

LA RADIOTERAPIA
PALLIATIVA CON
TECNICHE SPECIALI
DELLA MALATTIA
METASTATICA



# **ASPETTI GENERALI**

La scelta del trattamento nel malato con oligometastasi: criteri di selezione e valutazione dei risultati

E. Maranzano, *Terni - F. Trippa*, *Terni* 



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# **ASPETTI GENERALI**

- **4 OLIGOMETASTASES** definition
- **4** Patient selection
- **4** Evaluation of therapeutic results



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EDITORIAL 1995

Samuel Hellman Ralph R. Weichselbaum The University of Chicago Chicago, IL

Journal of Clinical Oncology, Vol 13, No 1 (January), 1995: pp 8-10

### **Oligometastases**

- •The *Halsted* theory ( $T \rightarrow N \rightarrow M$ ) proposed that cancer spread is orderly, extending in a contiguous fashion from the primary tumor through the lymphatic to the lymph nodes and then to distant sites (1907).
- A subsequent Hellman theory of systemic disease hypothesis (T → M) proposed that clinically apparent cancer is already a systemic disease (1980).
- A third *Hellman* & *Weichselbaum'* theory of spectrum hypothesis: cancer range between disease that remains localized and disease that is systemic at time of diagnosis → multistep nature of cancer progression (1995).

The <u>occasional success of surgical excision</u> of one or a small number of pulmonary mets, brain mets or hepatic mets (e.g., 25% of cure after hepatic resections for metastatic colorectal cancer) lets to hypothesize the theory of <u>oligometastases</u>

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**Oligometastases** 

### Hellman & Weichselbaum suggested that

for many cancers a few metastases exist at first, before the malignant cells acquire widespread metastatic potential.

### Consequently,

if radical intervention (Surgery or Stereotactic Body Radiotherapy - SBRT) could be delivered during an oligometastatic phase, the intervention could change disease progression in pts who would otherwise have been treated palliatively in most settings (the hierarchy theory)



# Solitary Metastases: Illusion Versus Reality

Philip Rubin, MD, Ralph Brasacchio, MD, and Alan Katz MD, MPH

"Suddenly a solitary horseman appeared on the horizon, then another, then another...in a few moments a whole crowd of horsemen swooped down upon him."—Leacock

### Can a solitary nodule truly be the only metastatic lesion? or is it the

first nodule/horseman on the horizon before the horde appears?







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M1: a solitary metastasis in a single organ

M2: oligometastases, designate number and limited to 1 organ

(5 nodules, 5 cm in total)

### Serum molecular markers as follows:

**S0**: not detectable

**S1**: detectable, low level

**S2**: intermediate level

S3: high level

A. no systemic signs: minimal 5% weight loss, minimal lab abnormalities.

**B. systemic signs**: 100% weight loss, cachexia, fevers nexplained, lab abnormalities, (i.e. altered lung function, abnormal liver enzymes)





# Solitary Metastases: Illusion Versus Reality

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### THE MAJOR METASTATIC ORGAN SITES

- Solitary lung mets
- Solitary liver mets
- Solitary brain mets
- Solitary bone mets
- **4** Other solitary site mets:

H&N, eye & orbit, ovary & vagina, heart, intestines

COMMENTS AND CONTROVERSIES

# Extracranial Oligometastases: A Subset of Metastases Curable With Stereotactic Radiotherapy

Kimberly S. Corbin, Samuel Hellman, and Ralph R. Weichselbaum, University of Chicago Medical Center, Chicago, IL

### **OLIGOMETASTASES**

### **Definition**

An intermediate state of cancer spread between localized disease and wide spread mets

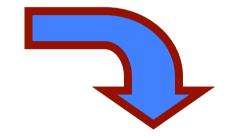
the implication is that oligometastatic disease may be cured with metastasis-directed therapy

## (SBRT)

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Kimberly S. Corbin, Samuel Hellman, and Ralph R. Weichselbaum, University of Chicago Medical Center, Chicago, IL

**OLIGOMETASTASES** 



Stereotactic Body Radiotherapy (SBRT)

## (SBRT)

# Extracranial Oligometastases: A Subset of Metastases Curable With Stereotactic Radiotherapy

Kimberly S. Corbin, Samuel Hellman, and Ralph R. Weichselbaum, University of Chicago Medical Center, Chicago, IL

### **OLIGOMETASTASES**

- Incidence has not been well established
- **Prognosis** can be derived from surgical reports (*lung mets from sarcoma*, *liver mets from colorectal cancer*, *oligometastasis from breast cancer*)
- → this subset of patient can be cured in about 15-20% of cases

### Diagnosis

- → Potential **differential genetic signature** between samples isolated from pts with few or many mets (e.g., microRNA 200c expression or microRNA signature).
- → Hierarchy to the appearance of metastatic sites (i.e., early mets may be of limited nature and early therapy could prevent future spread)
- → Potential increase of early identification of oligometastasis with PET-CT (e.g., NSCLC with mets to adrenal gland at diagnosis)

### (SBRT)

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# **Incidence of oligometastases**

**Lung** met from sarcoma

19% one met

Gadd MA et al: Ann Surg 1993

**Liver**-only mets from colorectal cancer

49% one met 38% ≤ 3 mets

Kienski et al: Ann Surg Oncol 2010

**Distant** failure from breast cancer

16% (median 1,7)

Dorn P et al: Int J Radiat Oncol 2011

**Distant** failure from lung cancer

19%

McManus MP et al: Int J Radiat Oncol Biol Phys 2001

## (SBRT)

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# Extracranial Oligometastases: A Subset of Metastases Curable With Stereotactic Radiotherapy

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**SBRT** treatment of limited metastases has shown promising local control rates for treated metastases, ranging from <u>67% to</u> 95%

Two- to 3-year survival rates have been reported in the range of 30% to 64%

SBRT results compare favorably with surgical results.

SBRT is less invasive than surgery and may be more broadly applicable to greater numbers of tumors in various organs

## (SBRT)

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- Diagnosis
- → Potential increase of early identification of oligometastasis with PET-CT (e.g., NSCLC with mets to adrenal gland at diagnosis)
- → Potential differential genetic signature between phenotype isolated in oligometastatic pts (e.g., microRNA 200c expression or microRNA signature).





**Review Article** 

# Oligometastases and Oligo-recurrence: The New Era of Cancer Therapy

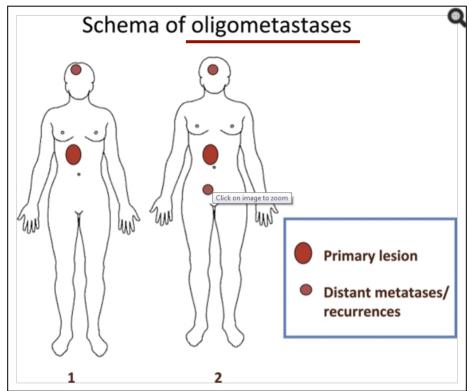
Yuzuru Niibe\* and Kazushige Hayakawa

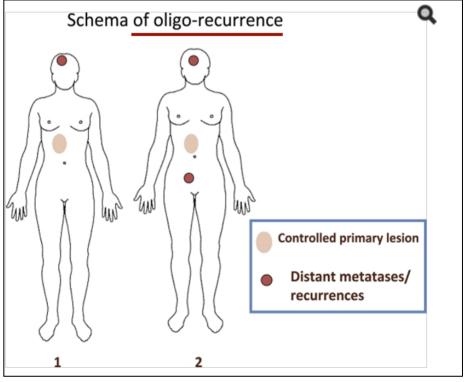
Department of Radiation Oncology, Kitasato University School of Medicine, Sagamihara, Kanagawa, Japan

### **OLIGO-RECURRENCE**

### Patient with a limited number of metastases and controlled primary tumor

Table 1.		
Oligometastases and oligo-recurrence		
	Oligometastases	Oligo-recurrence
Reference	Hellman and Weichselbaum	( <u>1</u> ) Niibe et al. ( <u>2,3,4</u> )
Primary lesion	Uncontrolled/controlled	Controlled
No. of distant/metastases/recur	rences One to several	One to several (one is better)







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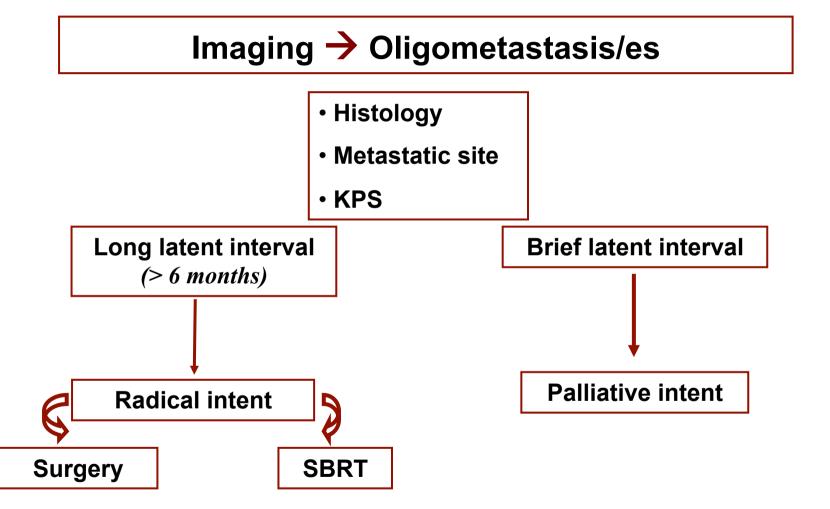
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- **4** Evaluation of therapeutic results

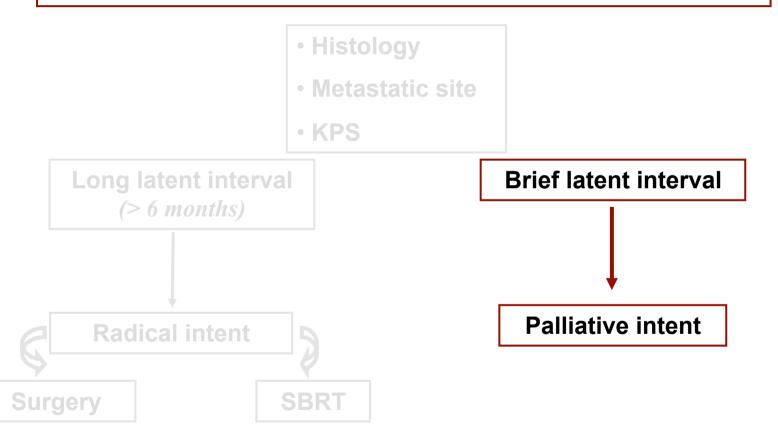
### 3 distinct cohorts of pts with oligometastases

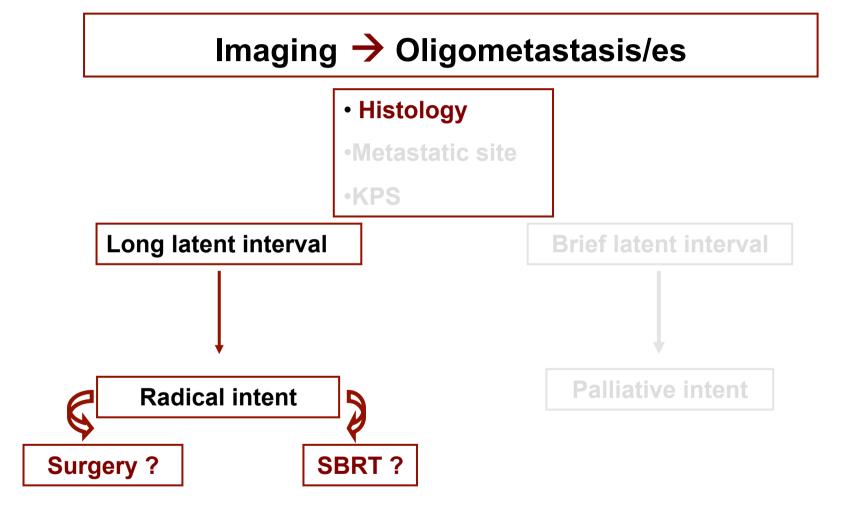
- those who <u>present with</u> oligometastatic disease
- those with induced oligometastatic disease <u>after cytoreductive</u>
  <u>therapy</u>
- those with relapsed oligometastatic disease <u>after curative</u> <u>locoregional therapy</u>

These different groups probably have <u>different prognoses</u>, so therapeutic <u>approaches might differ</u>



# Imaging → Oligometastasis/es





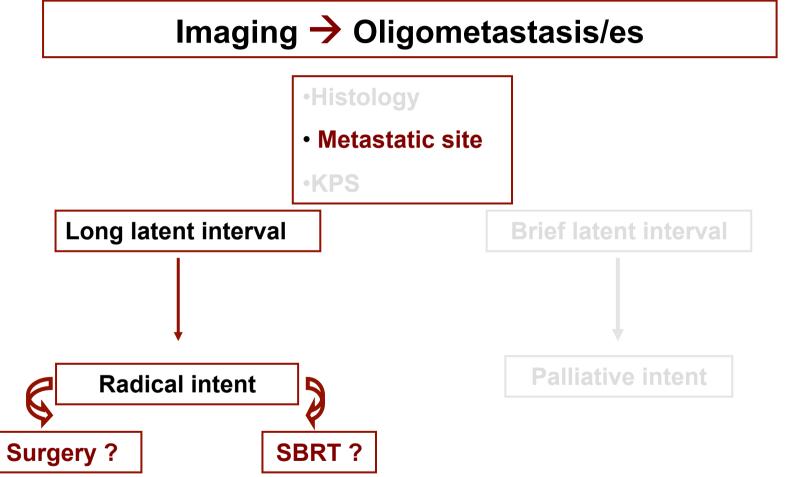
### On the basis of HISTOLOGY

- → Better survival for breast cancer with respect to colorectal & lung
- → Radioresponsive *vs* radioresistant histologies

# Surgery for (?)\*

- kidney clear cell tumor,
- melanomas, or
- sarcomas

<sup>\*</sup> Please note the ablative radiation effect that is not necessary related to radiosensibility!



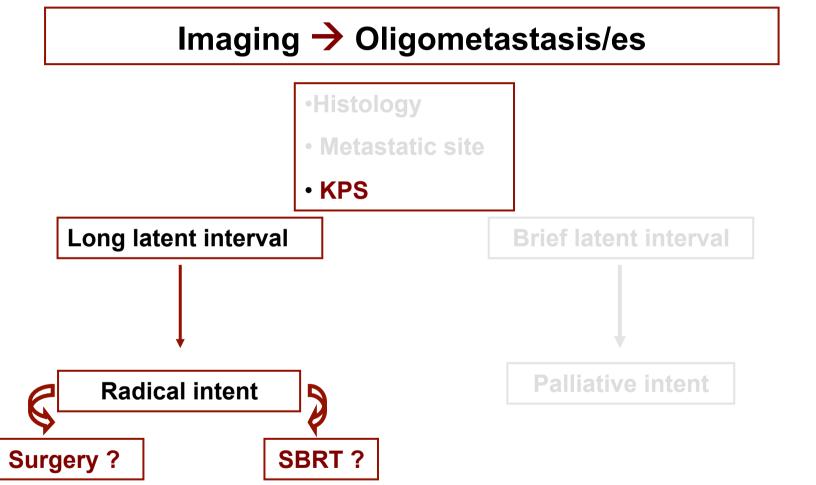
### On the basis of METASTATIC SITE

### Surgery

- for peripheral lung mets →
- for easy accessible liver masses →
- when spinal stabilization is necessary →
- to remove bone impingement to the spine →

### Radiation therapy

in other cases



### On the basis of PERFORMANCE STATUS

## **Surgery**

• for good PSK pts?



# **Radiation therapy**

in the other cases ?!



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# **ASPETTI GENERALI**

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### **EVALUATION OF THERAPEUTIC RESULTS**

- **4** Local control
- **♣ Disease-free survival** (& Freedom from widespread distant metastasis!)
- **4** Toxicity rate
- **QoL**

### **EVALUATION OF THERAPEUTIC RESULTS**

**Local control rate: 67-95%** 

Note that: most in-field recurrence occur (!)

- → 2-year in-field control rate (i.e., duration of response) ~ 20%
- Disease-free survival (& Freedom from widespread distant metastasis?)
- **4** Toxicity rate
- 4 QoL

### Oligometastases Treated With Stereotactic Body Radiotherapy: Long-Term Follow-Up of Prospective Study

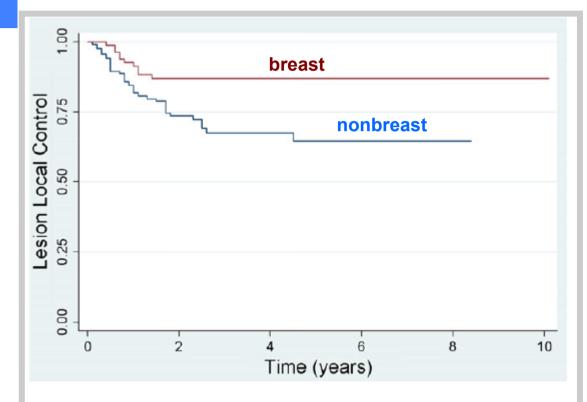
Michael T. Milano, M.D., Ph.D., \* Alan W. Katz, M.D., M.P.H., \*

Int J Radiation Oncol Biol Phys, Vol. 83, No. 3, pp. 878–886, 2012

### LOCAL CONTROL (LC)

The breast cancer patients had a 2-, 4-, and 6-year lesion LC rate of 87%;

the nonbreast cancer patients had a 2-, 4-, and 6-year lesion LC rate of 74%, 68%, and 65%, respectively.



**Fig. 2.** Kaplan-Meier actuarial lesion local control for breast cancer (red line) and nonbreast cancer (blue line) patients. A color

- **♣ CT** and MRI are routinely used to evaluate response to therapy, with RECIST criteria.
- **♣** Postsurgical and postradiotherapy <u>fibrosis</u> and <u>necrosis</u> of both malignant tissue as well as surrounding normal tissue, often making it difficult to differentiate between malignant and non-malignant tissues in surveillance CT and MRI imaging

Solanki et al. Radiation Oncology 2012, 7:216 http://www.ro-journal.com/content/7/1/216



RESEARCH Open Access

The utility of FDG-PET for assessing outcomes in oligometastatic cancer patients treated with stereotactic body radiotherapy: a cohort study

### **EVALUATION OF THERAPEUTIC RESULTS**

- **Local** control
- **♣ Disease-free survival** (& Freedom from widespread distant metastasis!)
- **4** Toxicity rate
- 4 QoL

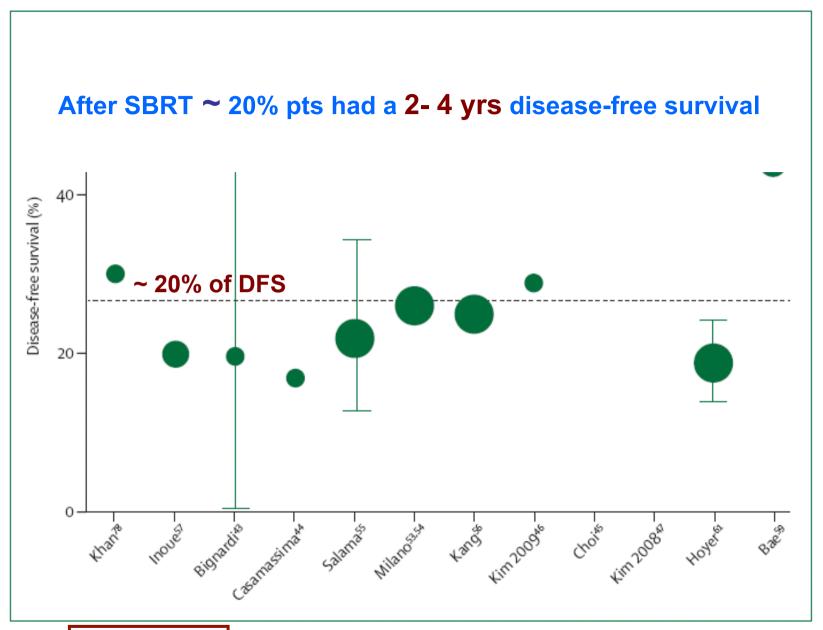


Figure 2 Disease-free survival in patients with oligometastatic disease at 17–48 months' follow-up

Dotted line represents mean proportion of patients who were disease free at the reported timepoint, weighted for number of patients in each cohort. Error bars represent 95% confidence intervals.

- **Local** control
- **↓ Disease-free survival (&** <u>Freedom from widespread</u> <u>distant metastasis</u>!)
- **4** Toxicity rate
- 4 QoL

## Oligometastases Treated With Stereotactic Body Radiotherapy: Long-Term Follow-Up of Prospective Study

Int J Radiation Oncol Biol Phys, Vol. 83, No. 3, pp. 878–886, 2012

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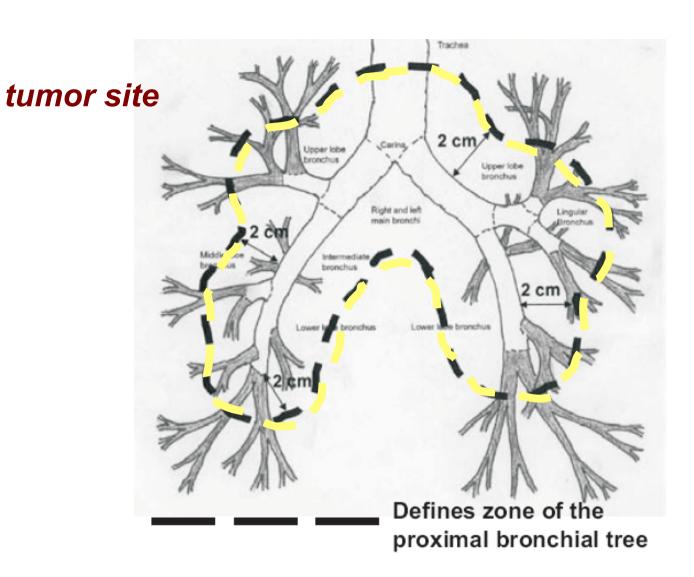
#### **EVALUATION OF THERAPEUTIC RESULTS**

## Freedom from widespread distant metastasis survival

Widespread distant metastases are defined as <u>distant progression</u> not amenable to resection or locally ablative therapy

(i.e., SBRT, stereotactic radiosurgery, radiofrequency ablation, embolization).

- **Local** control
- ♣ Disease-free survival (& Freedom from widespread distant metastasis!)
- **4** Toxicity rate
- **→**Toxicity is directly correlated to:
- administered dose,
- \* tumor volume,
- \* tumor site.
- 4 QoL



Reprinted with permission © 2008 American Society of Clinical Oncology. All rights reserved. Timmerman R, et al. J Clin Oncol 24, 2006:4833-4839.

- **4** Local control
- **♣** Disease-free survival (& Freedom from widespread distant metastasis!)
- **4** Toxicity rate
- → Toxicity is low if *radiation dose constraints* to normal tissues are respected
- 4 QoL



## NCCN Guidelines™ Version 1.2011 Non-Small Cell Lung Cancer

Table 5. Normal Tissue Dose Volume Constraints for SBRT\*

Table 5. Normal	lissue Dose voiu	ine constraints i	OI SBK I	
OAR	1 Fraction	3 Fractions	4 Fractions	5 Fractions
Spinal cord	14 Gy	18 Gy (6 Gy/fx)	26 Gy (6.5 Gy/fx)	30 Gy (6 Gy/fx)
Esophagus	15.4 Gy	30 Gy (10 Gy/fx)	30 Gy (7.5 Gy/fx)	32.5 Gy (6.5 Gy/fx)
Brachial plexus	17.5 Gy	21 Gy (7 Gy/fx)	27.2 Gy (6.8 Gy/fx)	30 Gy (6 Gy/fx)
Heart/ pericardium	22 Gy	30 Gy (10 Gy/fx)	34 Gy (8.5 Gy/fx)	35 Gy (7 Gy/fx)
Great vessels	37 Gy	39 Gy 13 Gy/fx	49 Gy 12.25 Gy/fx	55 Gy 11 Gy/fx
Trachea/ Large Bronchus	20.2 Gy	30 Gy (10 Gy/fx)	34.8 Gy (8.7 Gy/fx)	40 Gy (8 Gy/fx)
Rib	30 Gy	30 Gy (10 Gy/fx)	31.2 Gy (7.8 Gy/fx)	32.5 Gy (6.5 Gy/fx)
Skin	26 Gy	30 Gy 10 Gy/fx	36 Gy (9 Gy/fx)	40 Gy 8 Gy/fx
Stomach	12.4 Gy	27 Gy 9 Gy/fx	30 Gy (7.5 Gy/fx)	35 Gy 7 Gy/fx

- **Local** control
- **♣** Disease-free survival (& Freedom from widespread distant metastasis!)
- **4** Toxicity rate
- **QoL**

QoL

Since SBRT is associated with low toxicity rates with respect to surgery, the QoL benefit is probably greater

(Tree AC et al Lancet Oncol 2013)

#### Canadian-led International Development of a European Organization for Research and Treatment of Cancer Quality of Life Module for Malignant Spinal Cord Compression: Results of Phase I

Radiation Oncology
UNIVERSITY OF TORONTO Gunita Mitera<sup>1</sup>, Andrew Loblaw<sup>1</sup>, Arjun Sahgal<sup>1</sup>, Brita Danielson<sup>2</sup> 🎇 Sunnybrook



<sup>1</sup>Sunnybrook Odette Cancer Centre, Toronto, ON, Canada, <sup>2</sup>Cross Cancer Institute, Edmonton, AB, Canada

## **Objective**

To develop a relevant set of items assessing quality of life (QOL) issues in patients with malignant spinal cord compression (MSCC), not sufficiently covered by the European Organization for Research and Treatment of Cancer (EORTC) C15-PAL core questionnaire.

Table 1: Top 10 QOL issues ranked by patients (n=35)

Patients' Rank	QOL Issues	Freq (%)	HCPs' Rank
1	Have you had difficulty performing self- care (i.e. bathing, dressing)?	48.6	4
2	Did you have trouble controlling your bladder?	42.3	3
3	Did you have lower back pain?	42.3	7
4	Have you had difficulty in carrying out usual daily tasks (i.e. grocery shopping, housework)?	40.0	N/A
5	Have you worried about becoming dependent on others because of your illness?	40.0	6
6	Have you worried about becoming bed- bound because of your illness?	31.4	10
7	Did you have upper back pain?	28.6	N/A
8	Did you have to modify your daily activities because of your illness?	28.6	9
9	Have you worried about loss of mobility because of your illness?	28.6	7
10	Did you hope treatment would reduce pain as much as possible?	28.6	N/A

<sup>1 =</sup> Top priority issue for patients Items included within the red box are items ranked by both groups to be in the top 10

Table 2: Top 10 QOL issues ranked by health care providers (n=62)

HCPs' Rank	QOL Issues	Freq (%)	Patients' Rank
$\begin{pmatrix} 1 \end{pmatrix}$	Were you able to walk without assistance?	49.3	N/A
2	Did you have weakness of both legs?	47.9	N/A
3	Did you have trouble controlling your bladder?	45.2	2
4	Have you had difficulty performing self- care (i.e. bathing, dressing)?	39.7	1
5	Did you experience leakage of bowels?	35.6	N/A
6	Have you worried about becoming dependent on others because of your illness?	28.8	5
7	Have you worried about loss of mobility because of your illness?	28.8	9
8	Did you have lower back pain?	27.4	3
9	Have you had to modify your daily activities because of your illness?	27.4	8
10	Have you worried about becoming bed- bound because of your illness?	27.4	6

<sup>1 =</sup> Top priority issue for health care providers Items included within the red box are items ranked by both groups to be in the top 10

## Patients suggestions for questions to add:

- Do you have family support?
- **❖** Do you worry about your ability to drive in the future?
- Were you able to understand your procedures, treatments, & medications?

## Health care provider suggestions for questions to add:

- \* Have you experienced weakened relationships with family or friends?
- Do you feel like a burden to family/friends?
- ❖ Does MSCC have an effect on sexual function?
- Do you have control of your bowel or bladder?
- Are you worried of becoming dependant on others now?
- ❖ Are you more concerned about bodily pain or weakness/paralysis in the arms and/or legs?

## Canadian-led International Development of a European Organization for Research and Treatment of Cancer (EORTC) Quality of Life Module for Malignant Spinal Cord Compression

~RESEARCH PROTOCOL~

#### **ODETTE CANCER CENTRE, CANADA**

**Gunita MITERA** 

Andrew LOBLAW

Arjun SAHGAL

#### **CROSS CANCER INSTITUTE, CANADA**

Brita DANIELSON

Alysa FAIRCHID

#### Fundación Instituto Valenciano de Oncología, Spain

Estanislao ARANA

#### **LUMC, LEIDEN, NETHERLANDS**

Yvette van der LINDEN

#### TATA MEDICAL CENTER, KOLKATA, INDIA

Indranil MALLICK

#### TATA MEMORIAL CENTRE, MUMBAI, INDIA

Ashwini Budrukkar

#### **AUSTRALIA & NEW ZEALAND**

To be determined at April meeting

**GERMANY** 

Dirk RADES

**ITALY** 

**Ernesto MARANZANO** 

## CONCLUSIONS

#### JOURNAL OF CLINICAL ONCOLOGY

#### COMMENTS AND CONTROVERSIES

# Extracranial Oligometastases: A Subset of Metastases Curable With Stereotactic Radiotherapy

Kimberly S. Corbin, Samuel Hellman, and Ralph R. Weichselbaum, University of Chicago Medical Center, Chicago, IL

Trial Name or			
Number	Design	Eligibility	Intervention
SABR-COMET	Randomized	All metastatic sites treatable; maximum of three tumors to any single organ system; controlled primary tumor	Standard RT vs SBRT
UPCI 10-028	Phase II		
UPCI 10-027	Phase II		
NCT01565837	Phase II		
NCT01185639	Phase II		
PulMiCC	Randomized	Pulmonary metastases from colorectal cancer	Active monitoring vs Surgery

Ultimately, a <u>randomized trial</u> of ablative radiotherapy and/or surgery compared with the standard of care may be necessary to define the role of ablative modalities in oligometastases.

## Panel: Evidence-based practice for extracranial oligometastases

- Stereotactic body radiotherapy results in a high control rate of treated metastases (~80%)
- About 20% of patients are progression free at 2–3 years after stereotactic body radiotherapy
- Toxicity is low
- Stereotactic body radiotherapy should be considered in patients with isolated metastases, especially if the disease-free interval is longer than 6 months
- Randomised trials are needed to establish whether stereotactic body radiotherapy improves progression free and/or overall survival
- Patients most likely to benefit from stereotactic body radiotherapy have:
  - Long disease-free interval
  - Breast histology
  - One to three metastases



Giotto - Scrovegni - Christ among the Doctors