

## A Tale of Two Tails

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### Abstract

Neonatal Priapism is rare and usually self-limiting. We describe the first reported case in a father and son and describe the management.

### Case Report

A baby boy was born at term plus seven days to a fit and well 19 year old mother.

She had no antenatal problems and had one previous baby who was fit and well. On day one a midwife noticed a persistently erect penis. Doppler blood pressure was normal and a full blood count showed a haematocrit of 64% and haemoglobin of 24.7, excluding polycythaemia.

The boy's grandmother informed us that her son, the baby's father, also had a persistent erection at birth.

In adults priapism can cause long term damage. We performed a literature search to see to see if intervention was necessary.

### References

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Searching for priapism and restricting to neonatal age, 15 case reports were Identified<sup>1-14</sup>. One was excluded because it dealt with onset at 37 days<sup>1</sup>. Four cases were related to polycythaemia<sup>2-4</sup> but only one was managed with phlebotomy<sup>5</sup>. A case of idiopathic priapism was managed with IV ketamine<sup>7</sup>. All cases returned to normal between two and six days of age<sup>2-10</sup>. Since we saw the case, two more have been described. One did develop pyocavernositis<sup>11</sup> but the other resolved spontaneously<sup>12</sup>. Some authors recommend Doppler ultrasound studies to differentiate between ischemia (likely to be due to polycythaemia or sickle cell disease) and non-ischaemic<sup>1,3,11,12</sup>.

We watched and waited without Doppler ultrasound. The priapism resolved at five days and at four months had not recurred.

This is now the 18th case reported. We cannot explain the father son relationship in this presentation. Can any of your bright readers?