Pulmonary calcification following smallpox handler's lung

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Ross, P. J., Seaton, A., Foreman, H. M., and Morris Evans, W. H. (1974). Thorax, 29, 659–665. Pulmonary calcification following smallpox handler’s lung. A follow-up study of six nurses who developed smallpox handler’s lung during the 1962 South Wales smallpox epidemic and of the surviving smallpox patients is reported. The development of extensive punctate calcification in the chest radiographs of five of the nurses, together with the absence of abnormalities of lung function, is taken to support the view that the disease is a modified smallpox pneumonia rather than an allergic alveolitis.

A pneumonic illness occurring in vaccinated close contacts of smallpox patients was reported by Chadwick (1843) and has been described more recently by Howat and Arnott (1944) and Ratner and Khudyakova (1962).

Morris Evans and Foreman (1963) described several cases of the 'illness of contact' in nursing volunteers during the South Wales smallpox outbreak of 1962. They postulated that this was a pulmonary hypersensitivity reaction comparable to farmer's lung due to the inhalation of variola virus or scale dust from patients.

We report a follow-up of six of these nurses with smallpox handler's lung and of 10 surviving smallpox patients who did not develop pulmonary symptoms.

SUBJECTS

Thirty-two volunteer nurses had been involved in the care of 54 patients with smallpox in three isolation hospitals in South Wales in 1962. Of these nurses, 12 had developed pulmonary symptoms, five showing radiographic changes. This report records details of the follow-up of six of these nurses. Their ages in 1962 ranged from 38 to 50 years.

In addition, 12 patients with mild smallpox who had chest radiographs towards the end of their illness were recalled for repeat radiographs as part of the study. Two further surviving patients, who had not had chest radiographs in 1962, were also recalled.

METHODS

Standard six-foot postero-anterior chest radiographs were taken during the initial illness and at intervals afterwards. Radiographs taken before the illness were also available for all the nurses.

Radiographs were read in random order by two independent observers and classified according to the presence and type of peripheral opacities. Particular note was made of the distribution and size of the lesions and of the presence or absence of calcification.

Lung function tests were carried out in four nurses in 1962, in one in 1968, and in four in 1974. These tests included spirometry, using a lightweight waterless spirometer, lung volumes by the closed-circuit helium dilution technique, and carbon monoxide transfer factor by the single-breath technique. Predicted normal values were those of Cotes (1968).

Immunological testing for precipitating antibody to vaccinia virus, budgerigar, pigeon and canary serum and droppings, Microsporospora faeni, Thermactinomyces vulgaris, 'Air-conditioner pneumonia' organism, Aureobasidium pullulans, Candida albicans, house dust, and feathers was carried out by the Ouchterlony double diffusion technique.

CLINICAL FEATURES AND RESULTS

ORIGINAL ILLNESS IN THE NURSES  Typically, this began with progressive malaise nine to 12 days after the first exposure to a smallpox patient. This was followed four to six hours later by frontal headache, shivering, sweating, and generalized aching, especially in the lumbar area. The symptom-complex was described as 'influenza-like' but without the catarrhal features.

Some developed a dry irritating cough within three to four days with physical signs of prolonged expiration and scanty crepitations. Three
developed skin rashes, two resembling erythema multiforme and one with a diffuse erythematous flush. The febrile state lasted from three to 10 days and was intermittent for two to three days.

Of those with symptoms, five showed definite radiological lung abnormalities, three with marked changes. These three complained of slight exertional dyspnoea for a further six to eight weeks.

**SYMPTOMS ON FOLLOW-UP (1974)** There was no history suggesting a past episode of varicella pneumonia in any of the nurses interviewed. One (No. 4) had been treated for pulmonary tuberculosis in October 1962, approximately five months after the episode of smallpox handler’s lung. She reported mild dyspnoea on effort together with intermittent cough productive of white sputum in the last four or five years. She smoked 20 cigarettes daily. Two others (Nos. 1 and 2) reported recurrent winter cough often productive of purulent sputum. One of these had smoked for many years. Apart from No. 4 none reported any dyspnoea or symptoms of significant airways obstruction.

**RADIOGRAPHY** The two readers’ reports of the presence and type of parenchymal shadowing were identical in all films.

The chest radiographs of five nurses taken before the acute illness of 1962 showed no significant abnormality. The sixth showed evidence of calcified tuberculosis in both upper lobes and left hilar glands.

Radiographs taken during the acute illness of 1962 in five nurses showed nodular opacities. These varied in size from 2–3 mm (Nos. 2 and 3) to 5–10 mm (Nos. 4, 5, and 6). They were of low density, poorly defined, and distributed mainly in the upper and middle zones (Figs 1 and 2).

The radiograph of nurse No. 1 showed no significant abnormality, either during the illness or on follow-up. Serial chest radiographs in the other five nurses showed the subsequent appearance of diffuse nodular opacities 1–3 mm in size. These were of high density (calcified), sharply defined, and evenly distributed throughout both lung fields (Figs 3, 4, and 5).

The time interval before calcification appeared varied between four and seven years (Table 1). No relationship was noted between the size of the original ‘acute’ nodules and the size or time of appearance of the calcified nodules.

Follow-up chest radiographs of eight surviving smallpox patients taken in 1974 showed no change from those taken at the time of discharge from hospital in 1962. No calcification occurred in these radiographs or in those of the two other surviving patients not previously radiographed.

**LUNG FUNCTION** The results of lung function tests carried out are shown in Table II. There were no significant abnormalities found in 1962. Three of the four tested in 1974 were smokers and showed evidence of mild airways obstruction. A decreased transfer factor was noted in nurse No. 4 who smoked 20 cigarettes daily and had suffered from pulmonary tuberculosis in the past.

**IMMUNOLOGY** No precipitating antibodies to vaccinia virus, *Micropolyspora faeni*, *Thermoactinomyces vulgaris*, *Aurobasidium pullulans*, *Candida albicans*, house dust or feathers were found in 1962 or 1974. In addition, no precipitating antibodies to budgerigar, pigeon, and canary serum and droppings or ‘air-conditioner pneumonitis’ organism were detected in 1974.

**DISCUSSION**

Howat and Arnott (1944) reported an acute benign pneumonic illness in seven military hospital staff who had been in contact with smallpox patients in the Middle East. Symptoms of increasing malaise, headache, general aching, chills, and sweating developed 11 to 14 days after first contact with a patient. Cough was infrequent and unproductive, and appeared only after several days. Rash was absent. The febrile state lasted four to 12 days. Some crepitations were noted on auscultation. Radiological changes of fine diffuse mottling, usually in the lower lobes, and rounded opacities in the middle and lower zones were seen in six cases from the third day, and in two these persisted for six weeks. These authors considered the illness to be due to either modified variola or a virus of unknown origin.

Ratner and Khudyakova (1962) described an episode of fever and chest pain lasting up to four days in three vaccinated contacts of smallpox patients. This illness developed nine to 15 days after first contact. Chest radiographs showed single or several opacities which persisted up to eight months. No skin rashes were noted. They proposed the term ‘pulmonary form’ of smallpox to describe the illness.

In the cases we have studied, it is considered very unlikely that the illness was due to coincident virus, fungal or bacterial infection. This is indicated by the negative viral agglutination tests and
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examination of sputum carried out during the acute illness by Morris Evans and Foreman (1963). The affected nurses were working in separate isolation hospitals and there was no clinical history to suggest varicella pneumonia, Histoplasma infection, or miliary tuberculosis.

Morris Evans and Foreman (1963) commented at that time that actual invasion of the lung by the variola virus had never been proved and there was then no epidemiological evidence that this type of lung lesion was infectious. They therefore suggested the changes might have been due to a hypersensitivity reaction in the lungs of the nurses. The nature of the antigen was not certain but was thought to be the variola virus inhaled in droplets or scale dust from the patients during activities such as routine bed making.

The 'illness of contact' they and others described had previously been considered to be a highly modified smallpox infection, and in this case

TABLE I
SERIAL RADIOLOGY OF SIX NURSES WITH SMALLPOX HANDLER'S LUNG

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C—multiple calcified nodules seen on chest radiograph; NC—no calcification seen on chest radiograph; EC—early evidence of calcification in some lesions.
FIG. 3. Radiograph taken during the acute illness in nurse No. 4, showing nodular opacities in upper zones and calcified lesions of healed tuberculosis.

respect the erythematous rashes observed in three of our cases are interesting. They are seen as a prodromal manifestation of about 10% of all smallpox cases and are said to occur almost exclusively in the partially immune patient whose illness tends to run a benign course (Ricketts and Byles, 1908). There are several features of the acute illness which are unlike an allergic reaction such as farmer's lung. Dyspnoea was not a marked feature. There was no increase in serum gamma-globulin, and detailed studies of lung function performed while the lung lesions persisted were normal, in contrast to those seen in an allergic reaction such as farmer's lung where a restrictive defect with reduced transfer factor is commonly found (Hapke et al., 1968).

The initial radiographic lesions in our nurses appeared to be of two types. One, occurring in three nurses, was of a larger, blotchy, 'alveolar filling' lesion, while the other was of a smaller, interstitial type of nodularity. It is possible, however, that these represented two stages of the same process as both evolved in an identical manner over succeeding years. The most interesting feature on serial radiology was the appearance, after an interval of five or six years, of widespread punctate calcification. We know of no reference to calcification as a late feature of allergic alveolitis, but it is a well-known complication in survivors of varicella pneumonia (Mackay and Cairney, 1960; Abrahams, Evans, Knyvett, and Stringer, 1964). Although the viruses of varicella and variola are structurally different, the tissue reactions are known to be closely similar (Cappell and Anderson, 1971). This calcification is strongly suggestive of an infection by the inhaled variola virus.

The failure to grow variola virus from the sputum, nasopharyngeal washings, or skin scrapings of the rash from three of the nurses in 1962 may well have been due to the scarcity of the virus in the specimens and its modification by the defence mechanisms of the highly vaccinated host. Moreover, it is curious that none of the patients who survived the disease developed similar lung lesions. The differences between the nurses and the
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**FIG. 4.** Radiograph of the same subject as in Fig. 3 taken in 1974, showing the presence of miliary calcification in addition to that due to old tuberculosis.

**FIG. 5.** Chest radiograph of nurse No. 2 showing miliary calcification. Almost identical lesions were present in the other three nurses at follow-up.
TABLE II

RESULTS OF LUNG FUNCTION TESTS IN SIX NURSES WITH SMALLPOX HANDLER’S LUNG

<table>
<thead>
<tr>
<th>Nurse</th>
<th>Year</th>
<th>Peak Flow Rate (l/min)</th>
<th>Forced Expiratory Volume in one second (litres)</th>
<th>Vital Capacity (litres)</th>
<th>FEV FVC %</th>
<th>Residual Volume (litres)</th>
<th>Functional Residual Capacity (litres)</th>
<th>Total Lung Capacity (litres)</th>
<th>RV TLC %</th>
<th>Transfer Factor for Carbon Monoxide (ml/min/mmHg)</th>
</tr>
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<tr>
<td>1</td>
<td>1974</td>
<td>485 (121)</td>
<td>3.00 (122)</td>
<td>3.50 (117)</td>
<td>86</td>
<td>1.10 (59)</td>
<td>1.80 (78)</td>
<td>4.60 (91)</td>
<td>24 (91)</td>
<td>23 (92)</td>
</tr>
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<td>2</td>
<td>1974</td>
<td>240 (75)</td>
<td>1.30 (72)</td>
<td>2.20 (105)</td>
<td>59</td>
<td>1.05 (78)</td>
<td>1.45 (91)</td>
<td>3.25 (90)</td>
<td>32 (87)</td>
<td>22 (116)</td>
</tr>
<tr>
<td>3</td>
<td>1974</td>
<td>320 (93)</td>
<td>2.40 (120)</td>
<td>3.45 (144)</td>
<td>69</td>
<td>1.65 (110)</td>
<td>3.05 (142)</td>
<td>5.10 (124)</td>
<td>32 (88)</td>
<td>20 (95)</td>
</tr>
<tr>
<td>4</td>
<td>1968</td>
<td>2.08 (93)</td>
<td>2.04 (139)</td>
<td>3.21 (122)</td>
<td>62</td>
<td>2.00 (138)</td>
<td>2.83 (111)</td>
<td>5.19 (125)</td>
<td>39 (118)</td>
<td>25 (115)</td>
</tr>
<tr>
<td>5</td>
<td>1974</td>
<td>305 (87)</td>
<td>2.10 (102)</td>
<td>3.40 (139)</td>
<td>62</td>
<td>2.05 (132)</td>
<td>2.67 (111)</td>
<td>5.25 (122)</td>
<td>39 (115)</td>
<td>23 (110)</td>
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<tr>
<td>6</td>
<td>1962</td>
<td>1.9</td>
<td>2.68</td>
<td>1.73</td>
<td>71</td>
<td>1.72</td>
<td>2.66</td>
<td>4.18</td>
<td>41</td>
<td>17 (77)</td>
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Figures in parentheses indicate percentage of predicted normal value.
In Nurses 5 and 6 it was not possible to find a record of their height or weight at the time of testing.
patients must be explained by differences in the numbers of virus particles, the nurses presumably inhaling many more than any individual patient, and their different immunization status prior to exposure. It seems likely that smallpox handler's lung, a disease showing closely similar features in the three published reports, is a modified pulmonary infection with variola virus in highly immune subjects.

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REFERENCES


Requests for reprints to: Dr. A. Seaton, Sully Hospital, Sully, Penarth, Glamorgan CF6 2YA.
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