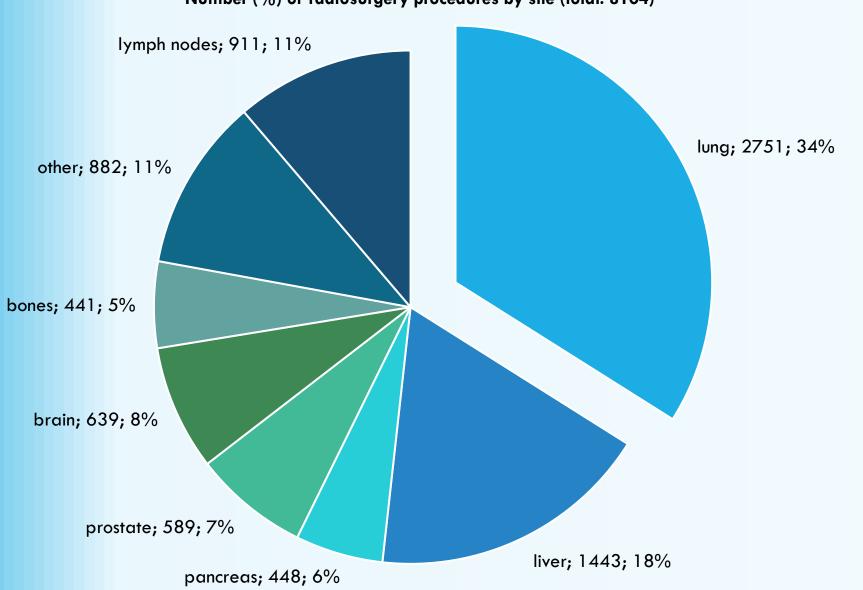


PATIENT RADIATION PROTECTION IN THE CONTEXT OF OLIGOMETASTATIC DISEASE MANAGMENT

Kosmina Domagoj





Number (%) of radiosurgery procedures by site (total: 8104)

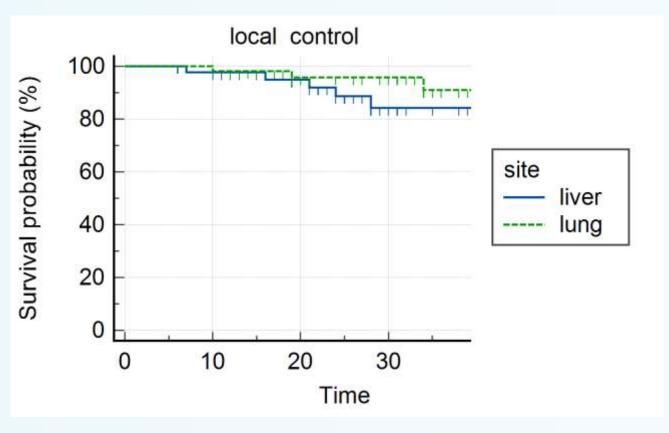
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OLIGOMETASTATIC DISEASE



A stage of cancer between local and systemic widespread disease

- •Typically defined as up to 5 lesions in up to 3 different organ systems
- As technology evolves more lesions can be treated
- Oligoprogression + systemic treatment

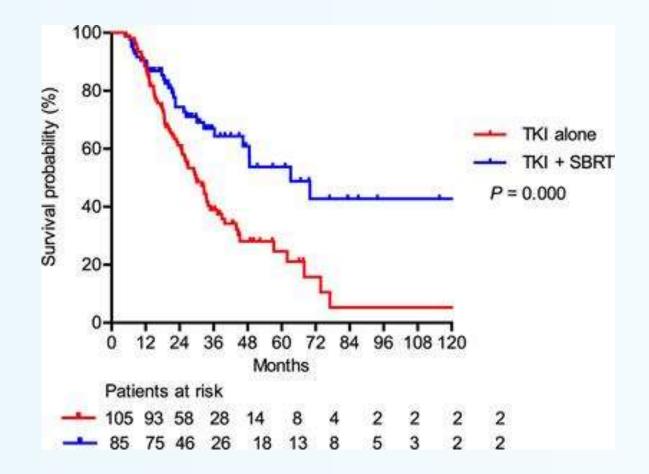


TREATMENT

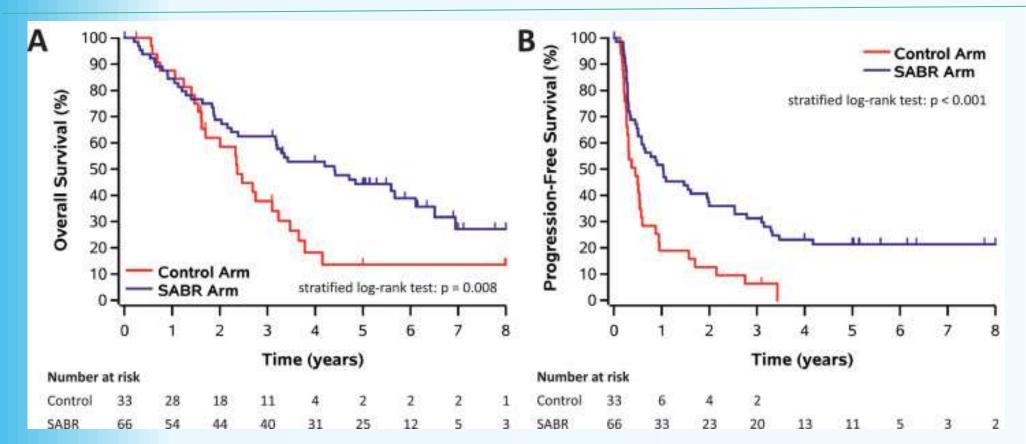


Stereotactic ablative body radiotherapy, SABR or SBRT, has emerged as a dominant treatment modality for metastatic disease

- •Ablative radiation doses to cancer lesion
- Outpatient procedure, can treat from 5 mm to >50 mm lesions
- Effective and low risk of toxicity
- Can only treat what we can "see"
- •Do we care about diagnostic dose?



TREATMENT



CLINICAL INVESTIGATION VOLUME 114, ISSUE 4, P611-616, NOVEMBER 15, 2022

Stereotactic Radiation for the Comprehensive Treatment of Oligometastases (SABR-COMET): Extended Long-Term Outcomes •<u>Stephen Harrow, MBChB, PhD, David A. Palma, MD, PhD, Robert Olson, MD, MSc,X. Melody Qu, MD, MPH,Andrew Warner, MSc, Suresh Senan, MBBS, PhD</u> Published:May 25, 2022DOI:https://doi.org/10.1016/j.ijrobp.2022.05.004



SECONDARY CANCER RISKS VS CANCER PROGRESSION

verywell

Secondary cancer risk increase with dose, to a relative increase of 50% for doses > 1 Gy

However mean time to secondary cancer measured in decades, example 22 years for breast¹

Stage 4 disease mean survival time is measured in years, sometimes months, colorectal cancer for example is ~ 1 year ²

Important to detect secondary lesions early

Non-Small Cell Lung Cancer: **5-Year Survival Rates** 100% 92% 83% 80% 68% 60% 60% 53% 36% 40% 20% 10% <1% 0% A1 A2 A3 B A B A B C Stage 1 Stage 2 Stage 3 Stage 4

Source: American Cancer Society 2017

1 Olsson H, Baldetorp B, Fernö M, Perfekt R. Relation between the rate of tumour cell proliferation and latency time in radiation associated breast cancer. BMC Cancer. 2003 Apr 9;3:11. doi: 10.1186/1471-2407-3-11. PMID: 12697074; PMCID: PMC155538.

2 Rodriguez-Bigas MA, Lin EH, Crane CH. Stage IV Colorectal Cancer. In: Kufe DW, Pollock RE, Weichselbaum RR, et al., editors. Holland-Frei Cancer Medicine. 6th edition. Hamilton (ON): BC Decker; 2003. Available from: https://www.ncbi.nlm.nih.gov/books/NBK13267/



MOTIVATION

Low dose protocols are being investigated for use in follow up after cancer, primarily in the lungs

LDCT have a significantly lower radiation dose, but the tradeoff is that their sensitivity is lower $(0.6-0.9)^{1,2,3}$

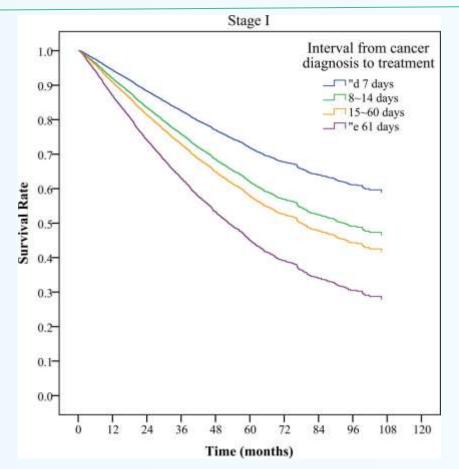
In the context of oligometastatic disease, the risk of not detecting a lesion is by far greater then the benefit of ionising dose reduction

1 Gobi K, Arunachalam VK, Varatharajaperumal RK, Cherian M, Periaswamy G, Rajesh S. The role of ultra-low-dose computed tomography in the detection of pulmonary pathologies: a prospective observational study. Pol J Radiol. 2022 Nov 4;87:e597-e605. doi: 10.5114/pjr.2022.121433. PMID: 36532248; PMCID: PMC9749781.

2 Gheysens, G., De Wever, W., Cockmartin, L. et al. Detection of pulmonary nodules with scoutless fixed-dose ultra-low-dose CT: a prospective study. Eur Radiol 32, 4437–4445 (2022).

3 Paks M, Leong P, Einsiedel P, Irving LB, Steinfort DP, Pascoe DM. Ultralow dose CT for follow-up of solid pulmonary nodules: A pilot single-center study using Bland-Altman analysis. Medicine (Baltimore). 2018 Aug;97(34):e12019. doi: 10.1097/MD.00000000012019. PMID: 30142849; PMCID: PMC6112944.

×



Tsai C, Kung P, Kuo W, et al

Effect of time interval from diagnosis to treatment for non-small cell lung cancer on survival: a national cohort study in Taiwan *BMJ Open* 2020;**10**:e034351. doi: 10.1136/bmjopen-2019-034351



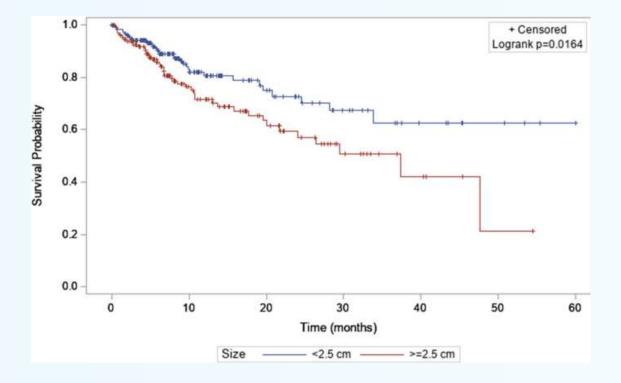
METASTASIS GROWTH AND APPEARANCE

Mean volume doubling time of secondary lesions is ~3 months

Secondary lesions can seed new secondary lesions, even reseed the primary tumor site

For cancer follow up, typical interval is 3 months

What happens if lesion is missed on 1 follow up and then detected on the next



METHODS AND RESULTS

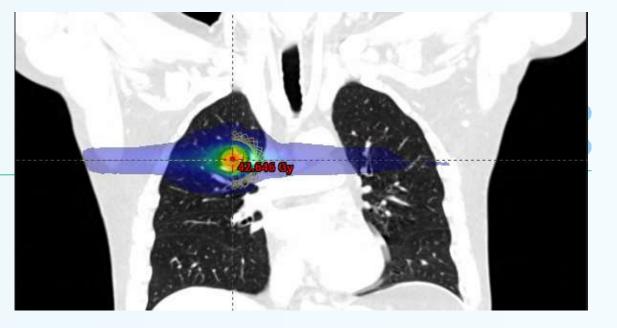
61 lung lesions from 20 patients, DIBH, single fraction

Lesions were isotropically expanded so their volume doubled

Forced density in new GTVs to correspond to original

Plans were calculated using knowledge based planning models, both original and modified

The 1 Gy volume was compared \sim 100x CTDI SDCT



	Original	Double GTV volume
Volume (mean)	0.1 – 2.1 ccm (0.6 ccm)	0.2 – 2.2 ccm (1.2 ccm)
1 Gy volume	1582 cc	2324 сс

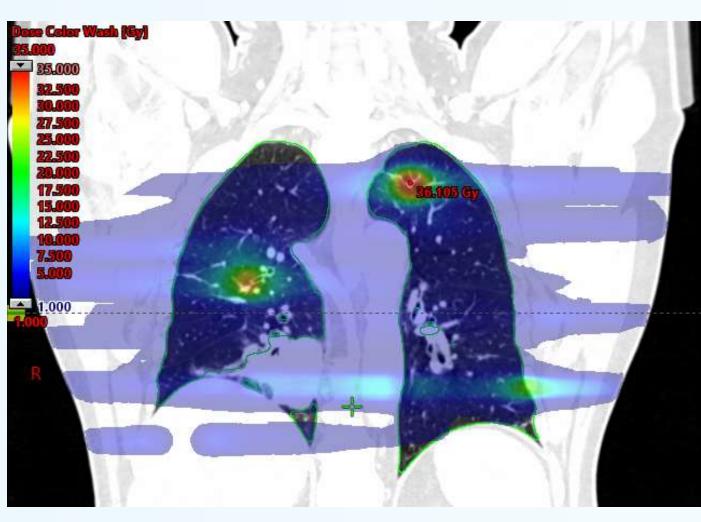
DRL DIAGNOSIS/DIAGNOSTIC



Optimize diagnostic procedure based on patient diagnosis/complaint

For cancer diagnosis focus on quality over lowering the diagnostic dose

Benefit does not outweight the risks for patients with a cancer diagnosis



CONCLUSION



Disease control is based on lesion detection – higher dignostic radiation doses improve detection

Aside from cancer biology risk factors, lower sensitivity of LD-CT can increase total patient radiation dose do to follow up SABR

HIGHER QUALITY DIAGNOSTIC IMAGING CRUTIAL FOR DISEASE MANAGMENT – smaller lesions, better disease control and less total dose to the patient

DRL's should take patient diagnosis into consideration – not just localization



OCCUPATIONAL RADIATION PROTECTION IN INTRAOPERATIVE RADIOTHERAPY

INTRAOPERATIVE RADIOTHERAPY

Low energy (50 kV) dose after tumor resection

Primary locations:

- Breast
- Brain
- Liver
- Rectum
- Spine

Is used for tumor control, iatrogenic dissemination Replaces some adjuvant radiotherapy



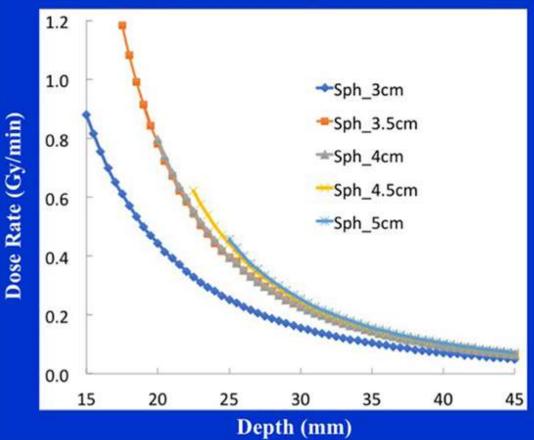
DOSES

50 kV energy

- Cca +30% additional DSB compared to MV external beam therapy
- Typical doses 20-25 Gy to the surface of the applicator
- 30-40 min treatments
- Fast dose fall off in water (soft tissue)
- Dose rate 0.6-0.8 Gy/min at aplicator surface
 - Soft tissue attenuates 99% of dose at 2 cm depth
- Treatment occurs in a shielded (2mm Pb shield walls and door) with no staff present in the room during treatment
- Successfully performed, treatment poses no occupational safety risk

X

Spherical Depth Dose



Intraoperative Radiotherapy With INTRABEAM: Technical and Dosimetric Considerations, Front. Oncol., 26 March 2018, Sec. Radiation Oncology Volume 8 - 2018

OCCUPATIONAL RISKS

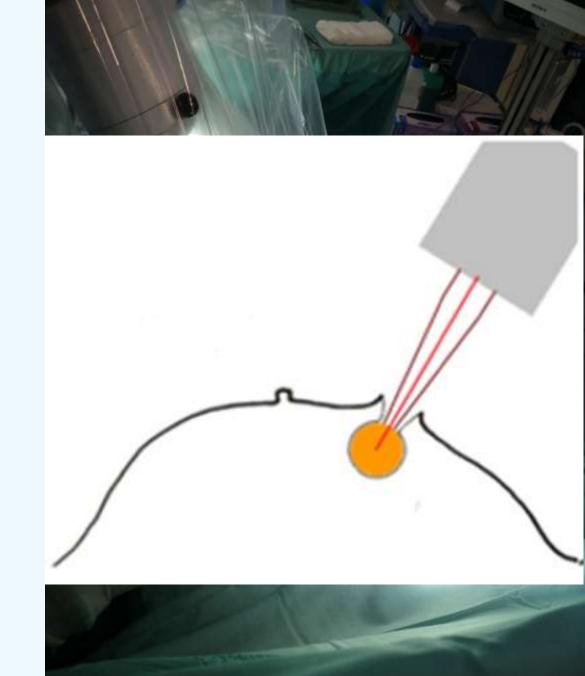
BUT operating theater is not always empty during IORT treatment:

REASONS:

- Anesthesia drip needs to be adjusted (3 min)
- Surgeon forgot phone (2 min)
- Anesthesiologist forgot her papers (4 min)

Is it safe?

- Treatment pause is possible, but do you prolong the patient treatment?
- Dose rates at ~1 m from patient range from 0.03 mGy/min to 2.8 mGy/min, depending on the site and depth of applicator
- High dose rates happen when breast applicator is used, and part of applicator is close to skin



OCCUPATIONAL DOSES

X

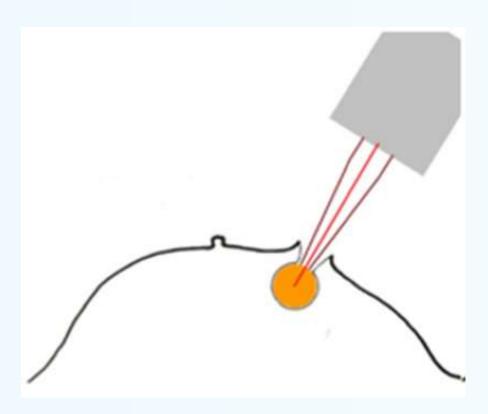
Staff can receive significant doses even with a short stay

A 3 min stay near the patient can lead to a additional dose of 140 µSv

While not significant, can accumulate over working lifetime, even if incidental

Recommendation:

- Always pause treatment when staff is entering theatre
- Staff entering to wear lead approns
- Identify potential risk patients (breast) and apply shielding to patient skin
 - 0.5 mm lead equvalent lead rubber shielding reduces dose rate by a factor of 20
 - Wet surgical gause wrapped around the breast in 1 cm thick layer reduces dose by factor of 6



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THANK YOU FOR YOUR ATTENTION

