

2022-RA-1075-ESGO

LONG TERM FOLLOW-UP AND OUTCOMES OF BORDERLINE OVARIAN TUMOURS – A TEN YEAR REVIEW OF THE SOUTH EAST WALES GYNAECOLOGICAL ONCOLOGY CENTRE (SEWGOC)

¹Hassan Zeinah, ¹Daniel Adama, ¹Elsie Tan, ²Upha Barclay, ¹Sadie Jones, ¹Ewelina Rzycka, ¹Kenneth Lim, ¹Robert Howells, ³Gareth Rowlands, ¹Anju Sinha, ¹Aarti Sharma. ¹The South East Wales Gynaecological Oncology Centre, Cardiff, UK; ²School of Medicine, Cardiff University, Cardiff, UK; ³Department of Pathology, University Hospital of Wales, Cardiff, UK

10.1136/ijgc-2022-ESGO.431

Introduction/Background Borderline ovarian tumours (BOT) are low malignant potential tumours. There is no consensus on how best to follow up those patients. The aim of this service evaluation project is to assess our current management and long-term outcomes and follow up in women diagnosed with borderline ovarian tumours over a ten-year period.

Methodology All women with confirmed histological diagnosis of BOT who underwent primary surgery at SEWGOC between 1st January 2007 to 31st December 2016 were included. Retrospective review of patients' electronic medical records was undertaken. Information regarding FIGO stages, management (fertility preserving surgery/pelvic clearance), follow up and recurrence were analysed.

Results Seventy-nine patients were diagnosed with BOT. The mean age was 48 years (range 18 – 86). Of these, 67 were stage I, 4 stage II and 8 stage III. Fertility sparing surgery (mean age 38) was performed in 32 patients (30 stage I, 2 stage III). Of these, 22 had follow-up. Four of 32 (12.5%) had recurrences. Pelvic clearance (mean age 55) was undertaken in 47 patients. Of these, 23 had follow up. Three of 47 (6%) patients presented with recurrence. All recurred within 5 years.

Conclusion This evaluation shows that recurrence in women who undergo fertility sparing surgery is doubled versus pelvic clearance. All patients with recurrence presented with symptoms within 5 years of initial surgery. Symptom-led follow up could be a suitable modality especially in those who underwent pelvic clearance.

2022-RA-1076-ESGO

LOW PRESSURE LAPAROSCOPIC PROCEDURES IN MORBIDLY OBESE GYNECOLOGICAL PATIENTS USING A NEW SUBCUTANEOUS ABDOMINAL WALL-RETRACTION DEVICE: A SAFETY AND FEASIBILITY STUDY

¹Antonino Ditto, ¹Umberto Leone Roberti Maggiore, ²Vincenzo Granato, ¹Fabio Martinelli, ¹Valentina Chiappa, ¹Salvatore Lopez, ¹Mauro Signorelli, ¹Matteo Maruccio, ³Simona Palladino, ⁴Giulia Chiarello, ⁵Ludovica Spanò Bascio, ¹Francesco Raspagliesi. ¹Fondazione IRCCS Istituto Nazionale Tumori, Milano, Italy; ²Università dell'Insubria, Varese, Italy; ³Università degli studi di Verona, Verona, Italy; ⁴Università 'Aldo Moro' di Bari, Bari, Italy; ⁵Policlinico universitario di Modena, Modena, Italy

10.1136/ijgc-2022-ESGO.432

Introduction/Background Laparoscopic surgery for female patients with high BMI is still challenging despite it has been shown safe in obese patients. According to available literature, laparoscopic to laparotomic conversion rate in these patients is 57%, mostly for inadequate surgical exposure and anaesthetics indications. The aim of this prospective study is to assess

feasibility, laparotomic conversion rate and perioperative complications after low-pressure laparoscopy (LPL) using a new subcutaneous abdominal wall-retraction device (LaparoTenser) in morbidly obese patients with gynecological diseases.

Methodology

Inclusion criteria were patients aged > 18 years, with BMI >30 kg/m² who were eligible for laparoscopic surgery for gynaecological disease. We excluded patients with preoperative diagnosis of extra-uterine disease and contraindication to upfront general anaesthesia/mini-invasive surgery. Anamnestic, surgical, postoperatively complications and hospitalization related data were prospectively collected.

Abstract 2022-RA-1076-ESGO Table 1

Main baseline population characteristics	Value
Age	68 (40 – 83)
Obesity grade	
Grade I (BMI ¹ 30 to 34.9)	3/24 (12.5%)
Grade II (BMI ¹ 35 to 39.9)	9/24 (37.5%)
Grade III (BMI ¹ over 40)	12/24 (50.0%)
Type of surgery	
TLH ² -BSO ³	3/24 (12.5%)
TLH ² -BSO ³ -SNL ⁴ biopsy	17/24 (71%)
MSO ⁵	1/24 (4.1%)
BSO ³ and omentectomy	1/24 (4.1%)
TLH ² -BSO ³ -SNL ⁴ biopsy and complete peritoneal staging	2/24 (8.3%)
Surgical outcome	
Operative time (min)	175 (111 – 249)
Conversion to laparotomy	3/24 (12.5%)
Advanced disease	2/24 (8.3%)
Difficult visualization of the operative field	1/24 (4.1%)
Complete surgical staging	23/24 (95.8%)
Intra-operative complications	0/10 (0%)
Hospital stay (days)	4 (3 – 7)
30-days complications	6/24 (25.0%)
Clavien Dindo grade 1	5/24 (20.9%)
Clavien Dindo grade 2	1/24 (4.1%)
Clavien Dindo grade 3	0 / 24 (0%)
Clavien Dindo grade 4	0 / 24 (0%)

data are expressed as median (range) or number (percentage)

¹ Body Mass Index; ²Total Laparoscopic Hysterectomy; ³ Bilateral Salpingo-Oophorectomy; ⁴ Seminal Lymph Node; ⁵ Monolateral Salpingo-Oophorectomy

Results We enrolled 24 patients since October 2020 to May 2022. table 1 summarizes the main characteristics of patients included in the study. The operating field visualization was optimal in 23 out of 24 cases (95.8%) with a median (range) CO₂ pressure of 3 (3–5) mmHg. Conversion rate for inadequate exposure was 4.1% (1/24). 2/24 patients underwent laparotomic conversion to be radically treated because of advanced disease. Operative time, blood loss, and hospital stay were similar to standard laparoscopy. No intraoperative