MASSIVE PULMONARY EMBOLISM IN A BOY AGED 9

BY

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Massive pulmonary embolism is not normally a post-operative complication to be feared in a boy of 9 and no description of another case in childhood has been found in the literature: it must therefore be a rare event.

Case History

The patient, a boy aged 9, was admitted to the children's ward at the Royal Berkshire Hospital in March, 1956, as a case of suspected acute appendicitis. Three days earlier he had complained of feeling sick and giddy and also had some central abdominal pain. He vomited several times that day and during the next two days and the pain persisted. The day before admission he passed several loose stools and the night before admission had been feverish. He had had a head cold with a little cough 10 days earlier which had not cleared up completely.

On admission he was a boy of good physique who was flushed but did not look ill. He was febrile (temperature 102° F., pulse 110), the tongue was furred and the pharynx and tonsils were injected. The chest was clinically clear and heart sounds were normal. There was lower abdominal tenderness, mainly on the right with muscle guarding. Bowel sounds were present. There was some tenderness per rectum to the right. Urine was normal.

The lower abdominal tenderness and guarding increased during three hours' observation and appendicectomy was decided upon. At operation Mr. G. L. Bohn found a high paracolic appendix, the tip reaching...
FIG. 2.—Electrocardiograph four days after embolism. Steep inversion of T.V.1 and T.V.3, with displacement of transition zone to the left.

FIG. 3.—Electrocardiograph 26 days after embolism. Return to normal.
to the liver. It was difficult to remove but was not
inflamed.

No chemotherapy was given and over the next four
days the temperature and upper respiratory tract infec-
tion settled. A chest radiograph was clear. Culture
from stools grew no pathogens.

On the fifth post-operative day he sat up for tea and
on the following day he got up again in the afternoon.
He had been sitting in a chair for a few minutes when he
complained of sudden severe pain in the left side of the
chest and became dyspnoeic, pale and sweating. The
degree of collapse rapidly became very severe.

He was put in an oxygen tent and given 10 minims of
nepenthe and afterwards there was some improvement
but a state of severe shock, with a hardly perceptible
wrist pulse, persisted for about six hours. There was
some blood-tinged sputum.

The first electrocardiogram taken four hours after the
onset of symptoms showed a tachycardia and a prominent
S in lead I (Fig. 1).

As the clinical picture was that of pulmonary embolism
as seen in the adult, he was given 10,000 units of heparin
intravenously eight hourly for three doses. An initial
dose of 100 mg. of ‘dindevan’ was followed by 75 mg.
the next day, then 25 mg. twice daily, the dose thereafter
depending on prothrombin estimation. ‘Dindevan’ was
continued for three weeks, maintaining the prothrombin
efficiency at 35 to 50%. He was given intramuscular
penicillin in addition for the first five days.

He remained extremely ill for 48 hours and then made
a steady recovery. There was never any calf or leg
tenderness and there were no further embolic episodes.
No physical signs appeared in the chest.

An E.C.G. repeated four days after the onset showed
a steep inversion of T.V.1 and T.V.3 with displacement
of the transition zone to the left (Fig. 2) and this was still
present five days later. After a further week the tracing
was within normal limits (Fig. 3).

The E.C.G. appearances of transient acute cor pul-
monale are characteristically those of massive pulmonary
embolism and are thought to be due to sudden clockwise
rotation of the heart.

Since discharge from hospital one month after the
illness the patient has been well and is now back at school
and completely active. A chest film is clear and the
heart shadow normal.

A case of this sort immediately involves a number of
the hospital team and I am grateful to Dr. J. J. Kempton,
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