Case Report

Personalized selective internal radiation therapy in liver metastasis of thyroid cancer with impaired liver function: A case report

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ABSTRACT

Follicular thyroid cancer (FTC) is a less common form of differentiated thyroid cancer. Liver metastasis of differentiated thyroid cancer frequently occurs in the late onset of the metastatic disease, are often unresectable and noniodine avid, leading to a poor prognosis. A 69-year-old man with a 14-year history of multi-metastatic follicular thyroid cancer was treated iteratively with 131-Iodine allowing to maintain a stable disease. Upon a recent exponential increase of the thyroglobulin, a peritoneal mass and a voluminous hepatic metastasis were discovered, comorbidities and an insufficient future remnant liver function excluded liver surgical resection. The tumour board proposed a resection of the peritoneal mass followed by selective internal radiation therapy of the liver mass. Due to the already impaired liver function, personalized dosimetry allowed a safe treatment delivering low activity to the nontumoral liver followed by a clinical and imaging response of the liver mass at 3 months. At our knowledge, this is the first case of thyroid liver metastasis treated by selective internal radiation therapy.

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Introduction

Follicular thyroid cancer (FTC) is a less common form of differentiated thyroid cancer and spreads via haematogenous dissemination with distant metastases occurring in 10 to 15 percent of patients most commonly located in bone or lungs. According to the American thyroid association guidelines, treatment of FTC distant metastases requires $^{131}$I as long as iodine avidity is conserved. We report the first case of a patient presenting a progressive non resectable and iodine nonavid hepatic metastasis of FTC treated by radioembolization using hepatic selective internal radiation therapy (SIRT).

Case presentation

A 69-year-old man presented with follicular thyroid cancer (T4N1M1) initially treated by total thyroidectomy and ablation of one bone metastasis (cervical) followed by a first dose of radioactive 131-iodine ($^{131}$I). During 14 years follow-up, the patient presented repeatedly severe recurrences in bones, lungs, mediastinal lymph nodes and peritoneum treated iteratively with $^{131}$I (cumulative administered activity: 40.7 GBq in 6 sessions) each time achieving temporary disease control.

In 2018, after 3-years of loss of follow-up, the patient presented at the emergency unit for pyrexia. An increase of thyroglobulin (13076 μg/L) was noted. A diagnostic $^{131}$I-single photon emission computed tomography (SPECT/CT) (192 MBq) (Fig. 1a) was performed and showed a progressive disease with an intense uptake in a peritoneal mass without significant uptake of the liver mass. $^{18}$F-fluorodeoxyglucose positron emission tomography ($^{18}$F-FDG PET/CT) showed an increased glucose metabolism of the peritoneal mass, a mildly hypermetabolic liver metastasis and stable disease in lung and mediastinal lymph node (Fig. 1b). Magnetic resonance imaging (MRI) showed a 76 mm liver tumour in segments I, V, VI, VII, VIII (Fig. 1c) and a liver biopsy confirmed the presence of follicular carcinoma cells consistent with the known primary tumour. As part of preoperative work-up a hepatobiliary scintigraphy using $^{99m}$Technitium-mebrofenine ($^{99m}$Tc) was performed to determine hepatic function (HF) and assess future remnant liver function. Briefly, it consists in a dynamic planar acquisition followed by a SPECT/CT (Fig. 1d). Despite normal liver blood tests and no history of prior hepatic disease, the measured HF was 4.4 %/min/m² (normal value >7%/min/m², with the lowest acceptable HF cut-off limit for future liver remnant set at 2.7 %/min/m²) [1,2]. On this basis, the liver mass was considered as initially nonresectable, requiring preoperative liver volumes modulation. Due to the uncertainty of the benefit, the impaired liver function and the expected morbidity of such major resection, the thyroid and digestive multidisciplinary tumour board decided to perform a complete resection of the peritoneal mass followed by radioembolization of the liver metastasis. The radioembolization was realized with 90-Yttrium ($^{90}$Y) labelled resin microspheres, supra-selectively, in the segmental artery covering the lesion in order to keep as much as possible liver parenchyma out of the field of therapy.

SIRT simulation was performed with 2 supra-selective injections of $^{99m}$Tc-macro-aggregates of albumin respectively in the arteries vascularizing segment I and segments V to VIII (Fig. 1e). The field of treatment represented 85% of the whole liver and high selective uptake was noted (Tumour to nontumour uptake ratio: 3.54). No extrahepatic activity was observed. Personalized predictive voxel-based dosimetry was performed on macro-aggregates of albumin SPECT/CT images using a dedicated SIRT treatment planning system (Planet Onco dose, Dosisoft). Activity was prescribed to reach mean absorbed doses of 95 Gy and 35 Gy for the liver lesion and for the non-tumoural liver within the treatment field, respectively [3].

One week later, SIRT was performed at the exact same catheter positions as during the simulations without any immediate complication. A $^{90}$Y-PET/CT was acquired at day 1 post-SIRT, and post-treatment voxel-based 3D dosimetry was performed. Computed mean absorbed doses to the lesion and to nontumoural liver were 85 and 24 Gy, respectively.

Outcome and follow-up

Eight weeks post-RE, the patient was clinically doing well with no clinical or biological signs of radioembolization-induced liver disease (REILD). REILD is defined as a symptomatic post-radioembolization deterioration in the ability of the liver to maintain its (normal or preprocedural) synthetic, excretory, and detoxifying functions. It is characterized by jaundice and the development of or increase in ascites, hyperbilirubinemia, and hypoalbuminemia developing at least 2 weeks to 4 months after radioembolization, in the absence of tumor progression or biliary obstruction [4]. After 3 months, the size of the lesion was stable on MRI (Fig. 1f) with an overall increase in apparent diffusion coefficient values and a partial devascularization of the liver lesion. Thyroglobulin decreased from 4844 μg/L to 3667 μg/L, only partial remission as explained by the persistence of a residual extrahepatic FTC (mediastinal nodes and lung).

Discussion

Liver metastases of differentiated thyroid cancer (LMDTC), occurs in approximately 0.5% to 1.7% of the population with thyroid cancer [5,6]. LMDTC generally appears after the onset of metastases at other sites [5,7]. Because of a frequent dedifferentiation process (decreased expression and/or function of the sodium iodide symporter), most cases of LMDTC are not iodine-avid, limiting the effectiveness of radioiodine therapy [8]. Most patients are not eligible for surgery due to multiple LMDTC not accessible to complete surgical excision, disseminated metastases, or significant comorbidities. In this case the patient had already received a cumulative administered activity of 40.7 GBq of $^{131}$I, which could classify this disease as a $^{131}$I refractory disease [9]. Tyrosine kinase inhibitors have been reported as an effective option in patients with inoperable or refractory recurrent disease [6,9]. Due to the already impaired liver function, predictive activity prescription was indicated in order to optimize treatment efficacy and avoid...
the risk of REILD. The loss of liver function could be related to the multiple prior iodine-131 therapies. Should the classical modified body surface area (mBSA)-method have been used for SIRT activity prescription in this patient, the given activity would have been double (1450 MBq instead of 728 MBq) which would have resulted in a mean absorbed dose of the normal liver superior to 40 Gy associated with a very high risk of REILD [3,10,11].

**Conclusion**

To our knowledge, this is the first case of a thyroid liver metastasis successfully treated by SIRT using 90Y-microspheres as part of a multimodal treatment strategy. This case also highlights the importance of pretherapeutic work plan and predictive dosimetry to optimize personalized treatment to minimize the risk of REILD in an already impaired liver.

**Statement of ethics**

All data about the patient's privacy is blinded. The patient gave his informed consent for the publication of this case report.

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