Efficacy of trabectedin in metastatic solitary fibrous tumor

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Abstract

Solitary fibrous tumor is a rare tumor type and has an unpredictable course. Local recurrence rate varies between 9 and 19%, and rate of metastatic involvement between 0 and 36%. It is characterized by a typical architecture and immuno-histochemistry tests. The most important prognostic factor is the complete resection of primary tumor. Treatment of recurrences is not clearly established. If a solitary fibrous tumor is too advanced to allow surgical resection, radiotherapy and chemotherapy may be used. The most often used drugs are doxorubicine and ifosfamide. We report the case of a man with metastatic solitary fibrous tumor treated with trabectedin, administered at a dose of 1.5 mg/m² every 3 weeks. After 3 cycles, metastases had significantly decreased. Recurrence of the disease was demonstrated 8 months after the start of trabectedin. This case shows that trabectedin is a possible treatment option.

Introduction

Solitary fibrous tumor is a rare tumor type and has an unpredictable course. Treatment of recurrences is poorly codified. The surgery remains probably the reference. In the forms too advanced for a surgical solution, it is usual to use radiotherapy and chemotherapy. We describe a case report of a 39-year-old man with metastatic solitary fibrous tumor treated with trabectedin.

Case Report

Diagnosis

On July 2007, a 39-year old man without any relevant medical history had an assessment for chronic cough. Chest radiography revealed a left retro-cardiac thoracic mass. CT-scan found a voluminous solid, heterogeneous, well-encapsulated mass, with benign aspect, developed on lower half of left thorax. Lung biopsy concluded on solitary fibrous tumor (SFT) of the pleura. Surgical resection was performed on October 2007. Histological examination confirmed the diagnosis of SFT and surgical margins appeared free. In spite that this subtype of mesenchymatous tumor is widely considered as benign, most cases have the potential for local recurrence even after wide surgical excision. Typically, patient received regular follow-up examination including thoracic CT-scan. In February 2008, CT-scan revealed a unique 8 mm-lesion in the left lung and three other lesions on the right lung. Physical exam had not revealed any significant abnormalities associated with a good performance status. No complaints or functional limitations were described. These micronodules were totally nonspecific and asymptomatic. Their exact nature was uncertain. Subsequent CT-scan performed in Mars 2008 showed an increase in size and number of lung lesions, and doubtful liver images.

Lung biopsies were performed shortly afterwards. Pathological examination showed several very small-sized metastatic nodes in lung parenchyma (Figure 1). The round to spindleshaped cells had little cytoplasm and round, large-sized nuclei with relatively thin chromatin and small nucleoli. Vascular component was prominent with dilated vessels. Mitotic activity was 4 out of 10 high-power fields. Tumors cells were immunoreactive and immuno-histochemistry testing showed heterogeneous immunoreactivity for CD34, CD99 and bcl2 (Figure 2). focal and limited immunoreactivity was observed for P S100 and Cytokeratins. Less than 2% of cells expressed the proliferation marker Ki67. These examinations confirmed the diagnosis of SFT with metastatic evolution, based on the shape of lung micronodes and of doubtful liver images observed on CT-scan.

Treatment

Chemotherapy has been started according to the MAID (Mesna, Doxorubicine, Ifosfamide, Deticene) regimen, within framework of the PALSAR protocol from the FNCLCC and the French Sarcoma Group. After 6 cycles of MAID, best response was stable disease.

In January, 2009, CT-scan showed disease progression on lung metastases, which had increased in number and size. The biggest metastases were located in lingula and in left lower lobe of the lung (Figure 3). The hepatic lesions had become more visible. A second line of chemotherapy by trabectedin has been initiated with a 24-hour infusion at the dose of 1.5 mg/m² every 3 weeks, in a compassionate use program. Premedication consisted on a 30-minute IV infusion of 20 mg of dexamethasone before administration of trabectedin, as antiemetic prophylaxis and for liver protection, in accordance with good practice recommendations. After 3 cycles, lung metastases had significantly decreased (Figure 4). In addition, liver metastases had decreased in size, particularly at the segment VI level. After cycle 4 administration, trabectedin was definitely terminated due to a septic shock in the context of a grade 4 neutropenia. Subsequent CT-scan has been done in June 2009, i.e. 2 months after discontinuation of all chemotherapy agents. The pulmonary lesions were still in partial remission, particularly at the level of the right hilus (Figure 5). However, second recurrence of the disease was demonstrated in September of the same year, i.e. 8 months after the start.
of trabectedin and 5 months after the end of the chemotherapy.

Discussion

Solitary fibrous tumor was first described on 1931 by P. Klemperer and C. Rabin. SFT is a rare tumor type and often discovered in a fortuitous way. Initially observed in the pleura, SFT has also been described at many other sites, including soft tissue of orbit, of liver, thyroid, breast, gastro-intestinal tract, and finally of almost all organs. SFT can occur at all ages, with the highest incidence being between 60 and 70 years without evidence of unbalanced sex ratio.

As this tumor type has a slow growth, it often remains asymptomatic for a long time and can then become very voluminous.

For pathologists, SFT is well circumscribed, often partially encapsulated, of firm consistency, and has a pearly or yellowish-white color. It is described as a patternless architecture of spindle-shaped cells mimicking sometimes typical sarcomas like leiomyosarcoma, or synovial sarcoma. Moreover, it is characterized by a hemangiopericytoma-like architecture consisting of proliferation of medium-sized, well-formed vessels in the thick wall and hyaline, branching out. Immunohistochemically, SFT neoplastic tumor cells show expression of vimentin, CD34, CD99, bcl2, and less frequently EMA and smooth muscle actin. Focal and limited reactivity for protein S100, cytokeratins, and desmin has also been reported. Histological criteria to classify a SFT as malignant (indicating a risk of recurrence) include more than 4 mitoses out of 10 high-power fields, hypercellularity, nuclear atypia and presence of necrosis. Cytogenetic data on SFT are sparse and not coherent.

SFT has an unpredictable course. Local recurrence rates vary between 9 and 19 %, metastases rates (intra-thoracic, osseous, hepatic, cutaneous) vary between 0 and 36 %. The recurrences justify a long-term follow-up as some recurrences have appeared very lately (until 17 years after initial complete resection). The most important prognostic factor remains the complete resection of primary tumor. Treatment of recurrences is not clearly established and requires discussion at multi-disciplinary medical meetings. Surgery probably remains the standard treatment. If an SFT is too advanced to allow surgical resection, a parallel with soft tissue sarcomas is usually drawn. Radiotherapy and chemotherapy may be used. The most often used drugs are usually doxorubicin and ifosfamide. In our case, we have shown that trabectedin may be a possible option. Trabectedin has allowed tumor response for 8 months.

Figure 1. Lung biopsy: several metastatic micronodules of SFT (HES x 10).

Figure 2. Biopsy: SFT is characterized by spindle cells with vesiculare nuclei (HES x 40). All the tumoral cells are immunoreactive with CD34 (x40).

Figure 3. Computed tomography scan on January 2009, before trabectedin.

Figure 4. Computed tomography scan on March 2009, after 3 cycles of trabectedin.

Figure 5. Computed tomography scan on June 2009, 2 months later the end of chemotherapy.

Conclusions

Solitary fibrous tumor is a rare tumor type and has an unpredictable course. There are no

Trabectedin (Yondelis®) is a synthezised molecule, initially extracted from a marine animal (Ecteinascidia turbinata). It binds on the minor groove of the DNA and alkylates N2 of guanines. It may interfere with the DNA repair machinery and blocks cell in G2-M phase. The European Commission delivered a Marketing Authorization in September 2007. It is indicated for the treatment of patients with two types of cancers: i) Advanced soft tissue sarcoma, after failure of anthracyclines and ifosfamide, or who are unable to receive these agents. Trabectedin has been studied in one main study involving 266 patients with progressive advanced or metastatic soft tissue sarcoma. All of the patients had been treated previously with an anthracycline and ifosfamide. This study compared two different trabectedin regimens: 0.58 mg/m² every week for 3 weeks out of a 4-week cycle, or 1.5 mg/m² once every three weeks. Trabectedin was more effective when it was given once every three weeks. Median progression-free survival was 3.3 months versus 2.3 months respectively (hazard ratio, 0.755; 95% CI, 0.574 to 0.992; P=0.0418). Efficacy data are based mainly on liposarcoma and leiomyosarcoma patients. Results suggest that the drug may be particularly effective in patients with myxoid liposarcoma; ii) Platinum-sensitive recurrent ovarian cancer, in association with pegylated liposomal doxorubicin (PLD). Trabectedin (1.1 mg/m²) in combination with PLD (30 mg/m² every 3 weeks) was compared to PLD alone (50 mg/m² every 4 weeks) in one main study involving 672 women whose disease had occurred after previous treatment. Median progression-free survival was 7.3 months for the combination versus 5.8 months for PLD alone (hazard ratio, 0.79; 95% CI, 0.65 to 0.96; P=0.0190). The effect of trabectedin was more pronounced in the women whose cancer was platinum-sensitive (platinum-free interval of 6-12 months); iii) Others studies are ongoing in sarcoma, ovarian carcinoma, breast carcinoma and melanoma.

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guidelines for the treatment of recurrences. This case report suggests a new treatment option (trabectedin) and gives hope for this indication. Unfortunately, the absence of clinical study and published observation on this subject does not allow drawing of formal conclusion.

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