

Original article

Reprint

Magnetic resonance diffusion-weighted whole-body imaging (DWIBS) in the urinary bladder cancer diagnostics

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Abstract: The purpose of the article is to identify the most characteristic and significant changes of magnetic resonance indicators in patients with the urinary bladder cancer during diffusion-weighted whole-body imaging (DWIBS). **Materials:** From September 2009 till April 2011 98 patients have been examined: 61 (62.2%) with morphologically verified bladder cancer and 37 (37.8%) with cystitis. **Results:** The study has revealed that the sensitivity of DWIBS investigation in detecting bladder cancer is 98.36%, specificity is 10.81, and the efficacy is 65.38%. **Conclusions:** DWIBS is an informative noninvasive method for screening diagnostics of bladder cancer, as well as for identifying suspicious areas of regional and distant metastases.

Keywords: urinary bladder cancer (UBC), diffusion magnetic resonance imaging (DMRT).

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Introduction

Primary neoplasms of the urinary bladder account for 2-6% of all tumours [1]. Due and adequate diagnostics of the urinary tract diseases still remains a complex and actual problem of oncology [2].

Diffusion-weighted magnetic resonance tomography (DMRT) which provides functional information can be used in detection and characterization of pathological processes in the area of investigation including malignant tumours. The proposed Diffusion-Weighted Whole-Body Imaging with Background Body Signal Suppression (DWIBS) depicts the whole body volume diffusion. This new conception possesses unique characteristics, differs from standard DMRT and is capable to play an important role in screening diagnostics of oncological diseases [3].

The aim of the study is to identify the most significant magnetic resonance (MR) indicators and DWIBS investigation capabilities in diagnostics of the urinary bladder cancer (UBC).

Material and Methods

During the period from September 2009 till April 2011 there were examined 98 patients at the age of 45-81 suspected of the urinary bladder cancer: 78 males (80%) and 20 females (22%). All the patients were admitted to the Scientific Research Institute of Clinical and Fundamental Urology and Urological Clinic of Saratov State Medical University. 45 patients (45.9%) complained of painful and frequent urination, 53 patients (54.1%) demonstrated the signs of macrohematuria.

Depending on the results of histological investigation the patients were divided into two groups: the basic group consisting

of 61 patients (62.2%) with morphologically verified UBC (average age of 67±13 years), and comparison group consisting of 37 patients (37.8%) with cystitis (average age of 61±11 years). Depending on the UBC stage in accordance with TNM clinical classification designed by International Anticancer Alliance in 2002, the basic group was divided into four subgroups. Stage T1 of the pathological process was diagnosed in 33 patients (54.1%), stage T2 - in 18 patients (29.5%), stage T3 - in 7 patients (11.5%), stage T4 - in 3 patients (4.9%). All patients were performed cystoscopy and ultrasound scan.

MR investigation was performed on three apparatus PHILIPS of 1.5 T magnetic field power with application of the "Body-array" roll for the body. The protocol of MR investigation consisted of two consequent steps. During the first step there was performed standard investigation (T2WI, T1WI, T2WI with fatty tissue signal suppression in three mutually perpendicular planes and VISTA sequence in the axial plane) in order to detect preliminary tumour localization and visualize regional metastatic areas (Table 1).

Table 1. Features of standard MR investigation of pelvic organs

MR stage	MR plane	TR, ms	TE, ms	Fov, mm	Stage duration
T2WI	Sag, ax, cor	3500	80	320	01:38
T1WI	Sag, ax, cor	900	7	320	02:30
T2WI with f.t.s.s.	Sag, ax, cor	3500	7	320	00:56
VISTA	ax	2000	200	130	01:02

T2WI with f.t.s.s. is T2WI with fatty tissue signal suppression.

For adequate urinary bladder fulfilment an hour prior MR investigation the patients had to drink 250-300 ml of water and not to void urine for two hours before the investigation. With the catheter fixed in the urinary bladder directly prior the investigation 250-500 ml of sterile solution were injected into the urinary bladder and the catheter was overpressed.

During the second step there was performed DWIBS investigation including complete 530 mm FOV in RL direction for complete imaging at simultaneous use of less FOV in FH direction for better resolution. Parameters of DWIBS investigation are: fat suppression STIR; b-value (s/mm^2) 1.00; direction of MPGs Phase, frequency and slice phase; frequency and slice; TR (ms) 10,205; TE (ms) 70; TI (ms) 180 NA; parallel imaging factor 2; halfscan factor 0.6; EPI factor 47; slice orientation Axial; no. of slices 60; slice thickness/gap (mm) 4/0; FOV (mm) 400; RFOV (%) 70; matrix 160; scan percentage 70%; actual voxel size (mm^3) 2.5x3.6x4.0; calculated voxel size (mm^3) 1.6x1.6x4.0; no. of signals averaged 10; scan time 7 min. 8 s.

DWIBS investigation resulted in three blocks of native axial scans visualizing the region from the upper third of the thigh up to the head with the possibility of native data processing and three-dimensional inverted image constructing (Fig. 1).

Results

MR semiotics of the detected bladder wall changes in patients of the basic and comparison groups is given in Table 2.

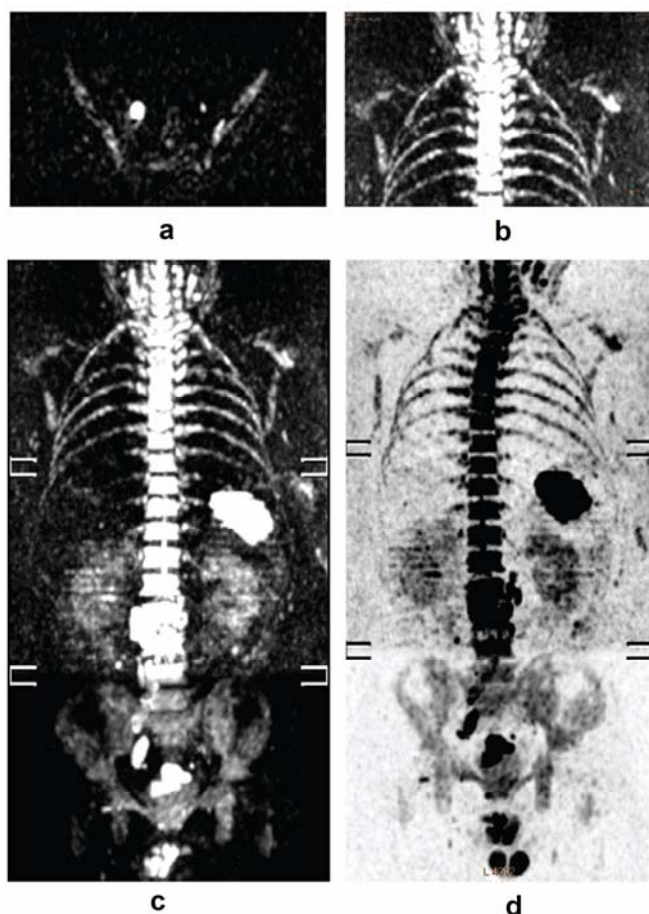


Figure 1. Sequence of formation DWIBS-image: native DWIBS-image in the axial plane (a), local 3D DWIBS-image (b), total 3D DWIBS-image (c), reversed total 3D DWIBS-image (d).

The analysis of the obtained data showed that independently of the pathological process stage lateral bladder walls were impaired more frequently; in stages T1, T3 and T4 - predominantly the right bladder wall: in 48.4%, 57.1% and 66.6% of the patients respectively. Diffuse bladder walls impairment was detected in cystitis.

Frequency of ureter ostia involvement was comparable in patients with stages T1, T3 and T4. At the same time frequency of ureter ostia involvement in the pathological process in patients with stage T2 of UBC was significantly higher and reached 61.1%. Independently of the stage there prevailed solitary bladder wall formations.

With advancing of the stage of the disease frequency of occurrence of local bladder wall thickening in the area of the pathological formation increased: in T1 - 51.5%, in T2 - 94.4%, in T3 and T4 - 100%. Change of MR bladder wall signal was detected in 42.4% of patients with stage T1, in 94.4% patients with stage T2, in 100% of patients with stages T3 and T4 of the disease. The same tendency was detected when evaluating the formation structure homogeneity and enlarged regional lymphatic nodes detection frequency. Regularity of external bladder wall contours in stage T1 was 100%. In stages T3 and T4 external bladder walls in all cases were irregular which corresponded to the verified stages of the pathological process. False signs of spread into the paravesical fat were detected in 6 patients (33.3%) with stage T2. However, in routine MR investigation the possibility to evaluate distant changes in patients with UBC was lacking.

Later comparison and native T2WI with high resolution and images of DWIBS investigation were carried out. Detection of areas with increased signal on native images and with decreased signal on three-dimensional inverted images was the criterion of DWIBS images evaluation. MR signal increase during native DWIBS investigation was normally detected from the brain structures, salivary glands, tonsils, spleen, gallbladder and small intestine (Fig. 2).

The data obtained during DWIBS investigation are given in Table 3. MR signal increase both from the pathological urinary bladder formation and regional and distal lymphatic nodes was detected in 60 patients (98.4%) of the basic group.

In the comparison group signal increase from the urinary bladder wall was not observed in 4 patients (10.8%). During DWIBS investigation of the patients with stage T2 of UBC irregular external bladder wall contours in the area of the malformation (false MR sign of the formation spread into the paravesical fat) were detected in 1 patient (5.6%), while during MR investigation - in 6 patients (33.3%) (Fig. 3).

The results of the carried out investigation demonstrate that DWIBS study sensitivity in detecting urinary bladder cancer is 98.4%, specificity - 10.8%, diagnostic effectiveness of the method - 65.4%.

Discussion

For a long time diffusion magnetic resonance tomography (DMRT) was used only for investigation of the brain.

Extracranial DMRT was complicated by multiple artefacts from the movement and magnetic receptivity which resulted in the loss of diagnostic significance of the given investigation [2]. Increase of gradients and appearance of new multichannel rolls almost solved that problem, but up to the recent time extracranial DMRT was obligatorily carried out at the breath holding, because respiratory movements were the obstacles for DWI due to the shifts during the internal organs respiration [4].

Table 2. MR semiotics of UBC and cystitis in routine MR investigation of the pelvis organs

No.	MR sign	Scale	UBC				Cystitis, n (%) (n=37)
			T1, n (%) (n=33)	T2, n (%) (n=18)	T3, n (%) (n=7)	T4, n (%) (n=3)	
1	What bladder walls were impaired?	right	16 (48.4)	6 (33.3)	4 (57.1)	2 (66.7)	0 (0)
		left	10 (30.3)	5 (27.8)	1 (14.3)	0 (0)	0 (0)
		anterior	2 (6.1)	2 (11.1)	0 (0)	1 (33.3)	0 (0)
		posterior	3 (9.1)	3 (16.7)	2 (28.6)	0 (0)	0 (0)
		urocystic triangle	3 (9.1)	2 (11.1)	0 (0)	0 (0)	0 (0)
2	Ureter ostia involvement	no	20 (60.6)	7 (38.9)	4 (57.1)	2 (66.7)	37 (100)
		right	6 (18.2)	6 (33.3)	3 (42.9)	1 (33.3)	0 (0)
		left	7 (21.2)	4 (22.2)	0 (0)	0 (0)	0 (0)
		both	0 (0)	1 (5.6)	0 (0)	0 (0)	0 (0)
3	Number of bladder wall formations	diffuse changes	2 (6.1)	2 (11.1)	0 (0)	0 (0)	37 (100)
		one thing	23 (69.7)	15 (83.3)	6 (85.7)	3 (100)	0 (0)
		two	3 (9.1)	0 (0)	1 (14.3)	0 (0)	0 (0)
		three	3 (9.1)	0 (0)	0 (0)	0 (0)	0 (0)
		more than three	0 (0)	1 (5.6)	0 (0)	0 (0)	0 (0)
4	Bladder wall thickening	yes	17 (51.5)	17 (94.4)	7 (100)	3 (100)	37 (100)
		no	16 (48.5)	1 (5.6)	0 (0)	0 (0)	0 (0)
5	Change of MR signal from bladder wall	yes	14 (42.4)	17 (94.4)	7 (100)	3 (100)	37 (100)
		no	19 (57.6)	1 (5.6)	0 (0)	0 (0)	0 (0)
6	External contours of wall formations	smooth	33 (100)	12 (66.7)	0 (0)	0 (0)	0 (0)
		rough	0 (0)	6 (33.3)	7 (100)	3 (100)	0 (0)
7	Regional lymph nodes (an increase of over 0.8 cm)	no	33 (100)	11 (61.1)	3 (42.9)	1 (33.3)	35 (94.6)
		single	0 (0)	7 (38.9)	4 (57.1)	2 (66.7)	2 (5.41)
		conglomerate	0 (0)	0 (0)	0 (0)	0 (0)	0 (0)
8	Homogeneity of the structure of bladder wall formations	yes	28 (84.8)	10 (55.6)	3 (42.9)	0 (0)	37 (100)
		no	5 (15.2)	8 (44.4)	4 (57.1)	3 (100)	0 (0)

Table 3. MR semiotics of UBC and cystitis in DWIBS investigation

No.	MR sign	Scale	UBC				Cystitis, n (%) (n=37)
			T1, n (%) (n=33)	T2, n (%) (n=18)	T3, n (%) (n=7)	T4, n (%) (n=3)	
1	MR signal increase from urinary bladder wall	yes	32 (97.0)	18 (100)	7 (100)	3 (100)	33 (89.2)
		no	1 (3.0)	0 (0)	0 (0)	0 (0)	4 (10.8)
2	MR signal increase from regional lymph nodes	yes	33 (100)	18 (100)	0 (0)	3 (100)	37 (100)
		no	0 (0)	0 (0)	7 (100)	0 (0)	0 (0)
3	MR signal increase from distant lymph nodes	yes	33 (100)	18 (100)	0 (0)	3 (100)	37 (100)
		no	0 (0)	0 (0)	7 (100)	0 (0)	0 (0)
4	External contours of urinary bladder wall	smooth	33 (100)	17 (94.4)	0 (0)	0 (0)	37 (100)
		rough	0 (0)	1 (5.6)	7 (100)	3 (100)	0 (0)

T. Takahara et al. in 2004 reported about a new unique conception of the whole body DWI called "diffusion-weighted imaging of the whole body with the background body signal suppression" (DWIBS) [5]. Most researchers report that DWIBS investigation is a means for detecting lymphatic nodes independently of their histological composition [5, 6]. The present study supports this viewpoint.

The results of the carried out study show that sensitivity of DWIBS investigation in detecting bladder cancer is high (98.4%) but low specificity parameters (10.8%) allow to judge rather about screening role of the given technique in complex diagnostics of the urinary bladder cancer. Rather significant results of the

investigation, to our opinion, are relevant differences in detecting MR signs during DWIBS investigation and routine MRT of false spread of the bladder wall formation into the paravesical fat, which allows to differentiate stages T2 and T3 of the urinary bladder cancer.

Evaluation of the bladder neoplasm, possible regional and distant metastases allows to consider another way of potential DWIBS application - as the means of radiation- and chemotherapy effectiveness evaluation. DWIBS investigation makes it possible to visualize the signs and characteristics of the detected areas of altered diffusion and to reveal pre-structural functional changes associated with the tumour [7].

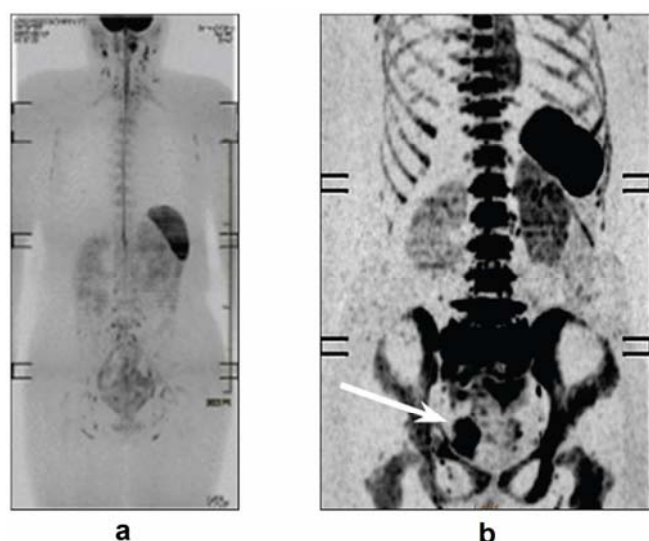


Figure 2. Reversed total 3D DWIBS-image in healthy subject (a) and UBC patients (b). The arrow indicates area of low power signal corresponding to UBC.

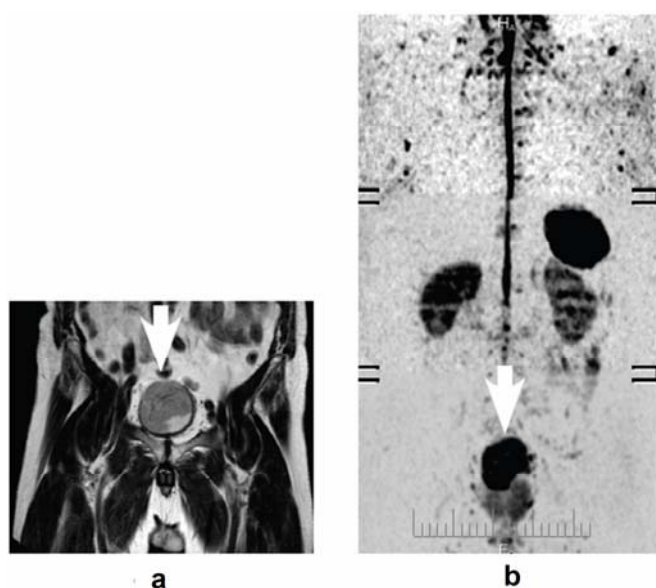


Figure 3. MR investigation: T2 WI-image (a) and DWIBS-image (b). The arrow indicates bladder wall formations, which has a homogeneous structure (T2 stage according TNM). External contours of wall formations is rough on T2 WI –image (signs distribution in perivesical tissue) and is smooth on DWIBS-image (no signs distribution in perivesical tissue).

Conclusions

DWIBS investigation in a number of cases verifies stages of the pathological process, i.e. allows to differentiate stages T2 and T3 of the urinary bladder cancer. Considering high sensitivity and low specificity of DWIBS investigation the described noninvasive informative technique could be used, basically, in screening diagnostics of UBC and in determining indications for expansive anatomical magnetic resonance tomography.

Conflict of interest: none declared.

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