Transcatheter intraarterial perfusion MRI approaches to differentiate reversibly electroporated penumbra from irreversibly electroporated zones in rabbit liver

Abstract

Rationale and Objectives: To investigate whether transcatheter intraarterial perfusion (TRIP) magnetic resonance imaging (MRI) can differentiate reversible electroporation (RE) zones from irreversible electroporation (IRE) zones immediately after IRE procedure in the rabbit liver.

Materials and Methods: All studies were approved by the institutional animal care and use committee and performed in accordance with institutional guidelines. A total of thirteen healthy New Zealand White rabbits were used. After selective catheterization of the hepatic artery under X-ray fluoroscopy, we acquired TRIP-MRI at 20 minutes post-IRE using 3 mL of 5% intraarterial gadopentetate dimeglumine. Semi-quantitative (peak enhancement, PE; time to peak, TTP; wash-in slope, WIS; areas under the time-intensity curve, AUT, over 30, 60, 90, 120, 150, and 180 seconds after the initiation of enhancement) and quantitative (K_{trans}, v_e, and v_p) TRIP-MRI parameters were calculated. The relationships between TRIP-MRI parameters
and histological measurements and the differential ability of TRIP-MRI parameters was assessed.

**Results:** PE, AUT\textsubscript{60}, AUT\textsubscript{90}, AUT\textsubscript{120}, AUT\textsubscript{150}, AUT\textsubscript{180}, K\textsuperscript{trans}, and v\textsubscript{e} were significantly higher in RE zones than in IRE zones (all \( P < 0.05 \)), and AUC for these parameters ranged from 0.91 (95% CI, 0.80, 1.00) to 0.99 (95% CI, 0.98, 1.00). There was no significant difference in AUC between any two parameters (\( Z, 0 \text{-} 1.47; P, 0.14 \text{-} 1.00 \)). Hepatocyte apoptosis strongly correlated with PE, AUT\textsubscript{60}, AUT\textsubscript{90}, AUT\textsubscript{120}, AUT\textsubscript{150}, AUT\textsubscript{180}, K\textsuperscript{trans}, and v\textsubscript{p} (the absolute value \( r, 0.6 \text{-} 0.7, \text{ all } P < 0.0001 \)).

**Conclusion:** AUT\textsubscript{150} or AUT\textsubscript{180} could be a potential imaging biomarker to differentiate RE from IRE zones, and TRIP-MRI permits to differentiate RE from IRE zones immediately after IRE procedure in the rabbit liver.

**Keywords:** Irreversible electroporation, liver, rabbit, reversible electroporation, transcatheter intra-arterial perfusion MRI
Abbreviations:

TRIP = transcatheter intraarterial perfusion

IRE = irreversible electroporation

RE = reversible electroporation

TUNEL = transferase dUTP nick end labeling

TIC = time-intensity curve

PE = peak enhancement

TTP = time to peak

WIS = wash-in slope

AUT = area under the time-intensity curve
Introduction

Image guided ablation techniques are standard care for early stage liver cancer patients with surgically unresectable tumors or on waiting list for transplantation (1). However, thermal ablation techniques can damage healthy adjacent tissues due to a lack of clear margins between treated and untreated tissues, irreversible electroporation ablation can generate distinct margins between treated and untreated tissues by delivering electrical pulses to induce cell death through permanent cell membrane permeabilization (2), allowing improved targeted treatment of tumors while minimizing adverse effects. Irreversible electroporation (IRE) is a new non-thermal ablation technique with significant advantages over conventional thermal ablative approaches, such as preserving the integrity of blood vessels and bile ducts and not being affected by the local blood flow (3-6). Strong IRE pulses produce both a central ablation zone containing complete tissue necrosis (defined as irreversibly electroporated zone, IRE zone) and a surrounding zone containing viable tissue (defined as reversibly electroporated zone, RE zone) (7-9). These findings have been confirmed by histologic results (7-9). As RE zone remains viable, there is a risk of tumor recurrence or local tumor progression, which presents significant concerns for more widespread clinical application of this treatment (6).
Advanced MRI methods permit a more accurate prediction of IRE volumes and allow immediate assessment of tissue response for patient-specific optimization of the treatment protocol (7, 10, 11). However, previous studies focused only on correlating imaging findings and IRE histology to discriminate between tumor and normal tissue regions and not on predicting the margins of IRE and RE zones.

Transcatheter intraarterial perfusion (TRIP)-MRI is used clinically to measure local tumor perfusion by selective intraarterial (IA) catheter directed contrast agent injections (12, 13). Continuous iterative volumetric imaging over time of TRIP-MRI incorporates the benefits of high spatial and temporal resolution with intraprocedural monitoring for objective perfusion quantification (14).

The purpose of our study was to investigate whether TRIP-MRI can differentiate RE zones from IRE zones immediately after the IRE procedure in the rabbit liver.

**Materials and Methods**

All experiments were approved by our institutional animal care and use committee and performed in accordance with National Institutes of Health guidelines.
Animal Model

Thirteen healthy New Zealand White rabbits (Covance, Denver, PA) were used. Basic anesthesia was induced by intramuscular injection of ketamine (100 mg/kg; Fort Dodge Animal Health, Fort Dodge, IA) and xylazine (5 mg/kg; Abbott Laboratories, North Chicago, IL). Rabbits were mechanically ventilated and maintained by inhalation of isoflurane (2-3% isoflurane with 100% oxygen, 3L/min) through a breathing mask during digital subtraction angiography (DSA), IRE, and MRI.

Conventional DSA

Conventional DSA was performed using a C-arm unit (PowerMobil; Siemens Medical Solutions, Erlangen, Germany). The femoral artery access was obtained through surgical cutdown. A 2-F catheter (JB1; Cook Medical, Bloomington, IN) was advanced over a 0.018-inch diameter guidewire into hepatic arteries. DSA of hepatic arteries was performed using an iodinated contrast agent (iohexol, Omnipaque 350; Amersham, Princeton, NJ). All of the rabbit hepatic arteries were successfully catheterized with DSA.

IRE Procedures

The left median lobe of the liver was exposed after median laparotomy. Two paralleled
MRI-compatible needle electrodes with 1-cm spacing (platinum-iridium materials; Microprobes, Gaithersburg, MD) were inserted into the center of the left median lobe with 1-cm depth. Next, the electrodes were connected to the IRE generator (ECM830; Harvard Apparatus, Holliston, MA) for application of 2000-V square wave pulses (number of pulses, 10; each pulse duration, 100 μs; the interval between pulses, 100 ms) to the targeted tissue (7, 15).

**MR Imaging**

MRI examinations were performed twenty minutes after IRE procedure using a 3T MRI scanner (MAGNETOM Skyra, Siemens Medical Solutions, Erlangen, Germany) with a 15-channel knee coil. Conventional MRI was performed before TRIP-MRI, and acquisition parameters are listed in Table 1. TRIP-MRI was repeatedly sampled for 250 times with 2.8 s sampling rate. 56 s after initiating TRIP-MRI, 3 ml of 5% gadopentetate dimeglumine (Magnevist; Bayer Schering Pharma, Whippany, NJ) was manually injected via the intra-arterial catheter over 6 s followed by a 3-ml saline flush at the same flow rate. Following TRIP-MRI, post-contrast T1-weighted (T1W) imaging was performed using the same sequence as pre-contrast T1W imaging.
**Histologic Evaluation**

Immediately after MRI, rabbits were euthanized by intravenous injection of Euthasol (200 mg/kg, Virbac AH Inc., Fort Worth, TX), and liver tissues were harvested. The sections, including the entire ablated zone (IRE and RE zone) and the surrounding normal liver tissue, were sampled along an orientation parallel to the electrode array. Liver samples were stained with hematoxylin-eosin (H&E) and terminal deoxynucleotidyl transferase dUTP nick end labeling (TUNEL) which was used as a marker of hepatocyte apoptosis. The RE and IRE zones were first identified on HE-stained slices. On the TUNEL-stained slices, hepatocyte apoptosis was quantified as the percentage of TUNEL-positive cells over the total number of cells in 10 random regions (× 200 magnification) on each slice.

**Image Analysis**

All the MRI images were analyzed by consensus of two radiologists using Jim software (version 7.0, Xinapse Systems, Essex, UK). Using post-contrast T1W images as reference (8, 11), three regions of interest (ROIs) were drawn within the hypo-intensity center (the IRE zone), the peripheral hyper-intensity region (the RE zone) on the axial section parallel to the electrode array, and in the normal liver parenchyma of the right lobe on TRIP-MRI images (7-9). A time-
intensity curve (TIC) was generated for each ROI, and the semi-quantitative parameters were calculated: peak enhancement (PE, the maximal signal intensity minus baseline signal intensity), time to peak (TTP, the time difference between PE and the initiation of enhancement), wash-in slope (WIS, the maximal upslope reaching PE), and the areas under the TIC (AUT) over seconds (AUT\textsubscript{30}, AUT\textsubscript{60}, AUT\textsubscript{90}, AUT\textsubscript{120}, AUT\textsubscript{150} and AUT\textsubscript{180}) after the initiation of enhancement (16). Single-input extended Tofts model was used to perform quantitative measurements (17). The maps of the following parameters were generated: $K^{\text{trans}}$, the volume transfer constant from the plasma to the extra-vascular extra-cellular space (EES); $v_c$, the EES volume fraction; $v_p$, the blood plasma volume fraction (18). The mean values of each quantitative parameter ($K^{\text{trans}}$, $v_c$, and $v_p$) were measured in the same ROIs that were drawn for the TIC analysis.

Statistical Analysis

Statistical analysis was performed using SPSS (version 22.0; SPSS, Chicago, IL). The results are presented as mean ± standard deviation (SD). Differences in TRIP-MRI parameters and hepatocyte apoptosis among the normal liver parenchyma, the RE and IRE zones were determined by the Kruskal-Wallis test followed by pairwise comparisons. The diagnostic
performance of TRIP-MRI parameters in differentiating RE from IRE zones was assessed by calculating the area under the receiver operating characteristic (ROC) curves (AUC), optimal cut-off values, sensitivity, specificity, positive and negative predictive values. The AUC for TRIP-MRI parameters were compared by Z test. The relationship between TRIP-MRI parameters and hepatocyte apoptosis was assessed by Pearson correlation coefficients ($r$). $P < 0.05$ was considered to indicate a significant difference in all statistical tests.

Results

MR Imaging

Representative MRI images of the liver after IRE procedure and the corresponding TICs derived from TRIP-MRI for the normal liver parenchyma, the RE and IRE zones were shown in Figure 1. The entire ablated regions were slightly hypo- or iso-intense on pre-contrast T1W images and hyper-intense on T2-weighted (T2W) images. On post-contrast T1W images, the entire ablated regions showed heterogeneous enhancement with the exception of a non-enhancing center and peripheral rim enhancement. The TICs and the maps of $K^{\text{trans}}$, $v_c$, and $v_p$ clearly depicted heterogeneous perfusion of the entire ablated zone.

TRIP-MRI parameters for the normal liver parenchyma, the RE and IRE zones were listed
in Table 2, Figure 2, and Figure 3. The results of the ROC curve analysis for TRIP-MRI parameters differentiating RE from IRE zones were summarized in Table 3. There was no significant difference in AUC between any two parameters (Z, 0-1.47; P, 0.14-1.00). $\text{AUT}_{150}$ and $\text{AUT}_{180}$ had the greatest ability to differentiate RE zone from IRE zone, with AUC of 0.99 (95% CI, 0.98, 1.00) for both, and an optimal cut-off value of 1066 for $\text{AUT}_{150}$ and 1343 for $\text{AUT}_{180}$. Both $\text{AUT}_{150}$ and $\text{AUT}_{180}$ had 96% (25/26) accuracy, 100% (13/13) sensitivity, and 92% (12/13) specificity for all.

**Histologic Findings**

Representative H&E- and TUNEL-stained images of the normal IRE treated, and untreated liver parenchyma are shown in Figure 4.

On TUNEL-stained images, the average hepatocyte apoptosis (%) in the normal liver parenchyma, the RE zone, and IRE zone were $0.1 \pm 0.08$, $3.0 \pm 1.3$, and $23.4 \pm 6.7$, respectively. There were significant differences in hepatocyte apoptosis among the normal liver parenchyma, RE zone and IRE zone (RE vs IRE, $P = 0.01$; IRE vs normal liver, $P < 0.0001$; RE zones vs normal liver, $P = 0.01$). The hepatocyte apoptosis strongly correlated with PE ($r = -0.74$, $P < 0.0001$), $\text{AUT}_{60}$ ($r = -0.63$, $P < 0.0001$), $\text{AUT}_{90}$ ($r = -0.65$, $P < 0.0001$), $\text{AUT}_{120}$ ($r = -0.68$, $P < 0.0001$), and $\text{AUT}_{150}$ ($r = -0.71$, $P < 0.0001$).
0.0001), AUT_{150} (r = -0.69, P < 0.0001), AUT_{180} (r = -0.71, P < 0.0001), K^{\text{trans}} (r = -0.63, P < 0.0001), and v_p (r = -0.66, P < 0.0001). There was moderate correlation of hepatocyte apoptosis with TTP (r = 0.40, P = 0.01), WIS (r = -0.54, P < 0.0001), and AUT_{30} (r = -0.60, P < 0.0001). A weak correlation was observed between hepatocyte apoptosis and v_e (r = -0.36, P = 0.02).

**Discussion**

Our study demonstrated that TRIP-MRI not only depicts a reduction in liver tissue perfusion immediately after IRE procedures but also enables clear discrimination between IRE and RE zones immediately following IRE procedures in a rabbit liver.

Guo et al. evaluated contrast-enhanced MRI to predict the size of IRE zones in the rat liver tissues (7). The contrast agents were injected intramuscularly before IRE, and MRI examinations were performed 2 hours after IRE without dynamic phase acquisition. Felker et al. described a three-layer histologic structure of porcine liver tissues with corresponding MRI features obtained 50 hours after IRE (10). Our study focused on the immediate response of liver tissues after IRE using not only conventional MRI to obtain the morphologic information, but also TRIP-MRI to measure the functional changes. The entire ablated regions appeared iso-
intense on pre-contrast T1W and hyper-intense on T2W images. Post-contrast, heterogeneous enhancement was shown on T1W images, with a non-enhancing central core and a peripheral rim enhancement.

TRIP-MRI is a serial iterative volumetric perfusion technique and can intra-procedurally measure tissue perfusion changes (13). Previous studies compared TRIP-MRI with the conventional intra-venous contrast perfusion MRI in a rabbit VX2 liver tumor model and showed that TRIP-MRI at a 60% reduction in contrast agent dose identified more liver tumors than the IV contrast perfusion MRI. In our study, 5% gadopentetate dimeglumine was used for TRIP-MRI compared with 100% agent for intravenous contrast perfusion MRI (19, 20). In our study, the TICs showed that the wash-in and wash-out time was relatively short, which was consistent with the previous studies (19, 20). Our study also showed that PE and AUT over more than 60 s after enhancement initiation were significantly decreased from the normal liver parenchyma to RE-zones and from RE zones to IRE zones; these changes strongly correlated with histologic results. As for the quantitative parameters, $K_{\text{trans}}$ was highest in the normal liver parenchyma, followed by RE and IRE zones. $v_e$ in RE zones was significantly higher than in IRE zones or the normal liver parenchyma, likely due to the presence of interstitial edema and
congestion in RE zones. However, there was only a weak correlation between $v_e$ and hepatocyte apoptosis. Among the quantitative parameters, $K^{\text{trans}}$ had a good diagnostic performance in differentiating RE from IRE zones with 85% accuracy, 77% sensitivity, and 92% specificity.

It’s worth noting that, $\text{AUT}_{150}$ and $\text{AUT}_{180}$ had the greatest ability among all the TRIP-MRI parameters to differentiate RE from IRE zones with 96% accuracy, 100% sensitivity, and 92% specificity. Our results demonstrated that semi-quantitative parameter $\text{AUT}_{150}$ or $\text{AUT}_{180}$ could be a potential imaging biomarker to differentiate RE from IRE zones, and easy-to-implement in clinical practice as they can be calculated without using advanced pharmacokinetic modeling.

Our findings will provide fundamental preclinical validation for clinical translation.

The main limitation of our study is that TRIP-MRI was only performed at an immediate time point after IRE, and future studies will investigate delayed findings of TRIP-MRI and corresponding histologic changes after IRE to more comprehensively evaluate TRIP-MRI for differentiation of ablation zones after IRE.

In conclusion, TRIP-MRI permits to differentiate RE from IRE zones immediately after IRE procedure in the rabbit liver.
References


### Tables

**Table 1.** Sequences and parameters of conventional T1W, T2W, and TRIP MRI

<table>
<thead>
<tr>
<th>Parameters</th>
<th>T2W (TSE)</th>
<th>T1W (FLASH)</th>
<th>T2W (TSE)</th>
<th>TRIP MRI (GRE)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Orientation</td>
<td>Coronal</td>
<td>Axial</td>
<td>Axial</td>
<td>Axial</td>
</tr>
<tr>
<td>Trigger</td>
<td>Respiration</td>
<td>None</td>
<td>Respiration</td>
<td>None</td>
</tr>
<tr>
<td>TR/TE (ms)</td>
<td>1769–4341/39</td>
<td>200/2.93</td>
<td>2247–4384/39</td>
<td>2.60/1.2</td>
</tr>
<tr>
<td>FOV (mm²)</td>
<td>180 × 180</td>
<td>180 × 180</td>
<td>180 × 180</td>
<td>128 × 112</td>
</tr>
<tr>
<td>Matrix</td>
<td>192 × 192</td>
<td>224 × 224</td>
<td>192 × 192</td>
<td>128 × 112</td>
</tr>
<tr>
<td>Number of slices</td>
<td>17</td>
<td>17</td>
<td>17</td>
<td>20</td>
</tr>
<tr>
<td>Slice thickness/gap (mm)</td>
<td>2/0.6</td>
<td>2/0.6</td>
<td>2/0.6</td>
<td>2/0.6</td>
</tr>
<tr>
<td>Flip angle (degrees)</td>
<td>150</td>
<td>70</td>
<td>150</td>
<td>8</td>
</tr>
<tr>
<td>Bandwidth (Hz/pixel)</td>
<td>221</td>
<td>350</td>
<td>221</td>
<td>980</td>
</tr>
</tbody>
</table>

FLASH, fast low angle shot; FOV, field of view; GRE, gradient echo; TE, echo time; TR, repetition time; TRIP, transcatheter intra-arterial perfusion; TSE, turbo spin echo.
Table 2. Comparison of TRIP MRI parameters among IRE, RE zones and normal liver parenchyma

<table>
<thead>
<tr>
<th>Parameters</th>
<th>IRE Zone</th>
<th>RE Zone</th>
<th>Normal Liver</th>
<th>( P \text{ Value}^a )</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td>IRE vs RE</td>
</tr>
<tr>
<td>PE</td>
<td>15 ± 8</td>
<td>39 ± 13</td>
<td>110 ± 17</td>
<td>0.025</td>
</tr>
<tr>
<td>TTP</td>
<td>450 ± 86</td>
<td>541 ± 106</td>
<td>17 ± 5</td>
<td>0.29</td>
</tr>
<tr>
<td>WIS</td>
<td>0.04 ± 0.02</td>
<td>0.2 ± 0.2</td>
<td>6.3 ± 1.7</td>
<td>0.07</td>
</tr>
<tr>
<td>AUT(_{30})</td>
<td>58 ± 67</td>
<td>231 ± 145</td>
<td>2381 ± 532</td>
<td>0.06</td>
</tr>
<tr>
<td>AUT(_{60})</td>
<td>116 ± 126</td>
<td>702 ± 354</td>
<td>4637 ± 1065</td>
<td>0.02</td>
</tr>
<tr>
<td>AUT(_{90})</td>
<td>193 ± 237</td>
<td>1226 ± 564</td>
<td>6344 ± 1428</td>
<td>0.017</td>
</tr>
<tr>
<td>AUT(_{120})</td>
<td>281 ± 319</td>
<td>1843 ± 791</td>
<td>8049 ± 1772</td>
<td>0.014</td>
</tr>
<tr>
<td>AUT(_{150})</td>
<td>376 ± 421</td>
<td>2535 ± 1054</td>
<td>9616 ± 2163</td>
<td>0.012</td>
</tr>
<tr>
<td>AUT(_{180})</td>
<td>460 ± 500</td>
<td>3193 ± 1307</td>
<td>10962 ± 2484</td>
<td>0.012</td>
</tr>
<tr>
<td>K(_{\text{trans}})</td>
<td>0.1 ± 0.2</td>
<td>0.8 ± 0.6</td>
<td>2.9 ± 0.9</td>
<td>0.027</td>
</tr>
<tr>
<td>(v_e)</td>
<td>17 ± 17</td>
<td>58 ± 25</td>
<td>26 ± 14</td>
<td>&lt; 0.0001</td>
</tr>
<tr>
<td>(v_p)</td>
<td>2.4 ± 2.4</td>
<td>8.7 ± 7.4</td>
<td>22 ± 4</td>
<td>0.05</td>
</tr>
</tbody>
</table>

Data was represented as mean ± standard deviation.

AUT, area under the time-intensity curve; PE, peak enhancement, TTP, time to peak, WIS, wash-in slope.

* \( p < 0.05 \) was considered to indicate a significant difference.
Table 3. ROC Curve Analysis of TRIP MRI Parameters in Differentiating RE from IRE Zones

<table>
<thead>
<tr>
<th>Parameters</th>
<th>AUC*</th>
<th>Cut-off Value</th>
<th>Accuracy (%)†</th>
<th>Sensitivity (%)‡</th>
<th>Specificity (%)‡</th>
<th>PPV (%)‡</th>
<th>NPV (%)‡</th>
</tr>
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<tr>
<td>PE</td>
<td>0.95</td>
<td>(0.88, 1.00)</td>
<td>24.4</td>
<td>88.5 (23/26)</td>
<td>92.3 (12/13)</td>
<td>85.7</td>
<td>91.7</td>
</tr>
<tr>
<td>AUT60</td>
<td>0.96</td>
<td>(0.90, 1.00)</td>
<td>308.9</td>
<td>88.5 (23/26)</td>
<td>92.3 (12/13)</td>
<td>91.7</td>
<td>85.7</td>
</tr>
<tr>
<td>AUT90</td>
<td>0.98</td>
<td>(0.92, 1.00)</td>
<td>520.5</td>
<td>96.2 (25/26)</td>
<td>100 (13/13)</td>
<td>92.9</td>
<td>100</td>
</tr>
<tr>
<td>AUT120</td>
<td>0.99</td>
<td>(0.96, 1.00)</td>
<td>783.8</td>
<td>96.2 (25/26)</td>
<td>100 (13/13)</td>
<td>92.9</td>
<td>100</td>
</tr>
<tr>
<td>AUT150</td>
<td>0.99</td>
<td>(0.98, 1.00)</td>
<td>1066</td>
<td>96.2 (25/26)</td>
<td>100 (13/13)</td>
<td>92.9</td>
<td>100</td>
</tr>
<tr>
<td>AUT180</td>
<td>0.99</td>
<td>(0.98, 1.00)</td>
<td>1343</td>
<td>96.2 (25/26)</td>
<td>100 (13/13)</td>
<td>92.9</td>
<td>100</td>
</tr>
<tr>
<td>Ktrans</td>
<td>0.94</td>
<td>(0.85, 1.00)</td>
<td>0.28</td>
<td>84.6 (22/26)</td>
<td>76.9 (10/13)</td>
<td>90.9</td>
<td>80 (12/15)</td>
</tr>
<tr>
<td>ve</td>
<td>0.91</td>
<td>(0.80, 1.00)</td>
<td>45.2</td>
<td>80.8 (21/26)</td>
<td>69.2 (9/13)</td>
<td>92.3</td>
<td>90 (9/10)</td>
</tr>
</tbody>
</table>

AUC, area under the ROC curve; AUT, area under the time-intensity curve; NPV, negative predictive value; PE, peak enhancement; PPV, positive predictive value.

* Data in parentheses are 95% confidence interval.
† Data in parentheses are raw data.
**Figures**

**Figure 1:** Representative MRI images of the liver and corresponding time-intensity curves immediately after IRE in a single rabbit. The entire ablated zone shows hyper-intensity on the T2W image (a, arrowheads), slight hypo-intensity on the pre-contrast T1W image (b, arrowheads), and heterogeneous enhancement with a non-enhancement center and an obvious peripheral enhancement on the post-contrast T1W image (c). The maps of $K_{\text{trans}}$ (d), $v_e$ (e) and $v_p$ (f) overlapped on the post-contrast T1W image clearly depicted heterogeneous perfusion of the entire ablated zone. The time-intensity curves (g) derived from the normal liver parenchyma, the RE and IRE zone show a unique pattern for each zone.
Figure 2: The Tukey boxplot shows the areas under the time-intensity curves (AUT) over 30, 60, 90, 120, 150 and 180 seconds after the initiation of enhancement for the IRE (red, n = 13), RE zones (green, n = 13) and the normal liver parenchyma (blue, n = 13). AUT\textsubscript{60}, AUT\textsubscript{90}, AUT\textsubscript{120}, AUT\textsubscript{150}, and AUT\textsubscript{180} had the ability to differentiate RE from IRE zones. The box plot displays the full range of variation (from min to max), and the dots indicate discrete values. Statistically significant differences are marked by stars (* \( P < 0.05 \), ** \( P < 0.0001 \)).
Figure 3: Scatter dot plots show peak enhancement (a), time to peak (b), wash-in slope (c), $K_{\text{trans}}$ (d), $v_e$ (e) and $v_p$ (f) for the IRE zones ($n = 13$), RE zones ($n = 13$) and the normal liver parenchyma ($n = 13$). Peak enhancement, $K_{\text{trans}}$ and $v_e$ had the ability to differentiate RE from IRE zones. Statistically significant differences are marked by stars (* $P < 0.05$, ** $P < 0.0001$).

The larger horizontal bar indicates the mean value, error bars indicate standard deviations, and dots indicate individual values.
**Figure 4:** Representative histologic images of the IRE zone (a, d), the RE zone (b, e) and the normal liver parenchyma (c, f). In the IRE zone, there is obvious hepatocyte necrosis and hemorrhage on the HE-stained image (a), and apoptotic cells (brown stain) were about 22.7% of the total cells on the TUNEL-stained image (d). In the RE zone, there is hepatocyte swelling with vacuoles, occasional hepatocyte necrosis and congestion on the HE-stained image (b), and apoptotic cells (brown stain) were about 3% of the total cells on the TUNEL-stained image (e). In the normal liver parenchyma, there is no obvious abnormal feature on the HE-stained image (c), and no apoptosis is visible on the TUNEL-stained image (f). Scale bar, 100 μm for all the images.