Introduction

Chylothorax, lymphatic pleural effusion, is a rare but severe complication of cardiac surgery. The symptoms of chylothorax are the same as in any fluidothorax of other etiology, and so are the findings on images (X-ray, CT, ECHO, MRI). However, the biochemical finding is typical. The effusion is a milky cloudy fluid with elevated levels of triacylglycerols (twice to eight times as much as in serum), low level of cholesterol, elevated lymphocyte count and presence of chylomicrons.

Clinical signs may be poor at the beginning, but later, with greater volume of fluid in the pleural cavity, dyspnoea becoming a dominating complaint. With a longer course of chylothorax malnutrition sequelae may develop due to loss of proteins, immunoglobulins, lipids, electrolytes and water (5, 10, 12).

Causes of chylothorax (so-called spontaneous) include inborn developmental anomalies of lymphatic vessels (congenital chylothorax), certain tumors (lymphomas), infections (lymphadenitis, filariasis), specific inflammations (tuberculosis, cirrhosis), liver cirrhosis, thrombosis of the subclavian vein or superior vena cava, or lymphangionymomatosis.

Chylothorax (traumatic chylothorax) may also result from a complication of trauma from a blunt or penetrating injury to the chest, or from vigorous coughing or vomiting. Chylothorax has been described as a complication of surgery of the esophagus, larynx, lungs, mediastinum, heart or great vessels, as a complication of great vessel cannulation or translumbar angiography, or as a sequela of radiotherapy (1, 9, 10, 11).

Chylothorax is a rare but severe complication of cardiac surgery. The authors present the case of a 76-year-old woman suffering from ischemic heart disease, after coronary artery bypass grafting that included a left internal mammary artery pedicle graft. On the ninth postoperative day the left-sided fluidothorax developed. The results of biochemical analysis were consistent with the chyle. Combined treatment with pleural drainage and total parenteral nutrition was effective.

A Case Report

A 76-year-old female patient was admitted to the hospital with ischemic heart disease (triple vessel disease, CCS III, EF 55%). She underwent an on-pump myocardial revascularization using mammo-coronary bypass grafting from the pedicled left mammary artery to left anterior descending artery and two aorto-coronary grafts from the great saphenous vein to the left marginal artery (obtuse branch) and right coronary artery.

The operation and postoperative course were standard without complications. The patient was extubated on the operation day and transfered from the intensive care unit on the second postoperative day. Perioperatively inserted drains were removed on the second postoperative day. There was no fluid drainage noted at that time. The patient...
presented no clinical problems, no dyspnoea and no fever during the next few days.

On routine examination before planned discharge, on the 9th postoperative day, the follow-up chest X-ray showed a severe fluidothorax of the left pleural cavity (Fig. 1). After puncture, 1850 ml of cloudy fluid was evacuated from the pleural cavity. The sample of the evacuated fluid was sent to the department of microbiology. Unfortunately no biochemical analysis was performed at that time.

On the following day, an additional 1500 ml of milky effusion had to be removed by puncture from the pleural cavity. By biochemical analysis chylous fluid was diagnosed (triglyceride level of 14.52 mmol/l) (Tab. 1). Microbiology revealed no growth on the third day. A thoracic drain was inserted in order to relieve the chylothorax, and subsequently a further 1000 ml of chyle were evacuated.

Concurrently total parenteral nutrition (8500 kJ/day) was commenced (Table 2). The clinical condition of the patient remained stable, without subjective complaints. Biochemical laboratory data showed optimal concentrations of Na (137...139), K (4.8...5.4), Cl (101...104), urea (4.1...4.2), krea (76...63), bil (6...5), chol (3.66...4.1), tag (1.83...2.29), prot (56.8...68.4) and higher concentrations of ALT (0.72...0.77), AST (0.53...0.66) during the course of parenteral nutrition without any significant reaction to losses of chyle.

The losses from the thoracic drain gradually diminished (750...250...100...200...0...100...0 ml per day), and after 16 days it was possible to remove the pleural drainage. The total loss of chylous fluid reached 5750 ml. Follow up chest X-ray confirmed no reaccumulation of fluid. Complete diet was introduced gradually.

The patient was discharged home aftercare in a satisfactory condition on the 32nd day. On the follow-up after one year, the patient was cardiopulmonary compensated, without complaints, and without pathologic findings in the pleural cavity.

The cause of this patient’s chylothorax was not clear; it might have been due to damage to the thoracic duct, truncus mammarius or aberrant lymphatic vessels during preparation of the left mammary artery.

Discussion

Chylothorax is a rare complication of cardiac surgery. It usually forms two to five days after surgery when the patients start peroral diet (mainly high-fat diet). Early identification and prompt treatment may decrease both the early and the late morbidity and mortality. It was a bad mistake that no biochemical analysis was performed at the time of the first evacuation of pleural effusion in our patient.

The therapy of chylothorax is always combined – conservative and invasive.

The basis of conservative part of treatment of chylothorax is administration of a low-fat diet or, even better, total parenteral nutrition. The main goals are to minimize chyle formation and to prevent dehydratation, malnutrition, protein and electrolyte deficiencies, or immunosuppression (9, 10).

Excellent results are described after administration of octreotide, a synthetic long-acting somatostatin analogue which increases splanchnic arteriolar resistance and decreases the lymphatic flow. Octreotid administration may be used as the initial conservative therapy for moderate chylous drainage (2, 8).

Invasive therapy, an integral part of the treatment, is always based on evacuation of the pathological pleural fluid by means of puncture or by permanent chest drainage.

More aggressive surgical treatment is considered when the output remains high (>1000-1500 ml per day) despite the conservative management for a period more than 5 days, or if drainage or lung re-expansion is incomplete.
intervention is indicated by an assessment of each individual patient’s condition. In many papers there is advocated the conservative management for maximum of 2 weeks period, where the indication for surgery is absent (9, 10). However, surgery for chylothorax can lead to a longer hospital stay, extra morbidity failure of the procedure, and high costs.

Lymphatic structures at the site of the defect may be treated by surgery (after thoracotomy or by means of video-assisted thoracotomy) by closure of the thoracic duct above the diaphragm (clipping, ligation, application of tissue glue). Thoracoscopic surgery may be the first choice (8, 13).

If this approach is not effective defects of lymphatic ways can be closed by pleurectomy with pleurodesis and application of anti-inflammatory agents (bontalec, antibiotics) (3,12). A pleuroperitoneal shunt could be indicated in high risk patients, with excessive chylos leak (2 week’s duration, with output greater than 1000 mL/day) when preceding treatment was not successful (10).

Outstanding results have been described after application of percutaneous transabdominal embolization and blockage of lymphatic vessels (6). An effective result may be achieved after targeted radiotherapy (2). Others critically assess an importance of percutaneous strategy for diagnosing and treating chylothorax (3). These procedures should be attempted especially if patients are very ill, before more risk surgical thoracic duct closure is considered.

The applied combined therapeutic method – the evacuation of the fluid by permanent thoracic drainage and total parenteral nutrition was chosen in our patient. This therapeutic scheme resulted in decrease in chyle production, without any complications. We have chosen, to keep safe, this less invasive approach due to age of the patient, concomitant diseases and health status after the cardiac operation (less than 2 weeks). The patient is doing well without the recurrence 1 year after the procedure.

Though we already published an experience with 3 cases of chylothorax in the year 1988 (7), as we had met for the first time this complication after myocardial revascularization, we thought appropriate to dare to add even this rare case of chylothorax.

We recommend a combined treatment of chylothorax – a drainage of pleural cavity and total parenteral nutrition for maximum of 2 weeks period. After it a surgical treatment is considered. Surgery or other intervention are indicated when the output remains high (> 1500 ml per day) despite the conservative management for a period more than 5–7 days.

**Conclusion**

Chylothorax is a very rare complication of cardiac surgery. The real cause of its development in our patient after aorto-coronary bypass was not found. Presumably it was an injury to the lymphatic vessels (thoracic duct, truncus mammarius, aberrant pathways) at preparation of the mammary artery. The applied therapeutic method – the evacuation of the fluid by thoracic drainage and total parenteral nutrition was successful.

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**References**