

# THE ROLE OF DIFFUSION WEIGHTED MRI IN THE DIFFERENTIATION BETWEEN BENIGN AND MALIGNANT HEPATIC FOCAL LESION

#### **Thesis**

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By

#### **Mohammed Uday Hatim**

M..B., B.Ch Faculty of Medicine - University of Basrah

Under Supervision of

# A. Prof. Dr. Amr Mahmoud Abdelsamed

Assistant Professor of Radiodiagnosis Faculty of Medicine – Ain Shams University

#### A. Prof. Dr. Remon Zaher Elia

Assistant Professor of Radiodiagnosis Faculty of Medicine - Ain Shams University

Faculty of Medicine
Ain shams University
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#### **Abstract**

**Introduction:** Early detection and diagnosis of hepatic focal lesions are an important step in clinical work, which would allow effective surgical or mini-invasive therapy.

**Aim of the Work:** The use of MR Diffusion imaging with both low and high B values to detect and differentiate between benign and malignant hepatic focal lesion.

**Patient and Methods:** This study were includes 30 patients. They are El-Demerdash hospital patients with hepatic focal lesions. Patients underwent US or CT before MR examination.

**Result:** Thirty patients were included in this study, 20 males and 10 females. There were 24 primary hepatic focal lesions, (36.7% HCC, 3.3% focal nodular hyperplasia, 3.3% cysts, 13.3% hemangiomas, 6.7% cholangiocarcinoma, 16.7% regeneration nodule) and 6 metastatic lesions.

**Conclusion:** We hope to use DWI to be helpful in the characterization of focal liver lesions, especially if the lesions show non classic appearance of contrast enhancement in Triphasic CT study and in patients with renal insufficiency with inability to use contrast enhancement.

**Recommendations:** In our opinion, DWI is a useful adjunct to routine liver imaging (i.e. used as an additional sequence to the standard protocol study and not as a unique imaging series); it is fast, requires no intravenous contrast and is non-invasive.

**Keywords:** Diffusion weighted MRI, Malignant hepatic, Focal lesion



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# **Contents**

Subjects		Page
List of abbre	eviations	I
	2S	
L	1	
Aim of the w	vork	4
Review of L	iterature	
-	ter (1): Anatomy & MR Anatomy of the Liver	
_	ter (2): Pathology and MRI Appearance of Hepatic Focal Lesions	
_	ter (3): Technique of MR Examination of the Liver	59
Patients and	l Methods	78
L	82	
Illustrative (	Cases	88
∟111		
Summary a	nd Conclusion	118
L121		
∟122		
Arabic S	ummary	

### List of Abbreviations

**ADC** : Apparent diffusion coefficient

**CCA** : Cholangiocarcinoma

**DWI** : Diffusion-weighted magnetic resonance imaging

**FLLs** : Focal liver lesions

**FNH** : Focal nodular hyperplasia

**GI** : Gastrointestinal

**GRAPPA**: Calibrating partially parallel acquisition

**HCCs** : Hepato cellular carcinomas

**MR** : Resonance imaging

**OCP** : Oral contraceptive

**RARE** : Relaxation enhancement

**RF** : Radiofrequency

**ROI** : Region of interest

**SGE** : Spoiled Gradient-Echo

**T2WI** : T2-weighted imaging

**TE** : Echo time

**TR** : Repetition time

# List of Figures

No.	<u>Figure</u>	Page
1	Gross right and left anatomical lobes of the liver based on falciform ligament.	6
<u>2</u>	Gross anatomical lobes of the liver.	8
<u>3</u>	Liver segmental anatomy according to Couinaud classification.	10
<u>4</u>	Showing the relation of the liver: Top left, superior view; top right, posterior view; bottom left, anterior view; bottom right, inferior view.	13
5	Ligaments of the liver.	15
<u>6</u>	The portal vein and its tributaries (semi-diagrammatic). Portions of the stomach, pancreas and left lobe of the liver and the transverse colon have been removed.	16
7	Normal anatomy of the celiac Artery.	17
<u>8</u>	Arrangement of the hepatic venous territories. Multiple lower group veins may be present. Individual segments may drain into more than one hepatic venous territory.	19
<u>9</u>	Anatomy of the biliary system.	20
<u>10</u>	Normal hepatic veins.	21
11	Portal vein anatomy.	22
<u>12</u>	Sagittal MR images of the liver.	23
<u>13</u>	Coronal MR images of the liver.	24
<u>14</u>	Normal MR Liver signal intensity on T1-weighted non contrast axial image.	25
<u>15</u>	Normal MR Liver signal intensity on T2-weighted non contrast axial image.	26

No.	<u>Figure</u>	Page
<u>16</u>	Axial T2-weighted turbo spin-echo image with fat suppression shows a strongly hyperintense, unilocular.	29
<u>17</u>	Unenhanced axial T1-weighted gradient-recalled echo three-dimensional image showing an Echinococcus granulosus cyst.	31
<u>18</u>	Pyogenic abscess. Axial T2-weighted.	33
<u>19</u>	Pyogenic abscesses. Axial contrast- enhanced 3D T1-weighted VIBE MR image shows multiple small, ring-enhancing pyogenic abscesses caused by Streptococcus viridans.	33
<u>20</u>	Axial unenhanced T1-weighted in-phase image shows a large adenoma.	35
<u>21</u>	Axial T2-weighted image with fat suppression showing the adenoma.	36
22	Axial, unenhanced, T1-weighted image shows a FNH that is approximately 5 cm in diameter.	38
<u>23</u>	Focal nodular hyperplasia. Contrastenhanced spin-echo T1-weighted image obtained 4 min after bolus injection reveals persistent hypointensity of central scar (arrowhead).	39
24	Axial ADC map shows a 5-cm FNH which appears as a lobulated, hypointense lesion in the right liver lobe (subcapsular in segment 6).	39
<u>25</u>	Axial T1-weighted gradient-recalled echo three-dimensional image shows a moderate-sized hemangioma with lobular borders.	41

No.	<u>Figure</u>		
<u>26</u>	Coronal T2-weighted turbo spin-echo image with fat suppression shows a giant hemangioma with exophytic growth that bulges the liver surface (large white arrow) and has a central area of high signal intensity (black arrow), and a network of multiple fibrous septae of low signal intensity (small white arrows).	42	
<u>27</u>	Axial ADC image shows a hemangioma of approximately 4 cm in diameter in the right liver lobe in segment 7 subcapsularly (arrow), which appears hyperintense and inhomogeneous.	44	
<u>28</u>	Stepwise pathway of carcinogenesis for HCC in cirrhosis.	46	
<u> 29</u>	Typical hepatocellular carcinoma.	49	
30	Fibrolamellar HCC.		
<u>31</u>	Intrahepatic cholangiocarcinoma in a 69-year-old man.	54	
<u>32</u>	Hypervascular metastases in a 55-yearold man with a carcinoid tumor.	57	
<u>33</u>	A patient with a large liver metastasis (from colorectal cancer) and three cysts.	58	
<u>34</u>	Schematic illustrates water molecule movement.	67	
<u>35</u>	Transverse breath-hold (BH) versus respiratory-triggered (RT) fat-suppressed single-shot SEecho-planar diffusion acquisition in a 78-years old woman with liver cysts.	73	

# **List of Tables**

No.	<u>Table</u>	Page
1	Segments numbering of the liver.	10
<u>2</u>	Proposed Strategy to Improve Single-Shot	
	Echo-planar DW MR Imaging Quality of	75
	the liver.	

#### Introduction

Early detection and diagnosis of hepatic focal lesions are an important step in clinical work, which would allow effective surgical or mini-invasive therapy (*Yau et al.*, 2014; Wáng et al., 2015 and Loffroy et al., 2015).

With the advances in magnetic resonance imaging (MR) technology, diffusion-weighted magnetic resonance imaging (DWI) is now widely used as a standard imaging sequence in clinical work and shows its potential benefit in evaluation of the focal hepatic lesions (*Kwon et al.*, 2015 and Namimoto et al., 2015).

DWI with low value can suppress the intra hepatic vascular signal, creating the so-called black blood effect, which improves the detection of small focal liver lesions (FLLs) especially localized near small hepatic vessels. Meanwhile, DWI with low -value has higher imaging quality compared with single shot fast spin-echo sequences (*Takahara and Kwee*, 2012 and Bharwani and Koh, 2013).

A substantial number of studies have compared low B-value DWI with T2-weighted imaging (T2WI) for image quality and detection of FLLs. These studies generally showed better performance of DWI with low -value in

terms of lesion detection compared with T2WI (Wang et al., 2010).

DWI with higher b-value mainly reflects diffusion information of water molecules motion within the lesions, which help to improve the characterization of solid FLLs (*Namimoto et al.*, 2015).

Meanwhile, we found in practice that DWI with higher b -value also enables a better detection of lesions in liver compared with T2WI or other conventional sequences. For example, solid focal liver lesions such as focal nodular hyperplasia and hepato cellular carcinomas (HCCs) sometime can be difficult to be detected on T2WI or even DWI with low b -value due to either iso- or slightly hyper signal intensity to liver parenchyma (*Hussainet al.*, 2004 and Chen et al., 2015).

Diffusion is a physical process that results from the thermally driven, random motion of water molecules. In a container of water, molecules undergo free, thermally agitated diffusion (with a three dimensional Gaussian distribution) (*Taouli and Koh*, 2010).

The b-value represents the diffusion factor (measured in s/mm<sup>2</sup>) and the strength of the diffusion gradients. The ideal b-value for lesion characterization is a trade-off between signal attenuation and perfusion contamination.

This is generally possible using b-values between 400 and 1000 s/mm<sup>2</sup> for liver imaging. Pure diffusion contrast is obtained when using b-values above 1000 s/mm<sup>2</sup>. However, image quality can be limited by signal loss that occurs at such high b-values (*Tejas Parikh et al.*, 2008).

Two independent observers reviewed DW (*b* values of 0, 500, and 1000 sec/mm<sup>2</sup>) and T2-weighted images for FLL detection and characterization. Reference standard for diagnosis was obtained from consensus review by the two observers of DW, T2-weighted, pathological data, and follow-up imaging results. Apparent diffusion coefficient (ADC) was measured for FLLs identified at consensus review. DW and T2-weighted images were compared for FLL detection and characterization by using a binary logistic regression model. Receiver operating characteristic curve analyses was conducted to evaluate the utility of ADC for diagnosis of malignancy (*Tejas Parikh et al.*, 2008).

# **Aim of the Work**

The use of MR Diffusion imaging with both low and high B values to detect and differentiate between benign and malignant hepatic focal lesion.