Iodine-125 Seeds Combined With Biliary Stent Placement Versus Stent Placement Alone For Unresectable Malignant Biliary Obstruction: A Meta-Analysis Of Randomized Controlled Trials

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Abstract

Background and Aims: Malignant biliary obstruction is always caused by tumors which are unresectable so that palliative stent placement is conducted for drainage of bile duct tree. Recently, irradiation stent with 125I seeds has been used to improve the stent patency and survival time of patients. We conducted this meta-analysis to evaluate the therapeutic efficacy and safety of biliary stent placement with 125I seeds compared with stent placement alone in patients with malignant biliary obstruction.

Methods: We searched Pubmed, Web of Science, ClinicalTrials.gov, Cochrane Library, Embase and CNKI databases for all relevant studies up to 1 May 2020. Patient survival, stent patency, and adverse events were the primary outcome measured. Also, Review Manager 5.3 and Stata/SE15.0 were used to perform the analysis.

Results: Eleven randomized controlled trials with a total of 767 patients were included for meta-analysis. Stent combined with 125I seeds showed lower risk of stent occlusion at 3 month (Odds Ratios(OR) = 0.15; 95%CI: 0.05-0.49, P =0.002), 6 month (OR = 0.18; 95%CI: 0.08-0.44, P = 0.0001), 9 month (OR = 0.10; 95%CI: 0.05-0.20, P < 0.00001) and 1 year (OR = 0.15; 95%CI: 0.07-0.31, P < 0.00001) and better mean survival (MD = 125days; 95% CI 91–159 days; P < 0.00001) compared with stent placement alone. Also, reconstructed Kaplan-Meier data demonstrated improved survival in patients treated with stent plus 125I seeds (hazard ratio(HR)= 1.886; 95% CI: 1.609 to 2.210; P < 0.0001) Moreover, our analysis did not show significant difference between the two groups about the risk of adverse events including abdominal pain, hemobilia, pancreatitis, cholangitis and cholecystitis.

Conclusion: 125I seeds combined with stent demonstrated superior stent patency and improved survival time compared to stent alone with acceptable complications.

Key words: malignant biliary obstruction, 125I seed, biliary stent, stent patency, survival.

Introduction

Malignant biliary obstruction (MBO), which means stenosis and blockage of extrahepatic or intrahepatic bile ducts, is generally caused by various cancers including cholangiocarcinoma, pancreatic cancer, gallbladder cancer and cancer metastasis [1]. Due to the fact that its clinical performance is silent and unobvious, MBO is always diagnosed at an advanced stage when painless obstructive jaundice occurs. As a result, only 10%-20% of patients are resectable [2, 3]. Therefore, palliative therapy with...
stent implantation has been widely accepted and used for several decades. Clinically, covered or uncovered self-expanding metal stents (SEMSs) and plastic stents have been used via Percutaneous Transhepatic Cholangiography (PTC) or ERCP (Endoscopic retrograde Cholangiopancreatography) and the patency time of SEMSs is better than plastic stents about 3-4 months[4]. In addition, covered SEMS is superior to uncovered SEMS in terms of stent patency but it may influence the drainage of bile, causing some other complications and it may increase the rate of stent migration [5-8]. However, stent itself has no therapeutic effect on tumors, which always grow into the lumen through the stent mesh, resulting in occlusion of the stent. In addition, epithelial hyperplasia, biofilm deposition, biliary sludge, and formation of granulation tissue may cause stent occlusion as well [9, 10]. So, maybe stent placement alone was not enough for patients with MBO, and stent placement combined with inner or outside radiotherapy maybe a promising treatment option. Sofi et al. [11] showed that Radiofrequency Ablation (RFA) plus stent placement was superior to stent monotherapy (3) at least one of the following was reported: survival time or rate; stent patency or stent occlusion rate. (4) full text of studies are available. (5) patients in studies did not receive other radiotherapy. Studies were excluded when information about the survival or stent patency in the patient groups was not provided. Meanwhile, non-randomized controlled trials, animal experiments, narrative reviews were excluded too.

**Data extraction and Assessment of study quality**

Data were extracted by 2 authors (Y.L and Z.L) respectively based on the characteristics of studies. Detailed information included (1) basic information, such as author’s last name, year and type of publication, country; (2) patient characteristics and treatment information; and (3) efficacy outcomes and adverse events. Due to the fact that all the studies were RCTs, we used the Cochrane collaboration tool to determine the risk of bias, including six items: (1) sequence generation, (2) allocation concealment, (3) blinding, (4) incomplete outcome data, (5) no selective outcome reporting, and (6) other sources of bias. For each item, a low risk counts as a score of 1, with a total of 6 scores. Any disagreement was discussed with a third author(Y, X) and resolved with our consensus.

**Statistical analysis**

The primary end points were stent occlusion rates and overall survival time. Stent occlusion was dichotomous variables, so we extracted the number of events and total number of observed patients. Then, the odds ratios (OR) with 95% confidence intervals (CIs) were calculated. Also, survival was continuous variables, the means and standard deviations of observed patients were extracted from the included studies. Then, the pooled estimate of the mean difference (MD) with 95% confidence intervals (CIs) was calculated. Finally, the OR or MD of each study was pooled using either a fixed-effects or random-effects model. Moreover, when important information was not clear enough, we used digital software Engauge Digitizer (version 12.0) to extract information in the coordinates of the Kaplan-Meier.
curves from each of the included graphs, the survival analysis data points were extracted simultaneously. All the extracted information was exported into Microsoft Excel 2019, and then was imported into software Graphpad Prism (version 8) to analyze survival data information to reconstruct Kaplan-Meier curves and to obtain the HR (Hazard Ratio), median and mean survival time, stent occlusion rate and P value. It was considered valuable when the difference was less than 5% between the original and the extracted information. P values < 0.05 was defined as significant. A p value for Cochrane’s $\chi^2$ <0.1 or $I^2 > 50%$ was considered to have high heterogeneity. Random-effects models were used when the $I^2 > 50%$; otherwise, fixed-effects models were used. Publication bias was assessed by Egger’s test for asymmetry. The statistical analysis was performed using Stata (version 15) and RevMan 5.3 (Cochrane Collaboration, Oxford, U.K.).

Results

Study selection and characteristics of included studies

We strictly searched several databases for the articles based on the inclusion criteria, the number of articles was reduced from 1021 to 11 and all the included studies are RCTs (randomized controlled trials) with full text. The search process is shown in Figure 1. Within these eleven articles, a total of 767 patients were randomly distributed to either $^{125}$I seeds plus stent combination therapy group (377 patients) or stent monotherapy group (390 patients). Meanwhile, among these studies, all the stent implantation was performed by Percutaneous Transhepatic Cholangiography (PTC) rather than ERCP (Endoscopic retrograde Cholangiopancreatography). Technical success rate which means successful stent implantation and no shifts was almost 100%.

In addition, all the stents used in trials were ordinary metal stents like uncovered self-expandable metallic stents (USEMS) rather than plastic stents or covered self-expandable metallic stents. Moreover, seven studies [17-20, 24-26] used $^{125}$I seed strands and other four studies [16, 21-23] used $^{125}$I seed particles, which was analyzed in subgroup analysis. Detailed characteristics of studies were presented in Table 1, and basic characteristics of patients such as sex, age and etiology of malignant biliary obstruction were showed in Table 2. The Chi-square Test was conducted and all of the p value > 0.05, which means these characteristics between the two groups are well balanced.

Stent Occlusion

Stent occlusion between the $^{125}$I seeds groups and control groups were measured based on different time points including 3-month [16, 20-24, 26], 6-month [16, 20-24, 26], 9-month [16, 20, 23, 24, 26] and 1-year [16, 17, 19-26]. Combination therapy with $^{125}$I seeds demonstrated significant lower risk of stent occlusion at 3 month (OR = 0.15; 95%CI: 0.05-0.49, $P$ =0.002), 6 month (OR = 0.18; 95%CI: 0.08-0.44, $P = 0.0001$), 9 month (OR = 0.10; 95%CI: 0.05-0.20, $P < 0.00001$) and 1 year (OR = 0.15; 95%CI: 0.07-0.31, $P < 0.00001$) (Figure 2). Low heterogeneity ($I^2 = 0%$; $P = 0.88$) at 9-month stent occlusion and high heterogeneity at 3-month, 1-year stent occlusion were found ($I^2 = 63%$; $P = 0.01$, $I^2 = 64%$; $P = 0.01$, $I^2 = 54%$; $P = 0.02$ respectively) and we used random-effects models and fixed-effects models at the same time. Also, subgroup analysis and sensitivity analysis were further conducted below to find possible reasons.

Survival

9 of 11 studies [16-18, 20-25] reported data on survival. The pooled MD was significant (MD = 125 days; 95% CI 91–159 days; $P < 0.00001$), suggesting that $^{125}$I seeds groups significantly improved mean survival compared with control groups (Figure 3A). High heterogeneity ($I^2 = 58%$; $P = 0.02$) was found and we used random-effects models. Also, subgroup
analysis and sensitivity analysis were further conducted below to find possible causes. Moreover, the pooled overall survival from reconstructed Kaplan-Meier analyses demonstrated improved survival in patients receiving Iodine-125 seeds compared with patients undergoing biliary stent placement alone (hazard ratio, 1.886; 95% CI, 1.609 to 2.210; p<0.0001) as well (Figure 3B).

Table 1. Basic characteristics of the included trials

| Author (year) | Country | Period | Study design | Article type | Treatment | Control |
|---------------|---------|--------|--------------|--------------|-----------|---------|
| Hasimu (2016) | China   | July2011-June2014 | RCT | Full text | Interventi |
| Zhu (2012) | China   | November2008-October2010 | RCT | Full text | No. of patients | 28 | Stent with 125I seed strand | 27 | A nitinol self-expandable stent |
| Jiao (2017) | China   | January2013-January2015 | RCT | Full text | 12 | An outer self-expandable 125I seeds-loaded stent and an inner conventional self-expanding biliary nitinol alloy stent |
| Zhu (2017) | China   | October2013-March2016 | RCT | Full text | 31 | Stent with 125I seed strand | 30 | A Nitinol self-expandable stent |
| Chen (2012) | China   | March2009-January2010 | RCT | Full text | 152 | An inner conventional uncovered SEMS and an outer 125I seeds-loaded stent | 155 | Uncovered SEMS |
| Chen (2018) | China   | September2014-November2016 | RCT | Full text | 17 | 125I seed strand with stent | 17 | A Nitinol self-expandable stent |
| Zhou (2018) | China   | May2015-September2016 | RCT | Full text | 13 | 125I seed particle stent particle | 19 | Ordinary metal stent |
| Yao (2014) | China   | February2012-April2014 | RCT | Full text | 11 | Iodine-125 seed particle | 21 | Uncovered SEMS |
| Liu (2016) | China   | October2012-March2015 | RCT | Full text | 31 | 125I seed strand with stent | 31 | Uncovered SEMS |
| Zhao (2015) | China   | July2013-January2015 | RCT | Full text | 24 | 125I seed strand with stent | 28 | Uncovered SEMS |
| Ren (2015) | China   | March2012-February2013 | RCT | Full text | 31 | 125I seed strand with stent | 31 | Uncovered SEMS |

RCT: randomized controlled trial; 125I: iodine-125; SEMS: self-expandable metal stent.

Table 2. Basic characteristics of the included patients

| Ref | Intervention | Patient (no.) | Mean age (year) | Mean length of stricture(mm) | Sex male no. (%) | Etiology of MBO |
|-----|--------------|---------------|-----------------|-----------------------------|-----------------|----------------|
| Hasimu (2016) | 125I group | 28 | 70.95(8.56) | 60.89(9.54) | 11(59.3) | Mixed |
| Zhu (2012) | Stent group | 27 | 70.26(9.71) | 61.78(8.12) | 14(51.9) | Mixed |
| Jiao (2017) | 125I group | 12 | 62.5(17) | 42.5(7.2) | 7(58.3) | Mixed |
| Zhu (2017) | Stent group | 11 | 71(18) | 40.5(6.7) | 9(81.8) | Mixed |
| Zhao (2016) | 125I group | 31 | 60.4(8.8) | 29.5(6.8) | 6(19.4) | Mixed |
| Zhou (2015) | Stent group | 30 | 60.2(8.1) | 30.1(7.5) | 16(53.3) | Mixed |
| Chen (2012) | 125I group | 30 | 65(14) | 35(7.6) | 10(33.3) | Mixed |
| Chen (2018) | Stent group | 155 | 65.13 | 32(11.2) | 109(70.3) | Mixed |
| Zhou (2018) | 125I group | 17 | 61.2(14.5) | NA | 12(70.6) | Mixed |
| Zhao (2015) | Stent group | 17 | 63.9(9.3) | NA | 10(58.8) | Mixed |
| Liu (2016) | 125I group | 13 | 66 | NA | 12(63.2) | Mixed |
| Ren (2015) | Stent group | 19 | 68 | NA | 8(61.5) | Mixed |
| Yao (2014) | 125I group | 11 | NA | NA | 7(63.3) | Mixed |
| Zhou (2015) | Stent group | 21 | NA | NA | 13(61.9) | Mixed |
| Liu (2016) | 125I group | 31 | NA | NA | 18(61.9) | Mixed |
| Zhao (2015) | Stent group | 31 | NA | NA | 18(61.9) | Mixed |
| Ren (2015) | 125I group | 24 | NA | NA | 18(61.9) | Mixed |
| Zhu (2015) | Stent group | 28 | NA | NA | 18(61.9) | Mixed |

MBO: malignant biliary obstruction; Mixed: various kinds of cancers.
Adverse events

All the complications from studies primarily included abdominal pain, cholecystitis, pancreatitis, cholangitis and hemobilia, abdominal pain was the most common complication, but it did not reach statistical significance between 125I seeds + stent group and stent alone group (p=.68) and the same with the cholecystitis (p=.60), abdominal pain (p=.35), cholangitis(p=.91) and pancreatitis (p=.47) (Table 3).

Subgroup analysis and Sensitivity analysis

We made subgroup analysis and sensitivity analysis to find the possible reasons for high heterogeneity and intend to analyze possible factors that may influence stent occlusion and survival. We separated studies based on the different types of 125I seeds and seven studies [17-20, 24-26] used 125I seed strand and other four studies [16, 21-23] used 125I seed particles. Significant difference was found between 125I seeds groups and control groups about 6-month, 9-month and 1-year stent occlusion regardless of 125I seed strand or 125I seed particles used, but 125I seed strand groups showed no significant difference.
(p=0.07) about 3-month stent occlusion (Figure 4). Moreover, significant difference of survival between 125I seeds groups and control groups was found either 125I seed strand or 125I seed particles used (Figure 5). As for heterogeneity, within 1-year stent occlusion, either 125I seed strand group or 125I seed particles group showed no high heterogeneity (I²=0%, I²=13% respectively), but the pooled result showed high heterogeneity (I²=54%), which demonstrated that different types of 125I seeds may account for the high heterogeneity (Figure 4). However, the reason for high heterogeneity within 3-month, 6-month stent occlusion and survival were unclear, therefore, we conducted sensitivity analysis and we found Zhu et al[21] may account for the high heterogeneity that including this study or excluding it showed huge difference. When excluded the study by Zhu [21], I² changed from 63% to 40%, 64% to 51%, 54% to 0% and 58% to 38% respectively at 3-month, 6-month, 1-year stent occlusion and survival (Table 4). The possible reasons were analyzed in discussion.

Figure 4. Subgroup analysis of 3-month (A), 6-month (B), 9-month(C) and 1-year (D) stent occlusion between 125I seeds groups and control groups.

Figure 5. Subgroup analysis of survival between 125I seeds groups and control groups.
Quality assessment

All the eleven trials included were from China and described proper randomization, whereas several studies provided no details of patient allocation. Only one study was blinded.

Validity was assessed in detail (Figure 6), the score of the trials varies from 3-7.

Table 4. Outcome of sensitivity analysis

| 3-month stent occlusion | OR and 95%CI | P value |
|-------------------------|-------------|---------|
| Include Zhu(2018) | 63% | 0.15[0.05,0.49] | <0.002 |
| Exclude Zhu(2018) | 40% | 0.11[0.05,0.24] | <0.000001 |
| 6-month stent occlusion | OR and 95%CI | P value |
| Include Zhu(2018) | 64% | 0.18[0.08,0.44] | <0.000001 |
| Exclude Zhu(2018) | 51% | 0.14[0.05,0.36] | <0.000001 |
| 12-month stent occlusion | OR and 95%CI | P value |
| Include Zhu(2018) | 54% | 0.15[0.07,0.31] | <0.000001 |
| Exclude Zhu(2018) | 0% | 0.16[0.06,0.20] | <0.000001 |
| Overall survival | MD and 95%CI | P value |
| Include Zhu(2018) | 58% | 125[91,159] | <0.000001 |
| Exclude Zhu(2018) | 38% | 133[109,157] | <0.000001 |

Discussion

For patients with MBO who have unresectable tumors or who are unwilling to accept surgery, biliary stent placement is the preferred choice [27, 28]. However, stent occlusion of patients remains challenging clinical problems. This meta-analysis of eleven RCTs [16 -26] provides relatively adequate evidence in favor of 125I seeds combined with stent placement, which showed improved survival and lower risk of stent occlusion compared with stent placement alone. Moreover, 125I seeds implantation is generally safe in the treatment of patients with malignant biliary obstruction.

125I can emit 27.4–31.5 keV χ-rays and 35.5 keV γ-rays, with an effective radiation radius of 17-20 mm, initial dose of 7.7 cGy/h, and a radioactive half-life of 60 days and it can be kept within the tumor area to directly kill tumor cells and inhibit tumor growth into the mesh of the stent but with limited damage to surrounding normal tissues and adjacent organs and subsequently improved patients’ liver function and performance status [29, 30]. Furthermore, implantation of 125I may cause CD3+ and CD4+ cell activation, which may lead to antitumor immune responses [14]. That maybe the reason why 125I seeds plus stent showed decreased rate of stent occlusion and superior survival.

Subgroup analysis and sensitivity analysis

In our meta-analysis, seven studies used 125I seed strands and other four studies used 125I seed particles and we made subgroup analysis based on this
difference. Although significant difference between $^{125}$I seeds groups and control groups was found about 6-month, 9-month and 1-year stent occlusion and survival either $^{125}$I seed strand or $^{125}$I seed particles used, even $^{125}$I seed strand groups showed no significant difference about 3-month stent occlusion, we speculated that $^{125}$I seed strand maybe the better choice. Possible reasons maybe as follows. First, irradiation dose was the focus of $^{125}$I seeds brachytherapy [31] and the minimum threshold for effective brachytherapy treatment of adenocarcinoma was 30 Gy, which has the capacity to kill tumor cells effectively [32]. $^{125}$I seed strand was constructed by $^{125}$I seeds linearly arranged which may cumulate higher irradiation dose than $^{125}$I seed particles placed apart. Second, $^{125}$I seed strand was more easily to place with lower risk of migration than $^{125}$I seed particles technically because $^{125}$I seed strand was sealed as a single object but $^{125}$I seed particles had to be loaded on the stent one by one to form the irradiation stent. Third, $^{125}$I seed strand was easily to withdraw if radiation-related complications occurred but $^{125}$I seed particles could not. Last, economic payment should be considered that whether $^{125}$I seed strand was cheaper than $^{125}$I seed particles used clinically.

In addition, based on $^{125}$I seeds plus stent placement, whether different etiology of malignant biliary obstruction such as cholangiocarcinoma, pancreatic cancer or hepatocellular carcinoma had effects on survival of patients, whether different types of stent influenced the stent patency, whether extrahepatic and intrahepatic biliary obstruction showed different therapeutic prognosis, whether $^{125}$I seeds plus stent placement is superior to RFA combined with stent, ought to be analyzed. However, there was not enough information in the included studies and more studies should be conducted to figure out the precise and specialized treatment for patient with MBO.

Furthermore, we conducted sensitivity analysis and found that high heterogeneity of 3-month, 6-month and 1-year stent occlusion disappeared when the study by Zhu [21] was excluded. Possible reasons maybe as follows. First, number of patients in this study were large that almost occupied the half of the total number. Second, some patients who underwent stent occlusion received stent implantation again in this study. Third, some patients received chemotherapy after stent placement too.

Advantages and disadvantages

In our meta-analysis, we included eleven RCTs which provided relatively adequate evidence and we used digital software Engauge Digitizer to reconstruct Kaplan-Meier curves and obtain the HR, median and mean survival time as well. However, there were several disadvantages in our meta-analysis. First, high heterogeneity was found in 3-month, 6-month and 1-year stent occlusion and we performed subgroup analysis and sensitivity analysis to find possible causes. Second, several included studies were of high quality that complied with the rules of RCT strictly but several studies did not abide by the rules strictly enough. Third, all the 11 studies were conducted in China, which may diminish the adequate representation of the current study. Last, sample size maybe not huge enough and included patients had various kinds of cancers which may hinder the applicability of this analysis.

In conclusion, $^{125}$I seeds plus stent placement was significantly superior to stent placement alone with regard to stent patency and survival of patients. In addition, $^{125}$I seeds implantation was a safe and tolerable procedure with comparable complications to stent placement alone. It may be a promising therapy for patients with malignant biliary obstruction and more well-designed randomized trials are necessary to further confirm these conclusions.

Abbreviations

MBO: malignant biliary obstruction; RCT: randomized controlled trial; $^{125}$I: iodine-125; SEMS: self-expandable metal stent; PTC: percutaneous transhepatic cholangiography; ERCP: endoscopic retrograde cholangiopancreatography; OR: odds ratios; MD: mean difference; CI: confidence interval.

Author Contributions

Y.X. and W.W. conceived the idea. Y.X. and S.L. searched and included articles and wrote the manuscript text. Y.L. and Z.L. collected and analyzed the data. Y.X. and Y.L. reconciled disagreements. S.L. and Z.L. provided statistic support. All authors reviewed the manuscript.

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Competing Interests

The authors have declared that no competing interest exists.

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