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Fatal pertussis in infancy, Italy



KEYWORDS

Pertussis;
Infants;
Pulmonary hypertension

Dear Editor,

Outbreaks of pertussis have recently occurred in industrialized countries with high vaccination rates such as the US, Australia and many European countries, involving particularly adolescents, adults and young infants.¹ The reasons for “resurgence” of pertussis include waning of immunity from vaccination or natural infection over time, low coverage by acellular vaccine, genetic changes in *Bordetella pertussis* strains, greater awareness of the disease, and improved diagnostic ability through the spread use of new molecular techniques.² Due to waning of mother’s immunity over time, transplacental transfer of pertussis antibodies is not protective, and infants too young to have completed their vaccination schedules are particularly vulnerable. They have the highest rate of morbidity, hospitalization, complications, intensive care unit (ICU) admission, and mortality.³

The incidence of pertussis deaths among infants in developed countries has increased, with very high mortality rates in those less than 3 months of age.³ Surprisingly, a national surveillance study up to 2012 showed that deaths from pertussis have not been reported in Italy since 2002.⁴ In addition, no death occurred in Tuscany, Italy, over the period 2013–2014, despite a dramatic increase in hospitalization rate (up to 95.7 per 100,000) in infants.⁵

A recent article in this Journal has highlighted the emerging problems associated with *B. pertussis* infection in infants and strategies to ameliorate this.¹ We report a retrospective chart review of six fatal cases of pertussis in infants recently admitted to three Italian tertiary care pediatric centers. A death attributed to pertussis was confirmed if: i) *B. pertussis* was isolated by culture or

Table 1 Characteristics of the patients at admission.

Pts	Gestational age (weeks + days)	Age (days)	Underlying conditions	Pertussis immunization	Duration of symptoms at presentation	Referring modality	Clinical features	Ongoing antibiotics
CM	26 + 5	89	BPD	No	3	Pediatrician	Rhinitis, dry cough, wheezing	CLR
PN	41 + 1	51	No	No	3	Parents	Dry cough, tachypnoea	No
CA	40 + 2	95	No	1st dose	5	Pediatrician	Dry cough, tachypnoea, poor feeding	CLR
SA	40 + 3	23	No	No	7	General hospital	Fever, dry cough, tachypnoea	No
ZV	38 + 1	24	No	No	4	Parents	Dry cough, tachypnoea, poor feeding	No
AM	36 + 0	23	No	No	2	Pediatrician	Rhinitis, dry cough, wheezing	No

BPD, Bronchopulmonary dysplasia; CLR, clarithromycin.

Table 2 Laboratory data, infective source, radiological findings, intervention in ICU, and outcome.

Pts	WBC $\times 10^3$ /mL (percentage lymphocytes/neutrophils) at admission	WBC $\times 10^3$ /mL (percentage lymphocytes/neutrophils) at peak	B. pertussis identified by RT-PCR	Infective source	Chest X-Ray pneumonia	Pulmonary artery pressure (mmHg)	Additional intervention in ICU	Time to death from hospital admission (days)	Time to death from ICU admission (days)
CM	17,4 (31/54)	109,1 (46/41)	LTA	Unknown	Bilateral	38	iNO	6	2
PN	31,2 (72/17)	104,1 (43/49)	NPA	Unknown	Bilateral	32		5	1
CA	71,5 (34/59)	71,5 (34/59)	NPA, BAL	Unknown	Bilateral	35	RBC, PLS	6	1
SA	83,3 (21/70)	83,3 (21/70)	NPA,BLD	Brother	Bilateral	Normal	iNO, RBC, PLS	4	1
ZV	34,1 (61/33)	65,5 (38/55)	NPA, BLD	Mother	Bilateral	39	iNO, RBC, PLS, PLT	2	1
AM	15,2 (62/24)	78,7 (26/62)	NPA	Unknown	Bilateral	60	iNO	4	2

WBC, white blood cells; RT-PCR, real-time polymerase chain reaction; LTA, lung tissue autopsy; NPA, nasopharyngeal aspirate; BAL, bronchoalveolar lavage; BLD, blood; iNO, inhaled Nitric Oxide; ECMO, Extra Corporeal Membrane Oxygenation; RBC, Red Blood Cells; PLS, Plasma; PLT, Platelets.

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