



Research Article



The Association of Kidney's GFR and Serum Creatinine Levels with Renal Scarring through ^{99m}Tc-Glucoheptonate Radiopharmaceutical Scan in Patients with Urinary Tract Infection

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Abstract

Background: UTI (urinary tract infection) is a leading case of renal scarring. Most of nephrons are found in kidney cortex, so it is important for early diagnosis of scarring. In this study, we investigated if scarring affects GFR (glomerular filtration rate) and serum level of creatinine through dual multipurpose ^{99m}Tc-GH (^{99m}Tc-Glucoheptonate) scintigraphy in patients with UTI. **Methods:** During this study 21 patients with UTI were studied by ^{99m}Tc-GH scan. For performing scintigraphy, the patient was injected by 370-555 MBq of ^{99m}Tc-GH and studied through two steps including dynamic (immediately after injection) and static (2-3h after injection) phases. The results were evaluated by appropriate analytical methods. Moreover, five patients were studied by both ^{99m}Tc-DMSA/GH (^{99m}Tc-dimercaptosuccinic acid and ^{99m}Tc-Glucoheptonate) to show ^{99m}Tc-GH scan is beneficial for detection of additional problems.

Results: The results showed that there is an association between right/left kidney scarring and related GFR. Furthermore,, the odds of decreased right and left kidney GFR with scarring is 12.64 and 11.89 times more than normal kidney, respectively. Also, there is a significant association between renal scarring and serum level of creatinine. The study showed that the odds of increased serum level of creatinine in patients with scarring are 6.75 times more than patients without scarring. Moreover, the survey of five patients with both ^{99m}Tc-DMSA/GH scans showed ^{99m}Tc-GH could be helpful with diagnosis of some renal problems as well as scarring.

Conclusion: Through this study by ^{99m}Tc-GH, it was demonstrated that kidney glomerular filtration rate and serum level of creatinine are associated with renal scarring. Furthermore, it was shown that ^{99m}Tc-GH scintigraphy could detect dilatation and abstraction of collection system in addition to renal scarring through static/dynamic renal scintigraphy.

Introduction

Renal scarring as a serious renal problem, leads to essential problems like chronic renal failure and hypertension.¹ There are some important etiologies of renal scarring including UTI (urinary tract infection), pyelonephritis, nephrolithiasis, hypertension, vesicoureteral reflux and diabetes.² Due to highly localization of nephrons in kidney's parenchyma, early diagnosis of cortical lesions could prevent permanent scars as a leading cause of renal failure. Moreover, renal scarring would lead to proteinuria, hematuria, electrolyte imbalance and hypertension.³ Basically, the origin of renal scarring is determined by biopsy, but it isn't preferable because of being invasive

method. Generally, various diagnostic methods including urography, sonography, CT (computed tomography), MRI (Magnetic resonance imaging) and renal scintigraphy are used to detect renal scarring.⁴ Renal scintigraphy as a functional imaging is used for kidney clearance assay, renal tubular secretion and morphological studies.⁵ Mainly, renal scintigraphy is classified into dynamic (perfusion and function scintigraphy) and static (renal cortical scintigraphy) studies.⁶ There are different types of radiopharmaceuticals determined to targeted renal scintigraphy (Figure 1). Dynamic scintigraphy is performed to evaluate GFR (glomerular filtration rate) and tubular secretion. The common radiopharmaceutical for calculation

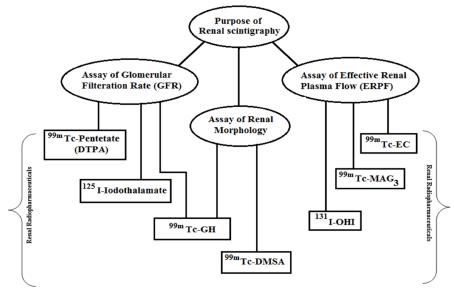


Figure 1. Different types of radiopharmaceuticals for renal scintigraphy.

of GFR is 99mTc-DTPA, but the 99mTc-GH could be also used. The 99mTc-GH (99mTc-Glucoheptonate) as a radiolabeled carbohydrate, is eliminated by glomerular filtration (>80%) and binds to kidney tubules (10%-15%). Therefore, 99mTc-GH as a multi-purpose radiopharmaceutical could be used to evaluate both renal function and cortical scar just via one-dose i.v. injection. It should be mentioned when the renal scintigraphy is just performed for functional study, the 99mTc-DTPA is preferable. For evaluation of both renal function and morphology, the 99mTc-GH is preferred in order to provide both dynamic and static scintigraphy. Renal cortical scintigraphy is performed to evaluate renal morphology, especially for diagnosis of renal scarring. 99mTc-DMSA (99mTc-dimercaptosuccinic acid) as common radiopharmaceutical is commonly used to perform cortical scintigraphy for diagnosis of renal scarring. After i.v. injection, the 99mTc-DMSA binds to a1-microglobulin in blood by sulfhydryl groups. The assembly (99mTc-DMSA-α1-microglobulin) aggregates in tubules of renal parenchyma by megalin/tubulin-mediated endocytosis that leads to parenchymal retention of 99mTc-DMSA by 50 percent. Free 99mTc- DMSA and trace amounts of assembly are secreted in tubular fluid.7 Generally, the study of renal function has been performed by glomerular radiopharmaceuticals, especially by 99mTc-DTPA with regard to GFR's calculation. The 99mTc-DTPA (99mTcdiethylenetriaminepentaacetic acid) was glomerularly filtered by %95 (about 5% of 99mTc-DTPA binds to plasma proteins), so it provides ability to calculate GFR and assay urine flow through pyelocalyceal system in order to detect obstructive problems.8 In comparison to renal scintigraphy, the CT scan can be used to study functional and morphological statues of unilateral chronic renal obstruction, but 99mTc-DTPA scan is preferable because of low radiation exposure dose. Furthermore, it was noted "comparative studies are needed to assess the role of CTbased parenchymal volume in evaluation of individual

renal functions in patients with acute obstruction, bilateral nephropathy and renal impairment".9 However, it is possible to study morphology and hemodynamics of kidney due to developments in ultrasound modalities, but the 99mTc-DTPA scan is necessary for assessment of clearance rate and accurate function of kidney.¹⁰ Although the renal magnetic resonance imaging could be used to perform the functional imaging by gadolinated tracers, it is accompanied by lack of linearity, therefore the 99mTc-DTPA scan still remains as a gold standard in this domain.¹¹ Moreover, from radiation protection point of view, radiation dose from 99mTc-GH is less than sums of 99mTc-DTPA and 99mTc-DMSA scans, because the recommended dose for 99mTc-GH scan is 370-555 MBq (the recent literature also recommended 296 MBq for 99mTc-GH (7)).12 Therefore, when there is necessity to perform both 99mTc-DTPA and 99mTc-DMSA scans, the 99mTc-GH scan could be just performed instead. Urinary tract infections (UTI) and acute pyelonephritis (APN) are the leading cause of renal scarring, especially in children. In general, 15-60% of children with UTI develop renal scarring.^{13,14} So, the early detection of scarring can prevent related problems as well as renal dysfunction due to parenchymal damage.15 Because patients with renal scarring have been routinely studied by 99mTc-DMSA scan, so the renal perfusion and GFR (that could be decreased) are missed during evaluation.

During this study, we attempted to detect renal scarring and evaluate association of that with GFR and serum level of creatinine by individual ^{99m}Tc-GH renal scan that could gather more data about morphology and function of kidney simultaneously. This study showed how missed ^{99m}Tc-GH radiopharmaceutical could be useful in diagnosis of urinary tract complications and kidney functions as well as renal scarring just through single scintigraphy instead of both separated ^{99m}Tc-DTPA and ^{99m}Tc-DMSA scintigraphy. Ultimately, ^{99m}Tc-GH scintigraphy as an alternative leads to reduce radiation exposure in patient candidate for both

^{99m}Tc-DTPA and ^{99m}Tc-DMSA scintigraphy.

Methods

During this retrospective study, 21 patients with UTI who had referred to our nuclear medicine division were studied during one year to find if renal scarring correlates with GFR and serum level of creatinine through 99mTc-GH renal scan. Ethical approval for this study was obtained from Research Ethics Committee of our institution. The participants were informed on data publishing.

Inclusion criteria were all patients with confirmed UTI by positive urine culture, and creatinine serum level of them had been determined prior to scintigraphy. All patients didn't have any systemic disease and hadn't consumed antibiotic recently. Exclusion criteria in this study were the pregnant patients.

The GH vacuumed cold kits were obtained from Pars Isotope Co. and radiolabeled with fresh 99mTcO, ⁹⁹Mo/^{99m}Tc (99m-Technetium pertechnetate) from generator according to kit instruction. All patients were evaluated by both dynamic and static scintigraphy after 370-555~MBq of $^{99\text{m}}\text{Tc-GH}$ i.v. injection. 16 The dynamic renal scan was subsequently performed after bolus i.v. injection of radiopharmaceutical at perfusion, parenchymal and excretory phases at posterior views. The split renal GFR was calculated via Gates method. The cortical scintigraphy was performed 2h after 99mTc-GH i.v. injection at Ant (anterior), Post (posterior), LPO (left posterior oblique) and RPO (right posterior oblique) views. The Siemens SPECT E.Cam Dual Head gamma camera with low-energy and high-resolution collimator was used in this study. It should be mentioned the normal serum level of creatinine was considered at 0.7-1.3 mg/dL.17 Also, the threshold for lowered GFR was considered as GFR 90≥ mL/min.¹⁸ For radiolabeling of GH cold kit, 1110-1850 MBq (30-50 mCi) of fresh 99mTcO₄ (up to 4 mL) was added to GH cold kit and shaken for 1 min then left in room temperature for 10 min to complete ligand/radioisotope incubation. For radiolabeling of DMSA cold kit, 1480 MBq of fresh 99mTcO₄ (up to 3 mL) was added to DMSA cold kit and shaken for 1-2 min then left in room temperature for 15-20 min to complete ligand/radioisotope incubation. The quality of radiolabeled kits were assayed by two systems of chromatography (by ITLC).19

Moreover, Schwartz formula (that it was considered as a confirmed standard way, Eq 1) was used to find out if the calculated GFR (based ^{99m}Tc-GH renal scan) is reliable.²⁰ In this Equation, the "k" is defined as 0.419 if males aged>13 years and otherwise considered at 0.373, the "Scr" and "L" are defined as serum level of creatinine and patient's height (cm) respectively.

Schwartz formula: eGFR = kL/Scr Eq. (1)

Through this investigation five patients were studied by either ^{99m}Tc-GH or ^{99m}Tc-DMSA renal scan after getting permission from vulnerable patients to show if ^{99m}Tc- GH

renal scan is beneficial for some patients in addition to renal scarring detection.

Determination of serum creatinine

The serum level of creatinine had been determined by autoanalyzer system (model: BS-800, Mindray factory, China) and enzymatic method.

Preparation of patient for scintigraphy

There is no special patient preparation for ^{99m}Tc-GH renal scan, but it is recommended that the patients should be hydrated by drinking 5-10 mL/kg of water 30-60 min prior to scintigraphy.²¹

Statistical analysis

The obtained data were analyzed by descriptive statistics methods as mean±SD and frequency (percent). For inferential statistics Fisher's exact test, Pearson correlation test and logistic regression analysis were used by SPSS 16 (SPSS Inc., Chicago, IL) Software. The Shapiro-Wilk test was used to evaluate normal distribution of data. In all analysis, p-values less than 0.05 (P<0.05) were considered as significant.

The intraclass correlation coefficient (ICC) has been advocated as a statistic for assessing agreement or consistency between two methods of GFR measurement. For calculation ICC, a two-way mixed-effect model based on single ratings assessed was used. Mean estimation along with 95% confidence intervals (CI) were reported for the ICC. Interpretation was as follows: <0.50, poor; between 0.50 and 0.75, fair; between 0.75 and 0.90 good; above 0.90, excellent.²²

Results

During this study, 21 patients with UTI, 11 female (52.4%) with mean of age 23.2±14.7 years and 10 (47.6%) male patients with mean of age 21.3±15.9 years were studied. The mean of GFR in patients with renal scarring (A) were calculated at 25.58±11.90 ml/min and 28.75±20.46 ml/min for right and left kidney respectively. The mean of GFR in patients without renal scarring (B) were calculated at 42.56±11.90 ml/min and 43.15±20.51 ml/min for right and left kidney respectively. Moreover, there was no significance difference between the mean serum level of creatinine for patients with (1.64±0.72 mg/dl) and without (1.5±0.94 mg/dl) renal scarring (p-value=0.751). Among 21 patients, 11 patients (52.4%) were demonstrated decreased right kidney's GFR. As Table 1 shows, 85.7% of patients with right renal scarring had decreased right kidney's GFR and 14.3% had normal GFR. Based on Fisher's exact test, there is a significant association between right kidney's GFR and renal scarring (p-value=0.031). Among all patients, 61.9% (11 patients) showed decreased left kidney's GFR. Also, 87.5% of patients with left renal scarring had decreased left kidney's GFR. Based on Fisher's exact test, there is also a significant association between left kidney's GFR and renal scarring (p-value=0.047).

Table 1. The rate of right/left kidney's GFR in patients with and without right renal scarring.

			Kidney's GFR		
Variable	Category	Decreased	Normal	Total	p-value*
		Number (%)	Number (%)	Number (%)	
	A	6 (85.7)	1 (14.3)	7 (100)	
Right kidney scarring	В	5 (35.7)	9 (64.3)	14 (100)	0.031
	Total	11 (52.4)	10 (47.6)	21 (100)	
	A	7 (87.5)	1 (12.5)	8 (100)	
Left kidney scarring	В	6 (46.2)	7 (53.8)	13 (100)	0.047
	Total	13 (61.9)	8 (31.8)	21 (100)	

^{*}Results were based on Fisher's exact test

Table 2. The effects of right /left kidney scar, gender and age on right and left kidney's GFR.

Variable	Category	OR*	95% CI**	p-value***		
		Right kidney's GFR				
Pight kidnov scar	-					
Right kidney scar	+	12.64	(1.12, 24.72)	0.043		
0	Female					
Sex	Male	1.27	(0.12, 13.54)	0.845		
Age (y)		1.05	(0.98, 1.13)	0.165		
		Left kidney's G	FR			
Left kidney scar	-					
	+	11.89	(1.09,22.91)	0.046		
Sex	Female					
	Male	3.55	(0.31, 40.56)	0.308		
Age (y)		1.04	(0.98, 1.11)	0.236		

^{*}Odds Ratio (OR), **Confidence interval (CI), *** Results were based on logistic regression

Table 3. The frequency (%) of creatinine serum level in patient with and without renal scarring.

Variable	Category	Increased Number (%)	Normal Number (%)	Total Number (%)	p-value*
	Α	8 (66.7)	4 (33.3)	12 (100)	
Renal scarring	В	2 (22.2)	7 (77.8)	9 (100)	0.030
	Total	10 (47.6)	11 (52.4)	21 (100)	

^{*}Results were based on Fisher's exact test

Moreover, after adjustments for sex and age, the results showed the odds of decreased right kidney's GFR in patients with right kidney scar is 12.64 times more than patients without that diagnosis (OR=12.64, p-value=0.043). Also, the results demonstrated that sex and age of the patients didn't affect right kidney's GFR (p-value_(sex)=0.845, p-value_(age)=0.165. Furthermore, the results indicated that the odds of decreased left kidney's GFR in patients with left kidney's scar is 11.89% times more than patients without that diagnosis (OR=11.89,p-value=0.046) through moderation of age and sex effects. Also, the results showed that sex and age of the patients didn't affect left kidney's GFR (p-value_(sex) = 0.308, p-value_(age) = 0.236) (Table 2).

Among all patients (n=21), 47.6% (11 patients) demonstrated increased serum level of creatinine. As Table 3 shows, 66.7% of patients with renal scarring associated with increased serum level of creatinine. According to Fisher's exact test, there is a significant association between renal scarring and serum level of creatinine

(p-value =0.030). Also, we developed a survey to find if there is correlation between right/left kidney's GFR and serum level of creatinine. According to Shapiro-Wilk test, normal distribution of variables including right kidney's GFR, left kidney's GFR and serum level of creatinine were concluded. Correlation between variables was assessed by Pearson correlation test. Furthermore,, there is a negative correlation between right/left kidney's GFR and serum level of creatinine (right kidney's GFR: r=-0.54, p-value=0.011; left kidney's GFR: r=-0.50, p-value=0.022) (Table 4). However, there is not significant correlation between left and right kidney's GFR (r=-0.01, p-value=0.982). Moreover, percent prediction (determination coefficient) of creatinine serum level by right/left kidney's GFR were calculated to be 25% and 29%, respectively. Following that, after adjustments of sex and age, the odds of increased serum level of creatinine in patients with renal scarring is 6.75 times more than patients without renal scarring (OR=6.75, p-value=0.032). Also, the results demonstrated

Table 4. The Correlation of right/left kidney GFR and serum level of creatinine.

Variable	Rig	Right kidney GFR		Left kidney GFR		Creatinine	
	r*	p-value	r*	P-value	r*	p-value	
Right kidney GFR	1		0.01	0.982	-0.50	0.022	
Left kidney GFR			1		-0.54	0.011	
Creatinine					1		

^{*}Pearson correlation coefficient

Table 5. The effects of right kidney scar, gender and age on creatinine serum level.

Variable	Variable Series	OR*	95% CI**	p-value***
Right kidney scar	- +	6.75	(1.31, 12.26)	0.032
Gender	Female Male	0.36	(0.03, 3.89)	0.362
Age (y)		1.05	(0.98, 1.13)	0.349

^{*}Odds Ratio (OR), ** confidence interval (CI), *** Results was gained upon logistic regression

that sex and age of patients didn not affect serum level of creatinine (p-value_(sex) = 0.362, p-value_(age) = 0.349) (Table 5).

Also, the intraclass correlation coefficient was computed to assess the consistency between two GFR measurement methods (calculated GFR from ^{99m}Tc-GH scan and estimated GFR from Schwartz formula). The ICC was between fair and good being 0.81 (0.56-0.92), for calculated GFR (by ^{99m}Tc-GH scintigraphy) and estimated GFR (by Schwartz formula).

Moreover, during this study, five patients that had been studied by both ^{99m}Tc-GH and ^{99m}Tc-DMSA scans, developed valuable results demonstrating ^{99m}Tc-GH renal scan could be beneficial for patients with some kidney problems as well as renal scarring. ^{99m}Tc-GH scan could evaluate renal perfusion-function in addition to renal cortical surveys (Figures 2-6). In all patients ^{99m}Tc-GH scan

was performed at least 48h after 99m Tc-DMSA scan in order to assay whether 99m Tc-GH shows renal cortical uptake similar to 99m Tc-DMSA scan.

Figure 2 (a-c) illustrates both ^{99m}Tc-GH and ^{99m}Tc-DMSA for a same patient. As shown, there is a decreased ^{99m}Tc-DMSA uptake in corticomedial parenchyma of left kidney that tends to be seemed as a scar. During ^{99m}Tc-GH scan, the early dynamic phases demonstrated the retention of ^{99m}Tc-GH in the mentioned region of left kidney. Following i.v. injection of diuretic at 10th min of study, completed clearance of radiotracer from corticomedial parenchyma of left kidney was shown. So it is concluded that the decreased parenchymal uptake is resulted from collecting system dilatation instead of renal cortical scar.

Figure 3 (a-c) illustrates a large cortical defect in the upper pole of right kidney in both ^{99m}Tc-GH (in static

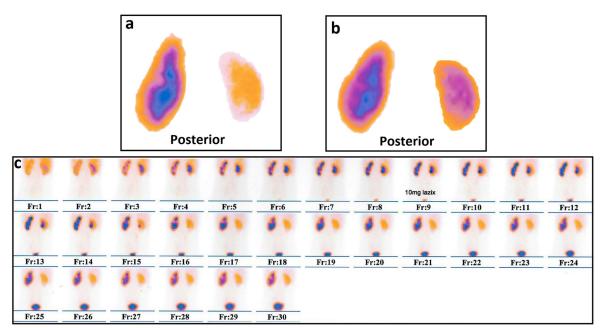


Figure 2. Both ^{99m}Tc-GH and 99mTc-DMSA scintigraphy for a same patient. (a): ^{99m}Tc-DMSA Renal Scintigraphy, (b): Cortical (static) renal scintigraphy by ^{99m}Tc-GH, (c): Dynamic functional renal scintigraphy by ^{99m}Tc-GH.

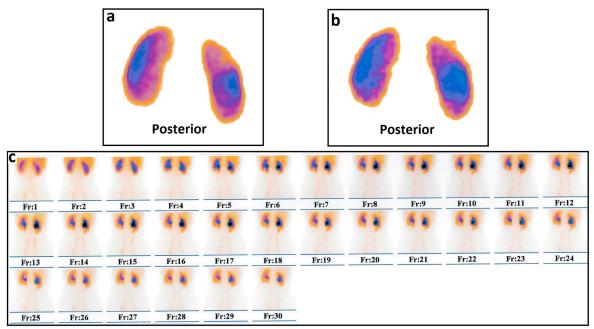


Figure 3. Both ^{99m}Tc-GH and ^{99m}Tc-DMSA scintigraphy for a same patient. (a): ^{99m}Tc-DMSA Renal Scintigraphy, (b): Cortical (static) renal scintigraphy by ^{99m}Tc-GH, (c): Dynamic functional renal scintigraphy by ^{99m}Tc-GH.

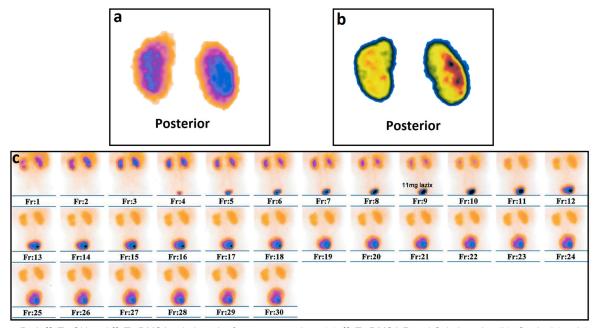


Figure 4. Both ^{99m}Tc-GH and ^{99m}Tc-DMSA scintigraphy for a same patient. (a): ^{99m}Tc-DMSA Renal Scintigraphy, (b): Cortical (static) renal scintigraphy by ^{99m}Tc-GH, (c): Dynamic functional renal scintigraphy by ^{99m}Tc-GH.

phase) and ^{99m}Tc-DMSA scans. Dynamic phase of ^{99m}Tc-GH scan showed accumulation of radiopharmaceutical in the mentioned defect that ruled out the probability of renal scarring. After Lasix injection, retention of ^{99m}Tc-GH in pelvis demonstrated obstructive problems like ureteropelvic junction stenosis (UPJS).

Figure 4 (a-c) shows small region of decreased cortical radiotracer uptake in the upper pole of left kidney in both static ^{99m}Tc-GH and ^{99m}Tc-DMSA scans that is suspicious of cortical scar. Both kidneys were also considered as normal at ^{99m}Tc-GH dynamic phase that ruled out renal scarring.

Therefore, it is concluded that such small defects could be due to partial dilatation of collecting system.

Figure 5 (a-c) illustrates normal uptake of radiopharmaceutical in kidneys cortex in both renal cortical scans (a,b). In addition, the ^{99m}Tc-GH dynamic phase demonstrated normal perfusion -function of right kidney and mild decreased perfusion-function and GFR for left kidney.

Figure 6 (a-c) indicates a large region of cortical defect in upper pole of right kidney in both ^{99m}Tc-DMSA and ^{99m}Tc-GH (in static phase) scans. Moreover, the ^{99m}Tc-GH

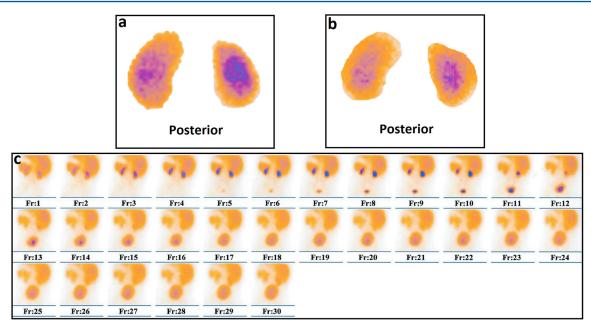


Figure 5. Both 99mTc-GH and 99mTc-DMSA scintigraphy for a same patient. (a): 99mTc-DMSA Renal Scintigraphy, (b): Cortical (static) renal scintigraphy by 99mTc-GH, (c): Dynamic functional renal scintigraphy by 99mTc-GH.

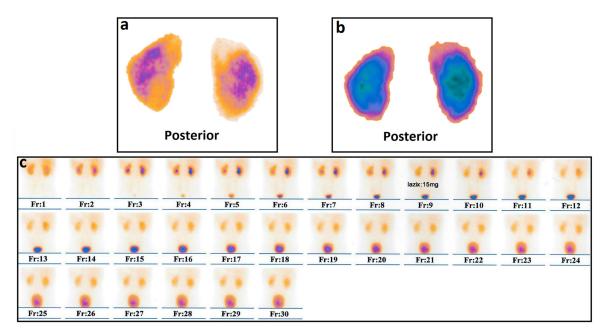


Figure 6. Both 99mTc-DMSA scintigraphy for a same patient. (a): 99mTc-DMSA Renal Scintigraphy, (b): Cortical (static) renal scintigraphy by 99mTc-GH, (c): Dynamic functional renal scintigraphy by 99mTc-GH.

dynamic phase showed normal perfusion-function of left kidney as well as non-obstructive partially dilatation of collecting system in right kidney. Consequently, there is correspondence between 99mTc-GH static phase and 99mTc-DMSA scans results.

Discussion

During this investigation it was concluded that 1) the mean of GFR in normal patients is supposedly higher than patients with renal scarring and also, the mean of creatinine serum level is elevated for patients with renal scarring, 2) the age and sex are not affected the GFR

and serum level of creatinine, 3) the separated GFR is associated with related kidney's scarring, and 4) the GFR and serum level of creatinine are not correlated with together. Similar to this findings, there are some similar studies. Because patients with UTI are susceptible to renal scarring,²³ there are numerous studies in this field. Bandari et al.24 studied the presence of renal scarring in children with UTI by 99mTc-DMSA scan. They reported the renal scarring associated closely with hydroureteronephrosis (100%), febrile UTI (34%), younger age group (67.5%) and E. coli UTI (86%). They also noted that cortical renal scintigraphy as a non-invasive method is preferable

for investigation of renal scarring in children with first and recurrent UTI. Shaikh et al.25 expressed the delayed antibiotic treatment of febrile UTI will be firmly associated with renal scarring. They mentioned that "understanding the association between the number of episodes of febrile UTI and the risk for renal scarring can help develop evidence-based management strategies for children with this frequently occurring problem". Najafi et al.26 did a meta-analysis to find the prevalence of kidney scarring caused by UTI in Iranian children. Their survey concluded that one-third of children with UTI involved with renal scarring and the odds of renal scarring in patients with urinary tract reflux is about four times. Recently, during a review article by Murugapoopathy et al.27 it was mentioned UTI as a common problem of hospitalized children could affect bladder and kidney if it is severe. The infection could destroy kidney's tissue by replacing of nephrons through deposition of extra cellular matrix as renal scar. Therefore, UTI could lead to renal scarring resulting in decreased GFR and elevated serum level of creatinine. Similar to our investigation, there are similar studies that noted the renal scarring affect kidney's function through decreasing GFR. Relevantly, Imamura et al.28 showed the renal scarring confirmed by 99mTc-DMSA scan leads to decrease GFR in related patients. Fattah et al.29 noted the nephrons losing cause to increase in single nephron glomerular filtration rate (SNGFR) as a compensatory mechanism to prevent decreasing of total GFR, but this mechanism could lead to hypertension or CKD (chronic kidney disease) if there is severe or long lasting of nephrons losing. Based on inherent characteristics of 99mTc-GH, both functional and morphological studies could be performed by 99mTc-GH scan. Related that, our study showed the calculated GFR by 99mTc-GH scan is in correlation with estimated GFR (from Schwartz formula). According to Schwartz equation, there is an inverse relationship between GFR and serum level of creatinine. So, the result could be considered as a validation for our study, and the odds of increased serum level of creatinine in patients with renal scarring was 6.75. Although no research found a relationship between renal scarring and serum level of creatinine directly, it seems that the renal scarring could affect serum level of creatinine through causing an injury to nephrons and decreasing GFR at least. Because the serum level of creatinine could be affected by several factors including age, gender, race, protein intake, muscle mass, infections, and inflammatory status, therefore the used of creatinine as real function's marker would be as a controversy especially in patients with acute kidney injury (AKI). Cystatin C proposed as a marker of renal function in critically patients with normal serum creatinine. The studies showed serum cystatin C is not superior to serum creatinine in the early detection of renal dysfunction.30,31 Also, during our study the participants did not have any systemic disease. However, 99mTc-DMSA scintigraphy is considered as a gold standard to detection of renal scarring,³² 99mTc-GH scintigraphy could show the kidney's cortex due to about 20% cortex retention too. Through our study by 99mTc-GH the valuable results were gained regarding the kidney's function, cortical disorders and obstructive problems. Therefore, this study suggests that it is better to investigate both kidney's cortex and perfusion in patients with risk factors of renal scarring rather than just cortical study. Generally, the strength of this study was determined to evaluate both GFR (for evaluating kidney function) and kidney's cortex (for evaluating renal scarring) just by 99mTc-GH scan instead of separated 99mTc-DTPA and 99mTc-DMSA scans, and we investigated the association of Kidney's GFR and serum level of creatinine with renal scarring just through one scintigraphy based on characteristics of 99mTc-GH radiopharmaceutical.

This study had some limitations. Firstly, the characteristics of patients were only sex and age, so other confounder factors which may affect renal function were not included in statistical analysis. Secondly, the creatinine serum level could be affected by several factors that were not considered during this study. Thirdly, the scintigraphy was limited in pregnant and breastfeeding patients.³³

Conclusion

In this study by 99mTc-GH renal scan some qualitative and quantitative findings were obtained. The study of patients with UTI by 99mTc-GH scintigraphy demonstrated that renal scarring has more effect on GFR. As this study shown, the GFR and serum level of creatinine is associated with renal scarring, also renal scarring and elevated serum level of creatinine correlated. Moreover, patients who were studied by 99mTc-GH developed both perfusion and cortical scans. During this study, limited survey of cortical study by both 99mTc-GH and 99mTc-DMSA scans in patients with UTI demonstrated that 99mTc-GH could show renal scarring if there is. Furthermore, dynamic phase of 99mTc-GH scan could detect additional problems like dilatation and obstruction of urinary tract. It is suggested that this kind of expanded renal scintigraphy via 99mTc-GH gives chance to detect perfusion-functional disorders that would cause serious kidney's problems in addition to cortical studies. Moreover, the patients are determined to be studied by both 99mTc-DMSA and 99mTc-DTPA scans could be replaced just by 99mTc-GH scan because of saving time and cost as well as decreased radiation exposure dose. Furthermore, reliability and consistency of GFR was calculated based on both 99mTc-GH renal scan and Schwartz formula methods.

Ethical Issues

The study had been approved by Tabriz University of Medical Sciences ethics committee (IR.TBZMED. REC.1396.480) and the necessary permission was obtained from patients to publish their findings.

Data Sharing

The data that support the findings of this study are available on request from the corresponding author.

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Author Contribution

AF designed the work and was a major contributor in drafting the work, EG interpreted the scintigraphy regarding the ^{99m}Tc-GH and ^{99m}Tc-DMSA renal scans, NM contributed in acquisition of all patient's documents and NG was done statistical analysis of the work, FFM participated in performing of scintigraphy and DO contributed in designing of project and revised the draft of manuscript.

Conflict of Interest

The authors report no conflicts of interest.

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