

2022-RA-758-ESGO

PRETREATMENT ¹⁸F-FDG PET/CT METABOLIC PARAMETERS AS PREDICTORS OF NON-COMPLETE CYTOREDUCTION IN PATIENTS WITH EPITHELIAL OVARIAN CANCER

¹Lidia Sancho Rodriguez, ²Luis Chiva de Agustín, ²Félix Boria Alegre, ²Luisa Sanchez Lorenzo, ²Teresa Castellanos Alarcón, ²Daniel Vazquez Vicente, ³Vicky Beteche Antar, ²Antonio Gonzalez Martin, ²Laura García Belaustegui, ²Teresa Iscar Galan, ³Angela Bronte Viedma, ³Enrique Maria Chacon Cruz, ³Jose Angel Minguez Milió, ²Andres Alcazar Peral, ²Guillermo Gallardo Madueño, ³Nabil Manzour Sifontes, ⁴Carmen Beorlegui Arteta, ²Felix Mauricio Cambeiro Vazquez, ²Jaime Espinos Jimenez, ²Monica Gutierrez Martinez, ²Jacobo Palma Delgado, ³Juan Luis Alcazar Zambrano, ³MJ Garcia-Velloso. ¹Nuclear Medicine, Clínica Universidad de Navarra, Madrid, Spain; ²Clínica Universidad de Navarra, Madrid, Spain; ³Clínica Universidad de Navarra, Pamplona, Spain; ⁴Universidad de Navarra, Pamplona, Spain

10.1136/ijgc-2022-ESGO.554

Introduction/Background The objective of this study was to analyse the utility of pretreatment ¹⁸F-FDG-PET/CT metabolic parameters to predict non-complete cytoreduction in patients with epithelial ovarian cancer.

Abstract 2022-RA-758-ESGO Table 1 Sample characteristics and relation to non-complete cytoreduction

Table 1 – Sample characteristics and relation to non-complete cytoreduction.		
VARIABLE	N (%)	X ² (Fisher)
FIGO		
≤ IIIB	21 (42)	0.092
IIIC / IV	29 (58)	
Histological type		
HGSOC (high grade serous ovarian cancer)	36 (72)	ns
Mucinous carcinoma	4 (8)	
Clear cell carcinoma	4 (8)	
Mesonephric adenocarcinoma	2 (4)	
Endometrioid carcinoma	4 (8)	
Type of debulking surgery		
Complete (R0)	39 (78)	ns
Non-complete (R1)	11 (22)	
Ascites		
Pathological ¹⁸ F-FDG uptake	10 (20)	0.004
Non-pathological ¹⁸ F-FDG uptake	40 (80)	
VARIABLE	Median (IQR)	Logistic regression
Age (years)	58 (47-62)	ns
Total metabolic active disease		
MTV	104.7 (14.1 – 313.4)	0.007
TLG	362.6 (65.9 – 1,281.4)	0.123
Infradiaphragmatic disease		
MTV	53.7 (14.1 – 281.6)	0.010
TLG	226.5 (63.1 – 1,125.7)	0.189
Total peritoneum		
MTV	107.4 (19.3 – 300.9)	0.027
TLG	307.8 (62.9 – 1,114.5)	0.055
Upper abdomen peritoneum		
MTV	28.3 (12.9 – 63.2)	0.420
TLG	82.9 (40.5 – 173.6)	0.412
Lower abdomen peritoneum		
MTV	63.9 (9.8 – 177.1)	0.104
TLG	189.5 (44.4 – 480.4)	0.134
Total pelvic disease (primary tumor and peritoneum)		
MTV	43.4 (9.8 – 213.0)	0.203
TLG	182.8 (42.1 – 859.9)	0.727
Infradiaphragmatic lymph nodes		
MTV	5.9 (3.8 - 8.4)	0.248
TLG	16.2 (10.2-25.9)	0.316

Methodology Transversal study on 50 patients with epithelial ovarian cancer at Clínica Universidad de Navarra who underwent pretreatment ¹⁸F-FDG-PET/CT and subsequent debulking surgery (R0 = complete, R1 = non-complete). The supra- and infradiaphragmatic metabolic active disease (primary tumor, peritoneal carcinomatosis and lymph nodes) visualized in the ¹⁸F-FDG-PET/CT was segmented using Syngo.via (automatic thresholding at 40% SUVmax and manual corrections). The extent and distribution of the peritoneal carcinomatosis

was evaluated globally and throughout abdominopelvic regions. The presence of pathological ¹⁸F-FDG uptake of the ascites was also evaluated. Metabolic parameters studied were metabolic active tumor volume (MTV) and total lesion glycolysis (TLG, defined as MTVxSUVmean), calculated for each segmented region and for the whole disease. Other variables studied were age, FIGO and histological tumor type. The dependent variable was non-complete cytoreduction. Data were described by median (IQR) and frequency (%). Chi-squared and median test were used to compare groups and ROC analysis to dichotomize continuous variables. Predictors of non-complete cytoreduction were analysed by multiple logistic regression.

Results Patient's characteristics are listed in table 1. Eleven patients (22%) showed non-complete cytoreduction, mostly associated to pathological uptake in ascites (60 vs 12.5%; OR= 10.5 95%CI: 2.2–50.7; p= 0.004), total MTV >192 (45.0 vs 6.7%; OR=11.5; 95%CI: 2.1–61.7; p=0.007; AUC=0.818) and MTV value of the whole infradiaphragmatic disease >209 (56.3 vs 5.9%; OR= 20.6; 95%CI: 3.6–116.8; p=0.010; AUC=0.818). Only the MTV of the whole infradiaphragmatic disease retains signification in the adjusted model.

Conclusion Despite the small sample size, this initial study highlights the possible role of some ¹⁸F-FDG-PET/CT metabolic parameters as predictors of non-complete cytoreduction in patients with epithelial ovarian cancer. Further validation in larger series is needed.

2022-RA-761-ESGO

ADVANCED OVARIAN CANCER IMITATING DEEP INFILTRATING ENDOMETRIOSIS. RADICAL RESECTION AND RECONSTRUCTIVE SURGERY OF THE ANTERIOR ABDOMINAL WALL

Piotr Lepka, Marcin Jedryka. Department of Gynecological Oncology, Lower Silesian Oncology, Pulmonology and Hematology Center, Wrocław, Poland

10.1136/ijgc-2022-ESGO.555

Introduction/Background Endometriosis is a disease affecting approximately 10–15% of the female population of reproductive age. Malignant transformation affects 0.7–1% of cases. In women diagnosed with ovarian cancer, endometriosis is present in up to 30% of patients.

Methodology The case of a 36-year-old patient initially diagnosed with pelvic and abdominal wall endometriosis with final diagnosis of advanced low-grade serous ovarian cancer in stage FIGO IVB. The diagnostic methods used and the extent of surgery with reconstruction of the anterior abdominal wall were described.

Results Patient with history of laparoscopic excision of benign ovarian cyst and caesarean section was presented to the gynecologist because of abdominal pain that worsened during menstruation. MRI (figure 1A) and core needle biopsy of the abdominal lesion and colonoscopy were performed. Histopathological report revealed low-grade serous carcinoma originating from the ovary. Complete surgical debulking was performed. Modified Ramirez surgical technique was used to approximate the borders of the fascia (figure 1B,C) and hernia mesh was applied (figure 1D).