

# Limited Resection Versus Pancreaticoduodenectomy for Duodenal Gastrointestinal Stromal Tumors? Enucleation Interferes in the Debate: A European Multicenter Retrospective Cohort Study

Clément Dubois, MD<sup>1</sup>, Frederiek Nuytens, MD<sup>1</sup>, Hélène Behal<sup>2</sup>, Caroline Gronnier, MD, PhD<sup>3</sup>, Gilles Manceau, MD, PhD<sup>4</sup>, Maxime Warlaumont, MD<sup>1</sup>, Alain Duhamel, PhD<sup>2</sup>, Quentin Denost, MD, PhD<sup>3</sup>, Charles Honoré, MD, PhD<sup>5</sup>, Olivier Facy, MD, PhD<sup>6</sup>, Jean-Jacques Tuech, MD, PhD<sup>7</sup>, Guido Tiberio, MD<sup>8</sup>, Cécile Brigand, MD, PhD<sup>9</sup>, Jean-Pierre Bail, MD<sup>10</sup>, Ephrem Salame, MD<sup>11</sup>, Bernard Meunier, MD, PhD<sup>12</sup>, Jérémie H. Lefevre, MD, PhD<sup>13</sup>, Muriel Mathonnet, MD, PhD<sup>14</sup>, Mohamed Sbai Idrissi, MD<sup>15</sup>, Florence Renaud, MD, PhD<sup>16,17</sup>, Guillaume Piessen, MD, PhD<sup>1,17</sup> on behalf of the AFC and FREGAT Working Group

<sup>1</sup>Department of Digestive and Oncological Surgery, University Lille, Claude Huriez University Hospital, CHU de Lille, Lille, France; <sup>2</sup>University Lille, CHU Lille, ULR 2694 – METRICS : Évaluation des technologies de santé et des pratiques médicales, Lille, France; <sup>3</sup>Department of Digestive Surgery, Haut Lévêque University Hospital, Bordeaux, France; <sup>4</sup>Department of Digestive Surgery, Pitié-Salpêtrière University Hospital, Paris, France: <sup>5</sup>Department of Surgery, Institut Gustave Roussy, Villejuif, France; <sup>6</sup>Department of Digestive Surgery, Dijon University Hospital, Dijon, France; <sup>7</sup>Department of Digestive Surgery, Charles Nicolle University Hospital, Rouen, France; <sup>8</sup>General Surgery, Department of Clinical and Experimental Sciences, University of Brescia, Brescia, Italy; <sup>9</sup>Department of Digestive Surgery, Hautepierre University Hospital, Strasbourg, France; <sup>10</sup>Department of Digestive Surgery, Brest University Hospital, Brest, France; <sup>11</sup>Department of Digestive Surgery, Tours University Hospital, Tours, France; <sup>12</sup>Department of Digestive Surgery, Pontchailloux University Hospital, Rennes, France; <sup>13</sup>Department of Digestive Surgery, Sorbonne Université, AP-HP, Hôpital Saint Antoine, Paris, France; <sup>14</sup>Department of Digestive Surgery, Limoges University Hospital, Limoges, France; <sup>15</sup>Department of Digestive Surgery, Clinique Claude Bernard RAMSAY, Ermont, France; <sup>16</sup>Department of Pathology, Lille University Hospital, Lille, France; <sup>17</sup>University Lille, CNRS, Inserm, CHU Lille, UMR9020-U1277 – CANTHER – Cancer Heterogeneity, Plasticity and Resistance to Therapies, Lille, France; <sup>18</sup>Department of Digestive Surgery, Amiens Picardie University Hospital, Amiens, France; <sup>19</sup>Department of Digestive Surgery, Angers, France; <sup>20</sup>Department of Digestive Surgery, Blois, France; <sup>21</sup>Department of Digestive Surgery, Hôpital Avicennes, Bobigny, France; <sup>22</sup>Department of Digestive Surgery, Haut-Levêque University Hospital, Bordeaux, France; <sup>23</sup>Department of Digestive Surgery, Hôpital Saint André, Bordeaux, France; <sup>24</sup>General Surgery, Department of Experimental and Clinical Sciences, University of Brescia, Brescia, Italy; <sup>25</sup>Department of Digestive Surgery, Cavale Blanche University Hospital, Brest, France; <sup>26</sup>Department of Digestive Surgery, Briey Hospital, Briey, France; <sup>27</sup>Department of Digestive Surgery, Coltea Hospital, Bucharest, Romania; <sup>28</sup>Department of Digestive Surgery, Polyclinique Synergia, Carpentras, France; <sup>29</sup>Department of Digestive Surgery, Beaujon University Hospital, Clichy, France; <sup>30</sup>Department of Digestive Surgery, Louis Mourier University Hospital, Colombes, France; <sup>31</sup>Department of Digestive Surgery, Dijon, France; <sup>32</sup>Department of Digestive Surgery, Douai Hospital, France; <sup>33</sup>Department of Digestive Surgery, Clinique de la Ligne Bleue, Epinal, France;

Clément Dubois and Frederiek Nuytens contributed equally to this work.

<sup>34</sup>Department of Digestive Surgery, Forez Hospital, Feurs, France; <sup>35</sup>Department of Digestive Surgery, Furiani Polyclinic, Furiani, France; <sup>36</sup>Department of Digestive Surgery, Grenoble, France; <sup>37</sup>Department of Digestive Surgery, Saint Odile Clinic, Haguenau, France; <sup>38</sup>Department of Digestive Surgery, Fondation Hôtel-Dieu Hospital, Le Creusot, France; <sup>39</sup>Department of Oncological and Digestive Surgery, Lille University Hospital, Lille, France; <sup>40</sup>Department of Digestive and Endocrine Surgery, Lille University Hospital, Lille, France; <sup>41</sup>Department of Digestive Surgery, Lille, Oscar Lambret Center, Lille, France; <sup>42</sup>Department of Digestive Surgery, Limoges, France; <sup>43</sup>Department of Digestive Surgery, Lyon-Sud Hospital, Lyon, France; <sup>44</sup>Department of Digestive Surgery, Croix-Rousse Hospital, Lyon, France; <sup>45</sup>Department of Digestive Surgery, Centre Leon Berard, Lyon, France; <sup>46</sup>Department of Digestive Surgery, Infirmerie Protestante, Lyon, France; <sup>47</sup>Department of Digestive Surgery, la Timone Hospital, Marseilles, France; <sup>48</sup>Department of Digestive Surgery, Saint Joseph Hospital, Marseilles, France; <sup>49</sup>Department of Digestive Surgery, la Conception Hospital, Marseilles, France; <sup>50</sup>Department of Digestive Surgery, Mende, France; <sup>51</sup>Department of Digestive Surgery, Legouest Hospital, Metz, France; <sup>52</sup>Department of Digestive Surgery, Saint François Clinic, Montluçon, France; <sup>53</sup>Department of Digestive Surgery, Val D'Aurelle Hospital, Montpellier, France; <sup>54</sup>Department of Digestive Surgery, Nancy University Hospital, Nancy, France; <sup>55</sup>Department of Digestive Surgery, Nantes University Hospital, Nantes, France; <sup>56</sup>Department of Digestive Surgery, Nice, France; <sup>57</sup>Department of Digestive Surgery, Pitié-Salpétrière University Hospital, Paris, France; <sup>58</sup>Department of Digestive and Visceral Surgery, Pitié-Salpétrière University Hospital, Paris, France; <sup>59</sup>Department of Digestive Surgery, Kremlin-Bicètre Hospital, Le Kremlin-Bicêtre, France; <sup>60</sup>Department of Digestive Surgery, Saint-Antoine Hospital, Paris, France; <sup>61</sup>Department of Digestive Surgery, Saint Louis Hospital, Paris, France; <sup>62</sup>Department of Digestive Surgery, Tenon Hospital, Paris, France; <sup>63</sup>Department of Digestive Surgery, Institut Curie, Paris, France; <sup>64</sup>Department of Digestive Surgery, Institut Mutualiste Montsouris, Paris, France; <sup>65</sup>Department of Digestive Surgery, Poitiers, France; <sup>66</sup>Department of Digestive Surgery, Reims, France; <sup>67</sup>Department of Digestive Surgery, Rennes, France; <sup>68</sup>Department of Digestive Surgery, Rouen, France; <sup>69</sup>Department of Digestive Surgery, Saint-Etienne, France; <sup>70</sup>Department of Digestive Surgery, New Civil Hospital, Strasbourg, France; <sup>71</sup>Department of Digestive Surgery, Strasbourg, CHU, Hôpital de Hautepierre, Strasbourg, France; <sup>72</sup>Department of Digestive Surgery, Foch Hospital, Suresnes, France; <sup>73</sup>Department of Digestive Surgery, Purpan Hospital, Toulouse, France; <sup>74</sup>Department of Digestive Surgery, Tours, France; <sup>75</sup>Department of Digestive Surgery, Trieste, Italy; <sup>76</sup>Department of Digestive Surgery, Valence, France; <sup>77</sup>Department of Digestive Surgery, Gustave-Roussy Institute, Villejuif, France

# ABSTRACT

**Background.** The optimal surgical procedure for duodenal gastrointestinal stromal tumors (D-GISTs) remains poorly defined. Pancreaticoduodenectomy (PD) allows for a wide resection but is associated with a high morbidity rate.

**Objectives.** The aim of this study was to compare the short- and long-term outcomes of PD versus limited resection (LR) for D-GISTs and to evaluate the role of tumor enucleation (EN).

**Methods.** In this retrospective European multicenter cohort study, 100 patients who underwent resection for D-GIST between 2001 and 2013 were compared between PD (n = 19) and LR (n = 81). LR included segmental duodenectomy (n = 47), wedge resection (n = 21), or EN (n = 13). The primary objective was to evaluate disease-free survival (DFS) between the groups, while the secondary objectives were to analyze the overall morbidity and mortality, radicality of resection, and 5-year overall survival (OS) and recurrence rates between groups. Furthermore, the short- and long-term outcomes of EN were evaluated.

**Results.** Baseline characteristics were comparable between the PD and LR groups, except for a more frequent

D2 tumor location in the PD group (68.3% vs. 29.6%; p = 0.016). Postoperative morbidity was higher after PD (68.4% vs. 23.5%; p < 0.001). OS (p = 0.70) and DFS (p = 0.64) were comparable after adjustment for D2 location and adjuvant therapy rate. EN was performed more in American Society of Anesthesiologists (ASA) stage III/IV patients with tumors < 5 cm and was associated with a 5-year OS rate of 84.6%, without any disease recurrences. **Conclusions.** For D-GISTs, LR should be the procedure of choice due to lower morbidity and similar oncological outcomes compared with PD. In selected patients, EN appears to be associated with equivalent short- and long-term outcomes. Based on these results, a surgical treatment algorithm is proposed.

Gastrointestinal stromal tumor (GIST) is the most common type of mesenchymal tumor of the gastrointestinal tract and is most frequently located in the stomach (60–70%) and small intestine (20–30%). Duodenal GISTs (D-GISTs) account for 5% of all GISTs.<sup>1</sup> The standard treatment for localized GIST is complete R0 surgical excision, avoiding tumor rupture.<sup>2–4</sup> Because of their

anatomic location, the optimal surgical procedure for D-GISTs remains poorly defined. Although limited resection (LR) such as a segmental duodenal resection, atypical lateral (wedge) resection, or enucleation (EN) may be technically feasible, anatomical considerations may render LR more difficult to perform due to the proximity of other critical structures, including the duodenal papilla and pancreas. As such, pancreaticoduodenectomy (PD) may be warranted in a subset of patients;<sup>3-10</sup> however, PD is associated with significant short- and long-term morbidity, especially for this specific indication.<sup>11–13</sup> In recent literature, several authors compared PD with LR for D-GISTs.<sup>14–24</sup> The results of these studies are in favor of offering an LR when possible rather than PD because of lower postoperative morbidity and equivalent long-term oncological results. However, in those small-size studies, the postoperative follow-up is often limited and patients are usually not comparable regarding comorbidities, tumor size, and tumor location within the duodenum.<sup>14–17</sup> To our knowledge, the role of EN for D-GISTs has not yet been evaluated in other studies. In a recent multicenter study of the French EsoGastric Tumors (FREGAT) Working Group network, the oncological safety of this approach has been suggested for esophageal GISTs of limited size (< 6.5 cm) without mucosal ulceration.<sup>25</sup> Whether or not the findings of this study could also apply to D-GISTs remains to be defined.

Consequently, the aim of this study was to evaluate (1) the postoperative course and oncological outcome of LR versus PD for patients with non-metastatic D-GISTs, and (2) the feasibility and short- and long-term outcomes of EN for D-GISTs.

# METHODS

#### Study Design and Population

In this observational cohort study, data from 1413 consecutive adult patients treated for histopathologically confirmed GIST in 61 French-speaking European centers between 2001 and 2013 were collected retrospectively through a dedicated website (http://www.chirurgie-viscera le.org). Data on patient demographics, clinical presentation, initial work-up, operative technique, histopathology, postoperative course, and oncological outcomes were gathered and analyzed. When missing, additional data were obtained by means of e-mail exchanges or phone calls with the collaborating centers. Patients were not included if the surgical and/or tumor data required for the analysis were missing.

Overall, 109 patients treated for a D-GIST were recorded in the database. The criteria for inclusion in this study were (1) D-GIST; (2) no distant metastasis; (3) with surgical treatment; and (4) no other progressive malignancy at the time of surgery. An overview of the study population and reasons for exclusion are shown in Fig. 1. Among the remaining population (n = 100), patients who underwent EN (n = 13), wedge resection (n = 21), and segmental resection (n = 47) were grouped together in an LR group (n = 81) and compared as a whole with the PD group (n = 19). Additionally, outcomes from the subgroup of patients who underwent EN were analyzed separately. This study was registered at researchregistry.com (UIN6164). Ethics Committee approval was not required due to the retrospective observational nature of this study.

#### Pretreatment Work-Up and Surgical Approach

Pretreatment investigations were standardized according to the guidelines of the European Society for Medical Oncology (ESMO) that were applicable at the time of surgical treatment.<sup>4</sup> The surgical approach (LR/PD) was determined according to the size and location of the tumor and/or involvement of the head of the pancreas, at the discretion of the operating surgeon. Reconstruction after PD was achieved using a pancreaticojejunostomy or pancreaticogastrostomy according to surgeon and center preferences. LR included segmental resection of the duodenum, local wedge resection, or EN (excision of the tumor without mucosal disruption). In general, following LR, the duodenal defect was closed primarily whenever possible or by means of a Roux-en-Y duodenojejunostomy if primary closure was not feasible. Margin status was ascertained based on the final pathologic assessment. Perioperative care was based on the usual practices of the individual surgeons.

# Postoperative Course

Postoperative morbidity was categorized into surgical complications (including anastomotic leak, intra-abdominal abscess, surgical site infection and bleeding necessitating blood transfusions, reoperation, and others) and medical complications (including urologic, pulmonary, cardiovascular, thromboembolic, neurologic complications, and others). The severity of complications was assessed according to the Clavien–Dindo (CD) classification and only grade II or higher complications were considered for the analysis of overall morbidity. Grade III/ IV complications (severe complications) were assessed likewise.<sup>13</sup>

# Histopathologic Analysis

The final diagnosis of GIST was based on histologic and immunohistochemical analysis, with selective use of



FIG. 1 Study population. GIST gastrointestinal stromal tumor

mutational analysis in doubtful cases. Tumor histopathology was studied to determine size (cm) and mitotic index (mitoses/5 mm<sup>2</sup>). Risk of recurrence was evaluated according to the modified National Institutes of Health Criteria, the Armed Forces Institute of Pathology (AFIP), and the TNM stage from the 7th edition of the International Union Against Cancer.<sup>12,26,27</sup> Resections were designated R0 if complete removal was obtained, both macroscopically and microscopically, and R1 in the case of a microscopically positive resection margin.

# Perioperative Treatment and Follow-Up

All cases were discussed during multidisciplinary team meetings. Decisions regarding the need for neoadjuvant and adjuvant tyrosine kinase inhibitors (TKIs) were made at the discretion of the local multidisciplinary teams in accordance with national and European guidelines.<sup>3,4</sup> Regular follow-up with clinical examination and computed tomography (CT) scan was recommended for at least 5 years, with frequency depending on the recurrence risk according to the European guidelines.<sup>4</sup> Disease recurrence was categorized as locoregional or distant. Mixed

recurrences included concomitant locoregional and distant relapses.

#### Study Endpoints

The primary objective of this study was to compare the disease-free survival (DFS) between LR and PD, while the secondary objectives were to analyze the overall morbidity and mortality, radicality of resection, 5-year overall survival (OS), and 5-year recurrence in the LR and PD groups. An additional secondary objective was to evaluate the oncological results of the EN subgroup, as well as the related overall postoperative morbidity and mortality.

# Statistical Analysis

Categorical variables were expressed as frequencies and percentages, and continuous variables were expressed as median and range. The distribution of the continuous variables was verified graphically and by means of the Shapiro–Wilk test. A Chi-square test or Fisher's exact test was used to compare categorical data between groups, a Cochran–Armitage trend test for ordered categorical

<b>TABLE 1</b> Baseline characteristics an	d therapeutic data
--	--------------------

	Overall $[n = 100]$	LR group $[n = 81]$	PD group $[n = 19]$	p value	EN group $[n = 13]$
Age < 60 years	52 (52.0)	40 (49.3)	12 (63.2)	0.28	4 (30.8)
Male	55 (55.0)	44 (54.3)	11 (57.9)	0.78	6 (46.2)
ASA score $> 2$	18 (18.0)	16 (19.8)	2 (10.5)	0.51	5 (38.5)
Tumor location				0.016	
D1	13 (13.0)	12 (14.8)	1 (5.3)		4 (30.8)
D2	37 (37.0)	24 (29.6)	13 (68.3)		6 (46.2)
D3	32 (32.0)	28 (34.6)	4 (21.1)		2 (15.4)
D4	18 (18.0)	17 (21.0)	1 (5.3)		1 (7.7)
Cause of diagnosis				0.78	
Symptomatic	71 (71.0)	58 (71.6)	13 (68.4)		3 (23.1)
Incidental	29 (29.0)	23 (28.4)	6 (31.6)		10 (76.9)
Biopsy	55 (55.0)	39 (48.1)	16 (84.2)	0.005	4 (30.8)
Mucosal ulceration on diagnostic endoscopy <sup>a</sup>	34/60 (56.7)	28/47 (59.6)	6/13 (46.2)	0.39	7/8 (87.5)
Neoadjuvant therapy	6 (6.0)	4 (4.9)	2 (10.5)	NA	0 (0.0)
Elective surgery	92 (92.0)	75 (92.6)	17 (89.5)	0.65	12 (92.3)
Laparotomy	95 (95.0)	76 (93.8)	19 (100.0)	0.59	11 (84.6)
Tumor breach or perforation	3 (3.0)	2 (2.5)	1 (5.3)	0.47	1 (7.7)
Median operating time, min (median [range])	224 [60-502]	190 [60-480]	350 [190-502]	< 0.001	180 [90–240]
Estimate blood loss, mL (median [range])	50 [0-1800]	0 [0-1600]	100 [0-1800]	0.050	0 [0–0]
Intraoperative transfusion	5 (5.0)	2 (2.5)	3 (15.8)	0.046	0 (0.0)
Adjuvant therapy	31 (31.0)	21 (25.9)	10 (52.6)	0.023	0 (0.0)

Data are expressed as n (%) unless otherwise specified

LR limited resection, PD pancreaticoduodenectomy, ASA American Society of Anesthesiologists, NA not applicable, EN enucleation

<sup>a</sup>Missing data in 10 patients, among 70 patients assessed by endoscopy

variables, and a Mann–Whitney test was used for nongaussian continuous variables. Follow-up data included the date of the most recent follow-up or 11 October 2016, the censoring date. The 3- and 5-year OS and DFS were estimated using the Kaplan–Meier method. The Cox proportional hazard regression model was performed to estimate the hazard ratio (HR) and its 95% confidence interval (CI) for PD versus LR group comparison. The DFS analysis was adjusted for D2 location and adjuvant therapy in the PD versus LR group comparison. All statistical analyses were performed using SAS software (version 9.4; SAS Institute, Cary, NC, USA) and a p value < 0.05 was considered statistically significant.

# RESULTS

# Pancreaticoduodenectomy (PD) Versus Limited Resection (LR)

*Baseline Demographics and Pre-Therapeutic Data* An overview of baseline demographics and pre-therapeutic data is illustrated in Table 1. Median age was 60 years

(range 28–107), and overall median follow-up after surgery was 63.2 months (interquartile range 40.6-86.5). Demographic characteristics and the American Society of Anesthesiologists (ASA) score did not differ significantly between the PD and LR groups. D-GISTs were discovered incidentally in 29% of cases, either by abdominal CT (n = 24) or by means of an endoscopy (n = 5). In symptomatic patients, the most common presenting symptoms were anemia (n = 48), gastrointestinal (most often externalized) bleeding (n = 31), abdominal pain (n = 24), asthenia (n = 23), weight loss (n = 12), anorexia (n = 10), and presence of an abdominal mass (n = 9); none presented with jaundice. Diagnostic modalities and the nature of presenting symptoms did not differ significantly between both groups (p = 0.39). The tumor location was significantly different between groups (p = 0.016); in the PD group, the tumor was most frequently located at the second part of the duodenum (68.3%), whereas in the LR group, the two most frequent locations were the third (34.6%) and second parts of the duodenum (29.6%). A diagnostic biopsy was performed in endoscopically 55% of cases. either (n = 50),

radiologically (n = 4), or surgically (n = 1). The number of patients who underwent a preoperative biopsy was significantly higher in the PD group (84.2% vs. 48.1%; p = 0.005). Among the 70 patients who underwent endoscopy as part of the preoperative assessment, data regarding endoscopic findings were available in 60 patients; mucosal ulceration was noted in 56.7% of these cases, with no significant difference between the PD and LR groups (46.2% vs. 59.6%; p = 0.39).

Therapeutic Data Details considering therapeutic data are shown in Table 1. Six patients received neoadjuvant TKI treatment, of whom two were included in the PD group. The median duration of neoadjuvant treatment was 8 months (1–18). Re-evaluation after neoadjuvant treatment demonstrated a partial response in three cases, stable disease in two cases, and disease progression in one case. Only five patients in the LR group underwent laparoscopic surgery, of whom two were converted to laparotomy. In the LR group, 47 patients underwent segmental resection (58%), 21 patients had a wedge resection (25.9%), and 13 D-GISTs were removed by EN (16%). Tumor breach was identified in two patients during surgical exploration. In one patient in the LR group, tumor perforation occurred intraoperatively during segmental duodenectomy. The median operating time was significantly longer in the PD group than in the LR group (350 min [190-502] vs. 190 min [60-480]; p < 0.001].Blood loss tended to be higher in the PD group (p = 0.050), with a required transfusion rate that was six times higher (p = 0.046) compared with the LR group. Intraoperatively, resection was considered curative in all patients. Adjuvant TKI therapy was more frequently prescribed in the PD group (52.6% vs. 25.9%; p = 0.023).

*Postoperative Outcomes* The in-hospital and 30-day mortality rate was 0% in both groups, while the 90-day mortality rate was 1% due to one patient dying of pneumonia, which occurred after discharge from the hospital on day 38. As shown in Table 2, the overall morbidity risk was significantly higher in the PD group (68.4% vs. 23.5%; p < 0.001). This increased complication risk in the PD group included both surgical (p < 0.001) and medical complications (p = 0.047). Patients in the PD group were affected by higher rates of reoperation (26.3% vs. 3.7%; p = 0.006) and severe complications (CD grade > 2; 26.3% vs. 6.2%; p = 0.020). The median length of stay was also significantly prolonged in the PD group (23 days [5–103] vs. 13 days [5–50]; p < 0.001).

*Histopathology Data* Details considering the histopathological data can be found in Table 2. Only four patients were found to have a microscopic incomplete

resection (R1), all of whom were in the LR group; two patients underwent an EN and two patients had a segmental resection. The number of R1 resections was too low to conclude any statistically significant difference between groups. In the overall population, the median tumor size was 4 cm (1-28), with a median mitotic index of 2 (0-50). In the PD group, tumor size, as well as the mitotic index, were not significantly higher compared with the LR group. There were no statistically significant differences in Joensuu score or modified AFIP scores, or in pTNM classification, between both groups (p = 0.21, p = 0.41, and p = 0.39, respectively). Molecular analysis of the resected specimen was performed in 40 patients; no mutation was detected in 9 patients, 29 patients had a KIT mutation (exon 11, 22 patients; exon 9, 6 patients; exon 13, 1 patient), and two patients had a mutation in platelet-derived growth factor- $\alpha$  (PDGFR $\alpha$ ; exon 12, 1 patient; exon 18, 1 patient).

Long-Term **Oncological** Results The long-term oncological results are summarized in Table 3. OS rates between the LR and PD groups were comparable, with a 3and 5-year OS rate of 95% versus 89% and 85% versus 89%, respectively (HR 1.28, 95% CI 0.36–4.54; p = 0.70) (Fig. 2a). The DFS rates of the LR and PD groups at 3 and 5 years were 95% versus 89% and 85% versus 89%, respectively. There was no significant difference in DFS (HR 1.24, 95% CI 0.47–3.28; p = 0.67] between the two groups, even after adjustment for location and adjuvant therapy (adjusted HR [aHR] 1.30, 95% CI 0.43-3.98; adjusted p = 0.64) (Fig. 2b). Disease recurrence occurred in 21 patients—17 patients in the LR group and 4 patients in the PD group. In the majority of cases, recurrence presented as metastatic disease (18/21, 86% of disease recurrence). Recurrence mode did not differ significantly between both groups (p = 0.61).

# Enucleation Subgroup

Of the 100 patients included in the study, 13 underwent EN. An overview of the baseline and treatment characteristics of these patients is shown in Table 1. None of the patients in the EN group received neoadjuvant treatment or adjuvant therapy. Tumor breach was noted during surgical exploration in one patient, but no intraoperative tumor rupture occurred. Global morbidity and surgical and medical complication rates in the EN group were comparable with the general LR group (Table 2). One case of postoperative leak occurred, which was treated conservatively. As illustrated in Table 2, R1 resection was found in 2 of 13 patients in the EN group. In both of these patients, mucosal ulceration was noted during preoperative endoscopy. In the EN group, only two patients had a lesion > 5 cm in

TABLE 2	Postoperative	outcome	and l	histopathology	data
---------	---------------	---------	-------	----------------	------

	Overall $[n = 100]$	LR group $[n = 81]$	PD group $[n = 19]$	p value	EN group [ <i>n</i> = 13]
Postoperative outcome					
Global morbidity	32 (32)	19 (23.5)	13 (68.4)	< 0.001	3 (23.1)
Surgical complications	22 (22)	11 (13.6)	11 (57.9)	< 0.001	2 (15.4)
Leak	14 (14)	6 (7.4)	8 (42.1)	0.001	1 (7.7)
Deep abscess	3 (3)	1 (1.2)	2 (10.5)	NA	
Bleeding	4 (4)	2 (2.5)	2 (10.5)	NA	
Medical complications	19 (19)	12 (14.8)	7 (36.8)	0.047	1 (7.7)
Urinary	2 (2)	0	2 (10.5)	NA	
Pulmonary	4 (4)	4 (4.9)	0	NA	
Cardiac failure	2 (2)	2 (2.5)	0	NA	
Thromboembolic	2 (2)	1 (1.2)	1 (5.3)	NA	
Neurologic	2 (2)	2 (2.5)	0	NA	
Reoperation	8 (8)	3 (3.7)	5 (26.3)	0.006	0 (0.0)
Clavien–Dindo grade $> 2$	10 (10)	5 (6.2)	5 (26.3)	0.020	0 (0.0)
Length of hospital stay, days (median [range])	13 [5-103]	13 [5-50]	23 [5-103]	< 0.001	9 [5–19]
Histopathology data					
Resection rate				NA	
R0	96 (96.0)	77 (95.1)	19 (100.0)		11 (84.6)
R1	4 (4.0)	4 (4.9)	0 (0.0)		2 (15.4)
Tumor size, cm				0.16	
$\leq 2$	8 (8.0)	7 (8.6)	1 (5.3)		3 (23.1)
2–5	52 (52.0)	45 (55.6)	7 (36.8)		8 (61.5)
5–10	31 (31.0)	22 (27.2)	9 (47.4)		2 (15.4)
> 10	9 (9.0)	7 (8.6)	2 (10.5)		0 (0.0)
Mitotic index, $n(/5 \text{ mm}^2)$				0.63	
<u>≤</u> 5	78 (78.0)	63 (77.8)	15 (78.9)		13 (100.0)
6–10	11 (11.0)	8 (9.9)	3 (15.8)		0 (0.0)
> 10	11 (11.0)	10 (12.3)	1 (5.3)		0 (0.0)
Joensuu risk of recurrence				0.21	
Very low risk	10 (10.0)	8 (9.9)	2 (10.5)		3 (23.1)
Low risk	47 (47.0)	41 (50.6)	6 (31.6)		8 (61.5)
Intermediate risk	0 (0.0)	0 (0.0)	0 (0.0)		0 (0.0)
High risk	43 (43.0)	32 (39.5)	11 (57.9)		2 (15.4)
Modified AFIP recurrence risk				0.41	
Very low risk	10	8 (9.9)	2 (10.5)		3 (23.1)
Low risk	48	42 (51.9)	6 (31.6)		8 (61.5)
Intermediate risk	17	11 (13.6)	6 (31.6)		2 (15.4)
High risk	25	20 (24.7)	5 (26.3)		0 (0.0)
pTNM				0.39*	
Ι	58	49 (60.5)	8 (42.1)		11 (84.6)
II	17	11 (13.6)	6 (31.6)		2 (15.4)
IIIA	4	3 (3.7)	1 (5.3)		0 (0.0)
IIIB	21	18 (22.2)	4 (21.1)		0 (0.0)

Data are expressed as n (%) unless otherwise specified

LR limited resection, PD pancreaticoduodenectomy, AFIP Armed Forces Institute of Pathology, NA not applicable, EN enucleation

\*Tested after pooled IIIA and IIIB

 TABLE 3 Long-term

 oncological results of local

 resection versus

 pancreaticoduodenectomy

	Non-adjusted HR (95% CI)	p value	Adjusted <sup>a</sup> HR (95% CI)	p value
Overall survival	1.28 (0.36-4.54)	0.70	NA	NA
Disease-free survival	1.24 (0.47-3.28)	0.67	1.30 (0.43-3.98)	0.64
Disease recurrence	1.33 (0.45–3.97)	0.61	NA	NA

HRs are expressed for the pancreaticoduodenectomy group, with the limited resection group as reference *HR* hazard ratio, *CI* confidence interval, *NA* not applicable due to the low number of events <sup>a</sup>Adjusted for D2 localization and adjuvant therapy



FIG. 2 Kaplan-Meier curve of the long-term oncological results of LR versus PD. a Overall survival; b disease-free survival. LR limited resection, PD pancreaticoduodenectomy, HR hazard ratio, CI confidence interval, adj. adjusted

diameter. Lesions in all other patients were  $\leq 3.5$  cm in diameter. According to both the Joensuu and AFIP classifications, 84.6% of EN patients were classified as having either low or very low risk of recurrence. Three deaths occurred in the EN group—two during the first year and one after 5 years; however, none were disease-related (pulmonary disease, cardiac insufficiency, and cerebrovascular incident). As such, the survival rate in the EN group was similar after 3 and 5 years (84.6%). Finally, in the EN group, no disease recurrence was noted.

# DISCUSSION

The surgical management of duodenal GIST is considered as complex due to the anatomical constraints related to the organ. Several surgical procedures have been described in this context. In addition to the radical gesture

of PD, other procedures with LR have been proposed, such as segmental and wedge resection, as well as EN. However, the different indications and results of these specific techniques remain poorly understood.<sup>10,11,15–17,28–36</sup> In this multicentric retrospective study, we observed that LR, when feasible, provides equivalent oncological results (OS, DFS, and risk of recurrence) compared with PD. Moreover, LR is associated with a reduced operating time and transfusion rate, as well as reduced morbidity and length of hospital stay. It should be noted that all R1 resections occurred in the LR group (n = 4). However, any statistical analysis was not applicable due to these low numbers. In our series, the tumor was most commonly located at the level of the second (37%) and third part (32%) of the duodenum, which is in accordance with other reports.<sup>16,17,35,37,38</sup> The majority of D-GISTs in our study, as well as in most other reports, had a low or very low risk of recurrence according to Joensuu's classification (57%), suggesting a better prognosis of GIST located at the level of the duodenum compared with the small intestine.<sup>16</sup> However, a case-control study by Zhang et al. showed that, after pairing, the prognosis between these two locations seems to be similar.<sup>14</sup> Nineteen percent of patients in this study underwent PD. This rate is among the lowest when compared with other reports (10–40%),<sup>18–25</sup> which might suggest a more conservative attitude of European surgeons in general and French surgeons in particular who participated in this study.

Regarding the pre-therapeutic data, the tumor was more frequently located in the second part of the duodenum in the PD group compared with the LR group. Indeed, the presence of the biliopancreatic junction and the close relationship between the pancreatic head and the duodenum at this level renders LR technically difficult to perform, except in cases of an antimesenteric location of the tumor. As expected, the preoperative biopsy rate was higher in the PD group due to the high morbidity rate and risk associated with the surgical procedure. As would be expected, the inherently more invasive technique of PD was associated with a significantly increased operating time and transfusion rate. In accordance with findings in other reports, PD was associated with a higher rate of postoperative complications, both medical and surgical in nature.<sup>10,11,16–24,28–36</sup> This in turn resulted in an extended length of hospital stay in the PD group. With regard to long-term oncological outcomes, LR was associated with an equivalent OS and adjusted DFS, without any measurable statistically significant difference in the R0 resection rate, or risk of recurrence between the two groups, and despite a higher rate of adjuvant TKI therapy in the PD group (p = 0.023). Moreover, since no differences were found in the mode of disease recurrence between PD and LR, the oncological safety of an LR technique for D-GISTs is demonstrated. However, it should be noted that despite the fact that this study can benefit from a relatively long median follow-up (63.2 months), it is known that a significant amount of disease recurrences (10-30%) are discovered more than 5 years after the initial surgical treatment.<sup>22</sup>

An overview of the literature comparing outcomes after PD or LR for D-GISTs is shown in Table 4. Regarding overall morbidity and survival, other studies including large numbers of patients reported results that are comparable with the findings in our study.<sup>16,17,35</sup> The majority of the authors report equivalent or inferior long-term oncologic outcomes in the PD group.<sup>14</sup> In these studies, the surgical technique could not be identified as an independent prognostic factor.<sup>16,17,30</sup> A recent meta-analysis by Shen et al. showed that PD was associated with a worse long-term prognosis.<sup>39</sup> However, Shen et al. emphasize that

this should be considered as a correlation and not as a causal relationship since patients who underwent PD were selected based on infaust prognostic factors (D2 localization, high-degree of mitosis, and high-risk classifications).

The strength of this study is that it comprises one of the largest study populations in the literature. Furthermore, we believe that the quality of this study lies in the fact that the DFS analysis was adjusted for D2 location and adjuvant therapy in the PD versus LR group comparison. As such, a potential selection bias of the surgical technique at baseline was maximally corrected. Despite the fact that this study is among the largest in the literature, an analysis by means of propensity score matching was not feasible because of the lack of statistical power.

EN has only rarely been discussed in the surgical management of D-GIST.<sup>40</sup> Our results suggest that EN could offer an interesting alternative for well-defined D-GISTs. In this study, EN was mainly proposed in elder and frail patients, probably because the surgical strategy was tailored in terms of the estimated medical risk. The majority of enucleated tumors were located in the proximal duodenum (46.2% D2; 30.8% D1) and could therefore offer a valid alternative for PD in cases of a tumor diameter < 5 cm. EN was associated with a low grade of morbidity without the need for surgical re-intervention, and a reduced length of hospital stay. Moreover, with no disease recurrence or disease-related deaths in the EN group, this technique appears to result in acceptable long-term outcomes. These findings have been incorporated in a surgical treatment algorithm for D-GISTs, as proposed in Fig. 3. A major French retrospective analysis and literature review by the FREGAT Working Group has demonstrated that EN is safe and feasible in esophageal GIST with a tumor diameter  $\leq 6.5$  cm and in the absence of mucosal ulceration on endoscopy.<sup>25</sup> In this study, in the two patients in the EN group who had an R1 resection, mucosal ulceration was identified during preoperative endoscopic evaluation. Mucosal ulceration has been described as a negative prognostic factor. In their study on esophageal GIST, Blum et al. identified two recurrences out of four ENs,<sup>41</sup> with both patients having signs of ulcerations on preoperative endoscopy. Although a low level of evidence is provided, mucosal ulceration is considered as a contraindication for EN in esophageal GIST.<sup>25</sup>

The main limitation of this study, besides its retrospective nature, is the fact that the rate of mutation analysis in this study is quite low at 40%. Nevertheless, this rate is still higher than the mutation analysis rate in other studies reporting on outcomes from comparable inclusion periods, and has to be seen within the context of an era when mutation analysis was not yet recommended in a standardized fashion.<sup>42</sup> A second limitation of this study is that although D2 location and tumor size are presumed to be the

Studies (year)	Time of inclusion	No. of patients	LR/ PD	Morbidity (LR/PD) (%)	Follow-up (months) (median [range])	OS (%)
Dubois et al. (current study)	2001–2013	100	81/19	23.5/68.4%	63.2 [40.6-86.5]	89/89 (5 years)
Zhang et al. <sup>19</sup>	1999–2016	52	37/14	$10.8/21.4 \text{ (CD} \ge 2)$	36 [6–138]	89.1 (5 years)
Lee et al. <sup>22</sup>	2000–2017	118	73/45	24.5/24.4 (in- hospital)	46 [3.9–167.9]	91.9/96.2 (5 years)
<b>•</b> • • • 18	2010 2016	27	22/0	1.9/15.6 (late)		37.4
Liu et al. <sup>10</sup>	2010-2016	37	22/9	NA	NA	NA
Shi et al. <sup>25</sup>	2005-2015	61	45/16	33.3/56.3	NA	NA
Lee et al. <sup>37</sup>	1994-2014	60	37/23	24/70	38 [21–72]	72/76 (5 years)
Chen et al. <sup>21</sup>	2005–2016	64	41/23	31.7/69.6	36 [5–102]	80.5/31.7 (5 years)
Sugase et al. <sup>20</sup>	1990-2014	25	16/9	31/33	NA	89/45 (5 years)
Crown et al. <sup>24</sup>	2000-2015	15	7/8	14/62	NA	NA
Chung et al. <sup>28</sup>	2001-2014	21	21/0	14.3	52 [5-125]	NA
Duffaud et al. <sup>17</sup>	1993-2002	105	82/23	18/26	36 [2–250]	86.5 (5 years)
Beham et al. <sup>29</sup>	NA	13	8/5	NA	NA	66 (months)
Zhou et al. <sup>30</sup>	2006-2012	48	34/14	11.8/35.7	36 [0-81]	NA
Hoeppner et al. <sup>31</sup>	2002-2011	9	8/1	NA	45 [6–111]	NA
Bourgouin et al. <sup>32</sup>	2000-2012	17	11/6	29/50	34	NA
Yang et al. <sup>33</sup>	1999–2011	22	13/9	15.4/88.9	67.5 [3–118]	80.6 (5 years)
Liang et al. <sup>34</sup>	1998-2006	28	18/10	NA	61 [23–164]	64.5 (months)
Colombo et al.35	2000-2011	84	56/28	9/36	42 [2–135]	89 (at 5 years)
Johnston et al. <sup>16</sup>	1994–2011	96	58/38	29.3/57.6	22 [4-81]	82 (5 years)
Kamath et al. <sup>36</sup>	1999–2011	41	30/11	20	18 [0–144]	74 (5 years)
Tien et al. <sup>11</sup>	2001-2008	25	16/9	12.5/44	18 [9–92]	NA
Goh et al. <sup>10</sup>	1992-2006	15	7/7	NA/43	42 [2–174]	NA

LR limited resection, PD pancreaticoduodenectomy, OS overall survival, CD Clavien–Dindo, NA not applicable, D-GIST duodenal gastrointestinal stromal tumor

main determinants for the decision of PD versus LR as the optimal surgical strategy, more details regarding the anatomical relationship of the tumor (relative to papilla, common bile duct, etc.) could not be retrieved from the database. Additionally, the authors acknowledge that although the rate of neoadjuvant TKI therapy is quite low at 6%, no further details regarding this patient group are known. However, to date, no randomized controlled trials have demonstrated any effect of neoadjuvant TKI therapy on the indication of surgical approach.<sup>43,44</sup> Moreover, since more patients in the PD group received neoadjuvant therapy, excluding this patient group would only further reinforce the results and message of this study.

# CONCLUSIONS

The results of this study suggest that in the surgical management of D-GIST, LR is preferred over PD whenever feasible considering the anatomical relations of the tumor. LR for D-GIST is associated with a reduced morbidity rate and provides long-term oncological results that are at least equivalent to PD. For D-GIST without endoscopic signs of mucosal ulceration, EN could serve as a potential alternative for PD, segmental resection, or wedge resection. However, future studies regarding this subject, including long-term results and larger patient cohorts, are required. Based on the findings of this study, we propose a surgical treatment algorithm for D-GISTs.



FIG. 3 Proposed algorithm for the treatment of resectable, non-metastatic duodenal GISTs. GIST gastrointestinal stromal tumor, ASA American Society of Anesthesiologists

**ACKNOWLEDGEMENTS** Participating Investigators (Collected Data) Department of Digestive Surgery, Amiens Picardie University Hospital, France: Francois Browet, MD; Charles Sabbagh, MD, PhD; Jean-Marc Regimbeau, MD. Department of Digestive Surgery, Angers, France: Emilie Lermite, MD PhD; Antoine Hamy, MD, PhD, Department of Digestive Surgery, Blois, France: Kevin Kraft, MD. Department of Digestive Surgery, Hôpital Avicennes, Bobigny, France: Richard Douard, MD, PhD; Philippe Wind, MD. Department of Digestive Surgery, Haut-Levêque University Hospital, Bordeaux, France: Hélène Gersen-Cherdieu, MD; Denis Collet, MD; Magalie Cabau, MD. Department of Digestive Surgery, Hôpital Saint André, Bordeaux, France: Christophe Laurent, MD PhD; Eric Rullier, MD. General Surgery, Department of Experimental and Clinical Sciences, University of Brescia, Brescia, Italy: Arianna Coniglio, MD. Department of Digestive Surgery, Cavale Blanche University Hospital, Brest, France: Charles-Henry Gancel, MD; Bogdan Badic, MD. Department of Digestive Surgery, Briey Hospital, France: Gilbert Ouedraogo, MD. Department of Digestive Surgery, Coltea Hospital, Bucharest, Romania: Mircea Beuran, MD. Department of Digestive Surgery, Polyclinique Synergia, Carpentras, France: Aude Brams, MD, PhD; Marc Kanor, MD; Christophe Louis, MD; Yves Russier, MD. Department of Digestive Surgery, Beaujon University Hospital, Clichy, France: Yves Panis, MD, PhD; Léon Maggiori, MD, PhD. Department of Digestive Surgery, Louis Mourier University Hospital, Colombes, France: Camille Caille, MD; Géraud Tuyeras, MD; Simon Msika, MD. Department of Digestive Surgery, Dijon, France: Luigi De Magistris, MD; T. Perrin, MD; Patrick Rat, MD; Pablo Ortega Deballon, MD, PhD. Department of Digestive Surgery, Douai Hospital, France: Philippe Meignie, MD. Department of Digestive Surgery, Clinique de la Ligne Bleue, Epinal, France: Jean François Bronner, MD; Jacques Moline, MD. Department of Digestive Surgery, Forez Hospital, Feurs, France: Claude Mondersert, MD. Department of Digestive Surgery, Furiani Polyclinic, France: André Caamano, MD. Department of Digestive Surgery, Grenoble, France: Catherine Arvieux, MD, PhD; Jean-Luc Faucheron, MD, PhD; Christian Letoublon, MD; Antoine Guillaud, MD. Department of Digestive Surgery, Saint Odile Clinic, Haguenau, France: Ségolène Lardenois, MD; Jean-Michel Nuss, MD. Department of Digestive Surgery, Fondation Hôtel-Dieu Hospital, Le Creusot, France: Patrick Chevillotte, MD. Department of Oncological and Digestive Surgery, Lille University Hospital, France: Edouard Vinatier, MD: Williams Tessier, MD. Department of Digestive and Endocrine Surgery, Lille University Hospital, Lille, France: Robert Caiazzo, MD, PhD; Francois Pattou, MD, PhD. Department of Digestive Surgery, Lille, Oscar Lambret Center, Lille, France: Nicolas Lamande, MD; Mehrdad Jafari, MD; Gauthier Decanter, MD. Department of Digestive Surgery, Limoges, France: François Paraf, MD. Department of Digestive Surgery, Lyon-Sud Hospital, Lyon, France: Mohamed Alyami, MD; Delphine Vaudoyer, MD; Guillaume Passot, MD, PhD; Olivier Glehen, MD, PhD. Department of Digestive Surgery, Croix-Rousse Hospital, Lyon, France: Hassan Demian, MD; Christian Ducerf, MD; Jean-Yves Mabrut, MD, PhD. Department of Digestive Surgery, Centre Leon Berard, Lyon, France: Marc Rivoire, MD, PhD. Department of Digestive Surgery, Infirmerie Protestante, Lyon, France: Vincent Garbit, MD; Thierry Leclercq, MD; Jean Loire, MD; Olivier Raspado, MD. Department of Digestive Surgery, la Timone Hospital, Marseilles, France: Rémy Le Huu Nho, MD; Mehdi Ouaissi, MD, PhD; Igor Sieleznef, MD, PhD; Bernard Sastre, MD. Department of Digestive Surgery, Saint Joseph Hospital, Marseilles, France: Bernard Pol, MD; Manuela Campanile, MD. Department of Digestive Surgery, la Conception Hospital, Marseilles, France: Sami Hamed, MD; Jean Hardwigsen, PD, PhD; Yves-Patrice Le Treut, MD. Department of Digestive Surgery, Mende, France: Gérard Carbonnel, MD. Department of Digestive Surgery, Legouest Hospital, Metz, France: Charlotte de Saint Roman, MD; M. Tréot, MD; Philippe Sockeel, MD. Department of Digestive Surgery, Saint François Clinic, Montluçon, France: Ghislain Tourreau, MD; Vasileios Baltzopoulos, MD. Department of Digestive Surgery, Montpellier, Val D'Aurelle Hospital, France: Anne Mourregot, MD; Philippe Rouanet, MD. Department of Digestive Surgery, Nancy University Hospital, France: L. Bresler, MD, PHD. Department of Digestive Surgery, Nantes University Hospital, Nantes, France: P. Senellart, MD;

Guillaume Meurette, MD, PhD; Paul Antoine Lehur, MD; Nicolas Regenet, MD. Department of Digestive Surgery, Nice, France: Vincent Casanova, MD; Anne Sophie Schneck, MD; Antonio Ianelli, MD, PhD; Jean Gugenheim, MD. Department of Digestive Surgery, Pitié-Salpétrière University Hospital, Paris, France: Mehdi Karoui, MD, PhD; Jean Christophe Vaillant, MD; Jean Baptise Bachet, MD, PhD; Laurent Hannoun, MD. Department of Digestive and Visceral Surgery, Pitié-Salpétrière University Hospital, Paris, France: Renato Lupinacci, MD; Francois Menegaux, MD, PhD; Christophe Tresallet, MD, PhD. Department of Digestive Surgery, Kremlin-Bicètre Hospital, France: Antoine Brouquet, MD, PhD; Stéphane Benoist, MD, PhD; Christophe Penna, MD, PhD. Department of Digestive Surgery, Saint-Antoine Hospital, Paris, France: Pierre Balladur, MD, PhD; Yann Parc, MD, PhD; François Paye, MD, PhD; Najim Chafai, MD; Emmanuel Tiret, MD. Department of Digestive Surgery, Saint Louis Hospital, Paris, France: Pierre Cattan, MD, PhD; Emile Sarfati, MD; Leonor Benhaim, MD. Department of Digestive Surgery, Tenon Hospital, Paris, France: F. Sista, MD; Sidney Houry, MD. Department of Digestive Surgery, Institut Curie, Paris, France: Pascale Mariani, MD. Department of Digestive Surgery, Institut Mutualiste Montsouris, Paris, France: Béatrice Ullua Severino, MD; Brice Gayet, MD; Thierry Perniceni, MD. Department of Digestive Surgery, Poitiers, France: Jean-Pierre Richer, MD; Jean Pierre Faure, MD; Jean Louis Kraimps, MD. Department of Digestive Surgery, Reims, France: Sophie Deguelte-Lardiere, MD; Olivier Bouche, MD, PhD; Reza Kianmanesch, MD, PhD. Department of Digestive Surgery, Rennes, France: Charles Le Clec'h, MD; Laurent Sulpice, MD; Bernard Meunier, MD; Karim Boudjema, MD. Department of Digestive Surgery, Rouen, France: Emmanuel Huet, MD; Michel Scotte, MD, PhD; Francis Michot, MD. Department of Digestive Surgery, Saint-Etienne, France: Thomas Sole, MD; Clément Costanza, MD; Charlotte Vermesch, MD; Santina Bruno, MD; Jack Porcheron, MD. Department of Digestive Surgery, New Civil Hospital, Strasbourg, France: M. Raharimanantsoa, MD; Patrick Pessaux, MD, PhD; Silvana Perretta, MD, PhD; Didier Mutter, MD. Department of Digestive Surgery, Strasbourg, CHU, Hôpital de Hautepierre, Strasbourg, France: H. Mercoli, MD; A. Klipfel, MD; E.H. Triki, MD; B. Romain, MD; S. Dragomir, MD; N. Chilintseva, MD; J.C. Olliern, MD; Serge Rohr, MD. Department of Digestive Surgery, Foch Hospital, Suresnes, France: Alexandre Rault, MD. Department of Digestive Surgery, Purpan Hospital, Toulouse, France: Charles Henri Julio, MD; Mael Chalret du Rieu, MD; Nicolas Carrere, MD, PhD; Bernard Pradère, MD. Department of Digestive Surgery, Tours, France: Perrine Senellart, MD: Baudoin Thébault, MD. Department of Digestive Surgery, Trieste, Italy: Nicolo De Manzini, MD. Department of Digestive Surgery, Valence, France: Christophe Henry, MD. Department of Digestive Surgery, Gustave-Roussy Institute, Villejuif, France: Sylvie Bonvalot, MD

AUTHOR CONTRIBUTIONS All authors meet the International Committee of Medical Journal Editors (ICMJE) authorship criteria. Clément Dubois: Analysis and interpretation of data, drafting the manuscript, final approval. Frederick Nuytens: Analysis and interpretation of data, drafting the manuscript, final approval. Hélène Behal: Analysis and interpretation of data, vouching for the data, revision of the manuscript, final approval. Caroline Gronnier: Acquisition of data, revision of the manuscript, final approval. Gilles Manceau: Acquisition of data, revision of the manuscript, final approval. Maxime Warlaumont: Acquisition of data, revision of the manuscript, final approval. Alain Duhamel: Analysis and interpretation of data, vouching for the data, revision of the manuscript, final approval. Quentin Denost MD: Acquisition of data, revision of the manuscript, final approval. Charles Honoré MD: Acquisition of data, revision of the manuscript, final approval. Olivier Facy: Acquisition of data, revision of the manuscript, final approval. Jean-Jacques Tuech: Acquisition of data, revision of the manuscript, final approval. Guido Tiberio: Acquisition of data, revision of the manuscript, final approval. Cécile Brigand: Acquisition of data, revision of the manuscript, final approval. Jean-Pierre Bail: Acquisition of data, revision of the manuscript, final approval. Ephrem Salame: Acquisition of data, revision of the manuscript, final approval. Bernard Meunier: Acquisition of data, revision of the manuscript, final approval. Jérémie H. Lefevre: Acquisition of data, revision of the manuscript, final approval. Muriel Mathonnet: Acquisition of data, revision of the manuscript, final approval. Florence Renaud: Acquisition of data, revision of the manuscript, final approval. Guillaume Piessen: Design of the study, decision on publication of the paper, analysis and interpretation of data, revision of the manuscript, final approval.

**FUNDING** No sources of funding were used to assist in the preparation of this study.

**DISCLOSURE** Clément Dubois, Frederiek Nuytens, Hélène Behal, Caroline Gronnier, Gilles Manceau, Maxime Warlaumont, Alain Duhamel, Quentin Denost, Charles Honoré, Olivier Facy, Jean-Jacques Tuech, Guido Tiberio, Cécile Brigand, Jean-Pierre Bail, Ephrem Salame, Bernard Meunier, Jérémie H. Lefevre, Muriel Mathonnet, Mohamed Sbai Idrissi, Florence Renaud, Guillaume Piessen have no conflicts of interest to declare.

# REFERENCES

- Joensuu H, Hohenberger P, Corless CL. Gastrointestinal stromal tumours. *Lancet*. 2013;382:973–83.
- Casali PG, Lost L, Reichardt P, Schlemmer M, Blay JY. ESMO Guidelines working group. Gastrointestinal stromal tumors: ESMO clinical recommendations for diagnosis, treatment and follow-up. *Ann Oncol.* 2009;35:739–45.
- Thésaurus National de Cancérologie Digestive. https://www.snf ge.org/sites/default/files/SNFGE/TNCD/tncd\_chap-12-gist\_2019 -11-25\_v2.pdf.
- ESMO/European Sarcoma Network Working Group. Gastrointestinal stromal tumours: ESMO Clinical Practice Guidelines for diagnosis, treatment and follow-up. *Ann Oncol.* 2014;25(suppl 3):21–6.
- Gervaz P, Huber O, Morel F. Surgical management of gastrointestinal stromal tumors. Br J Surg. 2009;96:567–78.
- Piessen G, Lefèvre JH, Cabau M, Duhamel A, Behal H, Perniceni T, et al. Laparoscopic versus open surgery for gastric gastrointestinal stromal tumors: what is the impact on postoperative outcome and oncologic results? *Ann Surg.* 2015;262:831–9.
- Woodall CE, Scoggins CR. Retroperitoneal and visceral sarcomas: issues for the general surgeons. Am Surg. 2007;73:631–5.
- De Matteo RP, Heinrich MC, El-Rifai WM, Demetri G. Clinical management of gastrointestinal stromal tumors. *Hum Pathol.* 2002;33:466–77.
- Gold JS, de Matteo RP. Combined surgical and molecular therapy. Ann Surg. 2006;244:176–84.
- Goh BK, Chow PK, Kesavan S, Yap WM, Wong WK. Outcome after surgical treatment of suspected gastrointestinal stromal tumors involving the duodenum: is limited resection appropriate? *J Surg Oncol.* 2008;97:388–91.
- Tien YW, Lee CY, Huang CC, Hu RH, Lee PH. Surgery for gastrointestinal stromal tumors of the duodenum. *Ann Surg Oncol.* 2010;17:109–14.

- Miettinen M, Lasota J. Gastrointestinal stromal tumors: pathology and prognosis at different sites. *Semin Diagn Pathol*. 2006;23:70–83.
- Clavien PA, Barkun J, de Oliveira ML, Vauthey JN, Dindo D, Schulick R, et al. The Clavien-Dindo classification of surgical complications: five-year experience. *Ann Surg.* 2009;250:187–96.
- Zhang Q, Shou C-H, Yu J-R, Wang JL, Liu XS, Yu H, et al. Prognostic characteristics of duodenal gastrointestinal stromal tumors. *Br J Surg.* 2015;102:959–64.
- Chok AY, Koh YX, Ow MY, Allen J, Goh B. A systematic review and meta-analysis comparing pancreaticoduodenectomy versus limited resection for duodenal gastrointestinal stromal tumors. *Ann Surg Oncol.* 2014;21:3429–38.
- Johnston FM, Kneuertz PJ, Cameron JL, Sanford D, Fisher S, Turley R, et al. Presentation and management of gastrointestinal stromal tumors of the duodenum: a multi-institutional analysis. *Ann Surg Oncol.* 2012;19:3351–60.
- Duffaud F, Meeus P, Bachet JB, Cassier P, Huyngh T, Boucher E, et al. Conservative surgery vs. duodeneopancreatectomy in primary duodenal gastrointestinal stromal tumours (GIST): a retrospective review of 114 patients from the French sarcoma group (FSG). *Eur J Surg Oncol.* 2014;40:1369–75.
- Liu Z, Zheng G, Liu J, Liu S, Xu G, Wang Q, et al. Clinicopathological features, surgical strategy and prognosis of duodenal gastrointestinal stromal tumors: a series of 300 patients. *BMC Cancer*. 2018;18:563.
- Zhang S, Tian Y, Chen Y, Zhang J, Zheng C, Wang C. Clinicopathological characteristics, surgical treatments and survival outcomes of patients with duodenal gastrointestinal stromal tumor. *Dig Surg.* 2019;36:206–2017.
- Sugase T, Takahashi T, Nakajima K, Hirota S, Masuzawa T, Nishida T, et al. Clinicopathological characteristics, surgery and survival outcomes of patients with duodenal gastrointestinal stromal tumors. *Digestion*. 2016;94:30–6.
- Chen P, Song T, Zhou H, Zhang T, Wu Q, Kong D, et al. Surgery for duodenal gastrointestinal stromal tumors: a single-center experience. *Dig Dis Sci.* 2017;62:3167–76.
- Lee S, Song K, Lee Y, Kim S, Hwang D, Lee J, et al. Clinicopathologic and optimal surgical treatment of duodenal gastrointestinal stromal tumor. *J Gastrointest Surg.* 2019;23:270–9.
- Shi HP, Huang ML, Wang ZQ, Zheng Y, Zhu Z, Sah B, et al. Clinicopathological and prognostic features of surgical management in duodenal gastrointestinal stromal tumors. *Dig Surg.* 2018;35:498–507.
- Crown A, Biehl TR, Rocha FG. Local resection for duodenal gastrointestinal stromal tumors. Am J Surg. 2016;211:867–70.
- Robb WB, Bruyere E, Amielh D, Vinatier E, Mabrut JY, Perniceni T, et al. Esophageal gastrointestinal stromal tumour: is tumoral enucleation a viable therapeutic option? *Ann Surg.* 2015;261:117–24.
- Joensuu H, Vehtari A, Riihimäki J, Nishida T, Steigen S, Brabec P, et al. Risk of recurrence of gastrointestinal stromal tumour after surgery: an analysis of pooled population-based cohorts. *Lancet Oncol.* 2012;13:265–74.
- International Union Against Cancer (UICC), Sobin LH, Gospodarowicz MK, Wittekind Ch. TNM classification of malignant tumors. 7th ed. New York: Wiley; 2010.
- Chung JC, Kim HC, Hur SM. Limited resections for duodenal gastrointestinal stromal tumours and their oncologic outcomes. *Surg Today*. 2016;46:110–6.
- Beham A, Schaefer IM, Cameron S, von Hammerstein K, Füzesi L, Ramasori G, et al. Duodenal GIST: a single center experience. *Int J Colorectal Dis.* 2013;28:581–90.

- Zhou B, Zhang M, Wu J, Yan S, Zhou J, Zhen S. Pancreaticoduodenectomy versus local resection in the treatment of gastrointestinal stromal tumors of the duodenum. *World J Surg Oncol.* 2013;11:196.
- Hoeppner J, Kulemann B, Marjanovic G, Bronsert P, Hopt UT. Limited resection for duodenal gastrointestinal stromal tumors: Surgical management and clinical outcome. World J Gastrointest Surg. 2013;5:16–21.
- Bourgouin S, Hornez E, Guiramand J, Barbier L, Delpero JR, Le Treut YP, et al. Duodenal gastrointestinal stromal tumors (GISTs): arguments for conservative surgery. *J Gastrointest Surg.* 2013;17:482–7.
- 33. Yang F, Jin C, Du Z, Subedi S, Jiang Y, Li J, et al. Duodenal gastrointestinal stromal tumor: clinicopathological characteristics, surgical outcomes, long term survival and predictors for adverse outcomes. Am J Surg. 2013;206:360–7.
- Liang X, Yu H, Zhu LH, Wang XF, Cai XJ. Gastrointestinal stromal tumors of the duodenum: surgical management and survival results. *World J Gastroenterol*. 2013;19:6000–10.
- Colombo C, Ronellenfitsch U, Yuxin Z, Rutkowiski P, Miceli R, Bylina E. Clinical, pathological and surgical characteristics of duodenal gastrointestinal stromal tumor and their influence on survival: a multi-center study. *Ann Surg Oncol.* 2012;19:3361–7.
- Kamath AS, Sarr MG, Nagorney DM, Que F, Farnell M, Kendrick M, et al. Gastrointestinal stromal tumour of the duodenum: single institution experience. *HPB*. 2012;14:772–6.
- Lee SY, Goh BK, Sadot E, Rajeev R, Balachandran VP, Gönen M, et al. Surgical strategy and outcomes in duodenal gastrointestinal stromal tumor. *Ann Surg Oncol.* 2017;24:202–10.
- Rutkowski P, Nowecki ZI, Michej W, Debiec-Richter M, Wozniak A, Limon J, et al. Risk criteria and prognostic factors for predicting recurrences after resection of primary gastrointestinal stromal tumor. *Ann Surg Oncol.* 2007;14:2018–27.
- 39. Shen Z, Cher P, Du N, Khadaroo P, Mao D, Gu L. Pancreaticoduodenectomy versus limited resection for duodenal gastrointestinal stromal tumors: a systematic review and metaanalysis. *BMC Surg.* 2019;19:121.
- 40. Miettinen M, Kopczynski J, Makhlouf HR, Sarlomo-Rikala M, Guyorffy H, Burke A, et al. Gastrointestinal stromal tumors, intramural leiomyomas, and leiomyosarcomas in the duodenum: a clinicopathologic, immunohistochemical, and molecular genetic study of 167 cases. *Am J Surg Pathol.* 2003;27:625–41.
- Blum MG, Bilimoria KY, Wayne JD, de Hoyos A, Talamonti M, Adley B. Surgical considerations for the management and resection of esophageal gastrointestinal stromal tumors. *Ann Thoracic Surg.* 2007;85:1717–23.
- 42. Verschoor AJ, Bovée JV, Overbeek LI, The PALGA group, Hogendoorn PC, Gelderblom H. The incidence, mutational status, risk classification and referral pattern of gastro-intestinal stromal tumours in the Netherlands: a nationwide pathology registry (PALGA) study. Virchows Archiv. 2018;472:221–9.
- 43. Rutkowski P, Gronchi A, Hohenberger P, Bonvalot S, Schöffski P, Bauer S, et al. Neoadjuvant imatinib in locally advanced gastrointestinal stromal tumors (GIST): the EORTC STBSG Experience. Ann Surg Oncol. 2013;20:2937–43.
- 44. Andtbacka R, Ng C, Scaife C, Cormier J, Hunt K, Pisters P, et al. Surgical resection of gastrointestinal stomal tumors after treatment with imatinib. *Ann Surg Oncol.* 2006;14:14–24.

**Publisher's Note** Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.