

HRN: ID Card:

OncoBREAST Dx - Order Form

Biochemistry

General:	<input checked="" type="checkbox"/> Creatinine	<input checked="" type="checkbox"/> ASAT	<input checked="" type="checkbox"/> ALAT
	<input checked="" type="checkbox"/> GGT	<input checked="" type="checkbox"/> Total Bilirubin	
Serum Tumor Markers:	<input checked="" type="checkbox"/> CA 15.3	<input checked="" type="checkbox"/> CEA	<input checked="" type="checkbox"/> EGFR
	<input checked="" type="checkbox"/> NGAL	<input checked="" type="checkbox"/> NSE	<input checked="" type="checkbox"/> 8-OHdG

Anamnesis

Personal Data

Age (years):

Comorbidities

Atherosclerosis	<input type="checkbox"/> Yes	<input type="checkbox"/> No	<input type="checkbox"/> Unknown
Chronic Liver Disease:	<input type="checkbox"/> Yes	<input type="checkbox"/> No	<input type="checkbox"/> Unknown
Diabetes Mellitus 1 or 2	<input type="checkbox"/> Yes	<input type="checkbox"/> No	<input type="checkbox"/> Unknown
Jaundice:	<input type="checkbox"/> Yes	<input type="checkbox"/> No	<input type="checkbox"/> Unknown
Renal Failure:	<input type="checkbox"/> Yes	<input type="checkbox"/> No	<input type="checkbox"/> Unknown

Lifestyle

Smoking:	<input type="checkbox"/> Never	<input type="checkbox"/> Currently	<input type="checkbox"/> Past
	<input type="checkbox"/> Unknown		

Order Information

Ordered by: Signature: Order Date:

Disclaimer

According to Annex VII of Medical Device Directive 93/42/EEC, Class I, Rule 12, OncoBREAST Dx is a non-invasive blood and urine test useful to suggest a possible diagnosis in women with suspected malignancy in the breast, reduce inappropriate diagnostic tests, days of hospitalization, as well as morbidity. It is specially designed for women with a suspicious lump found in the breast, as a complementary method in radiological testing for high-risk women or those with a level of BI-RADS 3 and 4A (image findings suspicious of Breast Cancer that should be biopsied to verify malignancy), as well as women with family history of Breast Cancer.

OncoBREAST Dx is based in a panel consisting of 6 serum Tumor Markers (CA 15.3, CEA, EGFR, NGAL, NSE and 8-OHdG), as well as clinical data to obtain the best performance (Sensitivity of 82.3% and Specificity of 85.7%), by reducing the number of false positives generated by benign comorbidities and other well-known causes.

