Initial Experience of Consecutive Robotic Pancreatoduodenectomies With Patient Reported Outcomes From a Single Center in the UK

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To the Editor

Robotic surgery has seen exponential growth in the UK during the last 5 years. In fact, in the country, only two centers (including ours) perform robotic pancreatoduodenectomies (RPDs). The aim of the present study was to evaluate and report the postoperative outcomes and patient satisfaction of the first 16 consecutive RPDs from a single center. Prospectively collected data for RPDs were analyzed for postoperative outcomes. All the operations were performed with the Da Vinci surgical system (Intuitive Surgical Inc., Sunnyvale, CA, USA). All operations were performed using the same technique. In particular, pancreatojejunostomies were performed according to modified Blumgart technique using 3-0 proline on 31 mm needle for the outer layer and 5-0 PDS on 17 mm needle for duct to mucosa anastomosis [1]. Hepaticojejunostomy was performed using 5-0 PDS continuous anterior and posterior layers when the duct was more than 1 cm and interrupted sutures in a smaller duct.

The International Study Group on Pancreatic Surgery (ISGSP) definitions of the post-pancreatectomy fistula, hemorrhage, delayed gastric emptying, and Dindo-Clavien classification were used to categorize the postoperative complications [2, 3]. A validated fistula risk score calculator was used to assess the risk for development of postoperative pancreatic fistula [4]. Patient satisfaction was assessed using the RAND 36-Item Health survey [5]. To prevent performance bias, a junior Doctor from a different team who had no prior knowledge of patient’s pre- and postoperative details conducted the interviews. The patients rated at discharge, 2 weeks and 6 months postoperatively.

All continuous data were presented as median or mean ± standard deviation (SD) and all categorical variables were presented as numbers and percentages.

Twelve of 16 patients were included in the analysis. Two operations were converted to open, one due to extend malignant portal vein involvement with a view to vein resection and the other due to failure to progress due to high body mass index (BMI). Another two operations were abandoned, one due to peritoneal metastases and the other with intraoperative possible diagnosis of liver cirrhosis.

Mean operative time was 547 ± 65 min and the mean intraoperative blood loss was 245 ± 119 mL. Docking and undocking time was included in the operative time and was about 30 min.

Eleven out of 12 patients belong to American Society of Anesthesiologists (ASA) tier II and the mean BMI was 24.9 ± 2.8. Obviously, the above parameters had a positive impact on the postoperative recovery. Only two out of 12 patients developed Dindo-Clavien IIIb complications. Although, four (33%) out of 12 patients were characterized as of high risk for occurrence of postoperative fistula; none of the patients developed grade B or C fistula, only one patient was diagnosed with grade A fistula (Table 1).

Only two (17%) out of 12 patients were diagnosed with R1 margins.

The low incidence rates of major complications and pancreatic fistula obviously reflected on the shorter hospital stay compared with open procedures performed by the same team. The mean hospital stay was 8.33 ± 2.8 days (Table 1).

Although the number of enthusiastic supporters of RPD has increased recently exponentially, there are limited data reported on patient satisfaction and the associated quality of life (QoL) following robotic surgery [1, 6-8]. The present study is one of the first reporting QoL. Notably, only one patient out of 12 complained of severe non-specific body pains at discharge and 2 weeks postoperatively (Table 2).

Both parameters of physical functioning score and general health score demonstrated an increase compared to baseline score; these findings provide evidence for better QoL. In addition, the patients score higher for social functioning. In par-
ticular, 10 patients reported that it took them up to 5 weeks to recover since operation and remaining two recovered within 5 - 10 weeks (Table 2).

To the author’s best knowledge, this is the first study in the UK reporting postoperative outcomes and QoL in patients undergoing RPD. It demonstrates the feasibility, safety and efficacy of the RPD; moreover, it demonstrates high patient satisfaction for major operation such as pancreatoduodenectomy. Results from comparative studies and data from international registries will certainly help health care providers further to support robotic pancreatic surgery in the future. However, the results of the present study should be interpreted in the context of its limitations. The data are from initial experience from a single center in the UK. Therefore, institutional and underpowered sample bias might have influenced the results.

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None to declare.

Table 1. Patient Characteristics, Intra- and Postoperative Outcomes, Postoperative Pain Scores

| Mean age (years) | 70 ± 9.5 |
|------------------|----------|
| Gender           |          |
| Male             | 7        |
| Female           | 5        |
| ASA tier         |          |
| II               | 11       |
| III              | 1        |
| BMI, mean (SD)   | 24.9 (2.8)|
| Indications for surgery | |
| Ampullary adenoma with high grade dysplasia | 4 |
| IPMN             | 1        |
| Ampullary adenocarcinoma | 4 |
| Cholangiocarcinoma | 2 |
| Mixed neuroendocrine tumor | 1 |
| R1 (positive resection margins) | 2 (17%) |
| Intraoperative blood loss (mL), mean (SD) | 245 (119) |
| Operating time (min), mean (SD) | 547 (65) |
| Complications (Dindo-Clavien grades of major morbidity) | |
| Grade II         | 3        |
| Grade IIIa       | 1        |
| Grade IIIb       | 2        |
| Pancreatic fistula |          |
| Grade A          | 1        |
| Grade B or C     | None     |
| Fistula risk score groups | |
| Low risk         | 3        |
| Intermediate risk | 5 |
| High risk        | 4        |
| Length of stay (days), mean (SD) | 8.33 (2.49) |
| Postoperative pain score | |

ASA: American Society of Anesthesiologists; BMI: body mass index; SD: standard deviation; IPMN: intraductal papillary mucinous neoplasm.

| Pain numerical rating | Mild (1 - 3) | Moderate (4 - 6) | Severe (7 - 10) |
|-----------------------|--------------|------------------|-----------------|
| Pain immediate postoperation | 3 | 7 | 2 |
| Pain at discharge     | 10 | 1 | 1 |
| Pain at 2 weeks postoperation | 11 | | 1 |
Table 2. Quality of Life Survey and Postoperative Pain Score

| SF-36 survey domains                        | Pre-operative scores, mean (SD) | Postoperative scores, mean (SD) |
|---------------------------------------------|---------------------------------|---------------------------------|
| Physical functioning                        | 83 (12)                         | 90 (9)                          |
| General health                              | 38 (7.2)                        | 70 (7.5)                        |
| Bodily pain                                 | 84 (9.5)                        |                                 |
| Social functioning                          | 96                              |                                 |

| Other condition-specific symptoms           |                                 |                                 |
| Feeling motivated                           | Yes: 11; No: 1                   |                                 |
| Feeling depressed                           | Yes: 1; No: 11                   |                                 |
| Full of energy                              | Most of the time: 4; Some of the time: 9; None of the time: 0 |                                 |
| Charge in emotional well-being              | Same: 7; Better: 4; Worse: 1     |                                 |

SD: standard deviation.

Financial Disclosure

None to declare.

Conflict of Interest

None to declare.

Informed Consent

Not applicable.

Author Contributions

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Data Availability

The authors declare that data supporting the findings of this study are available within the article.

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