Clinical Series

Liver Resections for Metastases from Intraabdominal Leiomyosarcoma

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This paper discusses liver resection for intraabdominal leiomyosarcoma metastases as a therapy for carefully selected patients. Of the 83 hepatectomies performed from 1992 to 1996, five were resections for liver metastases due to intraabdominal leiomyosarcoma, in 3 patients. The surgical indication was single liver metastases, without any evidence of extrahepatic disease. No mortality occurred during surgery and the longest survival was 38 months. We concluded that liver resection for leiomyosarcoma metastases can be performed, allowing a long term survival in an occasional patient.

Keywords: Leiomyosarcoma, liver metastases, liver resection, non-colorectal liver metastases

INTRODUCTION

Leiomyosarcomas are malignant tumors of smooth muscles, occurring most commonly in the uterus, retroperitoneal region and extremities. These lesions correspond to 7% of the soft tissue tumors, most often affecting adult, female patients [1].

Metastatic dissemination occurs by haematogenous spread, mainly to the lungs and liver, and only occasionally to regional lymph nodes [2]. Liver metastases are usually observed in the recurrence of visceral and retroperitoneal sarcomas they are often multiple and bilobar [3,4].

Some authors suggest that for certain non-colorectal tumors, including leiomyosarcoma, the resection of liver metastases might increase time of survival and/or promote substantial palliation [5–13].

This paper describes a group of patients with liver metastases due to leiomyosarcoma.

PATIENTS AND METHODS

During the period from 1992 to 1997, 83 hepatectomies were performed by a single surgical team in the Department of Surgery, Fundação Faculdade Federal de Ciências Médicas de Porto Alegre (FFFCMPA). Five of them were liver resections

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for metastases due to leiomyosarcoma, performed in three patients. This group consisted of two men and one woman whose ages ranged from 41 to 56 (mean, 46.6), all of them caucasian.

The site of the primary tumor, the time of diagnosis of liver metastases and their location are summarized in Table I.

All the patients were submitted to a preoperative assessment which consisted of routine lab tests, liver function tests and imaging (ultrasonography-US and computed tomography-CT).

Liver resection was indicated for a single metastasis from leiomyosarcoma, without any evidence of disease at another site.

Hepatectomy was performed by a bilateral subcostal incision, with afferent vascular exclusion in four resections and selective clamping in one hepatectomy (patient 3, the first resection). Subhepatic silicone drains were used in all patients. One patient required chest drainage (patient 3).

Surgical mortality was defined by death up to the 30th day post-surgery. Time of survival was computed in months (from date of liver resection up to the latest visit). Recurrence was defined by abnormalities in computed tomography, ultrasonography, biopsy or bone radio-nuclide scanning.

RESULTS

The presence or absence of recurrence after liver resection, the course and survival of patients are illustrated in Figure 1.

Two segmentectomies were performed in patient 1. Up to the present time, this patient,

FIGURE 1 Post-resection survival and evolution of liver metastases due to leiomyosarcoma 1-Patient alive, without recurrence at present, after re-hepatectomy at 25 months due to hepatic recurrence; 2-Died. Extra-hepatic recurrence at 4 months post-hepatectomy; 3-Died. Submitted to re-hepatectomy at 12 months due to hepatic recurrence and, at 24 months, presented new hepatic recurrence.

is alive and asymptomatic, having adjuvant chemotherapy after the second resection, with a survival of 37 months.

Left lobectomy was performed in patient 2, who was submitted to radiation therapy and chemotherapy five months after liver resection, due to bone metastasis.

Bi-segmentectomy was performed in patient 3. He survived for 38 months with recurrence in the liver after the second operation.

No mortality occurred in surgery, and the transfusion of blood products required ranged from 0 to 1000 ml (mean, 250 ml).

DISCUSSION

An assessment of risk factors for the development of metastases due to soft tissue sarcomas

TABLE I Study of primary tumor and metastases

Patients	Age (years)	Site of primary tumor	Type of metastasis	Location of metastases
1	41	S.B.*	1) Synchronous 2) Metachronous	1) Segment V 2) Segment VIII
2	56	Uterus	Metachronous	Segments II and III
3	43	Stomach	1) Synchronous 2) Metachronous	1) Segments V and VI 2) Segments VII and VIII

*S.B. – Small bowel.

is dependent on 3 factors: histological grade, histological type and primary tumor site [3].

Leiomyosarcomas constitute 7% of the histological types, often affecting the uterus, stomach and small bowel. The most undifferentiated tumors, with the worst prognosis, are found in the retroperitoneal sites [1,3].

Recent studies indicate the liver as a preferential site for metastases from visceral and retroperitoneal leiomyosarcomas [3,4,14]. Most of these metastases are multiple and bilobar [12], and 50% of the patients who developed them did so in the first year after diagnosis. This is the period during which one must carefully assess the liver of patients with intra-abdominal leiomyosarcoma [3,4]. Once the diagnosis of liver metastases has been established, the type and histological grade, as well as disease-free time after resection of the primary tumor do not appear to influence patient survival [3].

Chemotherapy, systemic or intra-arterial, does not have a substantial impact on the treatment of liver metastases from soft tissue sarcomas [3,4,15]. However, Mavligit *et al.* [4] report their experience with 14 patients who had gastrointestinal leiomyosarcoma, treated by hepatic chemoembolization, using a mixture of polyvinyl alcohol with cisplatin powder (150 mg), followed by intrahepatic arterial infusion of vinblastin (10 mg/m²). In this series, all the patients had unresectable liver metastases; however, one patient had a dramatic regression of the tumor, which enabled its resection.

From data obtained in series in which the time of survival after liver resection due to metastases from leiomyosarcoma was established, a mean time of 33 months was obtained, one patient presented a survival longer than 5 years [2–16]. Thus, it seems clear that the leiomyosarcomas constitute an indication for liver resection in the case of non-colonic metastases.

Based on these findings, we operated on 3 patients with liver metastases from leiomyosarcoma, and obtained a survival ranging from 24 to 38 months. Two patients presented with re-

ccurrence of the disease in the liver, and both of them were operated on again, corroborating the data from the literature which reports a high rate of recurrence of this type of tumor in the liver [3,6,13]. The worst prognosis, was in the case of the patient with a metastasis of non-digestive origin, who presented with an extrahepatic recurrence less than 6 months after hepatectomy.

Thus, we concluded that the resection of liver metastases due to leiomyosarcoma could be performed, allowing a long-term survival in an occasional patient.

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INVITED COMMENTARY

The incidence of isolated hepatic metastases from cancers of the colon or rectum is frequent enough to justify hepatic resection in selected patients. For other malignancies metastases to the liver only is quite a rare phenomenon, which explains why the literature contains relatively few and almost anecdotal reports of long-term survival following hepatic resection of metastases from non-colorectal malignancies, such as intra-abdominal leiomyosarcoma.

The present study has shown that long-term survival (at least for 3 years) is possible following

resection of metastatic intra-abdominal leiomyosarcoma. A few other authors have reported similar findings with an occasional 5-year survivor, although none of 10 patients with liver metastases from visceral leiomyosarcoma survived for 5 years in the largest single series [1]. It should be emphasized that spread to the liver is the main factor determining survival for visceral sarcomas but not for sarcomas originating at other locations, with the exception of retroperitoneal sarcomas that spread as often to the liver as to the lungs [1]. The pattern of spread, and the likelihood of having metastases only in the liver, is of course influenced by the portal venous drainage of gastrointestinal organs.

Definite indications do not exist, but it would appear that a patient with isolated and resectable hepatic metastases from gastrointestinal leiomyosarcoma should undergo resection [2]. A prerequisite is that the resection can be undertaken with low morbidity and low mortality (well under 5%). Considering the relative inefficiency of chemotherapy for leiomyosarcoma, it is also important to be able to ensure clear resection margins. The difficulty to be confident about indications and prognostic factors is true for hepatic resection of all other non-colorectal secondaries and should, hopefully, lead to an international registry for patients undergoing liver resection for non-colorectal secondaries.

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