

Oligometastatic NSCLC:

The changing role of radiotherapy

Professor Suresh Senan
VU University Medical Center



- The Department of Radiation Oncology at VUMC has a research agreement with Varian Medical Systems.
- S Senan has received speakers honoraria from Varian Medical Systems.



- Use of ablative radiotherapy (SRS, SABR/SBRT)
- Which patients are most likely to benefit?
- SABR/SRS versus other ablative treatments

Issues to address:

Clinical trials

Tumor biology

Immunology

Toxicity issues

SRS- stereotactic radiosurgery; SABR/SBRT – stereotactic body radiotherapy



An Individual Patient Data Meta-Analysis of Outcomes and Prognostic Factors after Treatment of Oligometastatic Non-Small Cell Lung Cancer

Allison B. Ashworth¹, Suresh Senan², David A. Palma¹, Marc Riquet³, Yong Chan Ahn⁴, Umberto Ricardi⁵, Maria T. Congedo⁶, Daniel R. Gomez⁷, Gavin M. Wright⁸, Giulio Melloni⁹, Michael T. Milano¹⁰, Claudio V. Sole¹¹, Tommaso M. De Pas¹², Dennis L. Carter¹³, Andrew J. Warner¹ and George B. Rodrigues¹.

Systematic review of the literature to identify reports.

- 757 NSCLC patients with 1-5 synchronous or metachronous metastases
- Median patient age at diagnosis was 61 years
- 98% of patients had a good performance status
- 2/3 had otherwise early-stage intra-thoracic disease staged IA-IIIB (after excluding metastatic disease)



Median OS of 26 months, 1-year OS 70.2%, and 5-year OS 29.4%.

Surgery was the most commonly used treatment modality for the primary (n=635, 83.9%) and for metastases (n=339 62.3%).

Predictors of OS: synchronous vs. metachronous metastases ($p<0.001$), N-stage ($p=0.002$) and adenocarcinoma histology ($p=0.036$)

Recursive Partitioning Analysis for risk groups;

Low-risk: metachronous metastases (5-year OS 47.8%);

Intermediate risk: synchronous metastases and N0 disease (5-year OS 36.2%);

High risk: synchronous metastases and N1/N2 disease (5-year OS 13.8%).



Table 2. Advantages of Surgery and Stereotactic Radiosurgery for Brain Metastases.

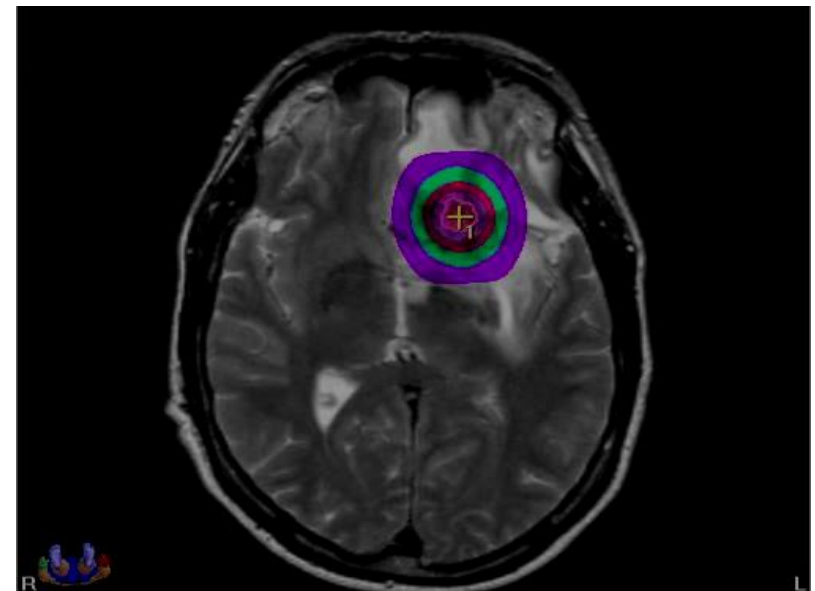
Surgery

Suh J, NEJM 2010

- Treatment of larger lesions (>4 cm in diameter)
- Rapid resolution of mass effect and edema
- Removal of cancer
- Histologic confirmation of cancer
- Rapid tapering of the dose of corticosteroids used to treat symptomatic lesions
- Less intensive follow-up
- Lower risk of radiation necrosis when combined with whole-brain radiation therapy

Stereotactic Radiosurgery

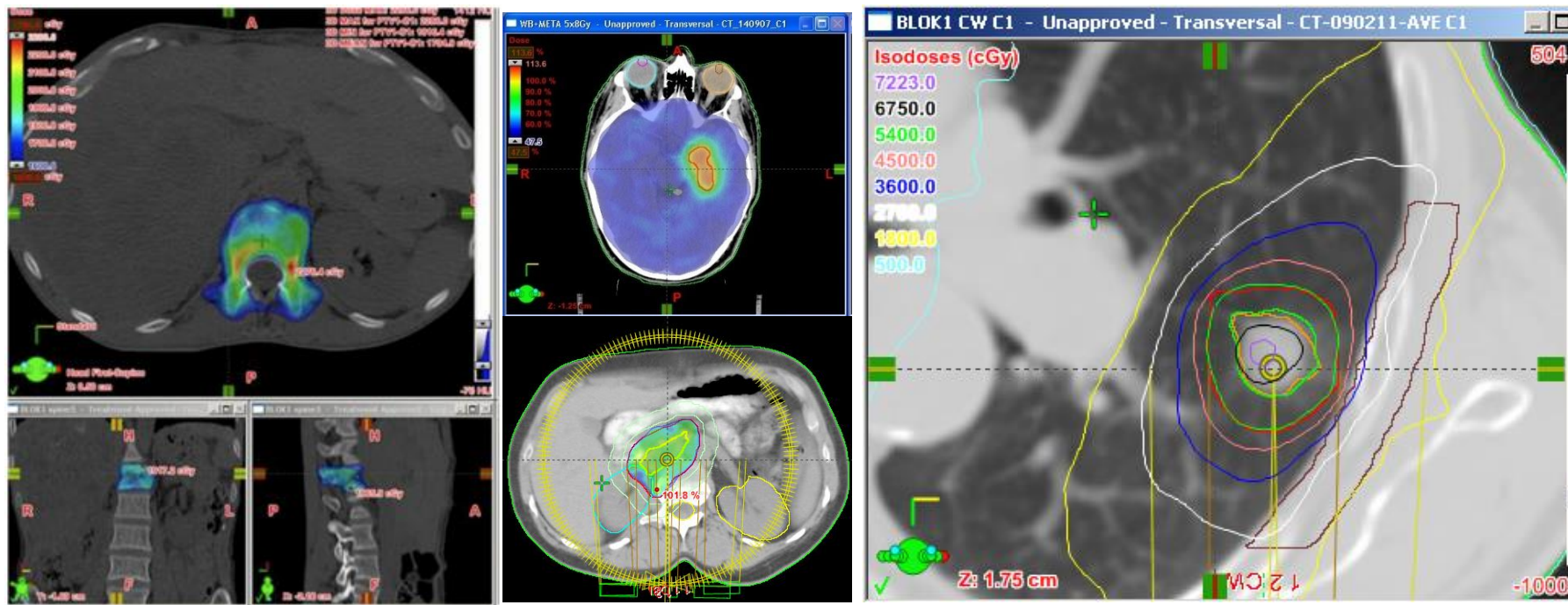
- Treatment of small, deep lesions or eloquent areas
- Minimally invasive or noninvasive approach
- General anesthesia not required
- Outpatient procedure
- Treatment of multiple lesions during same session
- Short recovery time (<1 wk)
- Potential avoidance of whole-brain radiation therapy
- Rapid initiation of systemic therapies



A technique for delivering external beam radiotherapy to an extra-cranial target

- (i) with a high degree of accuracy,
- (ii) using high doses of irradiation,
- (iii) delivered in 1-8 treatment fractions.

Senan, Guckenberger, Ricardi, IASLC textbook 2014



Extracranial Oligometastases: A Subset of Metastases Curable With Stereotactic Radiotherapy

Radiation Series	Year	No.		Local Control (%)	Survival (%)	Site
		Patients	Lesions			
Blomgran et al	1995	31	42	80	Not reported	Liver, lung, and retroperitoneum
Wulf et al	2004	41	51	80	33 ^a	Lung
Hoyer et al (colorectal cancer)	2006	64	141	86 ^a	38 ^a , 13 ^h	Lung, liver, and adrenal
Hof et al	2007	61	71	63 ⁱ	47.8 ⁱ	Lung
Rusthoven et al	2009	47	63	92 ^a	30 ^a	Liver
Rusthoven et al	2009	38	63	96 ^a	39 ^a	Lung
Kang et al (colorectal cancer)	2010	59	78	66 ^j	49 ^j	Multiple
Okunieff et al	2006	49	125	83 ^j	29 ^j	Lung
Katz et al	2007	69	174	57 ^k	24 ^{l,m}	Liver
Lee et al	2009	70	143	71 ^m	47 ⁿ	Liver
Milano et al	2011	121				Multiple ^p
Breast cancer		39		87 ^o	74 ^a , 47 ^o	
All others		82		65 ^o	39 ^a , 9 ^o	
Salama et al	2011	61	111	66.7 ^{q,r}	56.7 ^a	Multiple
Bae et al (colorectal cancer)	2012	41	50	64 ⁱ , 57 ^h	64 ⁱ , 38 ^h	Lung, liver, and lymph node
Norihisa et al	2008	34		90 ^a	84.3 ^a	Lung



- Use of ablative radiotherapy (SRS, SABR/SBRT)
- Which patients are most likely to benefit?
- **SABR/SRS versus other ablative treatments**

Issues to address:

Immunology

Clinical trials

Tumor biology

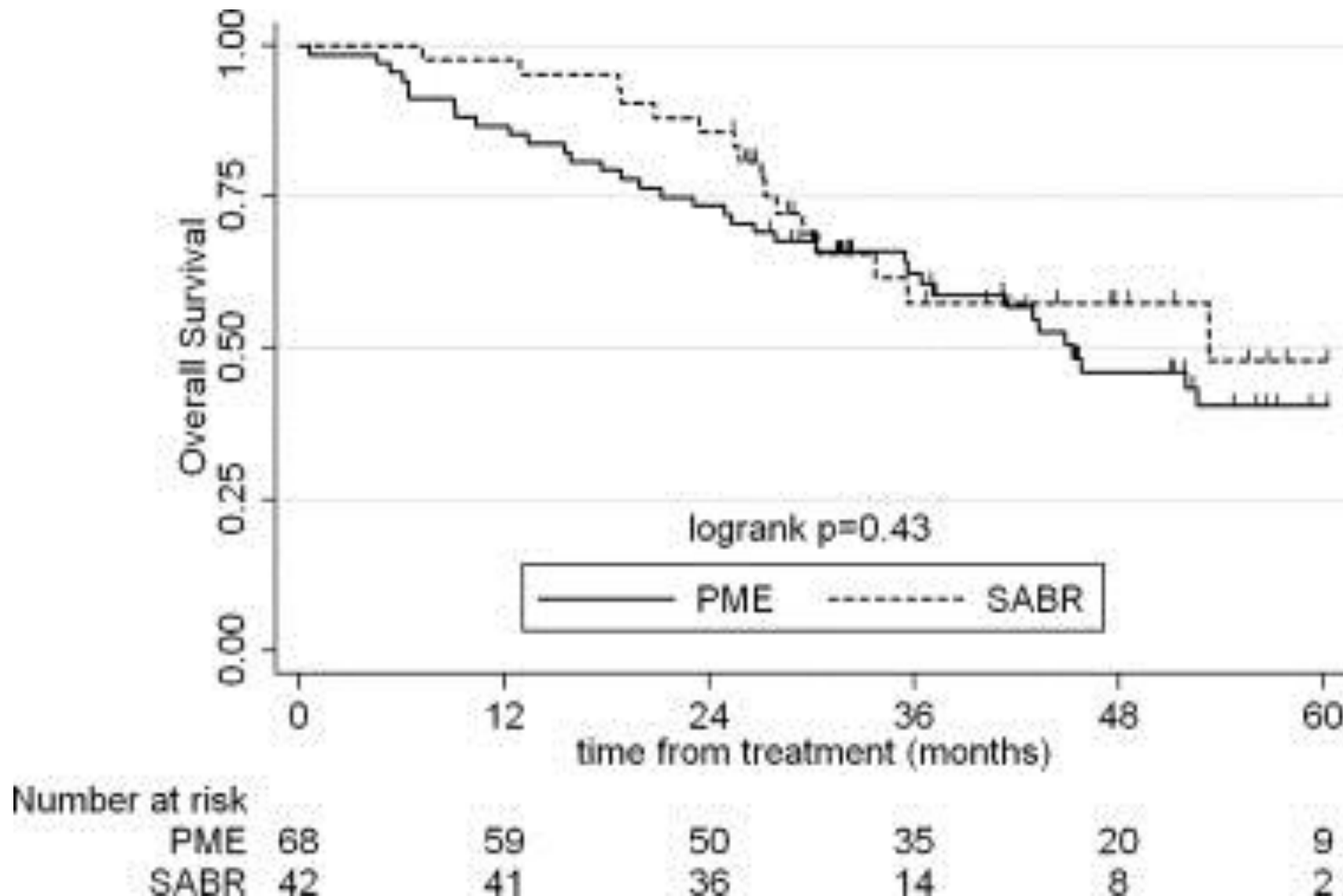
Toxicity issues

SRS- stereotactic radiosurgery; SABR/SBRT – stereotactic body radiotherapy



- Consecutive patients referred to a multidisciplinary team in a university-hospital from 2007-2010.
- Surgery was considered the first choice, and SABR otherwise
- 110 patients (surgery, n=68; SABR, n=42)
- Estimated OS rates at 1, 3 and 5 years:
 - 87%, 62%, and 41% for surgery, and
 - 98%, 60%, and 49% for SABR, respectively (logrank-test, $p=0.43$).
- Local control at two years was 94% (SABR) and 90% (surgery)
- Progression-free survival was 17% at three years





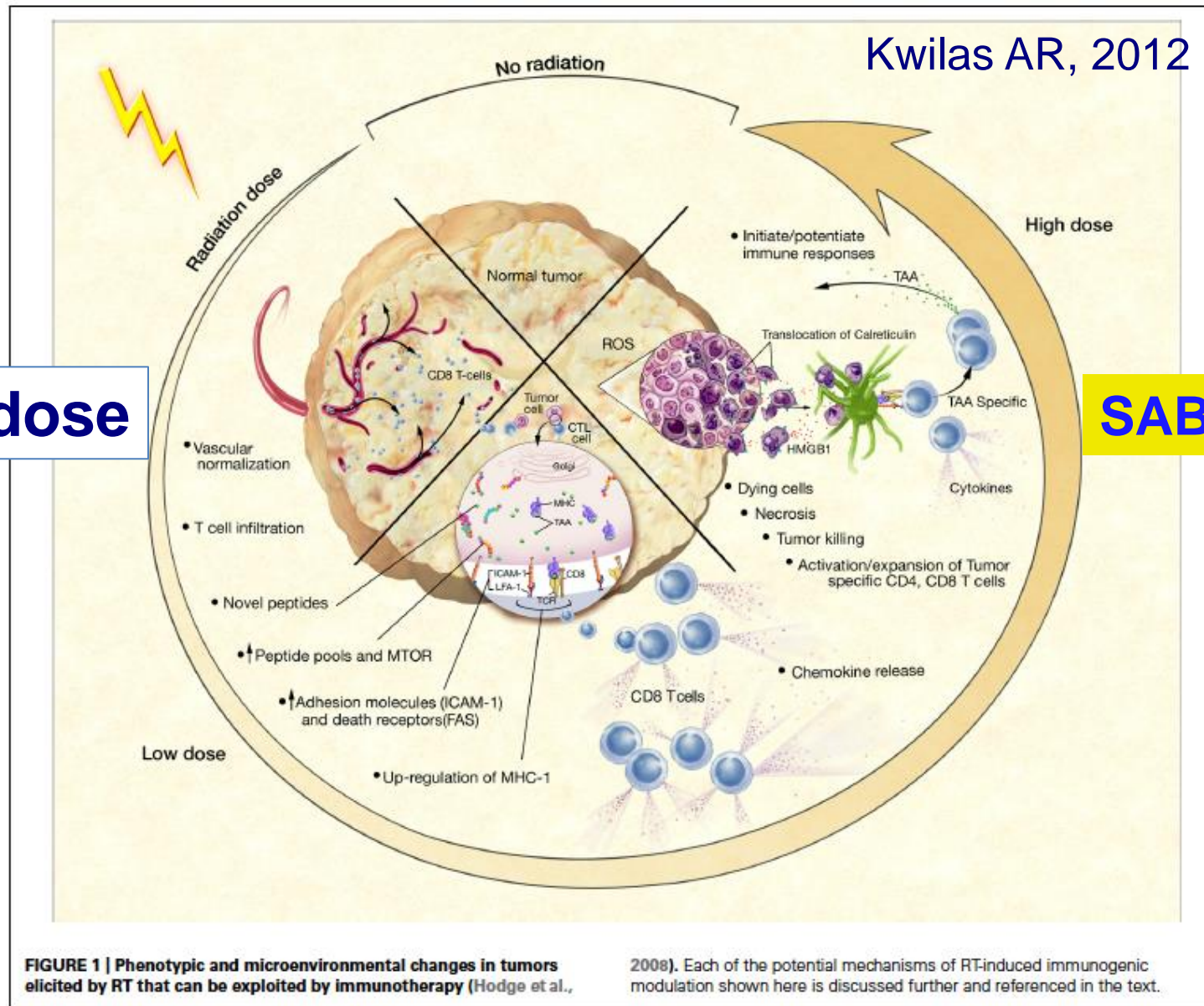
Overall survival, PME (pulmonary metastasectomy) versus SABR (stereotactic ablative radiotherapy).



Kwilas AR, 2012

Low dose

SABR doses



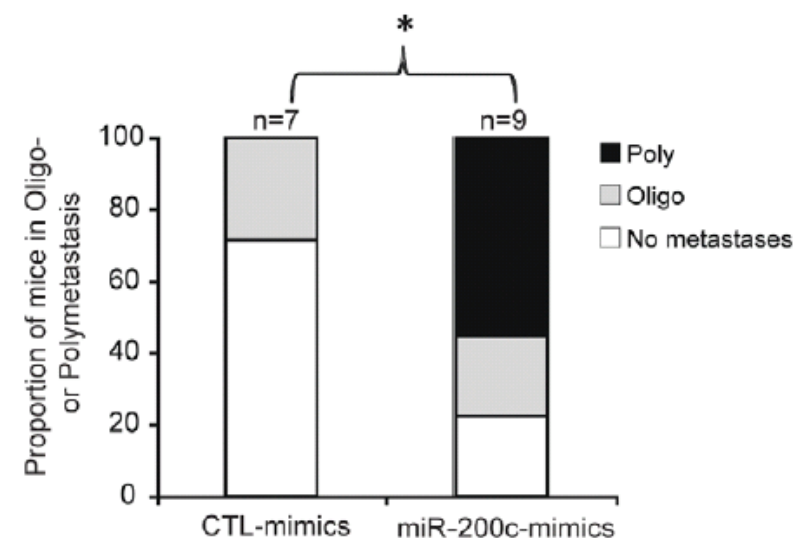
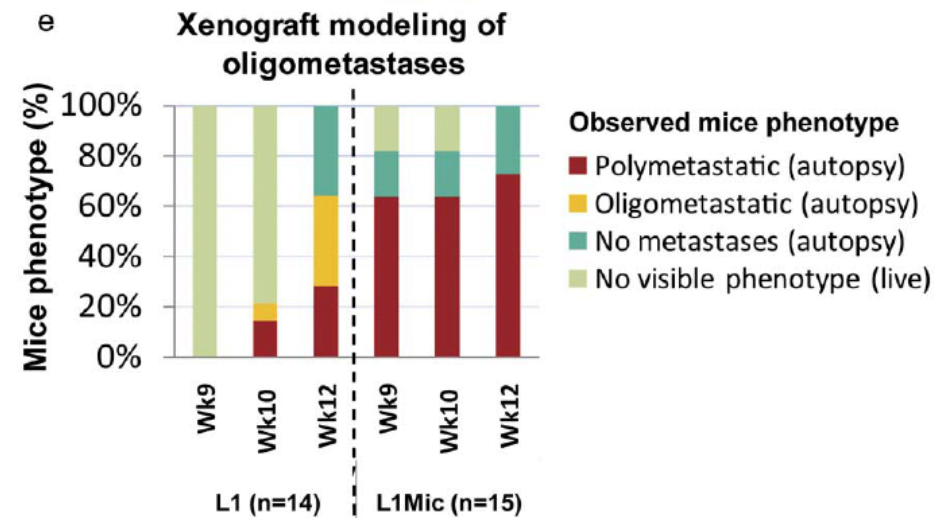
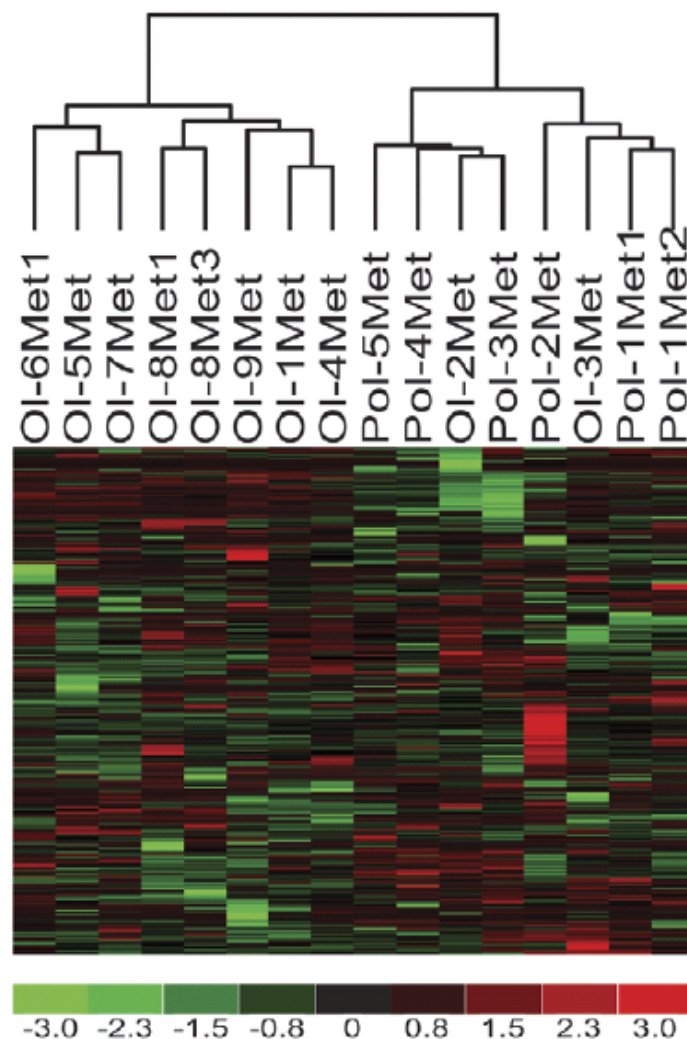
Radiation Therapy to Convert the Tumor into an In Situ Vaccine [Formenti SC, IJROBP 2012]



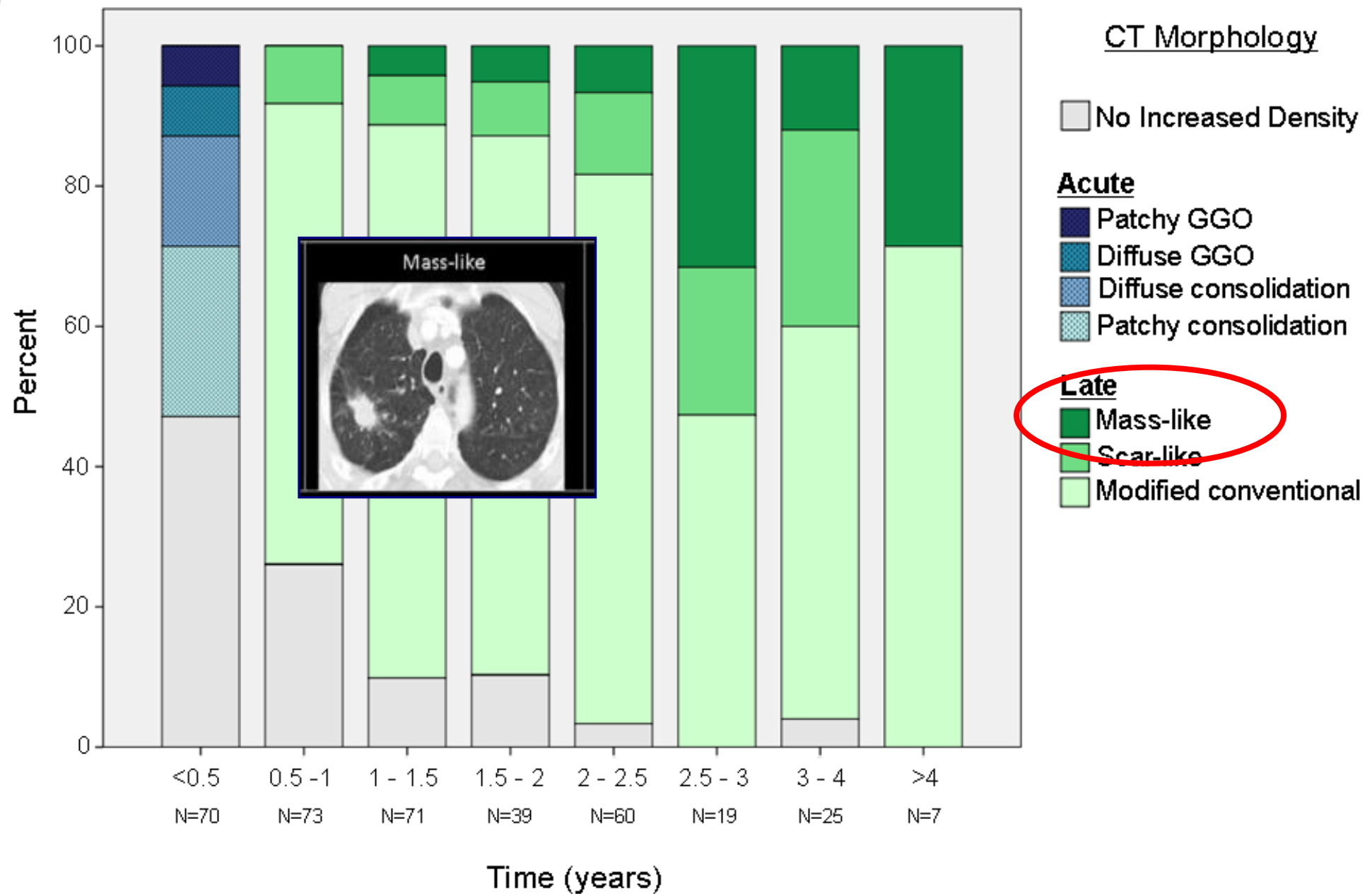


MicroRNA Expression Characterizes Oligometastasis(es)

Yves A. Lussier^{1,2,3,4,*}, H. Rosie Xing^{1,2,5,6,8}, Joseph K. Salama⁸, Nikolai N. Khodarev^{1,5}, Yong Huang^{1,3}, Qingbei Zhang^{3,6}, Sajid A. Khan⁷, Xinan Yang³, Michael D. Hasselle⁵, Thomas E. Darga⁵, Renuka Malik⁵, Hanli Fan⁶, Samantha Perakis⁵, Matthew Filippo⁵, Kimberly Corbin⁵, Younghee Lee³, Mitchell C. Posner⁷, Steven J. Chmura⁵, Samuel Hellman^{2,5}, Ralph R. Weichselbaum^{1,2,5,*}



Post-SABR radiological changes



Systematic review of literature on recurrences

High-risk features (HRF):

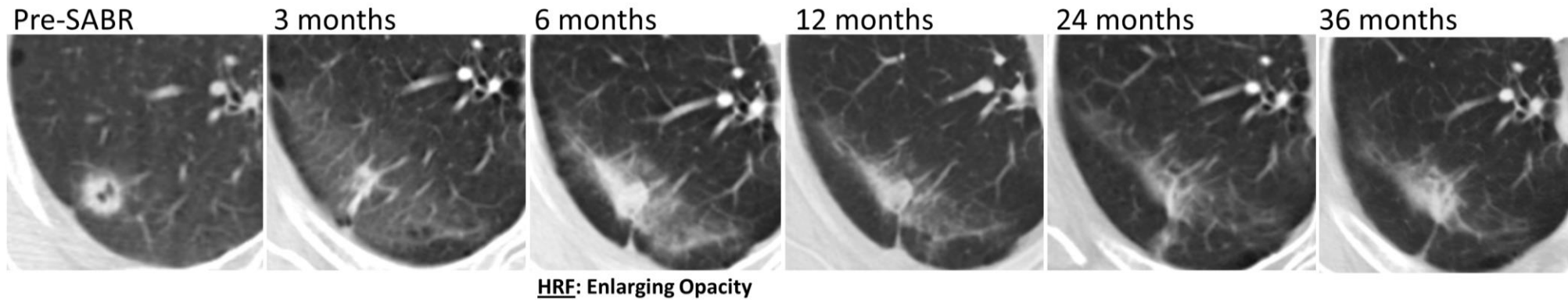
- enlargement of mass
- sequential enlargement on CT
- growing mass after 12 months
- bulging margin
- linear margin disappears
- air bronchograms disappear



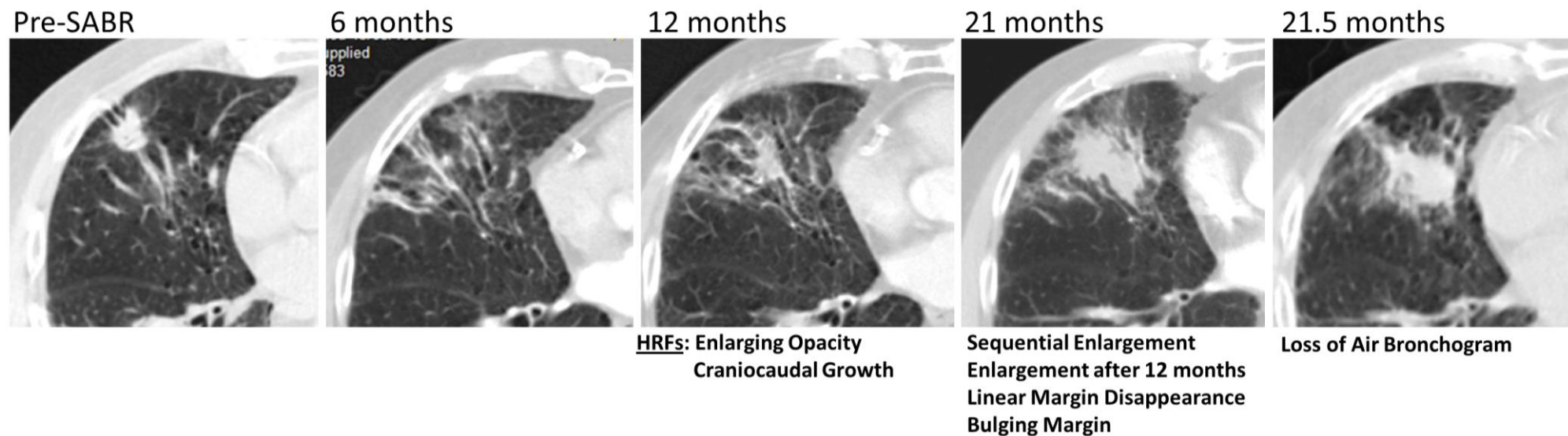
Fibrosis or recurrence after SABR?

Blinded scoring of 12 path. proven recurrences matched with 24 non-recurrences

A. No Recurrence



B. Recurrence



- Use of ablative radiotherapy (SRS, SABR/SBRT)
- Which patients are most likely to benefit?
- SABR/SRS versus other ablative treatments

Issues to address:

Immunology

Clinical trials

Tumor biology

Toxicity issues

SRS- stereotactic radiosurgery; SABR/SBRT – stereotactic body radiotherapy



Issues: treatment beyond progression, tumor flares, oligoprogression.

Weickhardt et al.

Journal of Thoracic Oncology • Volume 7, Number 12, December 2012

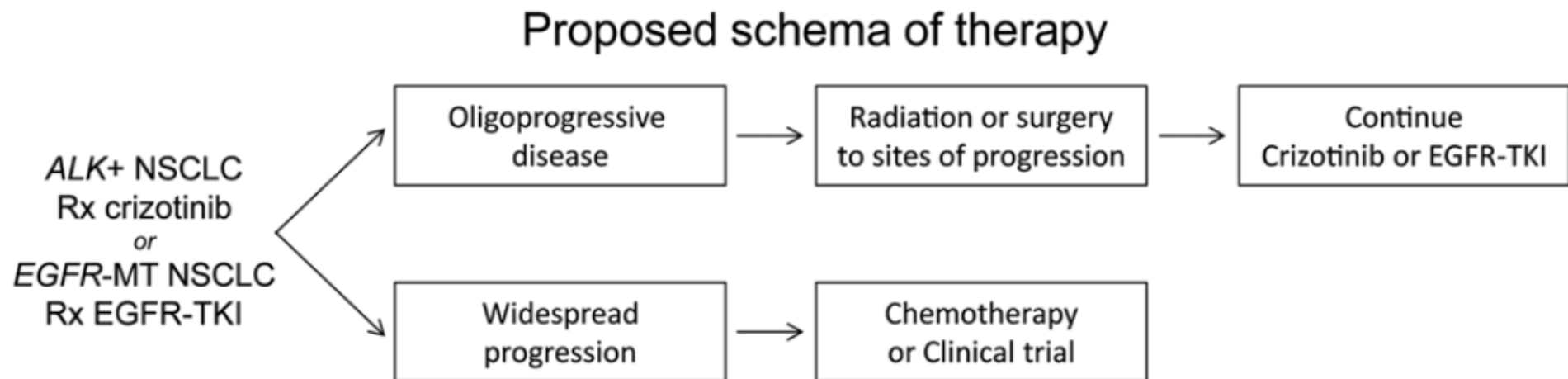


FIGURE 2. Proposed schema for incorporating local ablative therapy into therapy at time of first progression with *ALK+* or *EGFR*-MT NSCLC patients treated with TKI therapy. *ALK+*, anaplastic lymphoma kinase gene rearrangement; *EGFR*-MT NSCLC, epidermal growth factor receptor-mutant non-small-cell lung cancer; TKI, tyrosine kinase inhibitors.



How should oligometastatic progression during TKI be managed?

Local therapies including radiation, radiofrequency ablation, and metastasectomy are established treatment strategies in certain cancers including renal cell carcinoma, sarcoma, and colorectal cancer. Several experiences also support the use of local therapies (surgery, stereotactic radiation) with continued EGFR or ALK inhibition in cases of oligometastatic progression, resulting in minimal toxicity and in months to years of disease control [65].

Prior to proceeding with local therapy, patients should have a full evaluation of the extent of disease, including CNS imaging.

Recommendation 27: In case of oligometastatic progression during TKI treatment, use a local treatment (such as surgery or radiotherapy) and continue/resume TKI

Strength of recommendation: C

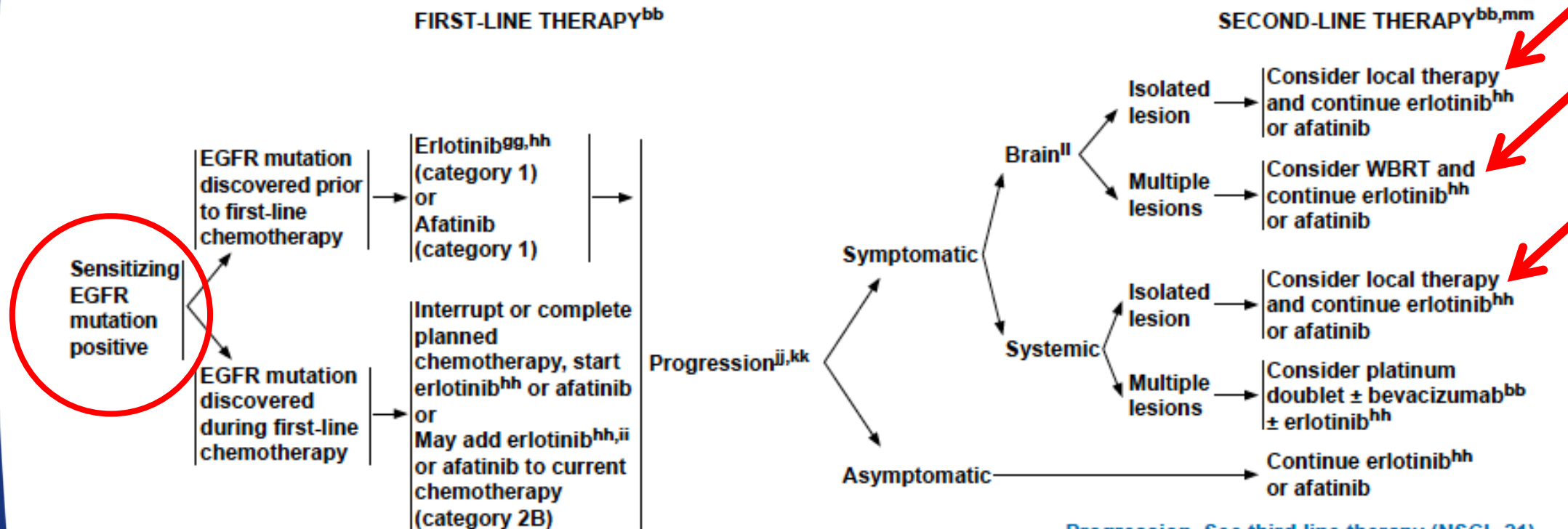
Level of evidence: V



ADENOCARCINOMA, LARGE CELL, NSCLC NOS: SENSITIZING EGFR MUTATION POSITIVE^a

FIRST-LINE THERAPY^{bb}

SECOND-LINE THERAPY^{bb,mm}



Progression See third line therapy (NSCL 21)



Clinical Investigation: Gastrointestinal Cancer

Increased Bowel Toxicity in Patients Treated With a Vascular Endothelial Growth Factor Inhibitor (VEGFI) After Stereotactic Body Radiation Therapy (SBRT)

Brandon M. Barney, MD,* Svetomir N. Markovic, MD, PhD,[†] Nadia N. Laack, MD,*
Robert C. Miller, MD,* Jann N. Sarkaria, MD,* O. Kenneth Macdonald, MD,[‡]
Heather J. Bauer, RN,* and Kenneth R. Olivier, MD*

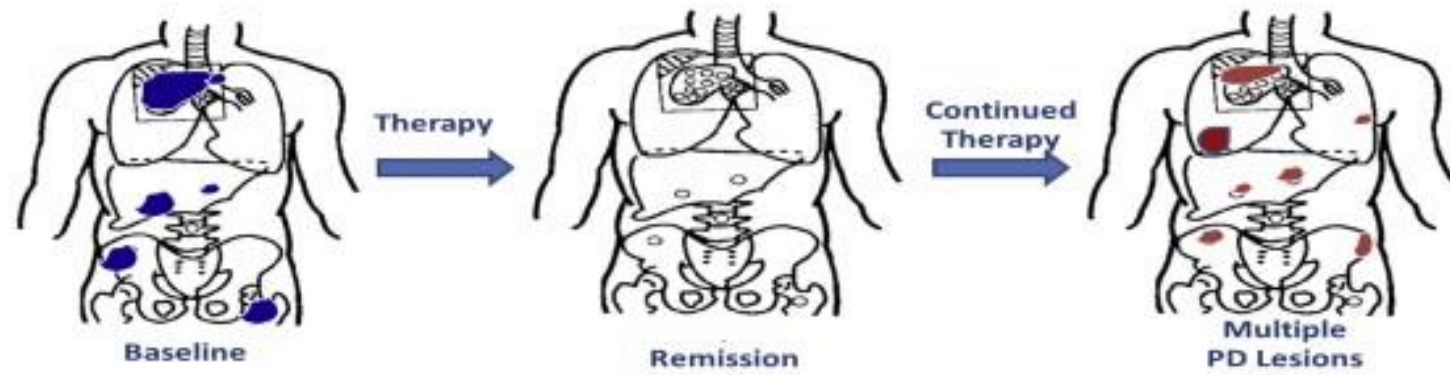
**Department of Radiation Oncology, Mayo Clinic, Rochester, Minnesota; [†]Division of Medical Oncology, Mayo Clinic, Rochester, Minnesota; and [‡]Therapeutic Radiologists Incorporated, Kansas City, Kansas*

Received Mar 29, 2013, and in revised form May 3, 2013. Accepted for publication May 5, 2013



PD Subtype

Systemic PD



- Timing of SABR (consider planned post-ablative systemic therapy; phased SABR)
- Registries; expert radiological assessment post-SABR
- Trial enrollment according to RPA groups (*Ashworth A*)
 - Low-risk: metachronous metastases (5-year OS 47.8%);
 - Intermediate risk: synchronous metastases and N0 disease (5-year OS 36.2%);
 - High risk: synchronous metastases and N1/N2 disease (5-year OS 13.8%).



Thank you for listening

