Comparison between cryoablation and irreversible electroporation of rabbit livers at a location close to the gallbladder

Jianying Zeng1,2, Zilin Qin1,2, Liang Zhou2, Gang Fang2, Jibing Chen2, Jialiang Li2, Lizhi Niu1,2, Bing Liang2, Kecheng Xu2

1 School of Medicine, Jinan University, Guangdong Province, Guangzhou, China
2 Fuda Cancer Hospital, Jinan University School of Medicine, Guangdong Province, Guangzhou, China


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Correspondence to: Lizhi Niu, Fuda Cancer Hospital, Jinan University School of Medicine, Guangzhou, China, 510665. E-mail: fudalab@163.com. Bing Liang, Fuda Cancer Hospital, Jinan University School of Medicine, Guangzhou, China, 510665. E-mail: 70404803@qq.com

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Jianying Zeng and Zilin Qin contributed equally to this work and share the first authorship.

Background. The ablation of liver tumors located close to the gallbladder is likely to lead to complications. The aim of this article is to compare the safety and efficacy of irreversible electroporation (IRE) and cryoablation in rabbit livers at a location close to the gallbladder.

Materials and methods. We performed cryoablation (n = 12) and IRE (n = 12) of the area of the liver close to the gallbladder in 24 New Zealand white rabbits in order to ensure gallbladder damage. Serum aminotransferase and serum bilirubin levels were measured before and after the ablation. Histopathological examination of the ablation zones in the liver and gallbladder was performed on the 7th day after the ablation.

Results. Seven days after the ablation, all 24 animals were alive. Gallbladder perforation did not occur in the IRE group; only mucosal epithelial necrosis and serous layer edema were found in this group. Gallbladder perforation occurred in four rabbits in the cryoablation group. Serum aminotransferase and serum bilirubin levels obviously increased in both groups by Day 3 and decreased gradually thereafter. The elevation in aminotransferase and bilirubin levels was greater in the cryoablation group than the IRE group. Pathological examination revealed complete necrosis of the liver parenchyma from the ablation center to the gallbladder in both groups, but bile duct and granulation tissue hyperplasia were observed in only the IRE group. Full-thickness gallbladder-wall necrosis was seen in the cryoablation group.

Conclusions. For ablation of the liver area near the gallbladder, IRE is superior to cryoablation, both in terms of safety (no gallbladder perforation in the IRE group) and efficacy (complete necrosis and rapid recovery in the IRE group).

Key words: cryoablation; irreversible electroporation; liver; gallbladder; ablation

Introduction

Hepatocellular carcinoma (HCC) is the sixth most prevalent malignancy worldwide, and a large proportion of patients with HCC are ineligible for tumor resection due to several factors, such as poor hepatic reserve (cirrhosis), multicentric tumors and extrahepatic disease. In HCC patients who are not candidates for surgery or liver transplantation, the National Comprehensive Cancer Network (NCCN) guidelines recommend the use of locoregional ablative methods, such as radiofrequency ablation (RFA), microwave ablation (MWA) and cryoablation. However, when the HCC lesion is located in close proximity to structures such as the major bile duct, gallbladder and diaphragm, ablation should be performed with caution in order to avoid damaging these structures. According to Lee et al., radiofrequency ablation of the area of the liver abutting the gallbladder can cause substantial...
complications, including gallbladder perforation, especially when the ablation is performed without maintaining a safe distance. Furthermore, it has been reported that ultrasound-guided percutaneous microwave ablation can be safely used for hepatic malignancies adjacent to the gallbladder, only when the ablation is performed under strict temperature monitoring. Cryoablation is achieved using the high-voltage-dependent release of argon and helium to induce a cycle of low temperature followed by thawing in order to cause physical damage and treat tumors. Because major vessels in close proximity to the tumor can absorb large amounts of heat during ablation (known as the “heat sink effect”), cryoablation can minimize vascular injury during the ablation of liver tumors. In addition, cryoablation is associated with good control, which allows the accurate targeting of the necrotic area and results in few side effects. This method might therefore be suitable for the ablation of liver lesions located near the gallbladder. Irreversible electroporation (IRE) is a novel ablation technology that utilizes short pulses of high-voltage electrical energy to induce tissue necrosis. IRE has many special advantages, including short ablation time, preservation of the internal structure of vital organs and lack of the heat/cold-sink effect. Although not yet included in the NCCN guidelines, IRE, owing to its intrinsic characteristics, might be a superior alternative to other ablation techniques for the ablation of tumors situated near the gallbladder. Therefore, IRE was selected for comparison with conventional cryoablation techniques. In this animal experimental study, we performed cryoablation and IRE of the liver area located 0.5 cm from the gallbladder in rabbits, and compared and evaluated the safety and efficacy of these two ablation techniques in order to identify the optimal ablation method for liver tumors located near the gallbladder.

Materials and methods

Experimental animals

In total, 24 healthy female New Zealand white rabbits, weighing (2.5 ± 0.2) kg each, were provided by the Animal Experimental Center of Jinan University. They were maintained in a clean, mechanically-ventilated environment at a constant temperature. The rabbits were randomly assigned to the cryoablation group (n = 12) and IRE group (n = 12). All animals were fasted for 24 h before surgery, and intravenous access was established for intraoperative drug administration. The study was approved by the Research Animal Care and Use Committee of Guangzhou Fuda Cancer Hospital (approval number, LL201402).

Cryoablation

A cryoablation system (Cryocare™, Endocare, Irvine, CA), composed of the main body, argon/helium gas container and cryoprobes (CRYO-42; Endocare), was used for cryoablation. Rabbits were anesthetized by the intramuscular injection of 44 mg/kg-bw ketamine, intubated and connected to a respirator for mechanically controlled respiration. Intraoperative anesthesia was maintained with 1.5% – 2% isoflurane.

General anesthesia was induced and maintained by the intramuscular injection of 44 mg/kg-bw ketamine. The surgeon had 17 years of experience in cryoablation. The rabbits were fixed in a supine position on the operating table and shaved. The skin was disinfected with iodine, and an abdominal incision was made to expose the liver and gallbladder under sterile conditions. The gallbladder was pressed on the liver, and a cryoprobe with a diameter of 1.4 mm was inserted into the liver until its tip was approximately 0.5 cm away from the gallbladder. The location of the tip was monitored using an ultrasound system (DP-50Vet, Mindray, Shenzhen, China; Figure 1A). When it was confirmed that the cryoprobe was connected to the main equipment, the argon gas was released to freeze the tissue. The cryoablation protocol was a double freeze-thaw cycle consisting of a 2-min freeze and 1-min thaw; this protocol ensured that part of the gallbladder was included within the ablation zone of the ice ball (Figure 1B). No animal died during the cryoablation process. After the cryoablation, the cryoprobe was retrieved. The liver was put back into the abdominal cavity, and conventional abdominal wall sutures were placed to complete the cryoablation. The cryoablation tract was filled in with a thrombin-soaked gelatin sponge and closed with a short suture.

IRE

General anesthesia was induced with an intramuscular injection of 44 mg/kg-bw ketamine and maintained with 1.5% – 2% isoflurane. Respiration was controlled using a respirator during the operation (tidal volume: 30 ml/time, respiratory rate: 30/min, oxygen concentration: 100%). Before the IRE operation, 0.12 mg/kg-bw pancuronium was
intramuscularly injected to block muscle contraction. IRE was performed using the AngioDynamics Nanoknife system (AngioDynamics, Latham, NY). The Nanoknife system consists of a main body and electrode probes (catalog no. 20400104, AngioDynamics), which were used for electroporation ablation. Monopolar probes were used, which could be changed to achieve 1 – 4 cm of electrode exposure. The size of the ablation zone created depended on the exposed length of the applicator as well as the distance between the probes. When the gallbladder was exposed, two monopolar probes were carefully inserted into the liver (Figure 2A). The probes were inserted to a depth of 0.5 cm into the liver parenchyma under ultrasound guidance (DP-50Vet, Mindray, Shenzhen, China), and were kept 1.5 cm apart and 0.5 cm away from the gallbladder (Figure 3). The monopolar probes were fastened with a spacer device. The ablation parameters consisted of nine groups of ten electrical...
pulses (total: 90 pulses), and each electrode pulse was 70-μs long; the output voltage was set at 1500 V/cm. This protocol ensured that the electric field covered part of the gallbladder (Figure 2B). Pulse delivery was synchronized with the R waves of the cardiac cycle in case of arrhythmia. One rabbit died during the IRE ablation because of tracheal intubation failure; we replaced this rabbit with another rabbit. After the IRE ablation, muscle relaxation was reversed with intramuscular injections of 5 μg/kg·bw neostigmine and 5 μg/kg·bw atropine.

Postprocedure care
The 24 experimental animals returned to consciousness under the monitoring of a senior veterinarian and were sent back to the specific pathogen-free experimental room, which was filled with fresh air and maintained at a constant temperature. All the rabbits were under meticulous care, and pain was managed with intramuscular buprenorphine (0.01 mg/kg) and oral meloxicam (0.4 mg/kg). In addition, the rabbits received daily intramuscular injections of 40 mg/kg·bw cephazolin for 3 days after the surgery in order to prevent infection. For 7 days after the surgery, the animals were observed by veterinarians, who assessed whether the rabbits had suffered any postoperative complications. Feeding was gradually restored over 3 days after the surgery in both groups of animals; however, both food and water intake were lower in the cryoablation group than in the IRE group.

Detection of serum aminotransferases and serum bilirubin
We collected blood (2 ml, from an ear marginal vein) 1 day before the operation and on days 1, 3, 5 and 7 after the ablation. The blood samples were centrifuged to obtain serum. Hepatic function was measured using an automatic biochemical analyzer (Hitachi-7100; Hitachi, Tokyo, Japan). Alanine aminotransferase/aspartate aminotransferase (ALT/AST) and serum bilirubin levels were determined with kits for in vitro diagnostic use (Biosino Biotechnology and Science Ltd, Beijing, China) by the velocity method.

Pathology
No animal died after the surgery. At 7 days after the ablation, all the animals were euthanized with an overdose of intravenous pentobarbital sodium (120 mg/kg, Sleepaway; Fort Dodge Animal Health, Fort Dodge, IA). The liver tissues from the ablation tract to the gallbladder were harvested, and the ablation lesions were cut and sectioned from the center. The maximum diameter was measured, and the lesions were photographed. Then, the tissue samples, including the abnormal and adjacent normal tissues, were fixed in 10% neutral buffered formalin, processed routinely for histology, embedded in paraffin, cut into 5-μm-thick slices and stained with hematoxylin and eosin. Pathological analyses were performed by an attending surgical pathologist.

Statistical analysis
All analyses were performed using GraphPad Prism 5 software (GraphPad Inc., San Diego, CA). The size of the ablation lesions and the serum ALT, AST and bilirubin levels were analyzed using the Student t-test. A P value of < 0.05 was considered to indicate a statistical difference, while P values of < 0.01 and < 0.001 indicated significant differences.

Results
Pathological appearance
In the cryoablation group, there were severe inflammatory adhesions between the ablation sites in the liver and gallbladder, the abdominal wall and the mesentery (12/12); the liver ablation zone was khaki-colored and oval. The lesion area had a narrow and sharp edge of light pink edematous tissue, and the ablation range obviously covered the gallbladder. There was a large gray area of necrosis adjacent to the gallbladder, and gallbladder perforation was found in four animals (Figure 4). In the IRE group, there were fewer inflammatory adhesions than in the cryoablation group (5/12 vs. 12/12).
The lesion area was yellow and oval with sharp edges. The ablated gallbladder tissue appeared grayish-white, but none of the 12 experimental animals showed gallbladder perforation (Figure 5). Details of the sizes of the ablation lesions are presented in Table 1; there were no differences in lesion size between the two groups. As gallbladder perforation was found in four animals in the cryoaablation group, we recorded the major diameter of the gallbladder lesions in only eight animals in this group.

There were no intact hepatic cells in the liver slices in both the cryoaablation and IRE groups. From the cryoaablation tract to the liver edge, the cryolesion appeared as a pink-stained area of disintegrating tissue that included hemorrhagic congestion and tissue necrosis, and it showed nuclear disappearance, no intact cellular structures, a band of local bleeding, and abundant inflammatory cell infiltration (Figure 6A). In the IRE group, the main pathological changes consisted of small bile duct, acute granulation tissue hyperplasia, rare congestion, hemorrhage band and a small amount of inflammatory cell infiltration (Figure 6B). The most conspicuous difference between the two groups was the appearance of the gallbladder: in the cryoaablation group, full-thickness gallbladder wall necrosis, muscular collapse and mucosal loss were found, and the necrotic area was large and uniformly stained red. In the IRE group, there was mucosal epithelial necrosis but no full-thickness gallbladder wall necrosis, and serous layer edema were largely concentrated in the area of the gallbladder wall near the liver.

Changes in serum aminotransferase and serum bilirubin levels

In both groups, the serum aminotransferase and bilirubin levels obviously increased on day 3, gradually decreased thereafter and recovered to normal in a week. At each time point, the serum aminotransferase and serum bilirubin levels were higher in the cryoaablation group than in the IRE group (Table 2).

<table>
<thead>
<tr>
<th>Ablation methods</th>
<th>Major diameter of liver lesions (cm)</th>
<th>P value</th>
<th>Major diameter (cm)</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Cryo</td>
<td>Gallbladder side (n = 12)</td>
<td>1.5 ± 0.4</td>
<td>3.8 ± 0.8</td>
<td>&gt; 0.05</td>
</tr>
<tr>
<td></td>
<td>Diaphragm side (n = 12)</td>
<td>3.8 ± 0.8</td>
<td>1.5 ± 0.4</td>
<td>n = 8</td>
</tr>
<tr>
<td>IRE</td>
<td>Gallbladder side (n = 8)</td>
<td>1.4 ± 0.4</td>
<td>4.1 ± 0.7</td>
<td>&gt; 0.05</td>
</tr>
<tr>
<td></td>
<td>Diaphragm side (n = 12)</td>
<td>0.8 ± 0.2</td>
<td>0.8 ± 0.2</td>
<td>n = 12</td>
</tr>
</tbody>
</table>

Table 1. Sizes of liver and gallbladder lesions at 7 days after the ablation (X ± s)
Discussion

In the study by Fairchild et al., the ice ball extended into the gallbladder lumen (up to a mean distance of 6 mm), but no cases of gallbladder perforation occurred, even when the iceball had extended as far as 1.8 cm into the gallbladder lumen. This may be explained by the theory that the bile within the gallbladder may dissipate the cooling effect of the cryoprobe around the gallbladder fossa. However, in our study, perforation occurred in one-third of the animals, and inflammatory adhesions occurred in all the animals in the cryoablation group. We attribute this difference to the fact that in our study, the ice ball extended to a far greater distance into the gallbladder lumen, and thus, the cryodamage to the gallbladder was more severe. Furthermore, as more of the gallbladder wall was cryoablated, the “dissipating effect” of the bile was extremely weakened, and a local self-limited inflammatory response resulted in gallbladder wall edema and reactive fluid accumulation, which in turn increased the severity of the gallbladder damage.

In contrast, IRE caused a color change in the gallbladder, but no perforation occurred 7 days after ablation. In addition, full-thickness wall necrosis occurred only in the cryoablation group; in the IRE group, the damage to the gallbladder was confined to the mucous layer. This is because IRE is a promising, non-thermal technology that takes advantage of electric fields instead of thermal energy to induce cell death. The most distinctive feature of IRE is that it has little influence on the adjacent large vessels and can preserve the collagen matrix of tissue structures. In addition, it does not lead to significant ductal damage if the electrodes are not placed very close to the ducts. In our study, the IRE probes were 0.5 cm away from the gallbladder, and the damage to the gallbladder caused by IRE was acceptable. Thus, our findings indicate that IRE is a potentially safe modality for the ablation of hepatic tumors adjacent to the gallbladder.

Furthermore, although the lesion sizes and changes in the serum aminotransferase and serum bilirubin levels were similar in the two groups, the damage to the liver cells was more severe in the cryoablation group. This phenomenon may be explained as follows: Cryodamage mainly leads to cell necrosis, and secondary injuries can be mediated by the resultant inflammatory reaction. According to current studies, thermal injury may require time to cause gallbladder perforation and perforations are usually found in the acute phase, not the immediate phase. In contrast, electrical injury (IRE) mainly causes cell apoptosis and tissue damage, and repair following IRE is rapid. At 7 days after cryoablation, tissue destruction was still ongoing (areas of cell disruption with hemorrhage), while at the same time point, tissue repair was obvious in the IRE group (areas of bile duct hyperplasia).

Collectively, the above findings indicate that IRE is unlikely to destroy the gallbladder wall and offers the advantage of faster repair of the damaged tissue. Thus, in this study, IRE was found to be superior to cryoablation in terms of protecting the gallbladder, and was a safer procedure than cryoablation.

Complete necrosis of tumor cells with no residual tumor is the main goal of ablation. During cryoablation, ice ball formation can be precisely monitored using ultrasonography or computed tomography. It is impossible to induce tumor cell necrosis in the periphery of the ice ball because the peripheral temperature of the ice ball is 0°C, and the temperature that is lethal to cells is at least –20°C. To ensure efficacious cryoablation, the edge of the ice ball is usually kept 1 cm beyond the edge of tumor, and the double freeze-thaw cycle protocol is used. In our study, the efficacy of

<table>
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<tr>
<th>Hepatic function</th>
<th>Ablation method</th>
<th>d-1</th>
<th>d-1</th>
<th>d-3</th>
<th>d-7</th>
</tr>
</thead>
<tbody>
<tr>
<td>ALT (U/L)</td>
<td>IRE</td>
<td>38.35 ± 3.22</td>
<td>167.05 ± 19.50</td>
<td>214.83 ± 10.37</td>
<td>119.60 ± 12.99</td>
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<td></td>
<td>Cryo</td>
<td>39.20 ± 3.90</td>
<td>195.03 ± 9.75**</td>
<td>289.58 ± 12.81**</td>
<td>142.35 ± 57.85*</td>
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<tr>
<td>AST (U/L)</td>
<td>IRE</td>
<td>34.10 ± 0.86</td>
<td>47.64 ± 0.69</td>
<td>60.21 ± 2.55</td>
<td>46.78 ± 1.71</td>
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<tr>
<td></td>
<td>Cryo</td>
<td>35.26 ± 2.45</td>
<td>60.11 ± 3.44**</td>
<td>79.12 ± 2.58**</td>
<td>52.80 ± 3.41*</td>
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<tr>
<td>Serum bilirubin</td>
<td>IRE</td>
<td>8.08 ± 2.19</td>
<td>26.01 ± 3.68</td>
<td>35.15 ± 5.70</td>
<td>29.09 ± 3.27</td>
</tr>
<tr>
<td>(μmol/L)</td>
<td>Cryo</td>
<td>7.59 ± 2.01</td>
<td>32.72 ± 4.08**</td>
<td>37.34 ± 6.14*</td>
<td>35.88 ± 4.28**</td>
</tr>
</tbody>
</table>

*P < 0.05 and **P < 0.01, compared with the IRE group.
cryoablation of the liver area near the gallbladder reached the expected levels, as determined based on the appearance (gray-red lesion spread to the gallbladder) as well as histopathological examination (no live cells between the cryozone and gallbladder) of the cryozone. IRE ablation has been reported to achieve nearly 100% necrosis in the ablation zone at 24 h in a swine liver model. The effectiveness of IRE ablation over a longer time is one of the focuses of this study. Seven days after IRE ablation, not only were the target cells destroyed completely, but the area of bile duct hyperplasia was far greater than the area of pink, disintegrating tissue. All in all, although there was a great difference between the abilities of the two therapies to destroy the liver cells, the long-term effects of both treatments were good.

The major limitation of this article is the lack of short- and long-term studies. Analysis of the pathological appearance of the gallbladder at different time points after ablation would help provide a stronger basis for the clinical application of the two ablation methods. Future experiments should use cholangiography to observe changes in the gallbladder and determine the integrity of the main bile duct.

In conclusion, we compared the safety and efficacy of cryoablation and IRE in rabbit livers, at a location close to the gallbladder. IRE provided complete ablation of the target tissue and completely retained the entire gallbladder structure. In the case of liver tumors situated close to the gallbladder, cryoablation must be performed carefully to avoid direct trauma to the gallbladder. The comparison in this article has a certain meaning in terms of the selection of instruments for liver tumor ablation. However, the clinical value of our findings and long-term curative effect of IRE remain to be confirmed.

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