

HRN: ID Card:

OncoOVARIAN Dx - Order Form

Biochemistry

General: Creatinine ASAT ALAT
 GGT Total Bilirubin
Tumor Markers: AFP β -hCG CA 19.9
 CA 125 CEA HE4

Anamnesis

Personal Data

Age (years):

Menopausal Status

Menopausal: Pre Post (less 1 year) Post (more 1 year)
 Unknown

Comorbidities

Ascites: Yes No Unknown
Cholestasis: Yes No Unknown
Chronic Liver Disease: Yes No Unknown
Jaundice: Yes No Unknown
Metrorrhagia: Yes No Unknown
Pancreatitis: Yes No Unknown
Pericardial/Pleural Effusions: Yes No Unknown
Renal Failure: Yes No Unknown

Order Information

Ordered by: Signature:
Order Date:

Disclaimer

According to Annex VII of Medical Device Directive 93/42/EEC, Class I, Rule 12, OncoOVARIAN Dx is a non-invasive blood test useful to suggest a possible diagnosis in women with suspected malignancy in the ovarians, reduce inappropriate diagnostic tests, days of hospitalization, as well as morbidity. It is specially designed for women with previous image findings suspicious of Ovarian Cancer that should be biopsied to verify malignancy.

OncoOVARIAN Dx is based in a panel consisting of 6 serum Tumor Markers (AFP, β -hCG, CA 19.9, CA 125, CEA and HE4), as well as clinical data to obtain the best performance (Sensitivity of 93.8% and Specificity of 94.4%), by reducing the number of false positives generated by benign comorbidities and other well-known causes.

