DBCG - PROTON TRIAL PHOTON RT PLANNING &THERAPY DANISH BREAST CANCER COOPERATIVE GROUP with boost Name - Patient ID Hospital ____ - _____ - ____ - ____ _ - _____ Month Year Day No. The upper part (Radiotherapy planning) of the sheet is filled in before RT. The lower part (Radiotherapy delivered) is filled in after completion of RT. If there is a deviation from the approved plan, the boxes for deviation are filled in with respect to the remaining fractions. Radiotherapy planning □CTVn L1 □CTVn L3 □CTVn L4 Indication for RT to □CTVn L2 □CTVn interpect □CTVn nodal boost levels according to □CTVn_IMN □CTVp_chest wall ESTRO guideline □CTVp_breast □CTVp_tumor bed boost (always SIB) Laterality Right Planned Revised plan CTVp-breast or chest wall Left 0=No V95% CTV_{p breast}-CTV_{p boost} (%) Breast implant 1=Yes Mastectomy: 0=No Mean dose CTV_{p breast}-CTV_{p boost} (Gy) Bolus on scar 1=Yes D25% CTV_{p breast}-CTV_{p boost} (Gy) Planned Revised plan V95% CTV_{p boost} (%) SIB dose (Gy) Mean dose CTV_{p boost} (Gy) Non-SIB dose (Gy) D2% CTV_{p_boost} (Gy) External receiving >105% but ≤107% **Fractions** of boost dose (ml) External receiving >107% but ≤110% 0=No Gating 1=Yes of boost dose (ml) External receiving >110% of boost CTVp_boost (ml) dose (ml) (Normofrac.: Max 10% may receive ≥20 Gy) Heart, V20/V17 (Hypofrac.: Max 10% may receive ≥17 Gy) (Normofrac.: Max 5% of heart may receive ≥40 Gy) Heart, V40/V35 (Hypofrac.: Max 5% of heart may receive ≥35 Gy) Mean heart dose (Gy) LADCA max dose (Gy) Ipsilateral lung, (Normofrac: Max 35% of ipsilat lung may receive ≥20 Gy) (Hypofrac: Max 35% of ipsilat lung may receive ≥17 Gy) V20/V17 CTVp_ breast / CTVp_chest wall (ml)

Form filled in by:
Name:(CAPITAL LETTERS)
Sign.:

Date	ddmmyy							
March 2015 RT PLANNING AND THERAPY – with SIB, page 2								
Patient ID Day Month Year No.								
Radiotherapy delivered								
Date of first RT	Form filled in by: Name:							
ddmmyy	(CAPITAL LETTERS)							
Date of last RT	Sign.:							
ddmmyy	Date ddmmyy							

Dose levels when therapy includes a simultaneous integrated boost, SIB, for tumour bed boost

SIB / Non-SIB in fr	V90%	V95%	Reference dose	D2% dose	Global		
	dose level	dose level	(mean dose)	level	max		
	(Gy)	(Gy)	(Gy)	(Gy)	(Gy)		
63 Gy / 51,52 Gy in 28 fr							
CTV _{p_breast} -CTV _{boost}	48,94		51,52				
CTV_{boost}		59,90	63,00	66,20	68,00		
57 Gy / 50 Gy in 25 fr							
CTV _{p_breast} -CTV _{boost}	47,50		50,00				
CTV_{boost}		54,20	57,00	59,90	61,60		
52,2 Gy / 42,3 Gy in 18 fr							
CTV _{p_breast} -CTV _{boost}	40,19		42,30				
CTV_{boost}		49,60	52,20	54,80	56,40		
45,75 Gy / 40 Gy in 15 fr							
CTV _{p_breast} -CTV _{boost}	38,00		40,00				
CTV_{boost}		43,50	45,75	48,00	49,40		

If the patient has a nodal boost, this is indicated in the tick off-box at the top of page 1. Only very few patients will have nodal boost, and data from these RT plans will be collected at a later point of time (most likely the RT plan with DICOM parameters will be collected).