

CLINICAL CASE REPORT

Chest pain and dyspnoea in a young primiparous woman following prolonged labour at a remote Australian community health centre

AUTHORS

Mohd Azlan Mohd Hashim¹ MBBS, Registrar

Timothy Linton² FRAGP, General Practitioner

Subash Heraganahally³ FRACP, Director, Respiratory and Sleep Medicine *, hssubhashcmc@hotmail.com

CORRESPONDENCE

*Dr Subash Heraganahally hssubhashcmc@hotmail.com

AFFILIATIONS

^{1, 3} Royal Darwin Hospital, 105 Rocklands Drive, Tiwi, NT, Australia

² Gove District Hospital, Gove, NT, Australia

PUBLISHED 18 May 2019 Volume 19 Issue 2

HISTORY RECEIVED: 15 October 2017

REVISED: 4 April 2019

ACCEPTED: 4 April 2019

CITATION

Mohd Hashim M, Linton T, Heraganahally S. Chest pain and dyspnoea in a young primiparous woman following prolonged labour at a remote Australian community health centre. Rural and Remote Health 2019; 19: 4615. https://doi.org/10.22605/RRH4615

Except where otherwise noted, this work is licensed under a Creative Commons Attribution 4.0 International Licence

ABSTRACT:

Reported here is a rare respiratory complication in a young primiparous woman in a remote community hospital in the Northern Territory, Australia. A Caucasian primiparous woman aged 23 years presented with chest pain following a very prolonged second stage of labour. On clinical examination a crunching and crackling sound was heard over the precordium, synchronous with the heartbeat (Hamman's sign). A chest radiograph confirmed the presence of pneumomediastinum. Complete resolution of symptoms and signs was achieved with supportive management.

Keywords:

Australia, chest pain, Hamman's sign, pneumomediastinum, primiparous.

FULL ARTICLE:

Introduction

Practising in remote community health centres can be challenging for healthcare professionals in dealing with medical emergencies, especially in areas where the resources are sparse and accessibility is limited. However, this provides an excellent opportunity to manage complex, unusual medical conditions first hand. Here we report a rare respiratory complication in a young primiparous woman following a very prolonged labour in a remote community hospital in the Northern Territory, Australia.

Case

A 23-year-old Caucasian primiparous woman who was at 40 weeks and 6 days of gestation presented to a remote community health center situated approximately 1000 km from the nearest tertiary care referral hospital for delivery of her first child after spontaneous rupture of membrane. She had an uneventful antenatal history and had no significant past medical or surgical comorbidities and was not on any regular medications. She had a BMI of 25 and had a few years of smoking history (about 10 cigarettes per day).

She was noted to have a very prolonged second stage of labour, lasting for about 3–4 hours. During that time she had been pushing with extreme efforts intermittently for prolonged periods. She was also noticed to have had multiple bouts of vomiting, presumably thought to be secondary to inhalation of medical nitrous oxide and oxygen. A healthy baby boy with a weight of 3.3 kg was delivered via normal vaginal delivery.

Six hours postpartum she complained of chest pain associated with mild dyspnoea and altered voice. Her vital signs were unremarkable and stable. She was examined by the on-call GP/obstetric physician who noted subcutaneous emphysema of the neck and chest on palpation. Her chest was clear with normal breath sounds on both lung fields. The physician also recognised crunching and crackling sound heard over the precordium, synchronous with the heartbeat (Hamman's sign). A chest radiograph revealed evidence of pneumomediastinum (Fig1). After consulting with the respiratory team at the tertiary referral hospital, she was managed with analgesia, supplemental oxygen and close observation at the community hospital. A repeat chest X-ray 24 hours later showed resolution of the pneumomediastinum, and the patient's symptoms improved. The subcutaneous emphysema resolved spontaneously after 3 days and she was discharged home. She was followed up by the respiratory team 2 months later with no further complications.

Discussion

Diagnosis of pneumomediastinum can be challenging, especially in remote community hospital settings, given that its most common symptom (chest pain) carries broad differentials¹. Clinical skills (physical examination) and decision making on management are extremely important where resources are limited. Awareness of this condition is crucial to differentiate between other conditions with higher morbidity and mortality such as pneumothorax, aortic dissection, oesophageal rupture, pulmonary or amniotic fluid embolism and myocardial infarction. The first report showing the occurrence of pneumomediastinum during labour was published by Simmons and later by Louis Hamman². Since then about 200 cases have been reported in the literature with an estimated incidence of 1 in 100 000 deliveries²⁻⁴. Although the clinical presentation can be dramatic, it is a benign condition with a self-limiting course.

The pathophysiology is thought to be secondary to a significant increase in intrapulmonary pressures, leading to disruption of alveolar walls and entry of air into the pulmonary interstitium. The negative pressure gradient of the mediastinum in relation to the pulmonary parenchyma will naturally cause air to flow into the mediastinum and vents out to the thorax and subcutaneous tissues, thus explaining the clinical signs associated with this condition⁵. In severe cases, tension pneumomediastinum can occur, which can prove fatal. Fortunately, this is a very rare event.

Management of this condition, unless complicated by tension pneumomediastinum or pneumothorax, is supportive as most cases resolve with conservative management alone. Supplemental oxygen may hasten reabsorption²⁻⁴. If detected during labour, steps to avoid repeated maternal expulsive efforts by using maternal forceps or converting to caesarean section under regional anaesthesia may be undertaken. A nitrous oxide and oxygen mix should be avoided as it may lead to expansion of trapped gas². Recurrence of pneumomediastinum in future pregnancies is not known to be common.



Figure 1: Chest X-ray of the patient, showing pneumomediastinum.

REFERENCES:

1 Gerazounis M, Athanassiadi K, Kalantzi N, Moustardas M. Spontaneous pneumomediastinum: a rare benign entity. *Journal of Thoracic and Cardiovascular Surgery* 2003; **126:** 774-776. https://doi.org/10.1016/S0022-5223(03)00124-7

2 Kandiah SI, Iswariah H, Elgey S. Postpartum pneumomediastinum and subcutaneous emphysema: two case reports. *Case Reports in Obstetrics and Gynecology* 2013; **2013:** 735154. https://doi.org /10.1155/2013/735154 PMid:23533868

3 Seidl JJ, Brotzman GL. Pneumomediastinum and subcutaneous

emphysema following vaginal delivery: case report and review of the literature. *Journal of Family Practice* 1994; **39:** 178-180.

4 Heffner JE, Sahn SA. Pleural disease in pregnancy. *Clinics in Chest Medicine* 1992; **13:** 667-678.

5 Maunder RJ, Pierson DJ, Hudson LD. Subcutaneous and mediastinal emphysema. Pathophysiology, diagnosis, and management. *Archives of Internal Medicine* 1984; **144:** 1447-1453. https://doi.org/10.1001/archinte.1984.00350190143024 PMid:6375617

This PDF has been produced for your convenience. Always refer to the live site https://www.rrh.org.au/journal/article/4615 for the Version of Record.