

## CLINICAL ALERT

# Intracranial complications of *Serratia marcescens* infection in neonates

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Even though *Serratia marcescens* is not one of the most common causes of infection in neonates, it is associated with grave morbidity and mortality. We describe the evolution of brain parenchymal affectation observed in association with *S. marcescens* infection in neonates. This retrospective case series details brain ultrasound findings of five neonates with hospital-acquired *S. marcescens* infection. Neonatal *S. marcescens* infection with or without associated meningitis can be complicated by brain parenchymal affectation, leading to cerebral abscess formation. It is recommended that all neonates with this infection should undergo neuro-imaging more than once before discharge from hospital; this can be achieved using bedside ultrasonography.

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*Serratia marcescens* infection has occurred sporadically from 2010 to 2014 in our 126-bed neonatal unit, with a tendency towards cluster infections. In each instance, staff at the Unit for Infection Prevention and Control, Tygerberg Hospital, Cape Town, South Africa employed their expertise to investigate, identify and manage appropriately the root cause of the cluster infection. On two separate occasions, colonised respiratory support equipment and suction points were identified as sources of infection; this situation was rectified. We recently noted a pattern of brain ultrasound findings consistently associated with the identification of *S. marcescens* septicaemia in neonates. These findings have not been noted in septicaemia secondary to other micro-organisms. The majority of neonates with such ultrasound findings died, despite intensive care and appropriate antimicrobial treatment having been instituted.

We describe a case series of five neonates with *S. marcescens* nosocomial infection in whom the abovementioned pattern was evident. In patient 1, autopsy characterisation of these findings was done.

Ethical approval for the case series and a waiver of individual informed consent were granted by the Human Health Research Ethics Committee of Stellenbosch University, protocol number N15/08/068.

## Case 1

Patient 1 was delivered by caesarean section because of maternal pre-eclamptic toxemia and fetal affectation at an estimated 30 weeks

of gestation (birth weight 1 460 g). Apart from antiretroviral, antihypertensive and antenatal corticosteroids medication, his mother had received no other treatment and had had a healthy pregnancy. He progressed well and was admitted to the neonatal high-care ward for further care because of a very low birth weight. He had a normal baseline full blood count and was treated with intravenous aminophylline and oral nevirapine for apnoea and HIV prophylaxis, respectively. He was commenced on formula feeds on his second day of life and required no respiratory support.

On day 6, he developed respiratory distress and progressed to hypoxic respiratory failure accompanied by convulsions, for which he had to be intubated and transferred to the neonatal intensive care unit (NICU) for ventilatory support. At this time, an infection screen was performed and he was commenced on meropenem and vancomycin antimicrobial therapy. A cranial ultrasound scan done on admission to the NICU showed a premature brain compatible with an estimated gestational age of 32 weeks, minimal vessel pulsation and predominant findings in the bilateral parieto-occipital area. The latter had well-defined hyperechogenicities with areas of cystic breakdown, predominantly on the left side. These findings were thought to represent ischaemic parenchymal injury and areas of white matter infarcts. The patient had a persistent low-voltage pattern on amplitude-integrated electro-encephalography, remained

in intractable hypotension and metabolic acidosis and succumbed to his illness within a day of admission to the NICU. His parents granted permission for an autopsy, and soon after he died, telephonic notification of Gram-negative bacilli on blood culture was received from the microbiology laboratory. This was later confirmed to be extended-spectrum beta-lactamase (ESBL)-producing *S. marcescens*.

Fig. 1 shows the predominant cranial ultrasound results of right parieto-occipital wedge-shaped hyperechogenicity in this patient; moderate lenticulostriate vasculopathy can also be

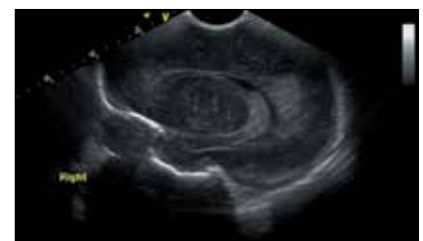


Fig. 1. Right parasagittal image.

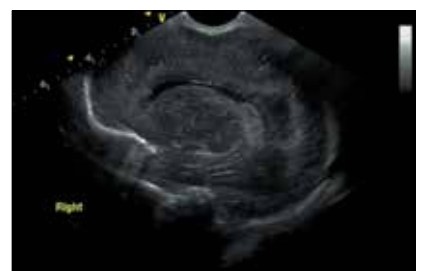


Fig. 2. Right parasagittal image approximately 12 hours later.

seen. Cranial ultrasonography was performed through the anterior fontanelle with an 8 MHz curvilinear probe using a Vivid S5 General Electric Healthcare ultrasound machine (USA).

Fig. 2 shows progressive change to cystic breakdown of the same area approximately 12 hours later.

In Fig. 3, the beginning of cystic breakdown within a hyperechogenic area is noted in the left posterior parietal region.

Autopsy findings included extensive white matter necrosis of the temporal lobes, haemorrhagic infarction of temporal and occipital lobes, leptomeningitis in sections of the parietal cortex, and haemorrhagic necrosis of the cerebellum. The brainstem was also noted to be disintegrated.

**Cases 2 - 5**

Details of four other patients with similar clinical and microbiological findings are given in Table 1, and the associated cranial ultrasound findings can be seen in Fig. 4.

The images of patient 2 were obtained with an 8 MHz sector probe using a Vivid S5 General Electric Healthcare ultrasound

machine. The images of patients 3, 4 and 5 were obtained using a 10 MHz sector probe (Antares Sonoline Siemens Medical Solutions, USA).

**Discussion**

*S. marcescens* is an Enterobacteriaceae micro-organism that is widespread in the environment, but not a common component of the human faecal flora. Most infections are acquired exogenously and spread in and among hospitals.<sup>[1]</sup> Hand-held transmission appears to be the primary mechanism of nosocomial spread. Environmental sources in hospitals have been found to include suction taps, colonised disinfectants or soap, and inadequately sterilised breast pumps and respirators. Disease in humans has been associated with infection of the urinary tract, respiratory tract, local wounds and central vein catheters. Colonisation in neonatal units has been associated with up to a tenfold increase in the rate of *S. marcescens* bacteraemia and meningitis.<sup>[2]</sup> Significant brain injury caused by ventriculitis, brain abscesses or porencephalic cysts is described in the majority of infants with meningitis. In our neonatal unit, we have noted intracranial complications in the presence or absence of meningitis.

NICUs in developed countries have reported serial infection outbreaks with this micro-organism over the past decade, although not all of these focused primarily on associated neurological morbidity. In a recently published report from Italy,<sup>[3]</sup> sepsis and pneumonia were the most severe clinical features described, with a mortality

rate of 7%. Only 1 of 18 affected neonates was found to have meningitis and a brain abscess. This neonate survived.<sup>[3,4]</sup> Published reports on neonatal *S. marcescens* infection in developing countries are limited with regard to availability and content. A global report on causes of neonatal sepsis in developing countries cited *Serratia* species as the fifth most common cause, accounting for 0.5% and 0.3% of early and late neonatal sepsis, respectively.<sup>[5]</sup> In their report on bloodstream infections in neonates in a developing country over a 12-month period, Ballot *et al.*<sup>[6]</sup> do not mention *S. marcescens* as one of the causative organisms. Unpublished data from our institution detailing infection patterns in very-low-birth-weight infants over an 18-month period, found that ESBL-producing *S. marcescens* accounted for 12% of late-onset neonatal sepsis. *S. marcescens* and ESBL-producing *S. marcescens* accounted for 5% and 8%, respectively, of infection-related mortality over a 24-month period in our unit.

Published case reports describe diagnosis of brain abscesses after an initial diagnosis of *S. marcescens* septicaemia and meningitis.<sup>[7,8]</sup> In our unit, three babies on whom cerebrospinal fluid (CSF) analysis had not been done died within 24 hours of onset of illness. Of the two survivors we describe, one had abnormal CSF analysis results of high protein, pleiocytosis and no organisms on culture, while the other had normal CSF results on two separate occasions. In both instances, the lumbar puncture and CSF analysis were done only after the diagnosis of septicaemia and commencement of

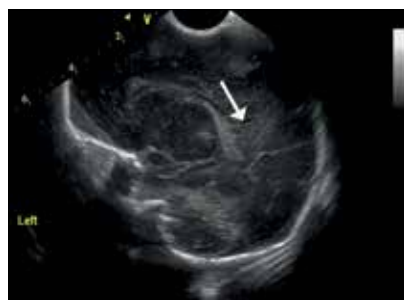


Fig. 3. Left parasagittal image.

**Table 1. Summary of details of patients with *Serratia marcescens***

Patient details	Age at <i>Serratia</i> diagnosis	Cranial ultrasound findings	Outcome
Case 2 GA 29 wks BW 1 608 g Positive	6 days CUS on day 6	Minimal cerebrovascular pulsation. Right-side moderate lenticulostriate vasculopathy. Bilateral extensive frontoparietal well-circumscribed hyperechogenicity with areas of echolucent breakdown	Died on day 6
Case 3 GA 31 wks BW 1 370 g Negative	7 days CUS on day 7	Bilateral frontoparietal parenchymal hyperechogenicity with areas of echolucent cystic breakdown. Bilateral basal ganglia/thalamic hyperechogenicity	Died on day 7
Case 4 GA 33 wks BW 1 550 g Positive	9 days CUS on days 9, 11 and 16	Initial bilateral anterior and posterior irregular hyperechogenicity. Progressed to echolucent cystic breakdown 48 hours later and to well-circumscribed thick-walled lesions with central echolucency 7 days later – suggestive of abscess formation	Alive on discharge
Case 5 GA 26 wks BW 840 g Negative	6 days CUS on days 33 and 47	Bilateral frontoparietal and parieto-occipital well-circumscribed thick-walled echolucent cystic lesions suggestive of abscess formation	Alive on discharge

GA = gestational age; BW = birth weight; CUS = cranial ultrasound findings; positive/negative relates to maternal HIV serology.





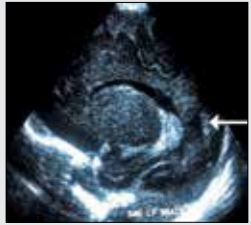



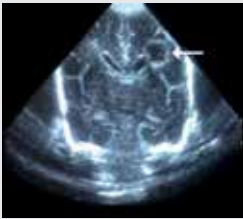


Patient	Image 1	Image 2
Case 2: right and left parasagittal views		
Case 3: coronal and right parasagittal views		
Case 4: left and right parasagittal views		
Case 4: 7 days later, left and right parasagittal views		
Case 5: coronal views		
Case 5: parasagittal left view		

Fig. 4. Patient cranial ultrasound images.

antibiotic therapy. With the exception of only one patient, we describe changes in the brain before the diagnosis of *Serratia* infection had been made in the babies with acute onset of illness. We also describe the progression and changing pattern

of these lesions. The progression seems to be that of ill-defined parenchymal hyperechogenicities, leading to cystic breakdown of these areas within 24 hours and culminating in areas of well-defined thick-walled cysts containing debris. The

sizes of these thick-walled cysts tend to remain the same on serial imaging. We postulate that the initial changes represent areas of ischaemic injury, which progress to infarction and liquefaction, and that the cysts represent abscess formation within the brain parenchyma.

We conclude that in a neonate who suddenly and unexpectedly becomes ill, cranial ultrasonography should be an urgent additional investigation. If changes such as inhomogeneous hyperechogenicities are noted, brain affection by a serious bacterial infection such as *S. marcescens* should be considered and appropriate antibiotic cover instituted without delay. Follow-up imaging within 24 - 48 hours and again within 7 and 14 days in survivors will aid in identifying progression of injury and planning for neurosurgical intervention and rehabilitation. We suggest that cranial ultrasonography should be performed more than once before discharge from hospital in all neonates diagnosed with *Serratia* septicaemia, regardless of the presence or absence of neurological symptoms or an abnormal CSF analysis. This is based on our subsequent observations of intracranial complications, including abscess formation in neurologically asymptomatic neonates with normal CSF findings, as in the case of patient 5. We also propose that good-quality bedside neuro-imaging can be undertaken with ultrasonography as often as clinically required in these critically ill neonates, without the need to move them from the NICU.

1. Donnenberg MS. Enterobacteriaceae. In: Mandell GL, Bennett JE, Dolin R, eds. Principles and Practice of Infectious Diseases, vol 2. 7th ed. Philadelphia, USA: Churchill Livingstone Elsevier, 2010.
2. Gruber WC, Fisher RG. *Serratia*. In: Feigin RD, Cherry JD, eds. Feigin and Cherry Textbook of Paediatric Infectious Diseases, vol. 1. 4th ed. Philadelphia, USA: Saunders, 1998.
3. Casolari C, Pecorari M, Della Casa E, et al. *Serratia marcescens* in a neonatal intensive care unit: Two long-term multiclonal outbreaks in a 10-year observational study. *New Microbiol* 2013;36:373-383.
4. Samuelsson A, Isaksson B, Hanberger H, Olhager E. Late-onset neonatal sepsis, risk factors and interventions: An analysis of recurrent outbreaks of *Serratia marcescens*, 2006 - 2011. *J Hosp Infect* 2014;86:57-63. DOI:10.1016/j.jhin.2013.09.017
5. Zaidi AKM, Thaver D, Ali SA, Khan TA. Pathogens associated with sepsis in newborns and young infants in developing countries. *Pediatr Infect Dis J* 2009;28:S10-S18.
6. Ballot DE, Nana T, Sriruttan C, Cooper PA. Bacterial bloodstream infections in neonates in a developing country. *ISRN Pediatrics* 2012;2012:508512. DOI:10.5402/2012/508512
7. Hirooka MT, de Vasconcellos Fontes RB, Diniz EM, Pinto FC, Matushita H. Cerebral abscess caused by *Serratia marcescens* in a premature neonate. *Arq Neuropsiquiatr* 2007;65(4-A):1018-1021.
8. Messerschmidt A, Prayer D, Olischar M, Pollak A, Birnbacher R. Brain abscesses after *Serratia marcescens* infection on a neonatal intensive care unit: Differences on serial imaging. *Neuroradiology* 2004;46(2):148-152.

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