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**Hepatic gastrointestinal stromal tumor: Systematic review of an exceptional location**

Manuel-Vázquez A *et al*.Hepatic GIST

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**Abstract**

***BACKGROUND***

A minor subset of primary gastrointestinal stromal tumors (GIST) can also arise outside the gastrointestinal tract, which is known as an extra-GIST (E-GIST). Primary GIST of the liver is an exceptional location.

***AIM***

To characterize epidemiological, clinical and pathological features and options of treatments.

***METHODS***

We performed a systematic review to search for articles on primary hepatic GIST.

***RESULTS***

This review shows that right hepatic lobe was the most frequent location. Regarding pathological and immunohistochemical features, mitotic count was ≥ 5/50 High Power Fields in more than 50%; and CD117 was negative in only 1 patient. More than 70% of patients had a lesion with high risk of malignancy.

***CONCLUSION***

The diagnosis of E-GIST must be considered in a liver mass. Rendering an accurate diagnosis is a challenge, as well as the confirmation of their primary or metastatic nature.

**Key words:** Gastrointestinal stromal tumors; Extra-gastrointestinal stromal tumor; Primary hepatic tumor; CD117; Primary hepatic gastrointestinal stromal tumor; Primary gastrointestinal stromal tumor of the liver

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**Core tip:** A great majority of primary gastrointestinal stromal tumors (GISTs) outside the gastrointestinal tract (GI) are metastases; however, a minor subset of primary GISTs can also arise outside the GI tract which is known as an extra-GIST (E-GIST). Among E-GIST, liver is an exceptional location. We systematically review the literature on primary GIST of the liver. Primary hepatic EGISTs have a male predominance and usually are incidental findings. The surgical approach is commonly performed, and the final diagnosis is made by pathological, immunohistochemical and molecular analysis. Primary hepatic EGISTs are often high-risk lesions. Literature is scarce and it is very difficult to establish guidelines for clinicians.

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**INTRODUCTION**

Gastrointestinal stromal tumors (GISTs) are a group of mesenchymal tumors characterized by the expression of KIT protein (CD117), with an overall incidence between 10-20 per million, which harbor different clinical behavior[1,2]. These are the most common mesenchymal neoplasm of the gastrointestinal tract, which represent 0.1%-3% of all gastrointestinal neoplasms[3,4].

Regarding the pathogenesis, GISTs are believed to originate from interstitial cells of Cajal (ICC), the pacemaker of gastrointestinal tract[5-7], or to a common precursor cell of ICCs and smooth muscle cells[8]. So, GISTs may arise anywhere along the gastrointestinal tract. Approximately 60%-70% of GIST occurs in the stomach, followed by 20%-30% in small intestine; colon and rectum (5%), esophagus (< 2%) and appendix are less frequent[9,10].

A great majority of GISTs outside the GI tract are metastases from GI GISTs; however, a minor subset of primary GIST can also arise outside the GI tract which is known as an extra-GIST (E-GIST). According to Miettinen *et al*[11], the frequency of EGISTs is no higher than 1% of all GISTs of defined origin, and they are characterized by the same morphological, immunohistochemical and molecular characteristic than conventional GIST[12].

Regarding embryology, the origin of EGISTs remains controversial, with certain hypotheses. Since some researchers have observed “ICC-like” cells with a similar structure and function to ICCs in organs outside of the GI tract[13,14], it is reasonable to presume that EGISTs originate from this common precursor cells of ICC. Other authors suggest that EGISTs may originate from a pool of undifferentiated pluripotent mesenchymal stem cells located outside GI tract, and then differentiate into ICCs[15-17].

Among E-GIST, mesentery, retroperitoneum and pancreas are the most frequent and this type of tumor has been reported in the omentum, bladder, gallbladder and uretus also; while, primary GIST of the liver is an exceptional location and its definitive diagnosis requires ruling out other types of liver mesenchymal tumors and other tumor types, such as sarcoma[6,12,15].

Most studies of E-GISTs are case reports, lacking enough information to clarify the disease; therefore, it is very difficult to establish guidelines for clinicians. In addition, if we refer to primary hepatic E-GIST, the literature is much scarcer due to this exceptional location.

This study is a systematic review of the literature on primary GIST of the liver. Our aim is to identify clinical and diagnostic features and treatment in this exceptional location of this type of mesenchymal tumor.

**MATERIALS AND METHODS**

We performed a search for articles on primary hepatic GIST in MEDLINE (PubMed), Tripdatabase, and Cochrane Library databases, with no restrictions on publication dates or author up until January 31, 2019.

The search items comprise the following MESH terms: “Extra-gastrointestinal stromal tumors” OR “extra-gastrointestinal stromal tumor” OR “extra-gastrointestinal stromal neoplasm” OR “extra-gastrointestinal stromal neoplasms” OR “E-GIST” OR “E-GISTs” and the following no MESH terms: “Primary malignant gastrointestinal stromal tumor of the liver” OR “Primary hepatic gastrointestinal stromal tumor” OR “Primary gastrointestinal stromal tumor of the liver”.

The articles were included or rejected based on the information obtained from the title and summary, and in case of doubt, after reading the complete article.

To evaluate the quality of the studies selected, we used the scale designed by Manterola *et al*[18] which evaluates each publication individually depending on the type of study, the sample size, and the methodology used. It has a range of 6 to 36 points, with a quality cutoff point of 18. We carried out a qualitative analysis of the studies included and their conclusions, based on the levels of evidence and degrees of recommendation proposed by Cook *et al*[19].

**RESULTS**

After the both initial searches, 420 articles were obtained. Only 23 (5.48%) met the search criteria, one of which were excluded because language (one in Romanian). The flowchart diagram is shown in Figure 1. We included 22 articles, including 23 patients[1,3,6,20-38].

The mean age was 56.18 years (± 15.4 SD), with a slightly male predominance (12/23). According country, 18 patients were from Southeast Asian, with 11 cases from China; while only 5 came from Western (France, Spain, Italy, United States). Right lobe was the most frequent location (12/23; 52.17%), and bilobar extension was present in 4 patients. Liu *et al*[28] report the coexistence of a hepatic and a pancreatic primary lesion. The median size was 15 cm (range: 2.2-27 cm). Among the symptoms, nearly 50% of patients (10/23) have no symptoms, being incidental finding during follow-up or extension study for gastric cancer in 2 of them (Table 1)

Regarding the diagnosis referred in Table 2, upper and lower endoscopy studies were performed in 9 patients, with no findings in 8 cases and an early gastric cancer in the other (biopsy: signet ring cell carcinoma). Biopsy of hepatic lesion was performed in 8 patients; one of them was surgical one.

Regarding the treatment of the selected patients, the management was surgical in 16 cases, ranging from limited surgical excision or anatomic resection to liver transplantation or extracorporeal resection and auto transplantation.

Two patients were received local treatment, with radiofrequency ablation (RFA) and microwave ablation. Luo *et al*[37] report a central hepatic lesion in a young patient. Liu *et al*[28] report a pancreatic primary E-GIST coexisting with another hepatic primary E-GIST, where both lesions were treated by RFA.

There were two refusals for treatment; in one case the patient refused surgery and received imatinib mesylate, and the other refused any action. In only one patient, the finding was a non-resecable hepatic lesion and the patient was treated with imatinib mesylate.

Regarding pathological, molecular and immunohistochemical findings, the most frequent cell type was spindle cells (17/23, 73.91%); molecular analysis was performed in 6 patients, with mutation in 5/6; mitotic count was ≥ 5/50 High Power Fields in 12 cases (52.17%); and CD117 was negative in only 1 patient. The risk of malignancy was classified according to Fletcher *et al*[2] and 17/23 (73.91%) patients had a lesion with high risk of malignancy (Table 4).

Mean follow-up was 14 mo (range: 3-252). During the follow-up, 10 patients were disease-free (follow-up: 3-30 mo) (Table 5).

**DISCUSSION**

Since Hu *et al*[6] reported the first primary hepatic GIST in 2003, we should consider that not all tumors of the liver with GIST features are metastasis and the liver could itself be the primary GISTs location.

Primary hepatic E-GISTs are extremely uncommon compared with their alimentary counterparts; thus, E-GISTs presenting in the liver raise a difficult diagnosis, management and prognosis[22].

In this review, primary E-GISTs of the liver have a slightly male predominance and the reported cases have a Southeast Asian predominance. Regarding symptoms, almost 50% have no clinical manifestations; among symptomatic patients, the symptoms are vague compared to GISTs, which commonly present with GI bleeding, abdominal pain, a palpable mass, weight loss, nausea and vomiting[12]. Also, the size of the hepatic lesion may become larger, mean 15 cm in our review.

Once we find a tumor with GIST characteristics presenting in the liver, the main challenge is determining whether this lesion is primary or metastatic, considering the liver is the most common site of distant metastasis for malignant GIST[21,28].

All of the studies included in this systematic review are case report, lacking enough information to clarify the disease and making difficult to establish protocols. The differential diagnosis is the main challenge for primary hepatic E-GISTs, but there are no consensus guidelines. According to Joyon *et al*[21], the diagnosis of primary E-GIST of the liver could be considered if these conditions are present: (1) Absence of GIST in the GI tract, with endoscopic and imaging studies; and absence of connection with the muscularis propia of the GI tract; (2) Absence of any past medical history which might suggest the resection of an overlooked or misdiagnosed GIST; and (3) Absence of GI tumor diagnosed during follow-up.

Thus, every hepatic GIST should be considered metastatic until no grossly nor histologically evidence of association with the muscularis propia have been shown and the remote medical history has been carefully explored, and no primary tumor could be found on long-term follow up[39]. Not all case included in this systematic review have been described enough information to be sure that all these requirements are fulfilled. In one hand, Kim *et al*[32]reported a patient with two synchronous lesions, an early gastric cancer and a primary hepatic GIST. Their preoperative diagnosis was a malignant hepatic lesion with an early gastric cancer, due to presence of signet ring cells in gastric carcinoma and radiological features of liver tumor. After surgery, the pathological study showed an early gastric cancer in the lesser gastric curvature, with signet ring cell carcinoma features, and spindle cells with a positive reactivity CD117 in hepatic tumor.Nagai *et al*[25] reported a hepatic primary E-GIST in a patient with a previous gastric cancer 7 years earlier. Histologically, gastric cancer was a poorly differentiated adenocarcinoma, with lymph node involvement. When the hepatic mass was founded, upper and lower gastrointestinal endoscopic studies were performed, with no findings. Microscopically, the hepatic tumor was composed of spindle cells, with positive results for KIT and CD34[25]. Ochiai *et al*[38] reported a patient with surgical resection of a primary hepatic GIST, based on the positive immunostaining for CD34 and c-kit, and recurrent hepatic lesion and submucosal gastric tumor 2 years after first operation. After surgical removal of both hepatic and gastric lesions, both specimens showed GIST features with expression of c-kit and CD34, but the different morphological and molecular findings (gastric lesion: spindle cells and mutation in exon of c-kit; hepatic tumor: round cells and no mutation at exon 11) were enough for the authors to conclude that the hepatic GIST and gastric tumor were independent. In these patients, the different histological and immunohistochemical findings were the reason to define the hepatic lesion as a primary E-GIST. On the other hand, Liu *et al*[28] reported two synchronous E-GISTs. The patient presented a 5 cm-pancreatic mass and a 2 cm-hepatic lesion, with the same radiological features, and no other lesions in upper/lower gastrointestinal endoscopic examinations. With the diagnosis of malignant pancreatic cancer with hepatic metastases, the authors performed a surgical fine-needle aspiration and pathological findings in the hepatic and pancreatic biopsy tissues indicated that the tumors were mitotic spindle cell with CD117+[28]. The authors conclude that they are two independent lesions due to the pancreas and liver are exceptional location for primary GIST.

EGISTs have the same morphological, immunohistochemical and molecular features than conventional GISTs, including metastatic ones[12]. The criteria for the histopathological diagnosis are now firmly established; tumor cells might present a spindle or an ephiteliod appearance and show a distinctive immunophenotype characterized by the expression of KIT (CD 117)[2,28,40]. Thus, the definitive diagnosis relies on the histopathological examination[41]; however, none of the pathological features is constant and required for a definitive diagnosis, including KIT mutation, which is no detectable in almost 5% of cases[29,34]. The concerning of KIT-negative GISTs could be solved after the discovery of novel mutations of the platelet derived growth factor receptor alpha oncogene as alternative pathogenetic mechanism[39].

In the majority of cases of E-GISTs, preoperative diagnosis is not possible; therefore, patients may be easily misdiagnosed with different types of cancer and surgery is performed to made a confirm diagnosis after histological examination.

The overall management of hepatic E-GISTs is generally based on the recommendations for GI GIST. A large spectrum of therapeutic options has been proposed depending on the initial presentation and clinical context. As with GISTs, complete surgical resection is the mainstay of treatment for E-GISTs, as long as the lesion is resectable[24,28].

A guided tumor biopsy must be considered for no-resectable tumors in order to assess the diagnosis and to offer another option for treatment such as radiofrequency, arterial embolization or chemoembolization may be considered[24,28,37]. The challenge is the risk of tumor rupture by the biopsy, well-known adverse prognostic factor in conventional GIST[42].

For E-GIST and as with GI GISTs, imatinib mesylate may be administered preoperatively in locally advanced tumor in order to minimizing the size, for adjuvant treatment for patients with a high recurrence risk or for palliative treatment in no-resectable lesions, which is similar to the guidelines for their alimentary counterparts[1,28,29,30,43]. Rediti *et al*[44] in 2014 and Wada *et al*[45] in 2012 reported the complete remission in a patient with greater omentum-mesentery E-GIST and a peritoneal E-GIST, respectively, who received only imatinib mesylate as a treatment.

On 2001, National Institutes of Health proposed a consensus classification system for defining the risk of malignant behavior, based on mitotic count and tumor size and now is widely in use, also for E-GIST[2]. Joensuu *et al*[46] proposed a new modified classification including primary tumor site and tumor rupture in the item for classifying the risk of GIST and the indication for adjuvant treatment.

Compared with GIST, E-GISTs have been reported to be accompanied by adverse prognostic factors, including a high proliferative index, large size and distant metastasis[15]; so E-GIST is considered to exhibit a worse prognosis, with a higher malignant potential and risk of recurrence following surgery compared with GISTs in the GI tract[2,11,15,24,26,42] Disease free-survival and disease specific-survival of hepatic GISTs are significantly worse than those of gastric and small intestine GISTs and the location is an independent prognostic factor[26]. There is a trend that E-GIST is an aggressive group with worse outcome. In this review, which includes only case reports, 10/23 patients are DF with a follow-up between 3 and 30 mo, and 8/23 patients had progression of E-GIST or metastasis, with not available data in 5 patients.

In conclusion, the diagnosis of E-GIST must be considered in a liver mass. Rendering an accurate diagnosis is a challenge, as well as the confirmation of their primary or metastatic nature. The optimal treatment is surgery and imatinib mesylate has a role as neoadjuvant treatment for locally advanced tumors, adjuvant treatment if the lesion has a high risk for recurrence, and palliative treatment if there is distant metastasis. Literature on hepatic E-GIST is scarce and further studies such as multicentric databases are needed to clarify diagnosis and treatment.

**Article Highlights**

***Research background***

A minor subset of primary gastrointestinal stromal tumors (GIST) can also arise outside the gastrointestinal tract, which is known as an extra-GIST (E-GIST).

***Research motivation***

Primary GIST of the liver is an exceptional location and this study aimed to characterize epidemiological, clinical and pathological features and options of treatments.

***Research objectives***

Our aim is to characterize epidemiological, clinical and pathological features and options of treatments.

***Research methods***

We perform a system review including all patients with hepatic GISTs.

***Research results***

This review shows that right hepatic lobe was the most frequent location, the median size was higher, there was a Southeast Asian predominance, and nearly 50% of patients have no symptoms. The most frequent treatment was surgery and more than 70% of patients had a lesion with high risk of malignancy.

***Research conclusions***

Literature on hepatic EGIST is scarce and further studies such as multicentric databases are needed to clarify diagnosis and treatment.

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**Figure 1** **Flowchart.**

**Table 1 Clinical characteristics and location in selected patients with extra-gastrointestinal stromal tumor of the liver**

|  |  |  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- | --- | --- |
| **Reference** | **Yr** | **Age/Sex** | **Country** | **Presentation** | **Location** | **Size (cm)** | **Multifocal** |
| Hu *et al*[20] | 2018 | 79/F | China | Epigastric discomfort | RL | 3.2 | No |
| Joyon *et al*[21] | 2018 | 56/M | France | Abdominal pain | Bilobar (Segments VII/VIII and LL) | 10 | Yes |
| 2018 | 59/F | France | Abdominal pain, weight loss | RL | 23 | No  |
| Carrillo *et al*[22] | 2017 | 41/M | Spain | Abdominal pain, weight loss | RL (S. V-VI) | 20 | No |
| Lok *et al*[23] | 2017 | 50/F | China | Abdominal pain | RL | 15 | Yes |
| Cheng *et al*[24] | 2016 | 63/M | China | No symptoms | RL | 15 | No |
| Nagai *et al*[25] | 2016 | 70/F | Japan | Follow-up gastric cancer, No symptoms | LL | 6 | No |
| Liu *et al*[26] | 2016 | NA | China | NA | NA | NA | NA |
| Wang *et al*[27] | 2016 | 61/M | China | No symptoms | Caudate lobe | 7.3 | No |
| Liu *et al*[28] | 2016 | 56/F | China | No symptoms | LL + Pancreas | 2.2 | No |
| Su *et al*[29] | 2015 | 65/M | Taiwan | Malaise, loss of apetite, abdominal pain | LL | 12 | No |
| Bhoy *et al*[3] | 2014 | 41/F | India | Abdominal pain, weight loss | RL (S.VI-VI)I | 15 | Yes |
| Lin *et al*[30] | 2015 | 67/F | China | No symptoms | RL | 7.4 | No  |
| Mao *et al*[31] | 2015 | 60/F | China | No symptoms | Bilobar (S I, IV, V, VIII) | 12.8 | No  |
| Kim *et al*[32] | 2014 | 71/M | Korea | Study for early gastric cancer, No symptoms | LL | 7 | No |
| Louis *et al*[33] | 2014 | 55/F | India | Abdominal pain, loss of apetite | Bilobar (SII, III, VI, VIII) | 14.5 | Yes |
| Zhou *et al*[34] | 2014 | 56/M | China | No symptoms | RL | 10 | No  |
| Li *et al*[35] | 2012 | 53/M | China | Abdominal discomfort | RL | 20 | No |
| Yamamoto *et al*[36] | 2010 | 70/M | Japan | Loss of apetite (12 years after gastric cancer) | LL | 20 | No |
| Luo *et al*[37] | 2009 | 17/M | China | No symptoms | RL | 5 | No |
| Ochiai *et al*[38] | 2009 | 30/M | Japan | Abdominal fullnes | Bilobar | 27 | No |
| De Chiara *et al*[1] | 2006 | 37/M | Italy | No symptoms | RL (SV) | 18 | No |
| Hu *et al*[6] | 2003 | 79/F | USA | Shortness of breath, pleuritic chest pain | RL | 15 | No |

M: Male; F: Female; RL: Right lobe; LL: Left lobe; NA: Not available.

**Table 2 Diagnostic characteristics in selected patients with extra-gastrointestinal stromal tumor of the liver**

|  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- |
| **Reference** | **Yr** | **Age/Sex** | **Endoscopy**  | **Imaging** | **Biopsy** |
| Hu *et al*[20] | 2018 | 79/F | Yes (no findings) | CT | No |
| Joyon *et al*[21] | 2018 | 56/M | NA | CT | Percutaneous |
|  | 2018 | 59/F | Yes (no findings) | CT | Guided |
| Carrillo *et al*[22] | 2017 | 41/M | Yes (no findings) | CT / MRI | No |
| Lok *et al*[23] | 2017 | 50/F | Yes (no findings) | CT | No |
| Cheng *et al*[24] | 2016 | 63/M | NA | CT | No |
| Nagai *et al*[25] | 2016 | 70/F | Yes (no findings) | CT / MRI | No |
| Liu *et al*[26] | 2016 | NA | NA | NA | NA |
| Wang *et al*[27] | 2016 | 61/M | NA | CT | No |
| Liu *et al*[28] | 2016 | 56/F | NA | CT | Surgical  |
| Su *et al*[29] | 2015 | 65/M | NA | CT | CT-guided |
| Bhoy *et al*[3] | 2014 | 41/F | NA | US/ CT | FNA |
| Lin *et al*[30] | 2015 | 67/F | Yes (no findings) | CT | No |
| Mao *et al*[31] | 2015 | 60/F | Yes (no findings) | CT / MRI | No |
| Kim *et al*[32] | 2014 | 71/M | Early gastric cancer | CT / MRI | No |
| Louis *et al*[33] | 2014 | 55/F | NA | CT | US-FNA/CT-FNA |
| Zhou *et al*[34] | 2014 | 56/M | Yes (no findings) | NA | No |
| Li *et al*[35] | 2012 | 53/M | NA | CT | US-FNAB |
| Yamamoto *et al*[36] | 2010 | 70/M | NA | CT | No |
| Luo *et al*[37] | 2009 | 17/M | NA | CT/US | US-FNA |
| Ochiai *et al*[38] | 2009 | 30/M | NA | CT/ MRI | No |
| De Chiara *et al*[1] | 2006 | 37/M | NA | CT | No |
| Hu *et al*[6] | 2003 | 79/F | NA | CT | No |

M: Male; F: Female; RL: Right lobe; NA: Not available; US: Ultrasound; CT: Computerized tomography; MRI: Magnetic resonance image; FNA: Fine needle aspiration.

**Table 3 Treatment and adjuvant treatment in selected patients with extra-gastrointestinal stromal tumor of the liver**

|  |  |  |  |  |
| --- | --- | --- | --- | --- |
| **Reference** | **Yr** | **Age/Sex** | **Treatment** | **Adjuvant imatinib mesylate** |
| Hu *et al*[20] | 2018 | 79/F | Curative surgical resection | Yes |
| Joyon *et al*[21] | 2018 | 56/M | OLT | No |
|  | 2018 | 59/F | Refused surgery: imatinib mesylate | No  |
| Carrillo *et al*[22] | 2017 | 41/M | Segmentectomy V-VI | Yes |
| Lok *et al*[23] | 2017 | 50/F | Right hepatectomy | Yes |
| Cheng *et al*[24] | 2016 | 63/M | Right hepatectomy | Yes |
| Nagai *et al*[25] | 2016 | 70/F | Left lateral segmentectomy  | No |
| Liu *et al*[26] | 2016 | NA | NA | NA |
| Wang *et al*[27] | 2016 | 61/M | Caudate lobe resection | No |
| Liu *et al*[28] | 2016 | 56/F | Microwave ablation | Yes |
| Su *et al*[29] | 2015 | 65/M | Irresecable: imatinib mesylate | No |
| Bhoy *et al*[3] | 2014 | 41/F | Right hepatectomy | Yes |
| Lin *et al*[30] | 2015 | 67/F | Surgical excision | Yes |
| Mao *et al*[31] | 2015 | 60/F | ECHRA | Yes |
| Kim *et al*[32] | 2014 | 71/M | Left lateral segmentectomy+excision of 1 intrabadominal nodule+total gastrectomy | NA |
| Louis *et al*[33] | 2014 | 55/F | Segmentectomy III and atypical resection (segments II, VI and VIII) | Yes |
| Zhou *et al*[34] | 2014 | 56/M | Anterior and median segmentectomy | No |
| Li *et al*[35] | 2012 | 53/M | Refused treatment | No |
| Yamamoto *et al*[36] | 2010 | 70/M | Left hepatectomy | NA |
| Luo *et al*[37] | 2009 | 17/M | RFA | NA |
| Ochiai *et al*[38] | 2009 | 30/M | L-Trisegmentectomy | Yes |
| De Chiara *et al*[1] | 2006 | 37/M | NA | No |
| Hu *et al*[6] | 2003 | 79/F | Right lobectomy | NA |

M: Male; F: Female; OLT Ortotopic liver transplantation; ECHRA Extracorporeal hepatic resection and autotransplantation; NA: Not available; RFA: Radiofrequency ablation.

**Table 4** **Pathological, immunohistochemical and molecular findings, and risk of malignancy according to Fletcher *et al*[2]**

|  |  |  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- | --- | --- |
| **Reference** | **Yr** | **Age/Sex** | **Cell type** | **Molecular analysis** | **Mitotic count (nº/50 HPF)** | **IH** | **Risk of malignancy** |
| Hu *et al*[20] | 2018 | 79/F | Spindle cells | NA | NA | CD117+, CD34+ | NA |
| Joyon *et al*[21] | 2018 | 56/M | Spindle cells | NA | 8 | CD117+, CD 34+ | High risk |
|  | 2018 | 59/F | Mixed (spindle and epithelioid) | 6 bp deletion in KIT exon 11 | 42 | CD117+ | High risk |
| Carrillo *et al*[22] | 2017 | 41/M | Spindle cells | 9 deletion in KIT | 5 | CD 117+, CD 34- | High risk |
| Lok *et al*[23] | 2017 | 50/F | Spindle cells | NA | 70 | CD117+, CD 34+ | High risk |
| Cheng *et al*[24] | 2016 | 63/M | Spindle cells | NA | >5 | CD 117 +, CD 34- | High risk |
| Nagai *et al*[25] | 2016 | 70/F | Spindle cells | NA | 40 | CD117+, CD 34+ | High risk |
| Liu *et al*[26] | 2016 | NA | NA | NA | NA | NA | NA |
| Wang *et al*[27] | 2016 | 61/M | Spindle cells | NA | NA | CD117+/CD34+ | High risk |
| Liu *et al*[28] | 2016 | 56/F | Spindle cells | NA | 2 | CD117+ | Low risk |
| Su *et al*[29] | 2015 | 65/M | Spindle cells | NA | 5 | CD117+, CD 34- | High risk |
| Bhoy *et al*[3] | 2014 | 41/F | NA | NA | NA | CD 117+ | High risk |
| Lin *et al*[30] | 2015 | 67/F | Mixed (spindle and epithelioid) | Mutation in exon 11 | 8 | CD117+, CD 34+ | High risk |
| Mao *et al*[31] | 2015 | 60/F | Spindle cells | Mutation in exon 11 | >10 | CD 117+, CD 34 - | High risk |
| Kim *et al*[32] | 2014 | 71/M | Spindle cells | NA | 30-32 | CD117+ | High risk |
| Louis *et al*[33] | 2014 | 55/F | Spindle cells | NA | 10 | CD117+ | High risk |
| Zhou *et al*[34] | 2014 | 56/M | Spindle cells | NA | <5 | CD117+/CD34+ | Intermediate |
| Li *et al*[35] | 2012 | 53/M | Spindle cells | NA | NA | CD117+, CD34+ | High risk |
| Yamamoto *et al*[36] | 2010 | 70/M | Epithelioid cells | mutation PDGFRA exon 12 | 1 | CD 117-/CD34+ | High risk |
| Luo *et al*[37] | 2009 | 17/M | Spindle cells | NA | 0 | CD117+, CD 34+ | Low risk |
| Ochiai *et al*[38] | 2009 | 30/M | Mixed (spindle and epithelioid) | No mutation at exon 11 | 75 | CD117+, CD 34+ | High risk |
| De Chiara *et al*[1] | 2006 | 37/M | Spindle cells | NA | 20 | CD117+, CD 34+ | High risk |
| Hu *et al*[6] | 2003 | 79/F | Spindle cells | NA | 20 | CD117+, CD 34+  | Low risk |

M: male; F: female; NA: not available; PDGFRA: Platelet derived Growth Factor Receptor Alpha; HPF: High Power Fields.

**Table 5** **Outcome and follow-up in selected patients with EGIST of the liver**

|  |  |  |  |
| --- | --- | --- | --- |
| **Reference** | **Yr** | **Outcome** | **Follow-up (mo)** |
| Hu *et al*[20] | 2018 | NA | NA |
| Joyon *et al*[21] | 2018 | Local recurrence (12 yr) | 252 |
|  | 2018 | DF | 18 |
| Carrillo *et al*[22] | 2017 | DF | 18 |
| Lok *et al*[23] | 2017 | Brain metastasis (6 mo) | 6 |
| Cheng *et al*[24] | 2016 | DF | 30 |
| Nagai *et al*[25] | 2016 | DF | 10 |
| Liu *et al*[26] | 2016 | NA | NA |
| Wang *et al*[27] | 2016 | DF | 12 |
| Liu *et al*[28] | 2016 | Abdominal metastasis | 17 |
| Su *et al*[29] | 2015 | Progression of disease (died 6 mo) | 6 |
| Bhoy *et al*[3] | 2014 | DF | 5 |
| Lin *et al*[30] | 2015 | Hepatic recurrence (24 mo) | 72 |
| Mao *et al*[31] | 2015 | DF | 12 |
| Kim *et al*[32] | 2014 | NA | NA |
| Louis *et al*[33] | 2014 | DF | 6 |
| Zhou *et al*[34] | 2014 | DF | 12 |
| Li *et al*[35] | 2012 | NA | NA |
| Yamamoto *et al*[36] | 2010 | NA | NA |
| Luo *et al*[37] | 2009 | DF | 3 |
| Ochiai *et al*[38] | 2009 | Hepatic recurrence, submucosal gastric tumor (24 mo) | 25 |
| De Chiara *et al*[1] | 2006 | Lung mestastasis (14 mo) | 39 |
| Hu *et al*[6] | 2003 | Portal lymph node mestastasis (16 mo) | 16 |

NA: Not available; DF: Disease free.