Case Report



Case report of ecstasy-induced renal venous thrombosis

Mohamed Tarek Eldehni¹, Ian S.D. Roberts², Ramesh Naik¹ and Emma Vaux¹

¹Department of Renal Medicine, Royal Berkshire NHS Foundation Trust, Reading, UK and ²Department of Cellular Pathology, John Radcliffe Hospital, Oxford, UK

Correspondence and offprint requests to: Mohamed Tarek Eldehni; E-mail: tarek.eldehni@nhs.net

Abstract

The use of 3,4-methylenedioxymethamphetamine, also known as MDMA or ecstasy, has been associated with vascular and end-organ damage. This case report describes, with histological evidence, the development renal venous thrombosis presenting with acute kidney injury following oral ingestion of 3,4-methylenedioxymethamphetamine (ecstasy).

Keywords: ecstasy; renal; thrombosis; venous

Introduction

This case describes the development of acute kidney injury following oral ingestion of 3,4-methylenedioxymethamphetamine (ecstasy) with histological demonstration of venous thrombosis.

Case report

A 22-year-old man presented with acute bilateral loin pain and was referred by his general practitioner for a urological consultation with suspected renal stones. He was on no regular medication, and at the beginning of his admission, he denied taking any recreational drugs. However, later on, he admitted taking ecstasy 1 day prior to his admission.

When examined, he was apyrexial but hypertensive with a blood pressure of 145/95 mmHg. His cardiovascular, respiratory and abdominal examinations were normal. Initial biochemical investigations showed evidence of acute kidney injury with a creatinine of 189 µmol/L. A creatinine of 73 µmol/L was documented 4 years previously. Other biochemistry results were Na 138 mmol/L, K 4.2 mmol/L and urea 7.7 mmol/L. Anti-neutrophil cytoplasmic antibodies (ANCA) and anti-nuclear antibodies (ANA) were negative; complement and immunoglobulins were normal. His urine dipstick was positive for both blood and protein. The protein–creatinine ratio was 92 mg/mmol. Renal ultrasound showed normal-sized kidneys and no renal stones.

Given the unexplained acute kidney injury with significant proteinuria and haematuria, he underwent ultrasoundguided renal biopsy. Over the next 3 days of admission, his renal function continued to improve, and his proteinuria started to disappear (Table 1).

Renal biopsy histology: A small vein at the corticomedullary junction was occluded by thrombus with evidence of early organization (Figure 1). Glomeruli, tubules and a small artery present appeared normal. There was no evidence of glomerulonephritis or tubulointerstitial nephritis.

A renal Doppler ultrasound did not show a thrombus in the main renal vein bilaterally. His renal function normalized, and his proteinuria resolved. Thrombophilia screen including antiphospholipid antibodies was negative. He was discharged with the diagnosis of acute kidney injury due to renal venous thrombosis induced by ecstasy.

Discussion

3,4-Methylenedioxymethamphetamine, also known as MDMA or ecstasy, is an amphetamine derivative. Its use has been associated with different organ failure such as acute kidney injury and fulminant hepatic failure [1]. The vascular effects of MDMA are poorly defined. Firstly, vasoconstriction caused by MDMA is described in animal models with effects such as increase in mean blood pressure [2], cutaneous vasoconstriction [3] and coronary artery vasoconstriction [4]. Moreover, necrotizing vasculopathy was described in a case report of a patient who used MDMA 10 days prior to presentation with acute kidney injury resulting in chronic renal failure [5]. Necrotizing vasculopathy was also reported in two transplanted renal grafts taken from the same donor who regularly used ecstasy; both grafts

Table 1. Progression of renal function with proteinuria during admission

Admission day	Creatinine (µmol/L)	Protein-creatinine ratio (mg/mmol)
1	189	92
2	254	101
3	215	39
4	128	
10	106	

© The Author 2010. Published by Oxford University Press on behalf of ERA-EDTA. All rights reserved. For permissions, please e-mail: journals.permissions@oxfordjournals.org

failed within the first week due to necrotizing graft vasculopathy [6]. Finally, there have been case reports of extra-renal venous thrombosis associated with the use of ecstasy such as a case of cerebral venous thrombosis in a 22-year-old woman who ingested one tablet of ecstasy prior to presentation with headache and photophobia [7]. Although

thrombus with evidence of early organization.

as a case of cerebral venous thrombosis in a 22-year-old woman who ingested one tablet of ecstasy prior to presentation with headache and photophobia [7]. Although MDMA can cause disseminated intravascular coagulation [1], venous thrombosis could also be a result of hyperthermia and dehydration induced by MDMA. We describe a case of acute kidney injury most likely caused by renal ve-

Fig. 1. A small vein at the corticomedullary junction was occluded by

nous thrombosis induced by ecstasy ingestion. The venous thrombosis involved small intra-renal veins, evident on renal biopsy, but did not include the main renal veins. This is supported by the degree of the acute kidney injury and the resolution of both the injury and the proteinuria.

Conflict of interest statement. None declared.

References

- Campbell GA, Rosner MH. The agony of ecstasy: MDMA (3,4methylenedioxymethamphetamine) and the kidney. *Clin J Am Soc Nephrol* 2008; 3: 1852–1860
- Vandeputte C, Docherty JR. Vascular actions of 3,4-methylenedioxymethamphetamine in alpha(2A/D)-adrenoceptor knockout mice. *Eur J Pharmacol* 2002; 457: 45–49
- Pedersen NP, Blessing WW. Cutaneous vasoconstriction contributes to hyperthermia induced by 3,4-methylenedioxymethamphetamine (ecstasy) in conscious rabbits. *J Neurosci* 2001; 21: 8648–8654
- Baker KE, Herbert AA, Broadley KJ. Vasoconstriction of porcine left anterior descending coronary artery by ecstasy and cathinone is not an indirect sympathomimetic effect. *Vascul Pharmacol* 2007; 47: 10–17
- Bingham C, Beaman M, Nicholls AJ et al. Necrotizing renal vasculopathy resulting in chronic renal failure after ingestion of methamphetamine and 3,4-methylenedioxymethamphetamine ('ecstasy'). Nephrol Dial Transplant 1998; 13: 2654–2655
- Hurault de Ligny B *et al.* Early loss of two renal grafts obtained from the same donor: role of ecstasy? *Transplantation* 2005; 80: 153–156
 Rothwell PM, Grant R. Cerebral venous sinus thrombosis induced by
- Kotnweit PM, Graft R. Cerebral venous sinus thrombosis induced by 'ecstasy'. J Neurol Neurosurg Psychiatry 1993; 56: 1035

Received for publication: 22.4.10; Accepted in revised form: 23.4.10

