

**A germline insertion in the tuberous sclerosis (*Tsc2*) gene gives rise to the Eker rat model of dominantly inherited cancer**

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Table 1 was inadvertently omitted. The table is shown below.

**Table 1 Comparison of the principal features of the Eker rat model and tuberous sclerosis**

	Eker rat	Tuberous sclerosis
Inheritance	Autosomal dominant	Autosomal dominant
Defective gene	<i>Tsc2</i>	<i>TSC1</i> and <i>TSC2</i>
Clinical		
Renal lesion	100%, renal cell carcinoma	Often, angiomyolipoma Rare, renal cell carcinoma
Skin lesion	No	Frequent, angiofibroma ungual fibroma
Neurological abnormalities	No	Frequent, convulsion mental retardation
Heart lesion	No	Often, rhabdomyoma
Pituitary adenoma	Often	No
Spleen tumour (probable hemangiosarcoma)	Often	No
Uterus tumour (probable leiomyosarcoma)	Often	No

**Establishing the identity of Anna Anderson Manahan**

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Due to a printing error, Table 1 and the order of authors were incorrectly published. The correct versions are shown below.

**Table 1 Short tandem repeat STR results**

	VWA	TH01	F13A1	FES/FPS	ACTBP2	AMELOGENIN
Tsar (Skeleton 4) <sup>a</sup>	15,16	7,9,3 <sup>b</sup>	7,7	12,12	11,32	X,Y
Tsarina (Skeleton 7) <sup>a</sup>	15,16	8,8	3.2 <sup>c</sup> ,5	12,13	32,36	X,X
Anna Anderson (intestine sample)	14,16	7,9,3	3.2,7	11,12	15,18	X,X

<sup>a</sup>Ref. 1.

<sup>b</sup>9,3 and 10 alleles are grouped and designated 9.3.

<sup>c</sup>3.2 allele was previously designated as a 3 allele under the old nomenclature.

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