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***Retrospective Study***

**Colorectal adenocarcinoma patients with M1a diseases receiving palliative primary tumor resection gained more clinical benefits than those with M1b diseases: A propensity score matching analysis**

Li CL *et al*. Palliative primary tumor resection in metastatic CRA

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**Abstract**

BACKGROUND

Surgical resection is regarded as the only potentially curative treatment option for patients with metastatic colorectal cancer (CRC). The National Comprehensive Cancer Network clinical practice guidelines do not recommend palliative surgery unless there is a risk of severe symptoms. However, accumulating evidence has shown that palliative surgery is associated with more favorable outcomes for patients with metastatic CRC.

AIM

To investigate the separate role of palliative primary tumor resection for patients with stage IVA (M1a diseases) and stage IVB (M1b diseases) of colorectal adenocarcinoma (CRA).

METHODS

CRA patients from 2010 to 2015 with definite M1a and M1b categories according to the 8th edition of American Joint Committee on Cancer staging system were selected from the Surveillance Epidemiology and End Results (SEER) database. To minimize potential selection bias, the data were adjusted with propensity score matching (PSM). Baseline characteristics, including gender, year of diagnosis, age, marital status, primary site, surgical information, race, grade, chemotherapy and radiotherapy, were recorded and analyzed. Univariate and multivariate analysis were performed to explore the separate role of palliative surgery for patients with M1a and M1b diseases.

RESULTS

A total of 19680 patients with metastatic CRA were collected from the SEER database, including 10399 cases of M1a diseases and 9281 cases of M1b diseases. Common independent prognostic factors for both M1a and M1b patients included year of diagnosis, age, race, marital status, primary site, grade, surgery and chemotherapy. After PSM adjustment, 3732 and 3568 matched patients in M1a and M1b group were included, respectively. Patients receiving palliative primary tumor resection had longer survival time than those without surgery (*P* < 0.001). For patients with M1a diseases, palliative resection could increase the median survival time by 9 mo; for patients with M1b diseases, palliative resection could prolong the median survival time by 7 mo. For M1a diseases, patients with lung metastasis had more clinical benefit from palliative resection than those with liver metastasis (15 mo for lung metastasis *vs* 8 mo for liver metastasis, *P* < 0.001).

CONCLUSION

CRA patients with M1a diseases receiving palliative primary tumor resection gained more clinical benefits than those with M1b diseases. Those patients with M1a (lung metastasis) had superior long-term outcomes after palliative primary tumor resection.

**Key words:** Colorectal adenocarcinoma; Palliative primary tumor resection; Distant metastasis; Propensity score matching; Surveillance Epidemiology and End Results; Overall survival

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**Core tip:** The National Comprehensive Cancer Network clinical practice guidelines did not recommend palliative surgery for metastatic colorectal adenocarcinoma (CRA). Using the Surveillance Epidemiology and End Results database, we found that patients with M1a diseases had a significant survival benefit compared to those with M1b diseases and patients with M1a (lung metastasis) got best long-term outcomes with median overall survival prolonged by 15 mo and with those without surgical treatment. These findings provided further evidence to support palliative surgical procedure to metastatic CRA and develop effective individualized treatment strategy.

**INTRODUCTION**

Colorectal cancer (CRC) is one of the three most common malignancies with 135430 individuals expected to be diagnosed in 2017 in the United States[1]. However, approximately 20% of new CRC patients are diagnosed with distant-stage tumors, resulting in poor long-term outcomes with a 5-year survival rate of 23.2%[2].

Surgical resection is regarded as the only potentially curative treatment option for this disease and could significantly improve the prognosis of patients with metastatic CRC[3]. Rees *et al*[4] reported that the 5-year cancer-specific survival (CSS) rate for metastatic CRC patients undergoing primary and hepatic resection was 36%. Abdalla *et al*[5]showed that patients receiving surgical resection of primary tumors and liver metastases had a 5-year survival rate of up to 58%. Similarly, curative surgical treatment could increase CRC patients with resectable lung metastasis 5-year survival rate of 32% to 68%[6,7]. Unfortunately, only one-fifth to one-quarter of metastatic CRC patients can receive curative surgical treatment[8], indicating that metastatic CRC patients were a heterogenous population.

According to the 8th edition of American Joint Committee on Cancer (AJCC) tumor-node-metastasis staging system, metastatic CRC are classified into M1a (metastasis confined to one organ or site) and M1b (metastases in more than one organ/site or the peritoneum). Complete resection is impossible for most metastatic CRC patients (especially for those with M1b diseases) even after neoadjuvant chemoradiation. The National Comprehensive Cancer Network clinical practice guidelines[9] do not recommend palliative surgery unless there is a risk of significant acute bleeding, obstruction, perforation or other severe symptoms based on comprehensive analysis of the literature[10-12]. However, accumulating evidence has shown that palliative surgery is associated with more favorable outcomes. For example, a pooled analysis including four randomized trials reported that patients receiving palliative primary tumor removal had prolonged overall survival (OS) compared with those not receiving operation.[13] Another population-based retrospective study reviewing 37793 metastatic CRC patients showed that palliative surgery was significantly related to better OS and CSS[14]. Finally, a systematic review consisting of 21 studies indicated that there was a survival benefit for palliative surgery in patients with metastatic CRC and criteria for palliative surgery should be extended on the basis of World Health Organization (WHO) performance status (PS) or tumor burden[15].

However, to our best knowledge, no studies have classified stage IV into subsets to assess the role of palliative surgery. Adenocarcinoma is the most common pathological type of CRC, accounting for approximately 90% of cases[16]. Thus, we subdivided colorectal adenocarcinoma (CRA) patient populations with stage IV disease on the basis of comorbidities from the Surveillance, Epidemiology, and End Results (SEER) database into stage IVA (M1a diseases) and stage IVB (M1b diseases). Outcomes of palliative surgery were then independently assessed.

**MATERIALS AND METHODS**

***Patient source***

Patient data, originating between 2010 and 2015, was collected from the SEER database, one of the largest cancer databases in the world[14]. The selection criteria were as follows: (1) patients 18 years old or older; (2) disease histologically diagnosed as adenocarcinoma; (3) treated for first primary tumor; (4) definite M1a or M1b diseases according to the 8th edition of AJCC staging system; (5) no surgery for metastatic sites (including distant lymph nodes); (6) surgical procedure or no surgical procedure to primary tumor (excluding tumor destruction or no pathologic specimen or unknown whether there was a pathologic specimen); and (7) active follow-up. Cases with unknown survival time, status or those coded as 0 mo were excluded. The entire cohort was divided into two groups based on the median age and calculation result of X-tile program (Yale University, 3.6.1, Supplementary Figure 1). After propensity score matching (PSM), 2935 patients with M1a diseases and 2145 patients with M1b diseases were excluded owing to lack of counterpart propensity scores. In survival analysis for M1a (liver metastasis) and M1a (lung metastasis), 2202 and 267 patients were further excluded, respectively. Follow-up time ranged from 1 to 71 mo.

***Statistical analyses***

Baseline characteristics of metastatic CRA patients, including sex, year of diagnosis, age, marital status, primary site, surgical information, race, grade, chemotherapy and radiotherapy, were recorded and analyzed by *χ2* test. The patient prognosis was assessed using univariate and multivariate Cox regression analysis with hazard ratios and 95% confidence interval. To minimize potential selection bias, 1:1 ratio PSM without replacement were used to investigate the effect of palliative primary tumor resection on metastatic CRA. After PSM adjustment, Kaplan-Meier methods were employed to analyze the OS for M1a and M1b patients. All statistical analyses were conducted using IBM SPSS Statistics 22.0. *P*-values < 0.05 were considered statistically significant.

**RESULTS**

***Baseline characteristics***

A total of 19680 patients with metastatic CRA were collected from the SEER database, which included 10399 cases of M1a diseases and 9281 cases of M1b diseases (Table 1). The entire cohort consisted of 11107 (56.4%) males and 8573 (43.6%) females with median age of 63 years (ranging from 18 to 108). Most patients were of White ethnicity (74.8%) and more than half of them had well or moderately differentiated tumors (grade I + II). Next, 15476 (78.6%) cases of primary tumors were located in colon and 4204 (21.4%) in the rectum. The prevalence of metastatic CRA between 2010 and 2012 was similar to that between 2013 and 2015. However, M1b diseases seemed to account for a larger proportion from 48.4% during 2010-2012 to 51.6% during 2013-2015 while M1a diseases showed the opposite prevalence trend.

Of the entire cohort, 14057 (71.4%) metastatic CRA patients received chemotherapy and 975 (5.0%) received radiotherapy. 9360 (47.6%) metastatic CRA patients received palliative primary tumor resection while 10320 (52.4%) did not. The proportion of patients with M1a diseases undergoing surgical procedure was much higher than that of M1b diseases (55.6% for M1a and 38.5% for M1b).

***Survival analysis***

Univariate and multivariate Cox regression analysis for OS for both M1a and M1b patients were performed (Table 2). The common independent prognostic factors in both M1a and M1b patients included year of diagnosis (2010-2012 *vs* 2013-2015), age (< 65 *vs* ≥ 65), race (white *vs* black), marital status (married *vs* others), primary site (colon *vs* rectum), grade (I + II *vs* III + IV) and surgery (yes *vs* no) and chemotherapy (yes *vs* no/unknown). Radiotherapy (yes *vs* no) was an independent prognostic factor for M1a patients but not for M1b patients.

After PSM adjustment, we obtained 3732 and 3568 matched patients in M1a and M1b group, respectively. Their survival curves were plotted by Kaplan-Meier methods (Figure 1A and D). Patients receiving palliative primary tumor resection had longer survival times than those without surgery (*P* < 0.001). For patients with M1a diseases, palliative resection could increase the median survival time by 9 mo; for patients with M1b diseases, palliative resection can prolong the median survival time by 7 mo (Table 3). Then M1a diseases were further subdivided by metastatic site. Patients with liver metastasis and lung metastasis were included in the further analysis, whereas patients with bone metastasis and brain metastasis were excluded because of small sample size. As shown in Figure 1B and C, patients with lung metastasis could obtain more clinical benefit from palliative resection than those with liver metastasis (15 mo for lung metastasis *vs* 8 mo for liver metastasis, Table 3, *P* < 0.001).

**DISCUSSION**

Metastatic CRC is a lethal disease with poor prognosis. While patients with metastatic CRC can obtain clinical benefits from curative surgery, there is still controversy with respect to the role of palliative primary tumor resection. To the best of our knowledge, this is the first population-based study subdividing stage IV into stages [IVA (M1a diseases) and stage IVB (M1b diseases)] to evaluate the effect of palliative primary tumor resection. Upon analysis it was determined that patients with M1a diseases could obtain more survival benefits than those with M1b diseases and patients with M1a (lung metastasis) got best long-term outcomes with median OS prolonged by 15 mo compared with those without surgical treatment. These findings provided further evidence to support palliative surgical procedure to metastatic CRA and develop effective individualized treatment strategy.

There were many predictors of OS in patients with unresectable metastatic CRC, including WHO PS, carcinoembryonic antigen level, number of metastatic sites and palliative surgery[13]. Li *et al*[17] showed that tumor location (right colon *vs* left colon *vs* rectum) was also an independent prognostic factor for metastatic CRC. The results were in line with our findings that patients with rectal cancer were at a lower risk of death than those with colon cancer, possibly owing to higher proportion of lung metastasis in patients with rectal cancer[18]. However, no studies focus on the effect of palliative surgery according to the number of metastatic sites or organs (M1a or M1b). Tarantino *et al*[14] reported that the survival difference between patients with palliative resection and those without palliative resection was anticipated to decrease due to the development of chemotherapeutic and molecule-targeted drugs. Actually, the significance of survival difference has persisted over time. This may be explained by the heterogeneity of stage IVA (M1a diseases) and stage IVB (M1b diseases). The development of systemic treatment could decrease the survival difference and increase surgery conversion indeed. The proportion of M1b diseases grew from 48.6% during 2010-2012 to 51.6% during 2013-2015 (Table 1, *P* < 0.001) and such patients were less likely to undergo surgical treatment than M1a diseases. This may also explain the decreased rate of patients undergoing primary tumor removal observed during 1998-2009 in Tarantino’s study[14].

Liver and lung metastasis are the most two common distant metastases from CRC[7], accounting for 50% and 10%-15% of CRC, respectively[19,20]. The prognosis of patients with liver or lung metastasis is usually better than those with brain or bone metastasis[21]. According to the published literature, median OS was 3-6 mo for patients with brain metastases and 5-7 mo for bone metastases[22-28]. For patients with unresectable liver metastases, who were treated with chemotherapy only, median OS was approximately 20 mo[29]. By comparison, patients with unresectable lung metastases, who achieved a complete or partial response to chemotherapy, could achieve a median OS of 27 mo[30]. From the perspective of epidemiology, the median time between the diagnosis of CRC and the emergence of liver metastases was shorter than that for lung metastases (17.2 mo for liver *vs* 24.6 mo for lung)[31], which indicated liver metastases possessed more aggressive malignant behavior to some extent. These survival findings were similar to the present results (22 mo for liver metastasis with palliative surgery *vs* 33 mo for lung metastasis with palliative surgery).

There are several limitations to this study that should be noted. PSM can adjust potential confounding variables and decrease selection bias as much as possible, increasing precision by creating a ‘quasi-randomized’ experiment[32]. However, we would like to admit the limitations of our study. First, some significant factors such as surgical complications, life quality, operative tolerance and laboratory parameters were not included. Second, detailed number of metastases in a single organ was not provided in the SEER database, which hampered further analysis for M1a diseases. Third, further classifications for M1 category in the AJCC staging system were not recorded before 2010, and only patient data between 2010 and 2015 were collected. The conclusions should be validated by more prospective data in the future.

**ARTICLE HIGHLIGHTS**

***Research background***

The National Comprehensive Cancer Network clinical practice guidelines do not recommend palliative surgery for metastatic colorectal adenocarcinoma (CRA) unless there is a risk of significant acute bleeding, obstruction, perforation or another severe symptom.

***Research motivation***

Accumulating evidence has demonstrated that palliative surgery for metastatic CRA patients was associated with more favorable outcomes. However, no studies further classified CRA patients with stage IV into subsets to assess the role of palliative surgery.

***Research objectives***

The purpose of this study was to investigate the separate role of palliative primary tumor resection for CRA patients with stage IVA (M1a diseases) and stage IVB (M1b diseases).

***Research methods***

CRA patient records with definite M1a and M1b categories were analyzed with adjusted propensity score matching. Patient prognosis was assessed by univariate and multivariate Cox regression analysis with hazard ratios and 95% confidence interval.

***Research results***

Patients with metastatic CRA receiving palliative primary tumor resection had a longer survival time than those who did not receive surgery (*P* < 0.001). Palliative resection increased the median survival time by 9 mo and by 7 mo for patients with M1a and M1b diseases, respectively. For M1a diseases, patients with lung metastasis had more Survival benefit from palliative resection than those with liver metastasis (15 mo for lung metastasis *vs* 8 mo for liver metastasis, *P* < 0.001).

***Research conclusions***

Palliative primary tumor resection improved survival for all CRA patients but more beneficial for those with M1a diseases than those with M1b diseases. Specifically, patients with M1a (lung metastasis) had the best long-term outcomes after palliative primary tumor resection.

***Research perspectives***

These findings provided further evidence to support the use of palliative surgical procedures treat metastatic CRA and develop effective individualized treatment strategies.

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**Footnotes**

**Institutional review board statement:** The study has been approved by the Bioethics Committee of the Affiliated Huaian No.1 People’s Hospital of Nanjing Medical University, China.

**Informed consent statement:** All patients from SEER database have agreed to participate in scientific researches.

**Conflict-of-interest statement:** All authors declare no competing financial interests.

**Data sharing statement:** No additional data are available.

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**Figure 1 Survival time for metastatic colorectal adenocarcinoma patients with or without palliative primary tumor resection.** A: Total patients with M1a diseases; B: Patients with M1a diseases (liver metastasis); C: Patients with M1a diseases (lung metastasis); D: Patients with M1b diseases.

**Table 1 Baseline characteristics of metastatic colorectal adenocarcinoma patients**

|  |  |  |  |  |
| --- | --- | --- | --- | --- |
| **Variables** | **Total (*n* = 19680) (%)** | **M1a (*n* = 10399) (%)** | **M1b (*n* = 9281) (%)** | ***P* value1** |
| Gender |  |  |  |  |
| Female | 8573 (43.6) | 4462 (42.9) | 4111 (44.3) | 0.050 |
| Male | 11107 (56.4) | 5937 (57.1) | 5170 (55.7) |  |
| Year of diagnosis |  |  |  | < 0.001 |
| 2010-2012 | 9835 (50.0) | 5342 (51.4) | 4493 (48.4) |  |
| 2013-2015 | 9845 (50.0) | 5057 (48.6) | 4788 (51.6) |  |
| Age |  |  |  | < 0.001 |
| < 65 | 10680 (54.3) | 5507 (53.0) | 5173 (55.7) |  |
| ≥ 65 | 9000 (45.7) | 4892 (47.0) | 4108 (44.3) |  |
| Race |  |  |  | 0.011 |
| White | 14715 (74.8) | 7856 (75.5) | 6859 (73.9) |  |
| Black | 3033 (15.4) | 1578 (15.2) | 1455 (15.7) |  |
| Others | 1932 (9.8) | 965 (9.3) | 967 (10.4) |  |
| Marital status |  |  |  | 0.661 |
| Married | 9774 (49.7) | 5180 (49.8) | 4594 (49.5) |  |
| Others | 9906 (50.3) | 5219 (50.2) | 4687 (50.5) |  |
| Primary site |  |  |  | 0.027 |
| Colon | 15476 (78.6) | 8114 (78.0) | 7362 (79.3) |  |
| Rectum | 4204 (21.4) | 2285 (22.0) | 1919 (20.7) |  |
| Grade |  |  |  | < 0.001 |
| I + II2 | 11321 (57.5) | 6525 (62.7) | 4796 (51.7) |  |
| III + IV3 | 3889 (19.8) | 1909 (18.4) | 1980 (21.3) |  |
| Others | 4470 (22.7) | 1965 (18.9) | 2505 (27.0) |  |
| Surgery |  |  |  | < 0.001 |
| Yes | 9360 (47.6) | 5787 (55.6) | 3573 (38.5) |  |
| No | 10320 (52.4) | 4612 (44.4) | 5708 (61.5) |  |
| Chemotherapy |  |  |  | 0.415 |
| Yes | 14057 (71.4) | 7402 (71.2) | 6655 (71.7) |  |
| No/unknown | 5623 (28.6) | 2997 (28.8) | 2626 (28.3) |  |
| Radiotherapy |  |  |  |  |
| Yes | 975 (5.0) | 670 (6.4) | 305 (3.3) | < 0.001 |
| No | 18705 (95.0) | 9729 (93.6) | 8976 (96.7) |  |

1*χ2* test. 2Well or moderately differentiated tumors. 3Poorly differentiated or undifferentiated tumors.

**Table 2 Univariate and multivariate cox regression analysis for overall survival for metastatic colorectal adenocarcinoma patients**

|  |  |  |  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- | --- | --- | --- |
|  | **M1a** | | | | **M1b** | | | |
| **Univariate analysis** | | **Multivariate analysis** | | **Univariate analysis** | | **Multivariate analysis** | |
| **HR (95%CI)** | ***P* value** | **HR (95%CI)** | ***P* value** | **HR (95%CI)** | ***P* value** | **HR (95%CI)** | ***P* value** |
| Gender |  |  |  |  |  |  |  |  |
| Female | Reference |  | Reference |  | Reference |  | Reference |  |
| Male | 1.100 (1.054-1.147) | < 0.001 | 1.025 (0.975-1.077) | 0.339 | 1.043 (1.002-1.086) | 0.042 | 1.020 (0.972-1.070) | 0.428 |
| Year of diagnosis |  |  |  |  |  |  |  |  |
| 2010-2012 | Reference |  | Reference |  | Reference |  | Reference |  |
| 2013-2015 | 0.972 (0.929-1.016) | 0.208 | 0.981 (0.929-1.036) | 0.486 | 0.965 (0.925-1.006) | 0.096 | 0.972 (0.924-1.023) | 0.277 |
| Age |  |  |  |  |  |  |  |  |
| < 65 | Reference |  | Reference |  | Reference |  | Reference |  |
| ≥ 65 | 1.675 (1.606-1.746) | < 0.001 | 1.413 (1.344-1.485) | < 0.001 | 1.527 (1.466-1.589) | < 0.001 | 1.319 (1.256-1.385) | < 0.001 |
| Race |  |  |  |  |  |  |  |  |
| White | Reference |  | Reference |  | Reference |  | Reference |  |
| Black | 1.138 (1.075-1.206) | < 0.001 | 1.114 (1.042-1.192) | 0.002 | 1.127 (1.068-1.191) | < 0.001 | 1.137 (1.064-1.215) | < 0.001 |
| Others | 0.910 (0.844-0.981) | 0.013 | 0.864 (0.792-0.942) | 0.001 | 0.953 (0.891-1.020) | 0.164 | 0.952 (0.878-1.031) | 0.227 |
| Marital status |  |  |  |  |  |  |  |  |
| Married | Reference |  | Reference |  | Reference |  | Reference |  |
| Others | 1.326 (1.272-1.383) | < 0.001 | 1.142 (1.086-1.201) | < 0.001 | 1.212 (1.165-1.262) | < 0.001 | 1.078 (1.026-1.132) | 0.003 |
| Primary site |  |  |  |  |  |  |  |  |
| Colon | Reference |  | Reference |  | Reference |  | Reference |  |
| Rectum | 0.799 (0.758-0.841) | < 0.001 | 0.759 (0.711-0.811) | < 0.001 | 0.839 (0.797-0.882) | < 0.001 | 0.779 (0.731-0.830) | < 0.001 |
| Grade |  |  |  |  |  |  |  |  |
| I + II | Reference |  | Reference |  | Reference |  | Reference |  |
| III + IV | 1.530 (1.450-1.614) | < 0.001 | 1.580 (1.484-1.682) | < 0.001 | 1.447 (1.375-1.523) | < 0.001 | 1.506 (1.417-1.601) | < 0.001 |
| Others | 1.702 (1.614-1.795) | < 0.001 | 1.149 (1.074-1.228) | < 0.001 | 1.544 (1.473-1.619) | < 0.001 | 1.178 (1.109-1.251) | < 0.001 |
| Surgery |  |  |  |  |  |  |  |  |
| Yes | Reference |  | Reference |  | Reference |  | Reference |  |
| No | 1.953 (1.859-2.051) | < 0.001 | 2.133 (2.011-2.262) | < 0.001 | 1.632 (1.552-1.716) | < 0.001 | 1.955 (1.843-2.074) | < 0.001 |
| Chemotherapy |  |  |  |  |  |  |  |  |
| Yes | Reference |  | Reference |  | Reference |  | Reference |  |
| No/unknown | 2.520 (2.395-2.651) | < 0.001 | 2.282 (2.164-2.405) | < 0.001 | 2.558 (2.430-2.692) | < 0.001 | 2.565 (2.432-2.705) | < 0.001 |
| Radiotherapy |  |  |  |  |  |  |  |  |
| Yes | Reference |  | Reference |  | Reference |  | Reference |  |
| No | 2.337 (2.069-2.639) | < 0.001 | 1.236 (1.085-1.408) | 0.001 | 1.695 (1.465-1.961) | < 0.001 | 0.946 (0.813-1.101) | 0.472 |

HR: hazard ratio; CI: Confidence interval.

**Table 3 Survival time for metastatic colorectal adenocarcinoma patients with or without palliative primary tumor resection**

|  |  |  |  |
| --- | --- | --- | --- |
|  | **Median survival time (95%CI)** | **1-yr survival rate (%)** | **3-yr Survival rate (%)** |
| M1a (total) |  |  |  |
| No surgery | 14 (13.275-14.725) | 54.4 | 13.9 |
| Surgery | 23 (21.977-24.023) | 70.4 | 32.6 |
| M1a (liver metastasis) |  |  |  |
| No surgery | 14 (13.320-14.780) | 53.8 | 13.4 |
| Surgery | 22 (20.955-23.045) | 69.2 | 29.8 |
| M1a (lung metastasis) |  |  |  |
| No surgery | 18 (15.692-20.308) | 62.2 | 18.6 |
| Surgery | 33 (28.014-37.986) | 77.5 | 45.4 |
| M1b |  |  |  |
| No surgery | 10 (9.401-10.599) | 42.2 | 8.6 |
| Surgery | 17 (16.209-17.791) | 60.0 | 20.4 |