Lack of risk factors predicting the development of genital mycotic infections among patients treated with dapagliflozin: the ABCD Nationwide Dapagliflozin Audit KY Thong¹, M Yadagiri², P Tong³, D Barnes⁴, D Morris⁵, T Chowdhury⁶, LL Chuah⁷, M Cohen⁸, A Kennedy⁹, T Robinson¹⁰, SC Bain¹¹, KA Adamson¹², REJ Ryder², ABCD Nationwide Dapagliflozin Audit contributors

University of Western Australia, Perth, Australia (1), City Hospital, Birmingham, UK(2), Qualigenics Medical, Central, Hong Kong(3), The Tunbridge Wells Hospital, Pembury, UK(4), Northgate Heatlh Centre, Shropshire UK(5), The Royal London Hospital, London, UK(6), Croydon Health Services NHS Trust, Croydon, UK(7), Barnet Hospital, Barnet, UK(8), Antrim Area Hospital, Antrim, UK(9), Royal United Hospitals, Bath, UK(10), Abertawe Bro Morgannwyg University Board, Swansea, UK (11), St John's Hospital, Livingston, UK(12)

ABSTRACT

Treatment of type 2 diabetes with sodium-glucose cotransporter 2 (SGLT2) inhibitors can result in genital mycotic infections. We investigated possible risk factors for developing such infections among patients treated with the SGLT2 inhibitor dapagliflozin. The Association of British Clinical Diabetologists (ABCD) nationwide dapagliflozin audit collected data on patients treated with dapagliflozin in routine clinical practice. Fifty nine diabetes centres participated. Using data collected, we assessed possible associations of patient's age, duration of diabetes, body mass index (BMI), glycated haemoglobin (HbA1c), renal function, sex, ethnicity, prior genital mycotic infection, urinary tract infection, urinary incontinence or nocturia with the occurrence of ≥ 1 genital mycotic infection within 26 weeks of treatment. A total of 1049 patients (476 women and 573 men) were analysed. Baseline characteristics were, mean (\pm SD), age 56.7 years (± 10.2) , BMI 35.5 kg/m² (± 6.9) and HbA1c 9.4% (± 1.5) , 76.2% Caucasian and 3.5% had prior history of genital mycotic infection. Only patient sex (13.2%) women vs 3.3% men) and prior history of genital mycotic infection (21.6% vs 7.3%) were found to be associated with the development of genital mycotic infections after dapagliflozin treatment (adjusted OR 4.22 [95%CI 2.48,7.19], p<0.001) and (adjusted OR 2.41 [95% CI 1.04,5.57], p=0.039), respectively. Women and patients with a history of genital mycotic infections had higher risks of developing genital mycotic infections with dapagliflozin treatment.

To investigate clinical factors associated with the development of genital mycotic infections (GMI) among patients treated with dapagliflozin.

- September 2016
- 59 diabetes centres participated
- 1049 patients analysed

- dapagliflozin

AIMS

SUBJECTS

• ABCD invited diabetes centres across UK to submit data on patients treated with dapagliflozin in routine clinical practice. Data presented are from October 2014 to

• Excluded from 1116 patients • <12 weeks of data (n=64) Previous SGLT2 inhibitor use (n=3)

ANALYSES

• Patients were divided into those with/without symptoms or confirmed GMI within 26 weeks of commencing

We compared baseline factors between patients with/without GMI using t-test and Chi-Square/Fisher's exact test.

RESULTS

	n	≥1 Genital mycotic infection		
		Yes	No	
Age (years)	1049	56 ± 11	57 ± 10	
Duration of diabetes (years)	1048	11 ± 6	12 ± 7	
HbA1c (%; mmol/mol)	1045	9.2 ± 1.5; 77 ± 16	$9.4 \pm 1.5;$ 79 ± 16	
Weight (kg)	1046	97.7 ± 25.1	101.2 ± 22.7	
Body mass index (kg/m²)	1023	34.0 ± 7.4	35.4 ± 6.9	
eGFR* (ml/min/1.73m²)	932	92 ± 18	90 ± 18	

		≥1 Genital my	F			
		Yes	No			
	Sex Female Male	63 (13.2) 19 (3.3)	413 (86.8) 554 (96.7)	·		
	Ethnicity (n=947) Caucasian Asian Afro-Carribean Mixed	51 (7.1) 20 (10.8) 3 (10.7) 1 (9.1)	671 (92.9) 166 (89.2) 25 (89.3) 10 (90.9)	0.11 (v 0.49 (v		
	Previous genital fungal infection <1 year Yes No	8 (21.6) 74 (7.3)	29 (78.4) 938 (92.7)			
	Previous urinary tract infection <1 year Yes No	2 (9.5) 80 (7.8)	19 (90.5) 948 (92.2)			
	History of urinary incontinence Yes No	2 (15.4) 80 (7.7)	11 (84.6) 956 (92.3)			
	History of nocturia Yes No	11 (13.6) 71 (7.3)	70 (86.4) 897(92.7)			

P value	
0.47	
0.27	
0.18	
0.23	
0.53	
0.32	

P value	
<0.001	
- /s Caucasian) /s Caucasian) -	
0.007	

0.27

0.06



RESULTS

Variables associated with GMI:

Patient sex (13.2% women vs 3.3% men) (adjusted OR 4.22 [95%CI 2.48,7.19], p<0.001)

Prior history of GMI (21.6% vs 7.3%) (adjusted OR 2.41 [95% CI 1.04,5.57], p=0.039)

CONCLUSION

Women and patients with a history of genital mycotic infections had higher risks of developing genital mycotic infections with dapagliflozin treatment

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