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Brief report

Mother as a vector of *Salmonella enterica* serotype Newport outbreak in a neonatal unit



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ABSTRACT

Introduction: In neonatal units, *Salmonella* infections have been attributed to food-borne sources and person to person transmission.

Methods: The outbreak described is the first reported by *Salmonella enterica* serotype Newport in a neonatal facility in Spain.

Results: The index case was an 8-day premature newborn that developed clinical sepsis with positive blood cultures. The outbreak source was the mother of the index case.

Conclusions: It is essential to improve infection control measures taking into account the parents, as they can be an important source of infection.

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Una madre como vector de un brote de *Salmonella enterica* serotipo Newport en una unidad neonatal

RESUMEN

Introducción: En unidades de neonatología, las infecciones por *Salmonella* han sido atribuidas a fuentes alimentarias y a transmisión de persona a persona.

Métodos: El brote descrito es el primero ocasionado por *Salmonella enterica* serotipo Newport en una unidad neonatal en España.

Resultados: El caso índice era un recién nacido prematuro que desarrolló sepsis clínica con hemocultivos positivos. La fuente del brote fue la madre del caso índice.

Conclusiones: Al implementar medidas de control de la infección nosocomial se debe tener en cuenta a los padres ya que pueden constituir una fuente importante de infección.

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Palabras clave:

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Introduction

Nontyphoidal *Salmonellae* are important causes of reportable foodborne infection. These bacteria are especially problematic in a wide variety of individuals especially infants, elderly and immunocompromised hosts and can lead to serious conditions

such as gastroenteritis, bacteraemia, septic arthritis, meningitis and pneumonia.¹ *Salmonella* as an etiologic cause of nosocomial infections is considered an infrequent agent. Health care-associated *Salmonella* outbreaks in neonatal units are rarely described but could be control challenging because of its persistence in the environment, the high prevalence of asymptomatic forms and prolonged carriage in infants.² In neonatal units, *Salmonella* infections have been attributed to food-borne sources, including contaminated infant formula and breast milk^{3,4} and person to person transmission by means of infected health care workers (HCWs)

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and fomites.^{5,6} In Europe, during 2008, *Salmonella enterica* serotype Newport only accounted for 0.7% of all salmonellosis reported cases according to the European Surveillance System (TESSy), with the fifth place among the other serotypes.⁷ We described an outbreak of *S. enterica* serotype Newport gastroenteritis in a neonatal unit in Spain.

Methods

Description of the outbreak

The outbreak occurred in the neonatal ward of the Hospital Universitari Vall d'Hebron (Barcelona, Spain). This is a 75-bed unit divided into 3 subunits: low-risk care (30 beds), intermediate care (20 beds) and intensive care unit (25 beds). The outbreak occurred in the low-risk care unit; this unit has 3 rooms communicated between them (room 4, 5 and 6). The outbreak period was defined from January 28th to February 15th, 2008. The index case was an 8-day premature newborn (34 weeks gestational-age) hospitalized since birth in room 4. On January 30th he developed gastroenteritis with dehydration and clinical sepsis, the blood and stool cultures were positive for *S. newport*. Empiric antibiotic treatment was initiated with ampicillin and gentamicin, and then changed to cefotaxime according to results of antimicrobial susceptibility. Contact isolation precautions were initiated and the epidemiological investigation was undertaken by the infection control team to trace the source of infection.

Epidemiological and microbiological investigation

A case of nosocomial *Salmonellosis* was defined as either the isolation of *S. newport* from blood or stool specimen or more than 1 episode of diarrhea and/or vomiting in an infant who was hospitalized in the neonatal unit during the outbreak period. The following variables were collected for each case: age, sex, gestational age, birth weight, date of birth, date of admission, hospitalization room, instrumentations, oral medications, type of milk or powdered formula, symptoms and date of onset, gastrointestinal symptoms in family members, results of cultures and outcomes.

Hospital staff members who worked in low-risk care unit during the outbreak period and parents of infant cases were interviewed to assess the presence of gastrointestinal symptoms and were requested to submit stool samples for *Salmonella* culture. Also, microbiological investigation was conducted in all family members of the index case. Active surveillance for salmonellosis among current hospitalized patients was initiated and stool specimens were collected from all symptomatic and asymptomatic newborns who shared the same rooms with the cases.

Microbiological study and antimicrobial susceptibility

Stool samples were plated on selective and differential media to study *Salmonella* (selenite enrichment broth, MacConkey agar, *Salmonella*-*Shigella* agar and xylose-lysine-desoxycholate agar). Cultures media were incubated at 37 °C in aerobic atmosphere. To bacterial identification we used the identification card and software AMS-GNI (Vitek, BioMerieux, Francia). *Salmonella* isolates were serotyped, based on the antigenic properties of their O (somatic), H (flagellar) and Vi (capsular) antigens, using agglutination test with specific antiserum in accordance with the existing Kauffmann-White serotyping scheme (WHO Collaborating Centre for Reference and Research on *Salmonella*, which is located at the Pasteur Institute in Paris, France).⁸

Antibiotic susceptibility tests were performed by disk diffusion method on Mueller-Hinton agar and Rosco disks (Neo-Sensitabs.

Rosco Diagnostica Taastrup, Denmark) following the interpretation of results by Clinical and Laboratory Standards Institute (CLSI) recommendations for *Enterobacteriaceae*. The antibiotics tested for the susceptibility study were: ampicillin, cefalotin, cefuroxime sodium, cefotaxime, ceftazidime, cefepime, ceftazidime, amoxicillin/clavulanate, piperacillin/tazobactam, aztreonam, gentamicin, tobramycin, amikacin, imipenem, nalidixic acid, ciprofloxacin, nitrofurantoin, fosfomicin and trimethoprim/sulfamethoxazole.

In order to establish the clonal relationships of the identified isolates of *Salmonella* the pulsed-field gel electrophoresis of restriction fragments (restriction enzyme: Xba1) of the chromosomal DNA were determined. Band patterns were visually interpreted following the criteria of Tenover et al.⁹

Infection control measures

All of the cases were under contact precautions and housed in nursing cohorts until discharge or until a negative stool sample was obtained. Hand hygiene was reinforced and more alcohol-based hand sanitizers were conveniently placed on all inpatient rooms. Health-care workers (HCWs) with positive culture were placed on sick leave until they had a negative stool sample.

Results

When the index case was reported, there were 20 newborns hospitalized in the low-risk care unit. Besides the index case, 40 people underwent microbiological testing: 11 neonates (2 symptomatic and 9 asymptomatic newborns hospitalized in rooms 4, 5 and 6), 27 HCWs and 2 parents. In total, six more cases were positive to *S. enterica* serotype Newport: 4 newborns (2 symptomatic and 2 asymptomatic cases), 1 parent and 1 nurse, the demographic and clinical characteristics of the cases are described in Table 1. The first secondary case was a premature 3-month infant with onset of gastroenteritis on January 31st (the day after the index case) and who had been hospitalized in room 5. She was discharged the 28th but remained under home visits by an ambulatory care nurse, who collected the stool sample that tested positive for *S. newport*. The next case was a 7-day full term newborn inpatient who was hospitalized also in room 5. The next two cases were asymptomatic infants who shared the same hospitalization rooms with the index case (room 4) or with the first secondary case (room 5). Only one case was detected among HCWs studied; it was an asymptomatic nurse who had attended three of the infant cases. All the infected newborns shared in some point of the outbreak period the rooms 4 or 5, they had common HCWs attending them but no medical or dietary risk factors were identified. The outbreak source was the index case's mother, a 30 year-old woman with Crohn's disease, without gastrointestinal symptoms at that moment but with a stool culture positive for *S. newport*. The seven isolated strains had the same antibiotic profile of resistance. All strains tested were sensitive to nalidixic acid, amoxicillin/clavulanate, ampicillin, aztreonam, cefepime, cefotaxime, ceftazidime, cefuroxime sodium, ciprofloxacin, trimethoprim/sulfamethoxazole, fosfomicin, imipenem, nitrofurantoin, piperacillin/tazobactam, and showed intermediate susceptibility to amikacin, cefalotin and gentamicin. These results and also PFGE showed it was the same strain (Fig. 1). All infected patients recovered completely without complications.

Discussion

This outbreak is the first reported in a neonatal facility by this serotype of *Salmonella* in Spain. The epidemiological investigation suggests as the origin of this outbreak, the index case's mother.

Table 1
Demographic and clinical characteristics of *Salmonella newport* cases.

| Case | Condition | Sex | Gestational Age at birth | Birth weight | Feeding | Clinical symptoms | Antibiotic treatment |
|------------|------------------------------|--------|--------------------------|--------------|-------------------------|----------------------------|----------------------|
| Index case | Prematurity | Male | 34 2/7 weeks* | 2320 g | Formula | Bloody diarrhea and sepsis | Cefotaxime |
| 1 | Prematurity | Female | 26 3/7 weeks | 840 g | Formula | Mild bloody diarrhea | None |
| 2 | Full-term with cervical mass | Male | 40 3/7 weeks | 3390 g | Breast milk | Mild diarrhea | None |
| 3 | Prematurity | Male | 33 1/7 weeks | 1580 g | Formula and Breast milk | Asymptomatic | None |
| 4 | Prematurity | Male | 34 2/7 weeks | 2190 g | Formula and Breast milk | Asymptomatic | None |
| 5 | Ambulatory nurse | Female | – | – | – | Asymptomatic | None |
| 6 | Index case's mother | Female | – | – | – | Asymptomatic | None |

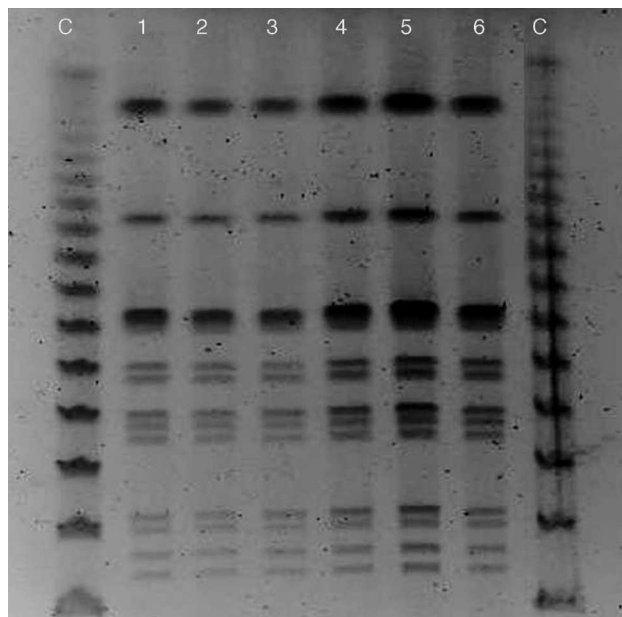


Fig. 1. Pulsed field gel electrophoresis (PFGE) of restriction fragments from 6 cases of the outbreak (except the nurse). The pulsotype profiles are almost identical and they can be attributed to the same strain of *Salmonella enterica* serotype Newport.

She was an asymptomatic carrier of *S. newport* and the bacteria was spread from person to person in the low-risk care unit. At the time of the index case detection, the occupancy in this unit was 87%, probably contributing to a disruption in hand hygiene compliance by HCWs. In France, Cartolano et al.¹⁰ have reported one case of *Salmonella brandenburg* from a premature newborn in a neonatal intensive care unit. This patient developed a septic shock syndrome on the 12th day of life and *S. brandenburg* was isolated in his gastric aspirate and a stool sample; according to the outbreak investigation the pathogen was almost certainly transmitted by contaminated hands because the mother had a positive stool specimen but her milk was free of *Salmonella*. In our outbreak, the route of transmission was probably the same, as the newborn was fed with artificial formula. Besides the four secondary cases occurred by horizontal transmission among the unit, the affected nurse probably was infected during domiciliary care of one of the infected newborns. As far as the clinical course is concerned, it was noticed that the

newborns being breastfed (mixed or exclusive) were asymptomatic in two cases or had mild diarrhea in one case. In contrast, the index case and one secondary case that had bloody diarrhea were exclusively fed with formula. These incidental findings could suggest a possible protective role of breastfeeding for severity of *Salmonella* infection already mentioned in other studies.¹¹ The rapidity of the diagnosis, the prompt application of control measures and reinforcement of hand hygiene helped to avoid transmission to more children and HCWs within our neonatal unit. Thus, it is mandatory to improve infection control measures with HCWs but also with the parents of hospitalized infants, specially reinforcing hand-washing, as they can be an important source of infection in this setting.

Conflict of interest statement

The authors declare that they have no conflicts of interest.

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