Purpose: To determine dose errors inherent in assuming that dose is fixed in the accelerator coordinate system for image guided planning of prostate cancer.

Methods: A 17 prostate patient cohort with 8-13 isocentrically aligned CTs/patient are used. For each patient, a 79.2 Gy, $18 \mathrm{MV}, 7$ beam IMRT plan is developed on the first/primary image reliant on RT0G 0126 objectives. Dose on the remaining images is evaluated in two ways: (A1) Dose is recalculated on each image. (A2) The planned/primary dose distribution is copied to each image. A2 assumes dose is invariant in the accelerator coordinate system. Effects of patient miss-alignment are simulated by evaluating dose with 27 patient shifts per image; 0 and $\pm 5 \mathrm{~mm}$ in left-right, anterior-posterior and superior-inferior directions. Per-image dose differences and dose-volume metrics (prostate-D90, bladder- and rectum-D25) are used to compare A1 and A2. With no shifts, 4D dose accumulated over all images is compared.

Results: The per-image root-mean-square-error percentage error (RMSPE) between A1 and A2 over all shifts and all patients is $1 \%$ for prostate-D90, $1.5 \%$ for rectum-D25, and $4.6 \%$ for bladder-D25. For accumulated dose, the RMSPE values are prostate-D $90=0.7 \%$, rectum$\mathrm{D} 25=12.0 \%$ and bladder-D25=0.7\%. 4 out of 17 patients had large variations. Excluding those patients RMPSE reduced from $12.0 \%$ to $3.0 \%$.

Conclusions: For the patient cohort studied, assumption of shift- and deformation-invariant dose distributions on average introduces $<2 \%$ error in evaluated dose-volume metrics in case of prostate and rectum. Further study is required for the bladder. Use of invariant dose distributions has a potential to reduce online re-planning time and permit pre-planning based on tissue deformation models. (Supported by NIH 5P01CA116602).

