Conclusion: There is evidence to suggest a dose volume effect between the PB and EP. Discriminatory PB dose-volume constraints were found to predict G2 EP. Further analysis is in progress to include patient reported outcomes related to EP.

Ref: (1) Wallner, IJROBP 2002 (2) Perna, Rad Onc, 2011 (3) Gay, IJROBP 2012

OC-0341
Anal dose reduction for radiotherapy of prostate cancer does not lead to less rectal incontinence
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Purpose or Objective: Radiation-induced rectal incontinence has a negative impact on Quality of Life in patients irradiated for prostate cancer. Several studies identified dose-effect relationships for the anal canal and lower rectum and hence, dose constraints for treatment planning have been implemented. We studied patient-reported rectal incontinence in a population treated with image-guided intensity modulated radiotherapy (IG-IMRT) and planned with a dose constraint for the anal canal, and compared it with a reference population treated with 3D-conformal radiotherapy (3D-CRT) with no dose constraint for the anal canal. For that purpose we analyzed data from two large prospective cohorts

Material and Methods: We selected patients treated to 78 Gy (39x2Gy) from two trials (CKTO 96-10 and CKTO 2006-08), who completed at least 2 follow-up questionnaires which included questions on pad use and fecal incontinence (IG-IMRT group n=242, 3D-CRT group n=189). In the IG-IMRT group, the mean dose to the anal canal was restricted to 58 Gy per protocol (more strict constraints depended on local planning guidelines). Grade >2 (G≥2) incontinence was defined as use of pads for uncontrolled loss of feces or mucus, Grade G1 incontinence was defined as any reported fecal incontinence regardless of use of pads. Prevalence and cumulative incidences of G2 and G1 incontinence were calculated. Cox regression was used to calculate Relative Risks (RR)

Results: Planned mean dose to the anal canal was on average 44.6 Gy (range 17-65) for 3D-CRT and 23.6 Gy (range 3-50) for IG-IMRT (p=0.001). Median follow up was 60 months. The 5y cumulative incidence of G2 incontinence was 15.2% for IG-IMRT vs 14.9% for 3D-CRT (RR=1.02, p=0.9). Prevalence of G1 incontinence was 5% at baseline and in the range of 30% - 40% in the years after treatment, with no significant differences between the groups (Figure 1). Within the 3D-CRT group, previous abdominal surgery was predictive for rectal toxicity (RR=2.9, p=0.05), whereas age >70 years at start RT (RR=2.9, p=0.01), diabetes mellitus (RR=2.4, p=0.04), and seminal vesicle dose 70 Gy vs 0 Gy (RR=9.2, p=0.03) were predictive in the IG-IMRT group. At multivariate analysis, adjusting for the significant baseline factors, RR of mean anal canal dose was 1.00 (p=0.9) for IG-IMRT patients and 1.05 (for each increase of 1 Gy) for 3D-CRT (p=0.04). Acute toxicity G2 (mainly proctitis) was predictive (p<0.01) in both groups with a RR of 3.1 (IG-IMRT) and 4.1 (3D-CRT). G1 incontinence at any time during follow-up was significantly associated with abdominal surgery in the 3D-CRT group, and with age >70 years, and diabetes mellitus in the IG-IMRT group

Conclusion: IG-IMRT with anal canal dose constraints did not reduce long-term incidence of rectal incontinence in prostate cancer patients, despite significantly reduced dose levels to the anal canal region. Further investigations are needed to understand the mechanisms of radiation damage causing rectal incontinence

OC-0342
Chemoradiotherapy in high-risk prostate cancer (QRT SOGUG trial): Preliminary report
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Purpose or Objective: To assess the toxicity and feasibility of concomitant radiotherapy with low doses of docetaxel plus standard hormonal treatment in patients with high risk localized prostate cancer

Material and Methods: Patients were randomly assigned to either arm A (LH-RH analogs every 3 months for 3 years and radiotherapy 74 Gy [20x3.7 fractions]) or arm B (LH-RH analogs every 3 months for 3 years, radiotherapy 73.8 Gy [1.8 Gy x 41 fractions] and concurrent weekly docetaxel at 20 mg/m2 for 9 weeks). Chemotherapy was started one week before of radiotherapy. Primary endpoint was PSA relapse according to the Phoenix definition. The planned number of patients was 130 to detect a 15% difference with a power of 80% and an alpha of 0.05 (two-sided).

Results: From 12/2008 to 9/2012, 130 pts were accrued (Arm A: 64, Arm B: 66). Median age was 68 years (61-73). Patients had T3-T4 (82.6%), Gleason Score 8 (76.3%), PSA > 20 ng/mL (26.9%) and pN+ (18.9%). All characteristics were well-balanced between arms. Median dose of radiotherapy was 74 Gy (72-74.8) in arm A, and 73.8 Gy (72.75.6) in arm B. 75.7% of patients received the planned 9 treatments of docetaxel and median number of cycles delivered per patient was 9. After a median follow-up of 29.6 months (9.6-40.2), most common grade 1/2 toxicities (arm A and arm B) were: cystitis (12.5% vs 8.3%), diabetes mellitus (12.5% vs 13.3%), rectal tenesmus (3.1% vs 23.3%), asthenia (23.4% vs 61.6%) and dysuria (28.1% vs 30.0%). Toxicity grade3/4, diarrhea was reported in 8.3% of patients in arm B and 0% in arm A. Grade3/4 lymphopenia occurred less often in arm A than in arm B (3.1% vs 23.3%). There was no toxicity-related death.

Conclusion: The QRT SOGUG phase IIb trial shows that standard doses of radiotherapy and concurrent weekly docetaxel can be administered without increasing toxicity profile.