HORIZONS



Understanding the impact of cancer diagnosis and treatment on everyday life

BASELINE OVARIAN CANCER CRF

(please also use for primary peritoneal and fallopian tube cancers)

FOR STAFF USE ONLY

CRF Completion Instructions

- This CRF is for completion by members of site staff NOT study participants
- Please complete the CRF when a patient has been recruited to the study
- Please complete as much of the CRF as possible
- If you have any queries, please contact the HORIZONS Coordinating Centre, email address <u>HORIZONS@soton.ac.uk</u>
- Please tick boxes when appropriate
- When you have completed the CRF, please keep a copy for your own records and return a copy to us, by post, fax or email along with the completed return cover sheet

Participant's Study ID / /				
Participant's date of birth	d d m m	у у у у		
Participant's weight kg Participant's height cms				
Participant's blood pressure which they were measured)	e (Please give the most r	ecently reported figures and th	e date on	
Systolic	mmHg	Date measure	d	
Diastolic	mmHg	d d m m	уууу	
Participant's tumour type (please tick one box)				
Туре	Sub-type			
Ovarian	Serous			
	Mucinous			
	Endometrioid			
	Clear Cell			
	Undifferentiated/unclas	ssifiable		
Primary peritoneal cancer				
Fallopian tube cancer				
Other (please describe)				
Not currently known				
Date of participant's current cancer diagnosis dd d m m m y y y y				

Participant's Study ID / /							
-	FIGO stage (please	tick one bo	x OR tick	the b	ox indi	icating t	the FIGO
Stage 1	Stage IA		FIGO stag	ge no	t curre	ently kn	own 🗍
	Stage IB						
	Stage 1C1						
	Stage 1C2						
	Stage 1C3						
Stage 2	Stage 2A						
	Stage 2B						
Stage 3	Stage 3A1						
Participant's	tumour grade (ple	ase tick one	box)			_	
Grade 1/lov	v grade/well differe	ntiated					
Grade 2/mc	oderate/intermedia	te grade					
Grade 3/hig	h-grade/poorly diff	erentiated					
Grade not c	urrently known						
Participant's	s pre-treatment ECC	OG status (pl	ease tick (one b	ox)		
ECOG 0 (the	patient has no sympto	ms)					
ECOG 1 (the	patient has symptoms	but is ambula	tory)				
ECOG 2 (the	patient is bedridden le	ess than half th	ne day)				
ECOG 3 (the	patient is bedridden h	alf the day or	longer)				
1	patient is chronically be activities of daily livir		requires as	sis-			
Is the partic	ipant pre or post m	nenopause?	(please tid	ck one	e box)		
Pre menopa	iuse						
Post menop	ause						
Unknown							

Participant's Study ID	
Has the participant had a prev	ious diagnosis of cancer (please tick one box)
Yes No	Unknown
	above question, please provide some information cancer(s) by completing the box(es) below
PREVIOUS DIAGNOSIS 1	
Type of cancer	
Date of diagnosis	
Treatment received	
Date treatment ended	
PREVIOUS DIAGNOSIS 2	
Type of cancer	
Date of diagnosis	
Treatment received	
Date treatment ended	
Has the participant been teste	ed for BRCA1 or BRCA2 (please tick one box)
Yes No	Unknown
If you answered "Yes" to the a	bove question, was the result (please tick one box)
Positive for a mutation in BRC	A1 or BRCA2
Negative for a mutation in BRG	CA1 or BRCA2
Ambiguous or uncertain	
Unknown	
Awaiting result	

Participant's Study ID / /				
Has the participant had any other genetic te	sts for inherited cancers?			
(please tick one box)				
Yes No	Unknown			
If you answered "Yes" to the above question, please provide some information about the participant's other genetic test(s) by completing the table(s) below				
Name of genetic test for cancer (1) Result of genetic test				
	Positive			
	Negative			
	Ambiguous/uncertain			
	Awaiting result			
	Unknown			
Name of genetic test for cancer (2)	Result of genetic test			
	Positive			
	Negative			
	Ambiguous/uncertain			
	Awaiting result			
	Unknown			
What is the participant's CA125 blood test result?				
Test result =				
Test was not carried out (please tick)				
Test result unknown (please tick)				

Participant's Stud	y ID	/			
Has a first degree relative of the participant (parent, sibling or child) been diagnosed with cancer? (please tick one box)					
Yes	No	Unknov	vn		
If you answered "	If you answered "yes" to the above question, what type of cancer and when was				
	Type of cancer	Age at diagnosis	Date of diagn	osis	
Relative 1					
Relative 2					
Relative 3					
Does the participa apply) Myocardial infarct	ant have any of the follo	owing co-morbidities	s (please tick a	ll that	
Angina/coronary a	rtery disease				
Congestive Heart Failure					
Cardiac Arrythmias					
Hypertension					
Venous Disease (PE/DVT)					
Peripheral Arterial	Disease				
Restrictive Lung Disease or COPD (chronic bronchitis, emphysema, asthma etc.)					
Liver Disease (portal hypertension, chronic/acute hepatitis, cirrhosis etc.)					
Stomach Ulcers or Inflammatory Bowel Disease					
Acute or Chronic Pa	ancreatitis				
End-stage Renal Dis	sease (chronic renal insufficion	ency, dialysis etc.)			
Thyroid problems (hyperthyroidism, hypothyroid	lism etc)			

Participant's Study ID / / /				
Participant's co-morbidities continued (please tick all that apply)				
Diabetes Mellitus Type 1				
Diabetes Mellitus Type 2				
Stroke/TIA				
Dementia				
Paralysis (paraplegia or hemiplegia)				
Neuromuscular Condition (multiple sclerosis, Parkinson's, myasthenia gravis, other chronic neuromuscular disorder)				
Clinical diagnosis of anxiety				
Clinical diagnosis of depression				
Other psychiatric Diagnosis (schizophrenia, bipolar disorder etc.)				
Osteoarthritis				
Rheumatoid Arthritis				
Other Rheumatological Disease (systemic lupus, mixed connective tissue disorder, polymyositis, rheumatic polymyositis, scleroderma etc.)				
HIV/AIDS				
Alcohol Abuse (or history of, must be accompanied by social, behavioural or medical complications)				
Drug/Substance Abuse (or history of, must be accompanied by social, behavioural or medical complications)				
Morbid Obesity				
Other (please give details)				
What is the participant's proposed treatment start date (main first-line treatment for ovarian, primary peritoneal or fallopian tube cancer)				
Please add your name and signature and the date that you completed this (CRF			
Name Signature				
Date d d / m m / y y y y				