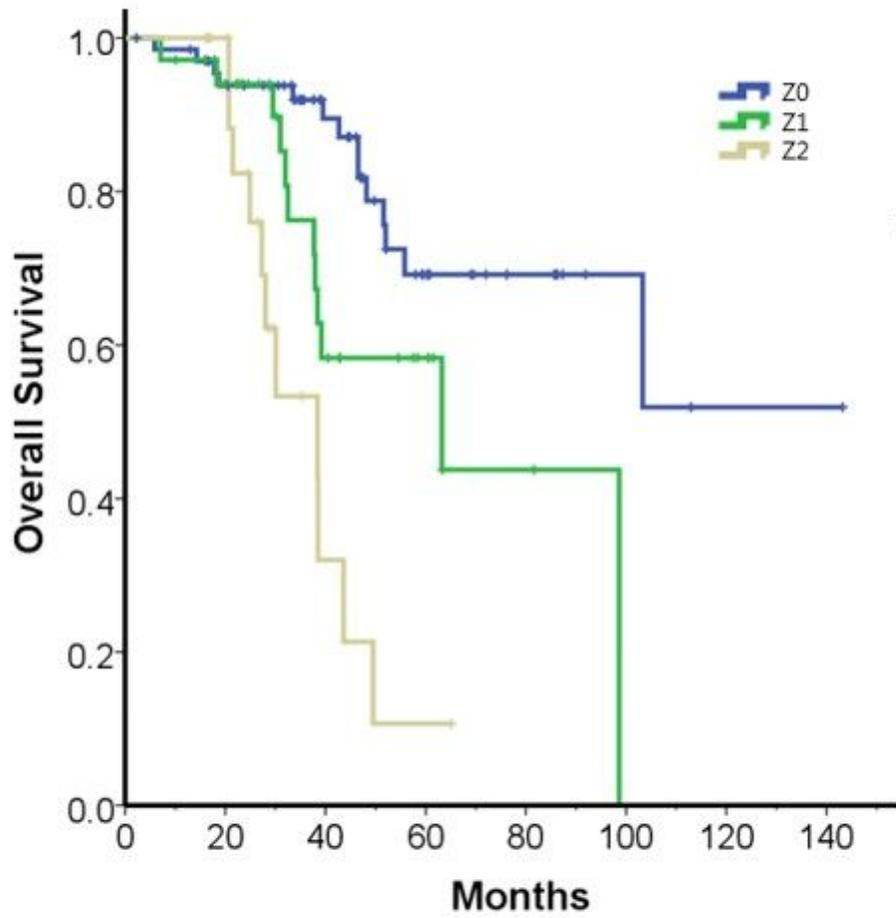
## Figure 2

(a) Distribution of centralities of all patients (b) Receiver operating characteristic curve analysis of centrality for positive resection margin. AUC = 0.720,  $P < 0.001$ , AUC: area under the curve



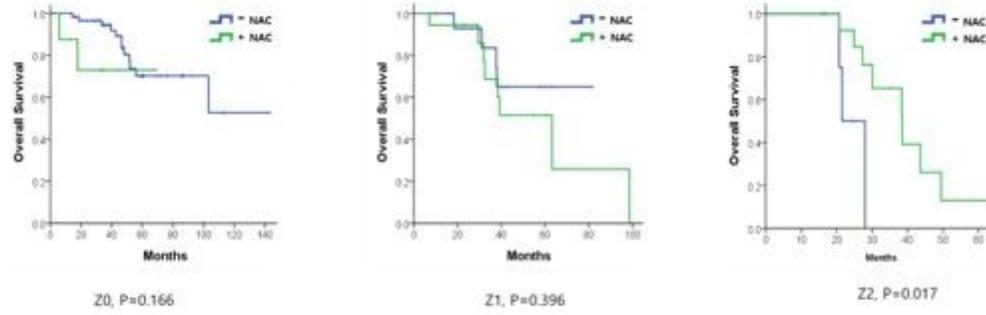
**Figure 3**

Distribution of patients according to centrality and number of liver metastases Z0: centrality > 1.5 cm and total number of metastases < 3, n = 68, Z1: centrality ≤ 1.5 cm or total number of metastases ≥ 3, n = 35, Z2: centrality ≤ 1.5 cm and total number of metastases ≥ 3, n = 21



**Figure 4**

Kaplan-Meier curves of overall survival of patients stratified by the number of risk factors observed. Significant differences in OS among the three groups were observed and Z2 showed worst OS (Z0 and Z1,  $P=0.026$ ; Z1 and Z2,  $P=0.016$ ; Z0 and Z2,  $P<0.001$ )



**Figure 5**

Kaplan-Meier curves of overall survival of subgroups stratified by neoadjuvant chemotherapy. Overall survival was statistically different only in Z2. P=0.166 in Z0, P=0.396 in Z1, and P=0.017 in Z2