

Review



Cryoablation versus Partial Nephrectomy for Clinical Stage T1 Renal Masses: A Systematic Review and Meta-Analysis

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Abstract

Introduction: Cryoablation has been considered as the most efficacious ablative alternative to partial nephrectomy (PN) for selected patients. Our objective is to assess the existing evidence relating to the safety and efficacy of cryoablation compared with PN for clinical T1 renal masses.

Materials and Methods: A comprehensive search of PMC, EMBASE, and the Cochrane Library was conducted to identify studies containing comparison of cryoablation and PN. By utilizing those included studies, a systematic review and cumulative meta-analyses were performed to assess the safety and efficacy between cryoablation and PN for T1 renal masses.

Results: 17 retrospective studies providing available data were included in our study. Significant differences were found about all oncological variables including all-cause death, cancer-specific death, metastasis and local-recurrence (p < 0.001, p = 0.03, p < 0.001, and p < 0.001, respectively) between the PN group and the cryoablation group. The mean difference between two groups for percent estimated glomerular filtration rate decrease and creatinine increase was -4.84 and 0.15 respectively (p < 0.001 and p = 0.006, respectively). The incidences of overall and postoperative complications in the PN group were significantly higher than that in the cryoablation group (p = 0.001 and p < 0.001, respectively), but the result about intraoperative complications didn't show a significant difference between the two groups (p = 0.53).

Conclusions: Comparing with PN, cryoablation for clinical T1 renal tumors is associated with poorer oncological outcomes, but the existing disadvantages are accompanied by lower rate of overall and postoperative complications and superior renal functional preservation. For patients with imperative indications for nephron-sparing surgery who can't risk more invasive PN, cryoablation could be an attractive option. Owing to the inherent limitations of eligible studies, conclusions drawn from our meta-analyses should be interpreted cautiously and be confirmed further with well-designed randomized controlled trials with extensive follow-up length.

Key words: cryoablation; partial nephrectomy; renal mass; clinical T1; meta-analysis

Introduction

An increase in newly detection of incidental small renal masses has reached to 2%-3.1% per year owing to routine application of multiple imaging modalities [1, 2]. Small renal tumors were more likely to originate in patients with elderly age and

comorbidities [3]. That's why sufficient renal function preservation and fewer complications were also important besides satisfactory oncological outcomes.

Partial nephrectomy (PN) has been adopted as standard treatment for stage T1a renal tumors.

Recently guideline has recommended PN rather than radical nephrectomy as management for selected T1b renal tumors [4]. PN can be conducted under open conditions, with laparoscopy or robotic assistance [5, 6].

Ablative therapies have been regarded as an alternative method to PN [7, 8]. Most frequent ablative technologies are cryoablation and radiofrequency ablation [9]. Cryoablation was considered as the most efficacious alternative to PN because of the lower possibility of local-recurrence progression metastatic compared with and radiofrequency ablation [9, 10]. Cryoablation can be performed with laparoscopy or percutaneous image guidance. Both approaches were reported to be safe and efficient to treat stage T1 renal tumors [11, 12].

More and more articles have reported that cryoablation could be applied for renal tumors up to T1b but not limited to small renal masses (usually defined as renal mass ≤ 4cm) [8, 13, 14]. A meta-analysis comparing laparoscopic cryoablation with laparoscopic PN for small renal masses has been reported [15]. However, no study systematically reviewed cryoablation for T1 renal tumors compared with PN. Several studies have compared cryoablation and PN applied to T1 renal masses [2, 3, 8, 10, 13, 14, 16-26], which may provide evidence that cryoablation could be an alternative challenging PN for surgical high-risk patients with T1 renal masses. Along with those series, a systematic review and meta-analysis was carried out to investigate the efficacy and safety of cryoablation for T1 renal masses compared with PN.

Materials and Methods

Data Sources and Searches

A comprehensive literature search utilizing the major electronic databases (PMC, EMBASE and the Cochrane Library) up to September 2018 was not restricted to regions or languages. ("partial nephrectomy" [Title/Abstract] or "nephron sparing surgery" [Title/Abstract]) and ("cryoablation" [Title/Abstract] or "cryotherapy" [Title/Abstract]) were combined as search terms. Related articles were also identified manually with screening the reference lists of all retrieved articles.

Inclusion and Exclusion Criteria

The following inclusion criteria were necessary: (1) Article has been published or accepted for full-length publication; (2) Article must be designed as a prospective or retrospective study that compared cryoablation and PN for T1 renal tumors; (3) At least one of the outcomes of interest below is evaluated and data is extractable. The following exclusion criteria were used: (1) A review or meta-analysis; (2) No available data could be extracted. (3) Editorials, letters, case report or animal experimental studies.

Outcome of Interest and Definitions

Following outcome variables were extracted to assess the efficiency and safety between cryoablation and PN.

1. Oncological variables: all-cause death, cancer-specific death, metastasis, local-recurrence.

2. Variables about renal function change: percent estimated glomerular filtration rate (eGFR) decrease, creatinine increase.

3. Variables about perioperative complications: overall complication rate, intraoperative complication rate, postoperative complication rate.

Data Extraction

Data from the included studies was extracted and calculated separately by two authors (Deng and Chen). Discussions were necessary until no doubt existed whenever there was a disagreement. Following information was obtained from every included article: first author's family name; year of publication; study design; number of included cases in both groups; cohort representative; surgical approach of each group; comparability between two groups; follow-up outcomes.

Quality assessment and statistical analysis

Criteria enacted by the Center for Evidence-Based Medicine was used to rate included studies for the level of evidence [27].

The modified Newcastle-Ottawa scale was used to assess the methodological quality of retrospective studies [28]. The scale evaluates risk in three factors of study design: patient selection; comparability between two groups; assessment of outcomes. Every study was scored from 0 to 9. High quality study was defined as study getting a score of \geq 7, otherwise studies were considered to be of moderate quality.

Hazard ratio (HR), mean difference (MD) and the odds ratio (OR) were used to measure the oncological variables, the renal function change and the complication rates, respectively. All outcomes were presented with 95% confidence intervals (95%CI). The method summarized by Tierney et al. [29] was used to calculate HRs and 95%CI when the data wasn't provided directly. When renal function change was presented as medians and ranges, the means provided by Hozo et al. [30] was used to convert the data into means with standard deviations. Yates' correction was used when no complication occurred in one of two groups [31]. Chi-squared test with significance set at p < 0.10 was used to assess the statistical heterogeneity between studies. The I² statistic was used to quantify heterogeneity. The fixed-effects (FE) model was applied to pool data if no significant heterogeneity existed between studies, or the random-effects model was applied [32].

In order to assess the stability of every outcome of interest, original analyses were repeated with exclusion of the study that got the lowest score to conduct sensitivity analyses. To explore the source of heterogeneity, subgroup analyses were performed according to the quality of included studies.

A funnel plot about all-cause death was inspected visually to assess the potential for publication bias.

Review Manager 5.3 (Cochrane Collaboration, Oxford, UK) was run to conduct all statistical analyses. A p value < 0.05 was deemed to mean statistically significant.

Results

Description of Included Studies

Table 1. Characteristics of included studies

Seventeen studies [2, 3, 8, 10, 13, 14, 16-26] fulfilling the inclusion criteria drafted in advance were singled out from the 1059 potentially relevant studies. Details of study selection were shown in flow-process diagram (Figure 1). All included studies were published in full with a retrospective observational study design. Level of evidence of all selected studies was 3b owing to their study design. More characteristics of the included studies were presented in Table 1.

Quality of Eligible Studies

No appropriate protocol was employed in advance to allocate patients and assign treatment in the retrospective studies, but just according to physician's discretion. Information about allocation concealment or the blinding method was not mentioned in studies. Comparable information about the included patients between the two groups varied from article to article. The mean follow-up period of every group in all studies was up to 6 months. 8 eligible studies got a score \geq 7 according to Newcastle-Ottawa Scale (Table S1) [8, 14, 16-18, 20, 24, 25], while the remaining ones were marked as moderate quality studies [2, 3, 10, 13, 19, 21-23, 26]. Most studies did not describe clearly about how to deal with missing data or intention-to-treat analyses.





Study	Country	Design	Patients, no		appro	oach	Matching	Follow-up, m		level of	Quality
			PN	CA	PN	CA	1,2,3,4,5,6,7,8,9,10	PN	CA	evidence	Score
Bhindi ²⁰¹⁸ [13]	USA	R	64	54	NA	Р	2,8, 10	47	47	3b	5
Caputo ²⁰¹⁷ [14]	USA	R	31	31	R	L+P	1,2,3,4,5,8,9,10	13.0	30.1	3b	8
Chehab ²⁰¹⁶ [16]	USA	R	128	37	R+O	Р	2,3,4,8	28.1	22.1	3b	7
Desai ²⁰⁰⁵ [17]	USA	RP	153	78	L	L	2,3,6,7	5.8	24.6	3b	7
Emara ²⁰¹⁴ [2]	UK	R	47	56	R	L	2,6,8	6	6	3b	6
Fraisse ²⁰¹⁸ [18]	France	R	470	177	R	Р	2,3,4,10	39.03	62.59	3b	7
Guillotreau ²⁰¹² [19]	USA	R	210	226	R	L	3,6,7	6	6	3b	5
Haber ²⁰¹² [20]	USA	RP	48	30	L	L	1,2,3,4,5,6,7,9,10	42.7	60.2	3b	8
Haramis ²⁰¹² [3]	USA	R	92	75	L	L	2,4,5,6,7,8	21.8	14	3b	6
Hruby ²⁰⁰⁶ [21]	USA	R	12	11	L	L	7	12	11.3	3b	4
Klatte ²⁰¹¹ [22]	Austria	R	82	41	0	L	1,2,4,5,10	33.6	33.6	3b	6
Lin ²⁰⁰⁸ [23]	USA	RP	14	13	L	L	2,3,6,9	38.5	24	3b	6
O'Malley2007[24]	USA	R	15	15	L	L	1,2,3,4,5,6,7,9	9.83	11.9	3b	8
Tanagho ²⁰¹³ [10]	USA	R	233	267	R	L+P	2,3,6	21.9	39.8	3b	4
Thompson ²⁰¹⁵ [8]	USA	RP	1383	240	0	Р	1,2,4,5,10	36	36	3b	7
Turna ²⁰⁰⁹ [25]	USA	R	36	36	L	L+P	1,2,3,4,7,9	6	6	3b	7
Zondervan ²⁰¹⁶ [26]	Netherland	R	124	137	O+L	L+P	2.3.10	44	43	3b	6

PN = partial nephrectomy; CA = cryoablation; m = months. R = retrospective; RP = retrospective design, prospective data collection; O = open; L = laparoscopic; R = robot-assisted; P = percutaneous.

Matching variables: 1 = age, 2 = gender, 3 = body mass index, 4 = tumor size, 5 = preoperative renal function, 6 = tumor side, 7 = tumor location, 8 = nephrometry (RENAL) Score, 9 = American Society of Anesthesiologists score, 10 = Charlson comorbidity index.



Figure 2. Forest plot and meta-analysis of all-cause death.



Oncological Outcomes

Available data provided by 8 of the 17 studies was pooled for all-cause death with nil heterogeneity [8, 10, 13, 14, 17, 20, 25, 26]. Original result indicated that stage T1 patients undergoing PN showed a significantly lower all-cause death compared with those undergoing cryoablation (HR, 0.52; 95%CI: 0.41 to 0.65; p < 0.001). Both the high quality study subgroup and the moderate quality study subgroup got as significant as the original result while the heterogeneity remained nil ((HR, 0.43; 95%CI: 0.31 to 0.61; p < 0.001) and (HR, 0.60; 95%CI: 0.44 to 0.82; p = 0.002), respectively) (Figure 2).

HRs on cancer-specific death could be extracted or calculated from 6 included studies [10, 13, 14, 20, 25, 26]. Significant higher possibility to cancer-specific death was found in the cryoablation group compared with the PN group (HR, 0.43; 95% CI: 0.21 to 0.91; p = 0.03). In the high quality study subgroup, the pooled HR was 0.14 (95% CI: 0.03 to 0.67; p = 0.01) without between-study heterogeneity. But no significant difference was found between the two treatment groups in the moderate quality study subgroup with slightly increased heterogeneity (HR, 0.60; 95% CI: 0.26 to 1.39; p = 0.23; $I^2 = 6\%$) (Figure 3).

4 studies [8, 13, 19, 20] including 2255 patients provided available details about the risk of metastasis. Statistical significance in the incidences of metastasis was found between the PN group and the cryoablation group (HR, 0.41; 95% CI: 0.26 to 0.67; p = 0.0004). The FE model was used to pool the result owing to the low heterogeneity. Results of the two subgroups remained to be significantly different between the two treatment groups, while the heterogeneity between the moderate quality studies increased to a high level (Figure 4).

Data on local-recurrence could be extracted from 11 studies [3, 8, 10, 13, 14, 17-20, 22, 26]. The FE model was used to pool the data on the basis of the

insignificant heterogeneity. The rate of local-recurrence in the cryoablation group was significantly higher than that in the PN group (HR, 0.35; 95%CI: 0.25 to 0.50; p < 0.001). Significant results were also concluded in the two subgroups while the heterogeneity between studies was decreased in the high quality subgroup and the opposite change was detected in the moderate quality subgroup (Figure 5).





Figure 5. Forest plot and meta-analysis of local-recurrence.

Renal Function Outcomes

Pooled data about percent eGFR decrease from 6 studies [10, 13, 14, 19, 20, 23] indicated a significantly better renal function preservation in the cryoablation group compared with the PN group (MD: -4.84; 95%CI: -7.02 to -2.36; p < 0.0001). The data was pooled without heterogeneity using the FE model. Results of the two subgroups were consistent with the original result (Figure 6).

There was no heterogeneity in 3 included studies [2, 20, 25] focusing on creatinine increase. The pooled WD for creatinine increase was 0.15 (95%CI: 0.04 to 0.26; p = 0.006) indicating that cryoablation protected renal function significantly better than PN for T1 renal

tumor cases. Subgroup analysis showed the same outcome as the original meta-analysis (Figure 7).

Complications Outcomes

There were significantly more overall complications and postoperative complications in the PN group than the cryoablation group ((OR: 1.86; 95%CI: 1.28 to 2.70; p = 0.001) and (OR: 2.97; 95%CI: 2.13 to 4.14; p < 0.001), respectively) (Figures 8 and 9). But no significant difference was found in intraoperative complication rate between two groups (OR: 2.02; 95%CI: 0.22 to 18.51; p = 0.53) (Figure 10). For three complication variables, pooled data of all subgroups were in accordance with those original results.



Figure 6. Forest plot and meta-analysis of percent eGFR decrease.

	partial nephrectomy			cryo	ablati	on		Mean Difference	Mean Difference
Study or Subgroup	Mean	SD	Total	Mean	SD	Total	Weight	IV, Fixed, 95% CI	IV, Fixed, 95% CI
2.2.1 high quality stud	ly						-		
Haber 2012	0.4	0.5	48	0.2	0.3	30	36.5%	0.20 [0.02, 0.38]	
Turna 2009	0.4	0.5	36	0.2	0.3	36	31.7%	0.20 [0.01, 0.39]	
Subtotal (95% CI)			84			66	68.1%	0.20 [0.07, 0.33]	
Heterogeneity: Chi ² = 0	.00, df = 1 (l	P = 1.00); I ² = 0	%					
Test for overall effect: Z	Z = 3.02 (P =	= 0.003)							
2.2.2 moderate quality Emara 2014 Subtotal (95% CI) Heterogeneity: Not app Test for overall effect: 2	/ study 0.1 licable Z = 0.41 (P =	0.49 = 0.68)	47 47	0.06	0.49	56 56	31.9% 31.9%	0.04 [-0.15, 0.23] 0.04 [-0.15, 0.23]	
Total (95% CI) Heterogeneity: Chi ² = 1 Test for overall effect: 2 Test for subgroup differ	.86, df = 2 (l Z = 2.72 (P = rences: Chi²	P = 0.40 = 0.006) = 1.86,	131); I² = 0 df = 1 (l	% P = 0.17	7), I² =	122 46.1%	100.0%	0.15 [0.04, 0.26]	-0.5 -0.25 0 0.25 0.5 Favours [experimental] Favours [control]

Figure 7. Forest plot and meta-analysis of creatinine increase.

	partial nephre	ctomy	cryoabla	ation		Odds Ratio	Odds Ratio
Study or Subgroup	Events	Total	Events	Total	Weight	M-H, Random, 95% CI	M-H, Random, 95% Cl
3.1.1 high quality stu	dy						
Caputo 2017	13	31	7	31	6.4%	2.48 [0.82, 7.47]	
Chehab 2016	9	128	4	37	5.6%	0.62 [0.18, 2.15]	
Desai 2005	49	153	6	89	7.9%	6.52 [2.66, 15.96]	
Fraisse 2018	104	470	28	177	12.0%	1.51 [0.96, 2.39]	
Haber 2012	15	48	3	30	5.1%	4.09 [1.07, 15.62]	
O'Malley 2007	3	15	4	15	3.6%	0.69 [0.12, 3.79]	
Turna 2009	21	36	5	36	6.1%	8.68 [2.74, 27.52]	
Subtotal (95% CI)		881		415	46.6%	2.45 [1.20, 4.99]	-
Total events	214		57				
Heterogeneity: Tau ² =	0.60; Chi ² = 20.4	3, df = 6	(P = 0.002); ² = 7	1%		
Test for overall effect:	Z = 2.47 (P = 0.0	1)					
3.1.2 moderate qualit	ty study						
Bhindi 2018	20	64	8	54	7.7%	2.61 [1.04, 6.55]	
Emara 2014	7	47	6	56	6.0%	1.46 [0.45, 4.68]	
Guillotreau 2012	49	210	27	226	11.4%	2.24 [1.34, 3.75]	
Haramis 2012	11	92	6	75	6.8%	1.56 [0.55, 4.44]	
Klatte 2011	14	82	8	41	7.4%	0.85 [0.32, 2.22]	
Lin 2008	3	14	4	13	3.5%	0.61 [0.11, 3.49]	
Tanagho 2013	22	233	23	267	10.5%	1.11 [0.60, 2.04]	
Subtotal (95% CI)		742		732	53.4%	1.54 [1.09, 2.17]	•
Total events	126		82				
Heterogeneity: Tau ² =	0.03; Chi ² = 7.00	, df = 6 (F	^o = 0.32);	² = 14%	þ		
Test for overall effect:	Z = 2.44 (P = 0.0	1)					
Total (95% CI)		1623		1147	100.0%	1.86 [1.28, 2.70]	◆
Total events	340		139				
Heterogeneity: Tau ² =	0.25; Chi ² = 29.3	5, df = 13	B (P = 0.00	6); l² =	56%		
Test for overall effect:	Z = 3.26 (P = 0.0	01)	10				0.05 0.2 1 5 20
Test for subaroup diffe	erences: Chi ² = 1	34 df = 1	(P = 0.25)	$ ^{2} = 2$	5.3%		partial hephrectomy cryoablation

Figure 8. Forest plot and meta-analysis of overall complication rates.

	partial nephrectomy		partial nephrectomy		partial nephrectomy cry			ation		Odds Ratio	Odds Ratio
Study or Subgroup	Events	Total	Events	Total	Weight	M-H, Fixed, 95% Cl	M-H, Fixed, 95% Cl				
3.3.1 high quality stud	У										
Caputo 2017	13	31	7	31	9.4%	2.48 [0.82, 7.47]					
Desai 2005	41	153	5	89	10.7%	6.15 [2.33, 16.23]					
Haber 2012	11	48	2	30	4.4%	4.16 [0.85, 20.30]	· · · · ·				
Turna 2009	21	36	5	36	4.8%	8.68 [2.74, 27.52]					
Subtotal (95% CI)		268		186	29.3%	5.09 [2.88, 8.99]	•				
Total events	86		19								
Heterogeneity: Chi ² = 2	67, df = 3 (P = 0	.45); l ² =	0%								
Test for overall effect: Z	= 5.60 (P < 0.00	0001)									
3.3.2 moderate quality	study										
Emara 2014	7	47	6	56	10.8%	1.46 [0.45, 4.68]					
Guillotreau 2012	42	210	27	226	48.1%	1.84 [1.09, 3.12]					
Tanagho 2013	18	233	6	267	11.9%	3.64 [1.42, 9.34]					
Subtotal (95% CI)		490		549	70.7%	2.09 [1.37, 3.18]	•				
Total events	67		39								
Heterogeneity: Chi ² = 1	92, df = 2 (P = 0	.38); 12 =	0%								
Test for overall effect: Z	= 3.41 (P = 0.00	006)									
Total (95% CI)		758		735	100.0%	2.97 [2.13, 4.14]	•				
Total events	153		58								
Heterogeneity: Chi ² = 1	0.53, df = 6 (P =	0.10); l ²	= 43%								
Test for overall effect: Z	= 6.40 (P < 0.00	0001)					0.02 0.1 1 10 50				
Test for subaroup differ	ences: Chi ² = 6.0)7. df = 1	(P = 0.01)), $ ^2 = 83$	3.5%		partial nephrectomy cryoablation				

Figure 9. Forest plot and meta-analysis of postoperative complication rates.

	partial nephrect	omy	cryoabla	ation		Odds Ratio	Odds Ratio
Study or Subgroup	Events	Total	Events	Total	Weight	M-H, Random, 95% C	M-H, Random, 95% Cl
3.2.1 high quality stud	У						
Desai 2005	8	153	1	89	25.1%	4.86 [0.60, 39.48]	+
Haber 2012	4	48	1	30	24.4%	2.64 [0.28, 24.79]	
Subtotal (95% CI)		201		119	49.5%	3.65 [0.79, 16.87]	
Total events	12		2				
Heterogeneity: Tau ² = 0	.00; Chi ² = 0.15, d	df = 1 (F	P = 0.69); I	² = 0%			
Test for overall effect: Z	= 1.66 (P = 0.10)						
3.2.2 moderate quality	study						
Guillotreau 2012	7	210	0	226	21.0%	16.70 [0.95, 294.15]	
Tanagho 2013	3	230	18	249	29.5%	0.17 [0.05, 0.58]	
Subtotal (95% CI)		440		475	50.5%	1.42 [0.01, 160.58]	
Total events	10		18				
Heterogeneity: Tau ² = 1	0.44; Chi ² = 9.22,	df = 1 (P = 0.002); l ² = 8	9%		
Test for overall effect: Z	= 0.14 (P = 0.88)						
Total (95% CI)		641		594	100.0%	2.02 [0.22, 18.51]	
Total events	22		20				
Heterogeneity: Tau ² = 3	.94; Chi ² = 14.48,	df = 3 (P = 0.002); ² = 7	9%		
Test for overall effect: Z	= 0.62 (P = 0.53)						0.001 0.1 1 10 1000
Test for subaroup different	ences: Chi ² = 0.14	4. df = 1	(P = 0.71)). ² = 0 ⁰	%		partial hephrectomy cryoablation
10. Forest plot and meta-a	nalysis of intraope	rative c	omplicatio	on rates			

Table 2. Results of meta-analysis and sensitivity analysis about comparison of partial nephrectomy and cryoablation

Outcomes of interest	Studies, no	PN patient, no	cryoablation patients, no	HR/WMD/OR (95%CI)	<i>p</i> value	Study heterogeneity			
						Chi ²	df	р	I²,%
Oncologic outcomes									
All-cause death	8	2072	873	0.52(0.41,0.65)+	< 0.001	7.05	8	0.53	0
Exclusion of [10]	7	1839	606	0.46(0.35,0.60)+	< 0.001	4.38	7	0.73	0
Cancer-specific death	6	536	555	0.43(0.21,0.91)+	0.03	4.81	5	0.44	0
Exclusion of [10]	5	303	288	0.36(0.14,0.92)+	0.03	4.42	4	0.35	9
Metastasis	4	1705	550	0.41(0.26,0.67)	0.0004	4.46	4	0.35	10
Exclusion of [13]	3	1641	496	0.29(0.15,0.57)	0.0003	2.22	3	0.53	0
Local-recurrence	11	2597	1356	0.35(0.25,0.50)	< 0.001	17.19	11	0.10	36
Exclusion of [10]	10	2364	1089	0.34(0.20,0.57)	< 0.001	16.25	10	0.09	38
Renal function outcomes									
Percent eGFR decrease	6	600	621	-4.84(-7.02,-2.66)	< 0.001	4.26	5	0.51	0
Exclusion of [10]	5	367	354	-4.10(-6.63,-1.57)	0.001	2.98	4	0.56	0
Creatinine increase	3	131	122	0.15(0.04,0.26)	0.006	1.86	2	0.40	0
Exclusion of [2]	2	84	66	0.20(0.07,0.33)	0.003	0.00	1	1.00	0
Complications									
Overall complications	14	1623	1147	1.86(1.28,2.70)+	0.001	29.35	13	0.006	56
Exclusion of [10]	13	1390	880	1.97(1.33,2.94)+	0.008	26.41	12	0.009	55
Intraoperative complications	4	641	594	2.02(0.22,18.51)+	0.53	14.48	3	0.002	79
Exclusion of [10]	3	411	345	5.93(1.59,22.07)+	0.008	1.04	2	0.60	0
Postoperative complications	7	758	735	2.97(2.13,4.14)+	< 0.001	10.53	6	0.10	43
Exclusion of [10]	6	525	468	3.15(1.75,5.69)+	0.0001	10.22	5	0.07	51

PN = partial nephrectomy; HR/WMD/OR = hazard ratio/weighted mean difference/odds ratio; CI = confidence interval; df = degrees of freedom; eGFR=estimated glomerular filtration rate;

hazard ratio; +Odds ratio.

Sensitivity Analysis

Sensitivity analyses were conducted by exclusion of the lowest scored study among included studies. All results except for the intraoperative complication rate were accordant with the primary outcomes. As to intraoperative complications, significant difference between the PN group and the cryoablation group was found after removing the study reported by Tanagho et al. [10] and the heterogeneity between studies decreased to nil (OR: 5.93; 95%CI: 1.59 to 22.07; p = 0.008; I² = 0%) (Table 2).

Publication Bias

A funnel plot of included studies reporting all-cause death was showed as Figure 11. All studies were located inside the 95%CI, which meant there was no obvious publication bias.



Discussion

PN has been recommended as management for T1 renal tumors in guideline [4]. For elderly patients or those with medical comorbidities, a higher surgical risk is accompanied with PN. Cryoablation has been regarded as an alternative to PN for T1 renal masses [8, 14]. However, whether cryoablation could lead to low levels of morbidities and surgical complications compared with those found in patients undergoing PN remained controversial and need to be proved further.

We report the first meta-analysis of studies comparing cryoablation to PN for clinical T1 renal masses, for which the role of cryoablation is still debated [10, 13, 14]. 17 eligible studies including a total number of 4666 patients can contribute to the ongoing discuss, providing a method for further research on this topic. Cryoablation for stage T1 renal tumors is a better choice to preserve nephron with less technically demand compared with PN. But the existing advantages are counterbalanced by the worse efficacy of cryoablation to PN in achieving effective oncological outcomes.

With regard to applying any new procedure, the safety is of primary significance to be taken into consideration. Both PN and cryoablation have been widely proved to be safe in the management of T1 renal masses [19]. PN which usually requires time-sensitive suturing with hilus clamping is more technically challenging than cryoablation [20]. According to our results a higher risk of incidences of both overall and postoperative complications was conveyed in PN group. Though no statistically significant difference about intraoperative complications between the two groups was found in original meta-analysis which was pooled with high heterogeneity, the result turned to significant in sensitivity analysis pooled without heterogeneity by eliminating the study reported by Tanagho et al. [10], in which many baseline characteristics were not comparable between two groups. The sensitivity analysis for intraoperative complications may hint the lower risk during the cryoablation procedure, which need to be proved further with more homogeneous studies.

Better oncological control is always pursued in management of tumors. PN achieved a better prognosis compared with cryoablation with respect to all variables including all-cause death, cancer-specific death, metastasis and local-recurrence. All results were pooled with nil or low heterogeneity. Selected patients who were considered more suitable to cryoablation were usually at elder age or with significant comorbidities [10, 13, 17, 19, 20, 26]. The selection bias contributed to the significant difference about oncological outcomes between two treatments. 4 of 11 studies providing information about oncological outcomes were at risk of a significant bias on tumor size [10, 17, 19, 26], but these results may be strengthened by the poorer prognostic feature that all mean tumor diameters in the PN group among these studies were significantly bigger. When matching these three factors, pooled results about all-cause death and cancer-specific death in the high quality study subgroup kept significantly different between the two treatments ((I²: 0%; HR: 0.43; 95%CI: 0.31 to 0.61; p < 0.001) and (I²: 0%; HR: 0.14; 95%CI: 0.03 to 0.67; p = 0.01), respectively), which implies that cryoablation brings poorer rates of overall survival and cancer-specific survival. Similar results about local-recurrence and metastasis have been summarized in study comparing laparoscopic cryoablation and laparoscopic PN reported by Tang et al. [15]. Rate of local-recurrence and metastases after renal cryoablation was deemed to be higher with more extended follow-up period [11]. Ultrasound was used to show the advancing edge of the ice ball without providing information on whether the lethal temperature has been achieved during cryoablation through laparoscopic or percutaneous approach [33], which may be the reason of poorer oncological outcomes after cryoablation.

To accomplish preservation of renal function was one of the main targets of nephron-sparing surgery [19, 34]. It is no doubt that PN and cryoablation are both effective in renal function protection [11, 35]. Cryoablation was associated with significantly less percent eGFR decrease and creatinine increase compared with PN. PN was usually conducted with hilum clamping, which was significantly associated with worse renal function [36]. Renal ischemia and actual parenchymal resection represent two separate mechanisms for potential renal function decrease during PN [37]. Cryoablation allows for direct visualization with laparoscopic approach or for real-time monitoring with percutaneous approach [13], which preserve functional nephron potentially [20, 33, 38].

Several limitations are worthy of discussion. The primary limitation of this systematic review and meta-analysis was that no randomized controlled trials were included, which made it hard to acquire enough data for meaningful results. Second, surgeries were performed by surgeons with different levels of surgical expertise and different choices of surgical approaches. Thirdly, patient allocation and treatment assignment were usually up to physician's attitude instead of randomized allocation, which led to a significant selection bias. Cryoablation was usually patient applied in elderly with significant comorbidities or those with a shorter life expectancy. Partial nephrectomy was chosen for renal masses with larger size. Next to last, the limited mean follow-ups in some studies resulted in follow-up bias. Finally, data about the incidences of complications was pooled with moderate or high heterogeneity using FE model.

Notable strengths of this systematic review are as follows. This is the first systematic review and meta-analysis comparing between cryoablation and PN for clinical T1 renal masses. The meta-analysis was conducted at an appropriate time, because more and more incidental renal tumors have been detected and enough studies have accumulated for inspection by meta-analytical methods. Studies were identified using multiple strategies, the methodological quality of the studies were evaluated according to strict inclusion and exclusion. Subgroup and sensitivity analysis were used to analyze the source of heterogeneity. Publication bias was not obvious with direct intuition.

Conclusions

Comparing with PN, cryoablation for clinical T1 renal tumors is associated with poorer oncological outcomes, but the existing disadvantages are accompanied by lower rate of overall and postoperative complications and superior renal functional preservation. For patients with imperative indications for nephron-sparing surgery who can't risk more invasive PN, cryoablation could be an attractive option. Owing to the inherent limitations of eligible studies, conclusions drawn from our meta-analyses should be interpreted cautiously and be confirmed further with well-designed randomized controlled trials with extensive follow-up length.

Abbreviations

PN: partial nephrectomy; eGFR: estimated glomerular filtration rate; HR: Hazard ratio; MD: mean difference; OR: odds ratio; CI: confidence interval; FE: fixed-effects.

Supplementary Material

Supplementary table. http://www.jcancer.org/v10p1226s1.pdf

Competing Interests

The authors have declared that no competing interest exists.

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