## The Role of PET CT In Comparison To Triphasic CT in Early Follow Up of Hepatocellular Carcinoma after Transarterial Chemoemoblization

### Thesis

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# By

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## List of Abbreviation

AFP	Alpha-fetoprotein	
BGO	Bismuth germanate	
CBD	Common bile duct	
СЕСТ	Contrast enhanced computed tomography	
СНА	Common hepatic artery	
CHD	Common hepatic duct	
СТ	Computed tomography.	
СТАС	Computed tomography attenuation	
	correction	
EASL	European Association for the Study of the	
	Liver	
FDG	Fluorodeoxyglucose	
GB	Gall bladder.	
GDA	Gastroduodenal artery	
GLUT	Glucose transporter	
GSO	Gadolinium oxyorthosilicate	
НСС	Hepatocellular carcinoma	
HCV	Hepatitis C virus	
HK enzyme	Hexokinase enzyme	
IMV	Inferior mesenteric vein	
IVC	Inferior vena cava	
kBq	Kolobecquerel	

Kev	Kiloelectron volt	
Kg	Kilogram	
LGA	Left gastric artery	
LHA	Left hepatic artery	
LHV	Left hepatic vein	
LPV	Left portal vein	
LSECs	Liver sinusoidal endothelial cells	
LSO	Lutetium oxyorthosilicate	
mCi	Millicurie	
MHV	Middle hepatic vein	
MIP	Maximum intensity projection	
MPV	Main portal vein	
mDECIST	Madified Degrange Evaluation Critaria in	
IIIKECISI	Modified Response Evaluation Criteria in	
IIIKECISI	Solid Tumors	
MRI	Solid Tumors Magnetic Resonance imaging	
MRI N/C ratio	Modified Response Evaluation Criteria in Solid TumorsMagnetic Resonance imagingNuclear/cytoplasmic ratio	
MRI N/C ratio Nal	Modified Response Evaluation Criteria in Solid TumorsMagnetic Resonance imagingNuclear/cytoplasmic ratioSodium iodide	
MRI N/C ratio Nal NASH	Modified Response Evaluation Criteria in Solid TumorsMagnetic Resonance imagingNuclear/cytoplasmic ratioSodium iodideNonalcoholic steatohepatitis	
MRI N/C ratio Nal NASH PET	Modified Response Evaluation Criteria in Solid TumorsMagnetic Resonance imagingNuclear/cytoplasmic ratioSodium iodideNonalcoholic steatohepatitisPositron Emission Tomography	
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MRI N/C ratio Nal NASH PET PMT PV	Modified Response Evaluation Criteria in Solid TumorsMagnetic Resonance imagingNuclear/cytoplasmic ratioSodium iodideNonalcoholic steatohepatitisPositron Emission TomographyPhotomultiplier tubePortal vein	
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MRI N/C ratio Nal NASH PET PMT PV PVTT RAPV	Modified Response Evaluation Criteria in Solid Tumors Magnetic Resonance imaging Nuclear/cytoplasmic ratio Sodium iodide Nonalcoholic steatohepatitis Positron Emission Tomography Photomultiplier tube Portal vein Portal vein tumoral thrombosis Right anterior portal vein	

	Tumors	
RFA	Radiofrequency ablation	
RHA	Right hepatic artery	
ROI	Region of interest	
RPPV	Right posterior portal vein	
RPV	Right portal vein	
RRA	Right renal artery.	
SA	Splenic artery	
SMA	Superior mesenteric artery	
SMV	Superior mesenteric vein	
SPD	Sum of the product diameters	
SUV	Standardized uptake value	
TACE	Transarterial chemoembolization	
WHO	World Health Organization	
18- FDG	18- fluorodeoxyglucose	

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## Introduction

Hepatocellular carcinoma (HCC) represents the commonest primary hepatic tumor of adults. It is the  $6^{th}$  most common tumor in the world and the third commonest cause of cancer related deaths (*Dai et al., 2014*).

Liver cancer represents about 11.85% of the malignancies of all GIT organs and 1.78% of the total malignancies among Egyptians (*Holah et al., 2015*).

HCC is caused by malignant transformation in hepatocytes due to chronic liver diseases resulting in cirrhosis (*Tsurusaki et al., 2014*).

From the selective treatment options of liver tumors, interventional procedures such as Trans arterial chemoembolization (TACE), has been widely used. The powerful cytotoxic effect of TACE by combined action of ischemia followed by chemoembolization of the tumor's feeding artery has been proved to result in therapeutic efficacy (*Song et al., 2013*).

Despite good results, this interventional procedure needs close monitoring to effectiveness of treatment because the rate of residual viable malignancy in tumors larger than 3 cm can reach 48% (*Tsurusaki et al., 2014*).

<sup>1</sup> 

Follow up of tumor response after TACE is important to determine whether the tumor is completely eradicated or additional treatment is required. Magnetic resonance imaging or computed tomography has been widely used for the assessment of treatment response after TACE. The determination of treatment response using size criteria, based on the Response Evaluation Criteria in Solid Tumors (RECIST), does not necessarily apply well to interventional therapy in such patients, so most radiologists have relied on the presence or absence of local contrast enhancement at the treated tumor in addition to changes in tumor size (*Kim et al.*, 2011).

The methods which are used to detect tumor viability depend on showing arterial enhancement for reporting treatment responses. However, this concept does not adequately consider the biological activity of HCC (*Song et al., 2013*).

Positron Emission Tomography (PET) is a noninvasive imaging tool that uses 18- fluoro-deoxy-glucose (18- FDG) as radioactive material showing difference in metabolism between tissues thus demonstrates the functional status of suspicious lesions (*Saif et al., 2010*).

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After interventional procedures, CT or MRI at one month are routinely performed to assess for residual tumors but there has been increasing evidence that PET can detect residual tumors earlier than CT and MRI (*Tsurusaki et al.*, 2014).

PET/CT is a new imaging tool, whose advantages are useful in clinical oncology. The combination of anatomical and functional image has been the true evolution in diagnosis (*Saif et al., 2010*).

So, PET/CT can be used in the assessment of hepatocelluar biological activity as an additional predictive tool (*Song et al., 2013*).

## Aim of the Work

The aim of this study is to emphasize the role of PET/CT in early follow up of HCC after transarterial chemoembolization in comparison to triphasic CT.