

Role of Dynamic Contrast Enhanced MRI& Diffusion Weighted Imaging in Evaluation of H.C.C After Chemo-Embolization

Thesis

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List of Abbreviations

Full term Abb. 3DThree-dimensional AASLD.....Association for the study of liver diseases ADCApparent diffusion coefficient AFP.....Alpha fetoprotein APF.....Arterioportal fistula APHEArterial phase hyper-enhancement BBEPI Black-blood echo-planar imaging BCLC.....Barcelona clinic liver cancer CNR......Contrast-to-noise ratio. DCE MRIDynamic contrast-enhanced mri DEB-TACE......Drug-eluting beads transarterial chemoembolization DMDiabetes mellitus DWI......Diffusion-weighted magnetic resonance imaging ETL.....Echo train length FL.....Falciform ligament FSE/TSE.....Fast/turbo spin echo GRAPPAGeneralized auto- calibrating partially parallel acquisition GRE Gradient echo H.C.C Hepatocellular carcinoma HASTE Half acquisition single shot turbo echo HIVHuman immunodeficiency virus IBR.....Institutional board review ICCIntrahepatic cholangiocarcinoma IP.....In-phase IVC.....Inferior vena cava

List of Abbreviations cont...

Full term Abb. IVIMIntravoxel incoherent motion LAVALiver acquisition with volume acceleration Lga.....Left gastric artery LhaLeft hepatic artery LHVLeft hepatic vein LI-RADSLiver imaging-reporting and data system LR-TIVLR-tumor in vein LR-TR.....Liver imaging reporting and data system treatment response LV.....Ligamentum venosum MDCTMulti-detector computed tomography Mha.....Middle hepatic artery MHV..... Middle hepatic vein MIP..... Maximum intensity projection MRCP......Magnetic resonance cholangiopancreatography MRI......Magnetic resonance imaging NEXNumber of excitations NPV Negative predictive value NSF......Nephrogenic systemic fibrosis OLTOrthotropic liver transplantation OPOut-of-phase PEI.....Percutaneous ethanol injection PI.....Parallel imaging PPV.....Positive predictive value RF.....Radiofrequency. RFARadiofrequency ablation Rha.....Right hepatic artery

List of Abbreviations cont...

Abb. Full term

RHV	Right hepatic vein
RT	Radiation therapy
R-T	Respiratory-triggered
SE	Single-shot spin-echo
Sma	Superior mesenteric artery
SNR	Signal to noise ratios.
SPAIR	Spectrally adiabatic inversion recovery
SPIO	Superparamagnetic iron oxide particle
SPSS	Statistical package for the social sciences
SSFSE	Single shot fast spin echo.
STIR	Short time inversion recovery sequences
TACE	Trans-arterial chemo-embolization
TAE	Trans-arterial embolization
TARE	Trans arterial radio-embolization
TE	Echo time
TR	Time of repetition
TVDT	Tumor volume doubling time
WO	Wash-out

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Introduction

INTRODUCTION

Hepatocellular carcinoma (HCC) is the most common primary malignant disease of the liver and is the third leading cause of death from cancer worldwide (*Bonekamp et al.*, 2013).

Only a minority of all patients with HCC are surgical candidates at the time of diagnosis (*Wang et al., 2014*).

Treatment options are divided into surgical therapies (i.e., resection, cryoablation and orthotropic liver transplantation (OLT), and nonsurgical therapies (i.e., percutaneous ethanol injection (PEI), radiofrequency ablation (RFA), trans-arterial chemo-embolization (TACE), radiation therapy (RT) and systemic therapy) (*Curley et al., 2017*).

Transcatheter arterial chemoembolization (TACE) is one of the most commonly used intra-arterial therapies to treat unresectable HCC, and several clinical trials have demonstrated that TACE has the potential to show survival benefits in patients with HCC. TACE is a frequently used technique and usually includes intra-arterial delivery of emulsions mixed with chemotherapeutic agents and lipiodol, followed by the administration of the embolic agent (*Wang et al., 2014*).

Assessing early response to therapy using objective criteria is paramount for clinical care. Identifying early responders could help individualize therapy and tailor future treatment strategies (*Bonekamp et al.*, 2013).

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Usually the presence of residual viable tumor or recurrence after TACE is assessed using multi-phasic multidetector computed tomography (MDCT) scanning, but this technique can be affected by artifacts produced by high concentrations of lipiodol, making it difficult to evaluate the characteristics of the lesion. Response may be better assessed by noting alterations in vascular tumor perfusion rather than size, and therefore functional measurements may be an appropriate method to assess tumor response (Ippolito et al., 2010).

MRI provides anatomic parameters, as well as functional and molecular parameters, for the assessment of the response to treatment. Non-contrast T1- and T2-weighted images provide information on morphological change, fluid content change and fibrosis. Dynamic contrast enhanced MRI can provide information on perfusion (Osama et al., 2013a).

Dynamic contrast-enhanced MRI (DCE MRI) can play a significant role as an imaging biomarker, especially in the detection of viable tumor foci, the differentiation between necrosis and viable tumor and early prediction of response (Lee et al., 2011).

Dynamic contrast-enhanced magnetic resonance imaging (DCE-MRI) also enables tumor vascular physiology to be assessed (Chen and Shih, 2014).