

Conclusion In a pooled analysis of 464 patients, MIRV monotherapy demonstrated ETB in ~10% patients. The safety profile consisted primarily of low-grade gastrointestinal and ocular events and reinforces MIRV's potential to become a new standard of care for FRα-positive ovarian cancer.

2022-RA-669-ESGO

PERIOPERATIVE MANAGEMENT OF ADVANCED OVARIAN (TUBAL/PERITONEAL) CANCER PATIENTS. A SURVEY FROM MITO-MANGO GROUPS

¹Francesca Falcone, ¹Maria Stella Gallo, ²Marco Cascella, ²Francesca Bifulco, ³Grazia Artioli, ⁴Paola Agnese Cassandrini, ¹Gennaro Casella, ⁵Maurizia Dalla Palma, ⁶Laura Falchi, ¹Cono Scaffa, ¹Felice Scala, ¹Serena Visconti, ¹Giuseppe Laurelli, ¹Rosaria Grimaldi, ¹Lucia Formisano, ⁷Antonio Frassoldati, ⁸Ettore Cicinelli, ⁹Emilio Stola, ¹⁰Giovanni Damiano Aletti, ¹Stefano Greggi. ¹Department of Gynecologic Oncology, Istituto Nazionale Tumori, IRCCS, 'Fondazione G. Pascale', Naples, Italy; ²Division of Anesthesia and Pain Medicine, Istituto Nazionale Tumori, IRCCS, 'Fondazione G. Pascale', Naples, Italy; ³U.O.C. Oncologia ed Ematologia Oncologica, ULSS 3 Serenissima, Mirano, Venice, Italy; ⁴U.O.C. Oncologia medica, 'Sacro Cuore-Don Calabria' Hospital of Negrar, Verona, Italy; ⁵U.O.C. Oncologia medica, ULSS 3 Serenissima, Venice, Italy; ⁶Obstetrics and Gynecology, 'San Giovanni Di Dio' Hospital, Florence, Italy; ⁷Unit of Clinical Oncology, S. Anna University Hospital, Ferrara, Italy; ⁸Department of Obstetrics and Gynaecology, Faculty of Medicine, University of Bari, Bari, Italy; ⁹Obstetrics and Gynecology, 'SS Annunziata' Hospital, Taranto, Italy; ¹⁰Department of Gynecologic Surgery, European Institute of Oncology, IRCCS, Milan, Italy

10.1136/ijgc-2022-ESGO.540

Introduction/Background The ESGO-quality indicators (QIs) for advanced ovarian cancer (AOC) have been assessed only by few Italian centres, and data are not available on the proportion of centres reaching the score considered for a satisfactory surgical management. There is great consensus that the ERAS approach is beneficial, but there is paucity of data concerning its application in AOC. This survey was aimed at gathering detailed information on perioperative management of AOC patients within MITO-MaNGO Groups.

Methodology A 66-item questionnaire, covering ESGO-QIs for AOC and ERAS items, was sent to MITO/MaNGO centres reporting to operate >20 AOC/year.

Results Thirty/34 questionnaires were analysed. The median ESGO-QIs score was 31.5, with 50% of centres resulting with a score ≥ 32 which provides satisfactory surgical management. The rates of concordance with ERAS guidelines were 46.6%, 74.1%, and 60.7%, respectively, for pre-operative, intra-operative, and post-operative items. The proportion of overall agreement was 61.3%, and with strong recommendations was 63.1%. Pre-operative diet, fasting/bowel preparation, correction of anaemia, post-operative feeding and early mobilization were the most controversial. A significant positive correlation was found between ESGO-QIs score and adherence to ERAS recommendations.

Conclusion This survey reveals a satisfactory surgical management in only half of the centres, and an at least sufficient adherence to ERAS recommendations. Higher the ESGO-QIs score stronger the adherence to ERAS recommendations, underlining the correlations between case volume, appropriate peri-operative management and quality of surgery.

2022-RA-672-ESGO

COMPARISON OF QUALITY OF LIFE IN PATIENTS WITH PLATINUM-SENSITIVE RECURRENT OVARIAN, FALLOPIAN TUBE AND PERITONEAL CANCER TREATED WITH TRABECTEDIN PLUS PEGYLATED LIPOSOMAL DOXORUBICIN (PLD) OR STANDARD PLATINUM-BASED THERAPY: DATA LOOK OF THE NOGGO S16/COMPASS TRIAL

^{1,2}Radoslav Chakerov, ³Mustafa Deryal, ⁴Bahriye Aktas, ⁵Robert Röhle, ²Annika Stürzebecher, ⁶Marco J Battista, ⁷Christian M Kurbacher, ^{8,2,9}Pauline Wimberger, ¹⁰Ralf Lorenz, ¹¹Jana Barinoff, ¹²Gunther Rogmans, ¹³Jens Kosse, ¹⁴Maximilian Klar, ¹⁵Tomas Kupec, ¹⁶Bernd Christensen, ^{17,2}Gülten Oskay-Özcelik, ¹⁸Thomas Illmer, ¹⁹Thorsten Götze, ^{1,2}Klaus Pietzner, ^{1,2}Jalid Sehouli. ¹Department of Gynecology with Center for Oncological Surgery, Charité-University Medicine of Berlin, Berlin, Germany; ²Nord-Ostdeutsche Gesellschaft für Gynäkologische Onkologie – NOGGO e.V., Berlin, Germany; ³Department for Gynecology and Obstetrics, CaritasKlinikum Saarbrücken St. Theresia, Saarbrücken, Germany; ⁴Klinik und Poliklinik für Frauenheilkunde, Universitätsklinikum Leipzig, Leipzig, Germany; ⁵Institute of Biometry and Clinical Epidemiology, Charité – Universitätsmedizin Berlin, corporate member of Freie Universität Berlin and Humboldt-Universität zu Berlin, Berlin, Germany; ⁶Department of Obstetrics and Gynecology, University Hospital Mainz, Mainz, Germany; ⁷Gynäkologisches Zentrum Bonn, Bonn, Germany; ⁸Technische Universität Dresden, Dresden, Germany; ⁹Nationales Centrum für Tumorerkrankungen (NCT), Dresden, Germany; ¹⁰Studien GbR Braunschweig, Braunschweig, Germany; ¹¹Sankt Gertrauden Krankenhaus, Berlin, Germany; ¹²ZAGO – am Helios Klinikum Krefeld, Krefeld, Germany; ¹³Sana Klinikum Offenbach, Offenbach, Germany; ¹⁴Klinik für Frauenheilkunde, Universitätsklinikum Freiburg, Freiburg, Germany; ¹⁵Klinik für Gynäkologie und Geburtshilfe RWTH Aachen, Universitätsklinikum Aachen, Aachen, Germany; ¹⁶Klinik für Gynäkologie und Geburtshilfe, ukbr – Universitätsklinikum Ruppert-Brandenburg, Neuruppin, Germany; ¹⁷Praxis für Krebsheilkunde, Berlin, Germany; ¹⁸Oncology Praxis – BAG, Dresden, Germany; ¹⁹Institut für Klinisch-Onkologische Forschung (IKF), Krankenhaus Nordwest, Frankfurt, Germany

10.1136/ijgc-2022-ESGO.541

Introduction/Background Despite recent progress regarding surgical and medical management of primary ovarian cancer, relapses are still frequent and one of the most critical challenges in the clinical routine. There is a broad consensus that quality of life (QoL) should be one of the most relevant goals of any therapy in relapsed ovarian cancer.

Methodology We report the results of a data look of the multicentre, randomized (1:1), active-controlled, open-label phase IV NOGGO-S16/COMPASS trial performed in patients with recurrent, platinum-sensitive, ovarian, peritoneal or fallopian tube cancer. The scope of this trial is to evaluate QoL with EORTC QLQ-C30 and QLQ-OV28 questionnaires during/after chemotherapy either with platinum/taxane-free combination of trabectedin (Yondelis®) plus PLD or with standard platinum-based chemotherapy comprising combination of carboplatin with PLD, gemcitabine or paclitaxel. The current data look serves to characterise the included patient population.

Results Data from 76 patients screened have been analysed. Patients have a median age of 63 years (range: 21–82), the performance status score of 0/1 was recorded in 75 patients (98.7%), and most are BRCA-negative (77.4%). They are diagnosed with primary ovarian carcinoma (83.6%), primary peritoneal carcinoma (9.6%) and fallopian carcinoma (6.8%), and 79% of patients have a grade 3 histopathological staging. Poly (ADP-ribose) polymerase inhibitors or bevacizumab were given as prior maintenance therapy to 15.3% and 76.4% of