

Phase III Randomized Trial of
Postoperative Adjuvant Conventional
Radiation (3DCRT) versus Image Guided
Intensity Modulated Radiotherapy (IG-IMRT) in Cervical Cancer (PARCER):
Final Analysis

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Disclosures

Research Funding PARCER Trial

- Department of Science and Technology, India
- Department of Atomic Energy, Clinical Trials Centre, India.





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Background

- Postoperative Radiation indicated for Cervix and Endometrial Cancers.
- Increase in GI symptom burden and toxicity in long term survivors after adjuvant radiation
- Phase II Studies
 - RTOG 0418/ RTCMIENDOMETRE demonstrated 27-28% acute GI toxicity with IMRT
 - No comparator arm
- Phase III Trial
 - NRG 1203: Improvement in patient reported outcomes at wk 5 and year 1 with IMRT as compared to 3DCRT.
 - No difference at 3 years with IMRT.
- Lack of Clarity on Long Term Impact of Postoperative IMRT

Hypothesis

IG-IMRT will improve late GI toxicity free survival in patients undergoing adjuvant RT for cervix cancer.

Conducted across 3 clinical sites of Tata Memorial Centre

NCT01279135/CTRI2012/120349

Study Eligibility

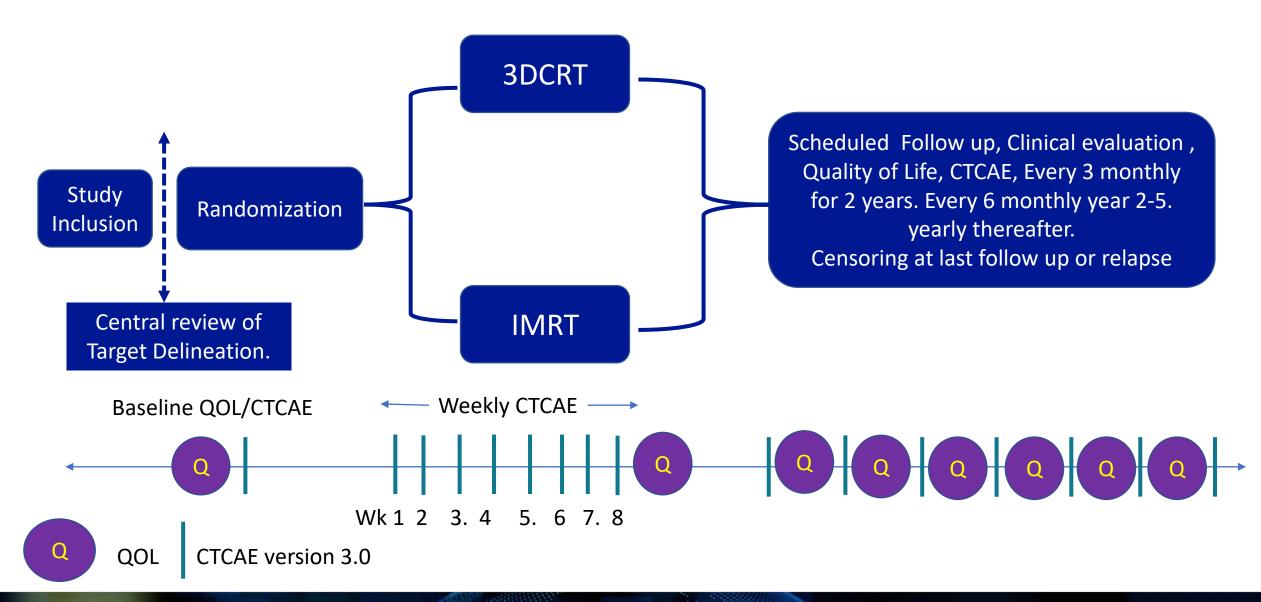
Inclusion Criteria

- Cervical Cancer
- Age >18 years
- Type III Hysterectomy with intermediate or high risk features
- Type I/II hysterectomy necessitating adjuvant CRT

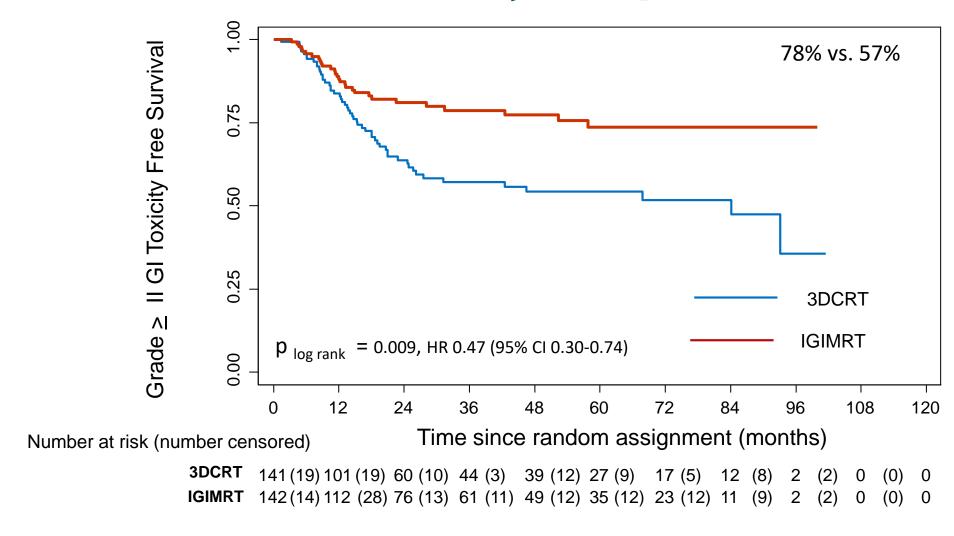
Exclusion Criteria

- Positive Para aortic nodes or indication for extended field RT.
- History of multiple previous abdominal surgeries/radiation
- Any medical condition predisposing to bowel toxicity

Trial Schema: PARCER

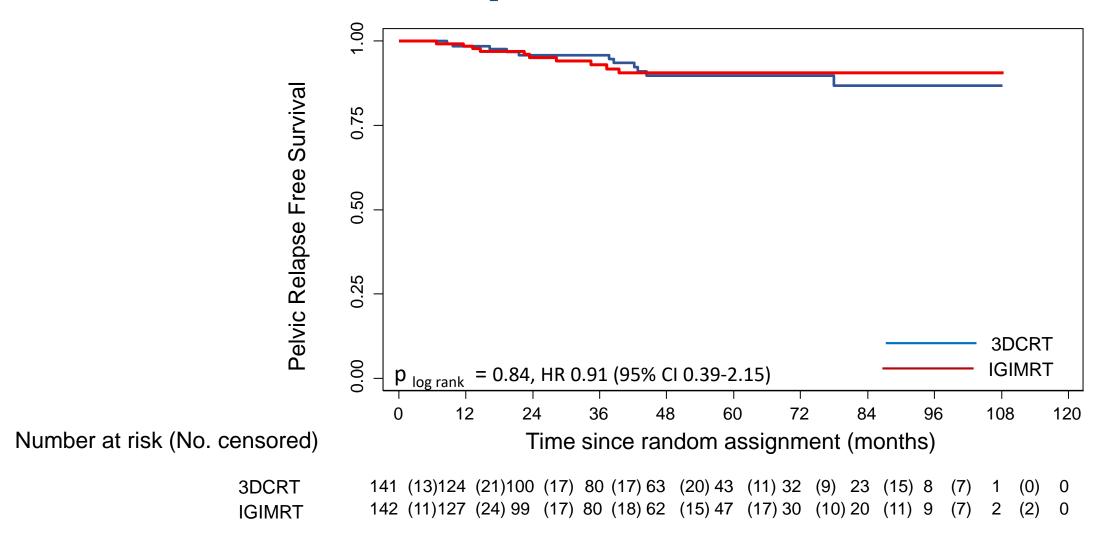


Primary Endpoint

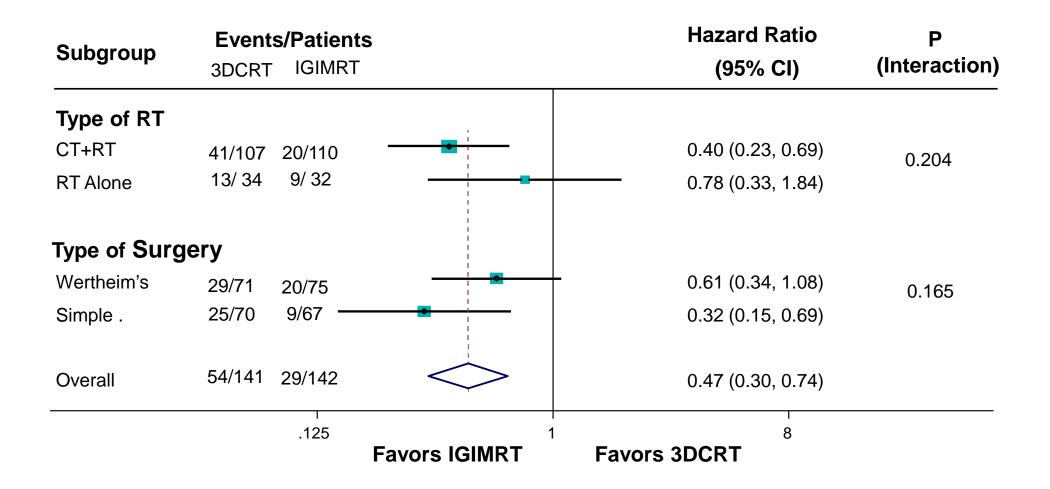


Adjusted for stratification factors- RT type and Sx type

Pelvic Relapse Free Survival



Planned Subgroup Analysis



Conclusions

- IG-IMRT is superior to 3DCRT in reducing Late GI toxicity in women undergoing postoperative pelvic RT.
- Greater Benefit of IG-IMRT in those receiving radio-sensitizing concurrent chemotherapy.
- No difference in tumour control rates in the pelvis with use of IG-IMRT
- IG-IMRT should represent the new standard of care for postoperative pelvic RT in women with gynecological cancers.