Systemic generalised lymphangiomatosis: unknown aetiology and a challenge to treat

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SUMMARY
We describe a case of a woman diagnosed at the age of 35 years with a generalised mediastinal and abdominal lymphangiomatosis associated with a protein losing enteropathy, who successfully improved when treatment with sirolimus was initiated.

BACKGROUND
Systemic generalised lymphangiomatosis is a very rare condition commonly appearing during childhood, characterised by proliferation of normal mature lymphatic ducts with formation of numerous cysts in varied body organs. These cysts can occur in any organ except for the brain that lacks lymphatics. Its aetiology remains unknown, but lymphangiomas are considered to be uncommon benign tumours. Different putative mechanisms of lymphangioma development are proposed including failure of the lymphatic system to connect with or separate from the venous system or abnormal budding of the lymphatic system from the cardinal vein. In adults, sequestration of lymphatic tissue secondary to inflammatory processes or surgical therapy has been incriminated. Recent studies suggest the potential role of specific lymphangiogenic growth factors. Increased interleukin-6, vascular endothelial growth factor c (VEGF-c), vascular endothelial growth factor receptor 3 (VEGFR-3) and Prospero Homeobox 1 (prox-1) have been correlated with the genesis of lymphangiomas. These new insights create new opportunities for treatment, including the use of inhibitors of angiogenesis.

Although lymphangiomas are benign lesions, they may cause significant mortality because of their large sizes, critical locations and the possibility of secondary infections. The lymphangiomatosis spectrum is very heterogeneous in its presentation, and can include microcystic and macrocystic isolated lymphatic malformations, thoracic and intra-abdominal diffuse lymphangiomatosis, and osseous and soft-tissue presentations known as Gorham-Stout disease.1

Lung involvement is paramount and the leading cause of death.2 Abdominal lymphangiomatosis is rare in adult patients. The most frequent symptoms are abdominal pain. Some patients with limited lymphangiomas are asymptomatic. Few case reports describe adult patients with lymphangiomatosis, gastrointestinal bleeding and hypoproteinemia, that may be life-threatening. Intestinal lymphangiectasia is one of the reasons of protein-losing enteropathy.2–5

Since the condition is rare there is no uniform treatment strategy for symptomatic patients and evidence-based therapies are lacking. Surgery has been attempted for localised and generalised gastrointestinal disease, but recurrence rates are high.6 Various other treatment modalities have been proposed, including radiotherapy, chemotherapy, interferon-alfa and systemic corticosteroids with varied success in small cohorts.7–10 A limited number of case reports suggests a potential benefit of angiogenesis inhibitors, such as VEGF inhibitors, mammalian target of rapamycin (mTOR) inhibitors and tyrosine kinase inhibitors.11–15

These treatments were successful at a young age of diagnosis but their efficacy in adults remains to be established.

Recent published case reports and a small patient cohort show an adult phenotype resembling lymphangiomatosis, but who are associated with phosphatidylinositol 3 kinase catalytic alpha polypeptide (PIK3CA) gene mutations. These PIK3CA-related overgrowth syndromes (PROS) can lead to asymmetric growth of lymphatics and mimic a systemic lymphangiomatosis. When confronted with patients suffering from PROS, efficacy has been established with sirolimus, and recent evidence shows promising results with BYL719, an inhibitor of PIK3CA.16

The present case illustrates the beneficial effect of sirolimus in a therapy resistant generalised lymphangiomatosis in an adult patient and highlights the importance of CT volumetry in the monitoring of the condition. Therefore, sirolimus therapy should be considered for diffuse lymphangiomatosis as it may be effective in improving the condition as illustrated in this patient, with persistent benefit 4 years after the initiation of treatment.

CASE PRESENTATION
A 35-year-old woman progressively developed abdominal pain, associated with severe lymphoedema of the legs, ascites and diarrhoea. She was not taking any medication and her medical history was unremarkable, except for a sphincterectomy and papillotomy for gallstones, a laparoscopic cholecystectomy and a left ovarian torsion. She did not have any familial antecedents of similar findings.

She did not smoke and there was no alcohol consumption.

Laboratory results showed a potassium level of 2.4 mmol/L, a calcium level 1.43 mmol/L, an albumin concentration 14 g/L, uric acid 1.8 mg/dL, gamma-globuline 1.7 g/L, transferrin saturation 28%, ferritine level 7 µg/L, haemoglobin 9 g/L, mean corpuscular volume (MCV) 72 fl, platelets 63 × 10^9/L, white cell count of 1.600 × 10^3/L, absolute lymphocytes 2.151 × 10^9/L and vitamin B₁₂ 0.18 µg/L.

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INVESTIGATIONS
An initial CT of the thoracoabdominal region showed a multilobar cystic mass in the mediastinum extending in the retroperitoneum and intraperitoneum. A total body MRI confirmed multiple hyperintense cystic lesions in mediastinum and retroperitoneum on T2-weighted images suggestive of lymphangiomatosis. A biopsy of the cystic lesions was performed under CT guidance, but biopsy was non-diagnostic, because there was not enough tissue sample.

DIFFERENTIAL DIAGNOSIS
The differential diagnosis should include metastatic ovarian cystic adenocarcinoma, cystic teratoma, lymphangioleiomyomatosis (LAM), PROS syndrome and benign lymphangiomatosis. The diagnosis of metastatic ovarian cystic adenocarcinoma was excluded with normal CA 125 and the atypical aspect on CT scan and MRI. Cystic teratoma usually presents as a complex cystic mass with a considerable solid component and calcifications, moreover the patient had normal alpha fetoprotein and beta-human chorionic gonadotropin (HCG) levels.16

PROS syndrome is a genetic disorder that results from somatic, mosaic gain-of-function mutations of the PIK3CA. Most patients present with major deformities and vascular masses that were not seen in the present case. LAM results from progressive proliferation of atypical smooth-muscle like cells. It often occurs as an inherited disorder called tuberous sclerosis complex (TSC), but it can be isolated and sporadic and primarily affects the lungs. Serum VEGF-D may be a useful biomarker in LAM.20

We proceeded with an upper gastrointestinal endoscopy that revealed multiple white spots in the duodenum and jejunum. Histopathology of the biopsy showed lymphangiectasia with no evidence of malignancy, nor the presence of LAM cells. Immunostaining for HMB45 was negative, but positive for CD31 and smooth muscle actin. TSC gene analysis was negative, and serum VEGF-D level was normal. Recently PIK3CA gene was analysed in a small skin vascular lesion and the mutation could not be identified.

The diagnosis was generalised lymphangiomatosis with a malabsorption due to protein losing enteropathy.

TREATMENT
The patient was put on a low-fat protein diet, with weekly infusion of albumin and immunoglobulin supplementation. Pan abdominal radiotherapy, with a total of 14.3 Gy in 11 fractions, was proposed but was stopped prematurely due to severe gastrointestinal side effects.

In the following years, different treatments, based on recent insights of the pathogenesis of this disease, were administered: sandostatine, low fat with medium chain triglyceride diet, followed by doxycline, beta-blockade and finally thalidomide. All treatments failed to induce significant changes in clinical, biological and radiographic findings. A treatment based on sirolimus 0.8 mg/m²/12 hours was initiated in January 2016. The dose of sirolimus was adjusted to keep trough concentrations between 6 and 30 µg/L.

OUTCOME AND FOLLOW-UP
Subsequent CT imaging showed a significant regression of the cystic lesions with an estimated volume reduction of 1226 cc to 447 cc (−64%) since the introduction of Sirolimus (Figure 1A–D). The cystic structures seen on the CT scans were manually segmented by the ‘volume tracing function’ (Intellispace Portal, Philips, The Netherlands), which implements a region growing algorithm that takes into account the density similarity of the cysts. Those densities were added to the segmentation. When all the cystic structures were selected, the algorithm calculated the content of the volume of interest.21 22

The patient continued infusion of albumin and immunoglobulin supplementation at a lower frequency with controlled serum albumin, and improvement of initial severe lymphopenia (white cell count of 3,400 × 10⁹/L, absolute lymphocytes 0.550 × 10⁹/L). The patient lost weight with regression of lymphoedema of lower limbs. She did not experience severe infections since treatment was started. This was indeed a major concern in the presence of severe lymphopenia and immunosuppression with sirolimus.

We postulate that regression of cystic lesions contributed to decreased lymphocyte sequestration and leakage resulting from diminished intestinal lymphangiectasias. Throughout the follow-up, the sirolimus levels were within therapeutic range.

DISCUSSION
Systemic generalised lymphangiomatosis is a rare disorder and usually becomes symptomatic at a young age. Although the condition is benign in nature, it may lead to severe symptoms and in some cases may have a debilitating course with high
mortality. Indeed, abdominal lymphangiomatosis can result in severe protein losing enteropathy and gastrointestinal bleeding due to the presence of diffuse intestinal lymphatic abnormalities, as described in the present case report. 4–9

The lymphopenia as observed in the present case is, however, exceptional and rarely reported.

Whether lymphangomas are true malformations or neoplastic in nature remains unsettled. Taking into account the unusual proliferative of lymphatics, mTOR inhibitors may represent a promising treatment strategy in the management of the observed lymphatic anomalies, although the level of evidence of efficacy is low and is essentially based on occasional case reports. 6–9

Patients treated with angiogenesis inhibitors were diagnosed at a young age. One can indeed presume that early intervention may be more promising by intervening in the process of lymphangiogenesis at an early stage. The efficacy at adult age of diagnosis remains uncertain.

In the broad differential diagnosis of lymphangiomatosis, patients should be screened for PIK3CA mutations, because several case reports and a recent published patient cohort have shown adult patients with a phenotype mimicking systemic lymphangiomatosis. 14–18

However, we could not detect this mutation in the patient.

The present observation adds support to the possible beneficial treatment of sirolimus in adult patients with diffuse lymphangiogenesis. Our patient was treated with sirolimus with dose adjustment according to measured trough levels with a favourable evolution of the associated protein losing enteropathy. This could not be achieved with previous therapeutic regimens, administered in parallel with supportive treatment. The treatment was not only efficient in controlling disease symptoms but also induced radiological regression of the cystic lesions. Volumetric CT as illustrated in the present case report constitutes a good measure to evaluate efficacy of treatment. 29–32

A similar observation has recently been made in the treatment of an adult patient with diffuse pulmonary lymphangiomatosis treated with sirolimus with a decrease in mediastinal masses and an improvement in pulmonary function. 32

Therefore, this case report highlights an important role of the angiogenesis inhibitors in the treatment of symptomatic and therapy resistant systemic generalised lymphangiomatosis.

Lymphopenia is not a limiting factor. Increase in absolute numbers of lymphocytes under sirolimus treatment might even be considered as a positive result with a possible benefit, including a decreased risk of secondary infectious complications.

Further insights in the pathogenesis of the disease and targeted treatments can result in a better outcome of patients with diffuse lymphangiomatosis.

**Learning points**

- Systemic generalised lymphangiomatosis is a benign disorder that may mimic malignancy with severe symptoms and aggressive behaviour.
- There is no uniform consensus on treatment of generalised lymphangiomatosis.
- Promising results can be expected with angiogenesis inhibitors.

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