

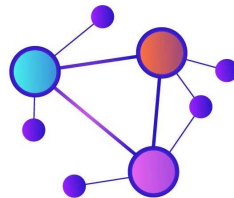
28eme congrès de Radiothérapie Tunisien

Sousse 2025

**Stéréotaxie
os et oligométastase**

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Centre Henri Becquerel
Quant.IF - AIMS



Liens d'intérêt

- Congrès: Merck
- Réunions scientifiques: AMGEN, BMS, MSD, Lilly, ASTRA ZENECA, MERCK
- Boards scientifiques: ASTRA ZENECA
- Projets scientifiques: Brainlab, Varian

Définition

Recommandations
ESMO ESTRO

Oligoprogression
= traitement

Oligoréccurrence
= absence de
traitement

A De-novo oligometastatic disease

Synchronous oligometastatic disease



- T0: first time diagnosis of primary cancer (green) and oligometastases (red) within 6 months

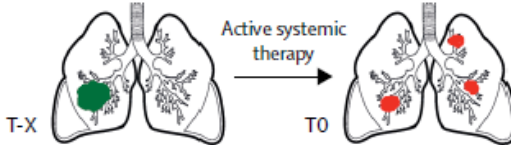
6 mois

Metachronous oligorecurrence



- T-X: diagnosis and treatment of primary cancer (green) in a non-metastatic state
- Systemic therapy-free interval
- T0: First time diagnosis of new oligometastases (red) >6 months after diagnosis of cancer

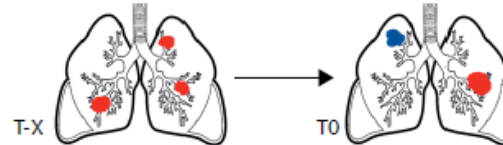
Metachronous oligoprogression



- T-X: diagnosis and treatment of primary cancer (green) in a non-metastatic state
- Under treatment with active systemic therapy
- T0: first time diagnosis of new oligometastases (red) >6 months after diagnosis of cancer

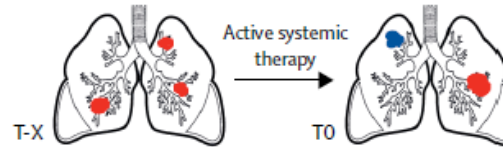
B Repeat oligometastatic disease

Repeat oligorecurrence



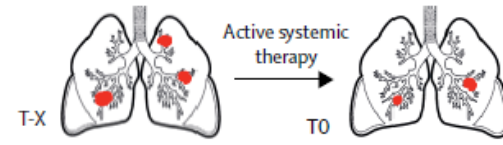
- T-X: diagnosis of oligometastases followed by local treatment or systemic treatment or both
- Systemic therapy-free interval
- T0: diagnosis of new (blue) and growing or regrowing (red) oligometastases

Repeat oligoprogression



- T-X: diagnosis of oligometastases followed by local treatment or systemic treatment or both
- Under treatment with active systemic therapy
- T0: diagnosis of new (blue) and growing or regrowing (red) oligometastases

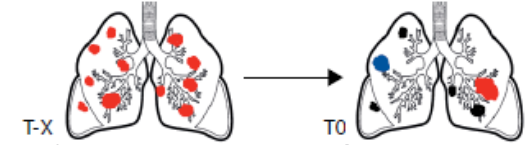
Repeat oligopersistence



- T-X: diagnosis of oligometastases followed by local treatment or systemic treatment or both
- Under treatment with active systemic therapy
- T0: diagnosis of persistent non-progressive (red) oligometastases

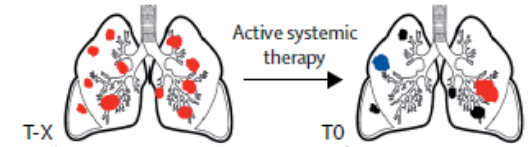
C Induced oligometastatic disease

Induced oligorecurrence



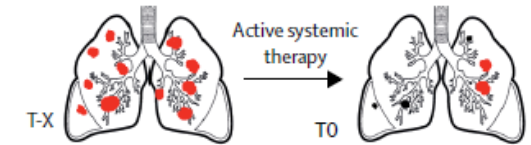
- T-X: diagnosis of polymetastatic metastatic disease followed by systemic treatment with or without local treatment
- Systemic therapy-free interval
- T0: diagnosis of new (blue) and growing or regrowing (red) oligometastases, possible residual non-progressive metastases (black)

Induced oligoprogression



- T-X: diagnosis of polymetastatic metastatic disease followed by systemic treatment with or without local treatment
- Under treatment with active systemic therapy
- T0: diagnosis of new (blue) and growing or regrowing (red) oligometastases, possible residual non-progressive metastases (black)

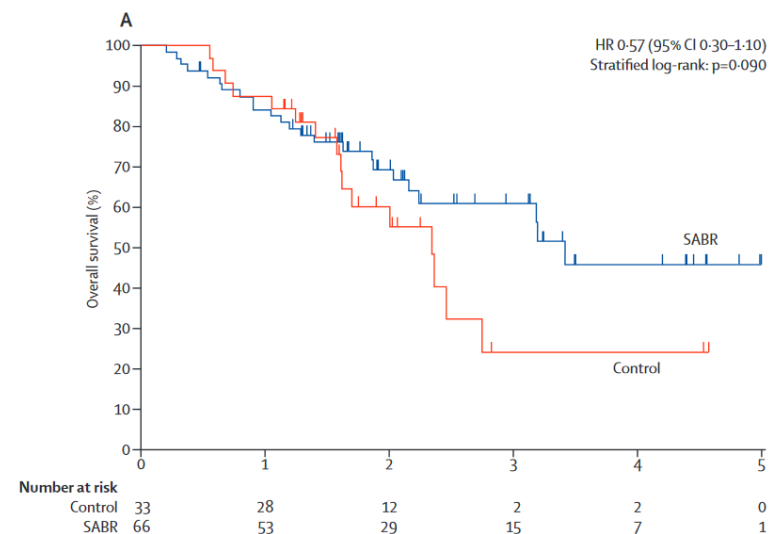
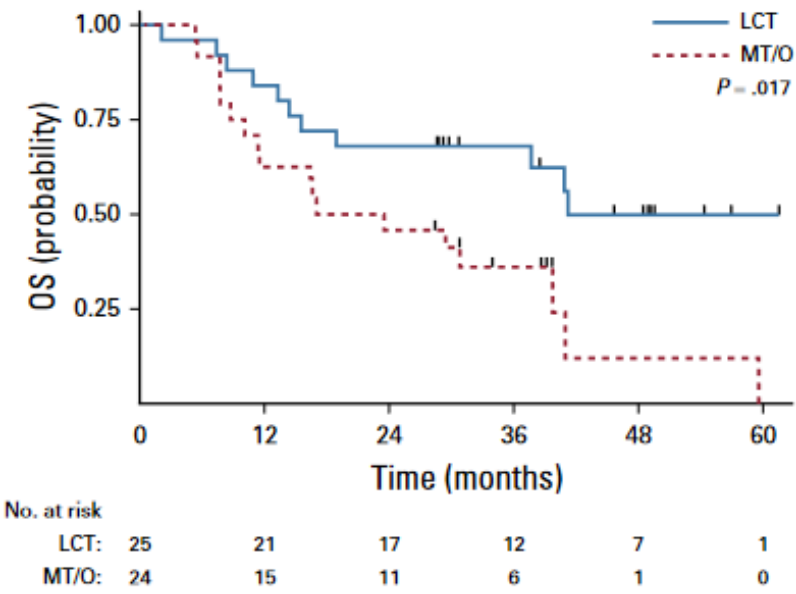
Induced oligopersistence



- T-X: diagnosis of polymetastatic metastatic disease followed by systemic treatment with or without local treatment
- Under treatment with active systemic therapy
- T0: diagnosis of persistent non-progressive oligometastases (red), where response is worse compared with other residual metastases (black)

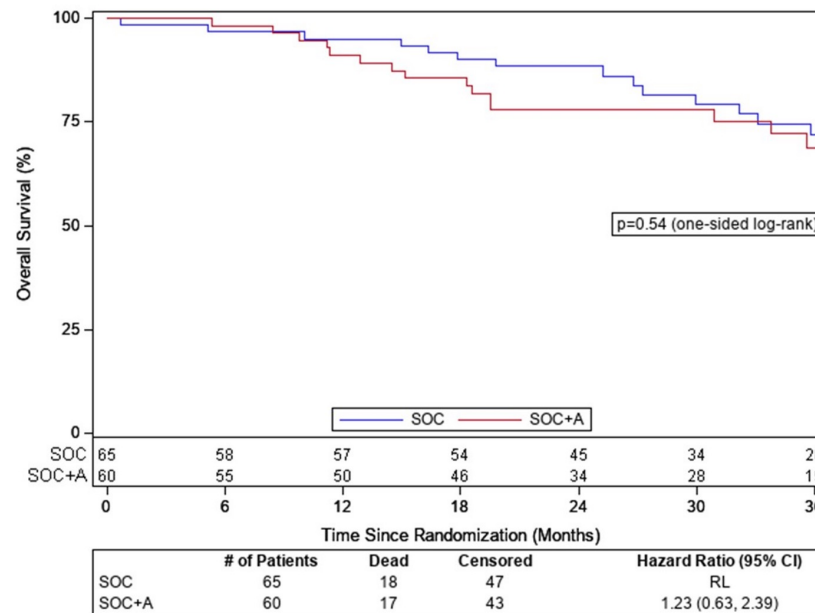
Guckenberger, Lancet Oncol, 2020

Phase II



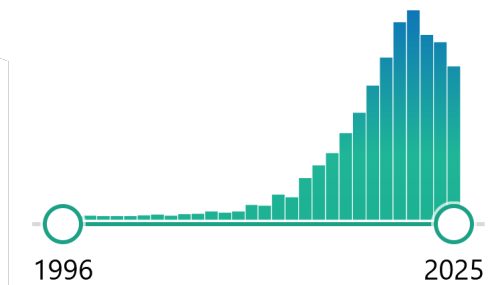
Etat des lieux

Phase III



Oligometastatic radiotherapy
2600 articles

RESULTS BY YEAR



Clinical trial
154

Randomized Controlled Trial
48

Palma, JCO, 2020 ; Gomez, JCO, 2019; Schmura, ASCO, 2022

Résultats

Meta-
analyse

Prostate

✓ SABR-COMET

✓ 99 patients

PFS

✓ EXTEND

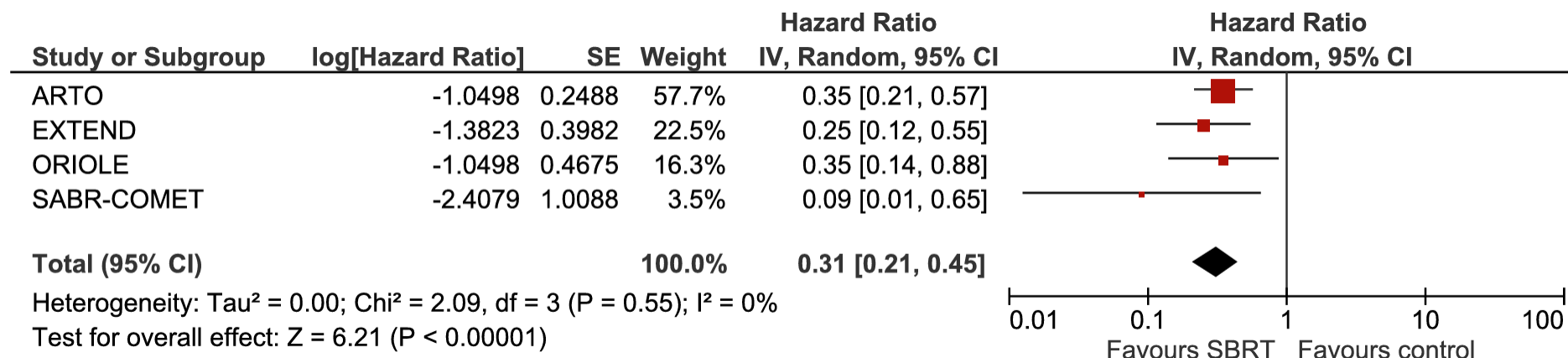
✓ 87 patients

✓ ORIOLE

✓ 54 patients

✓ ARTO

✓ 157 patients



Cas clinique

Mme B, 68 ans, OMS 0, absence de comorbidités

En 2018

CCI de 15 mm N- de grade 2 RH+ Her2- Ki 67 à 15%

Tumorectomie, Radiothérapie et Anti-aromatase

En 2025

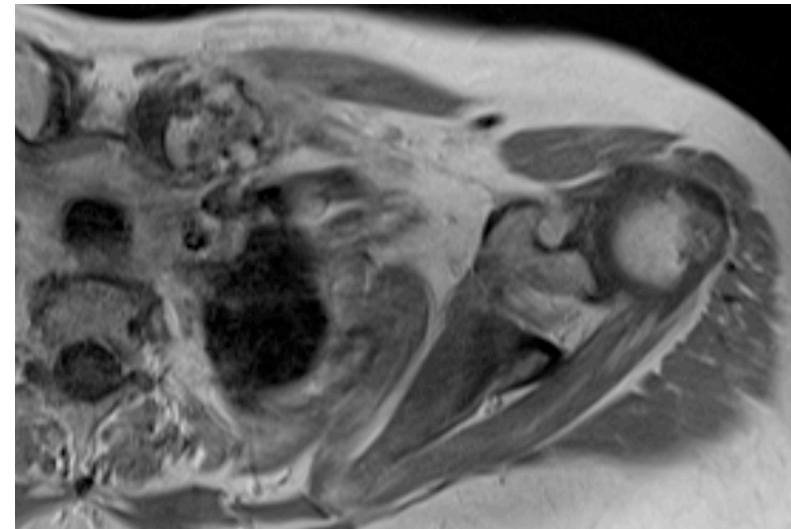
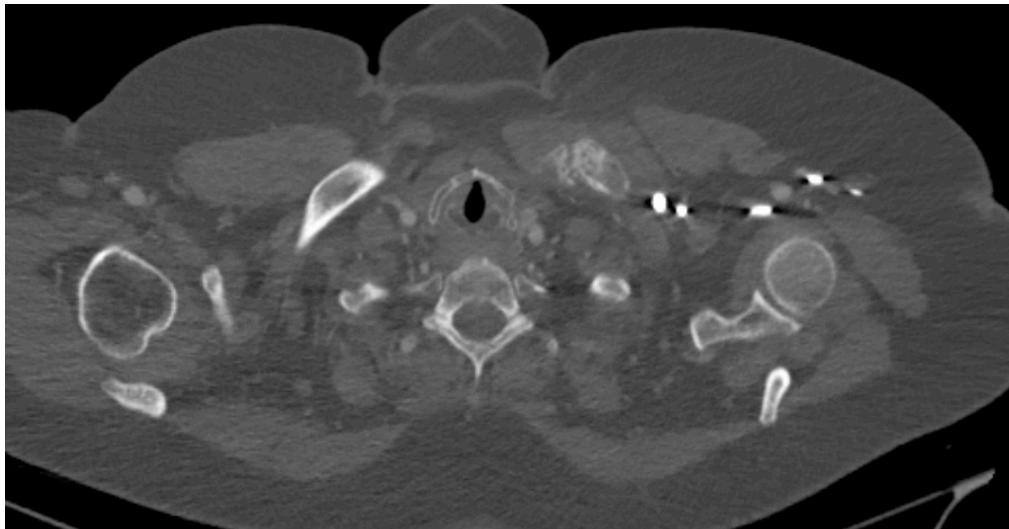
Douleurs de l'épaule Gauche

Bilan par TDM injectée et Scintigraphie Osseuse

Lésion de la clavicule Gauche et de la voute crânienne

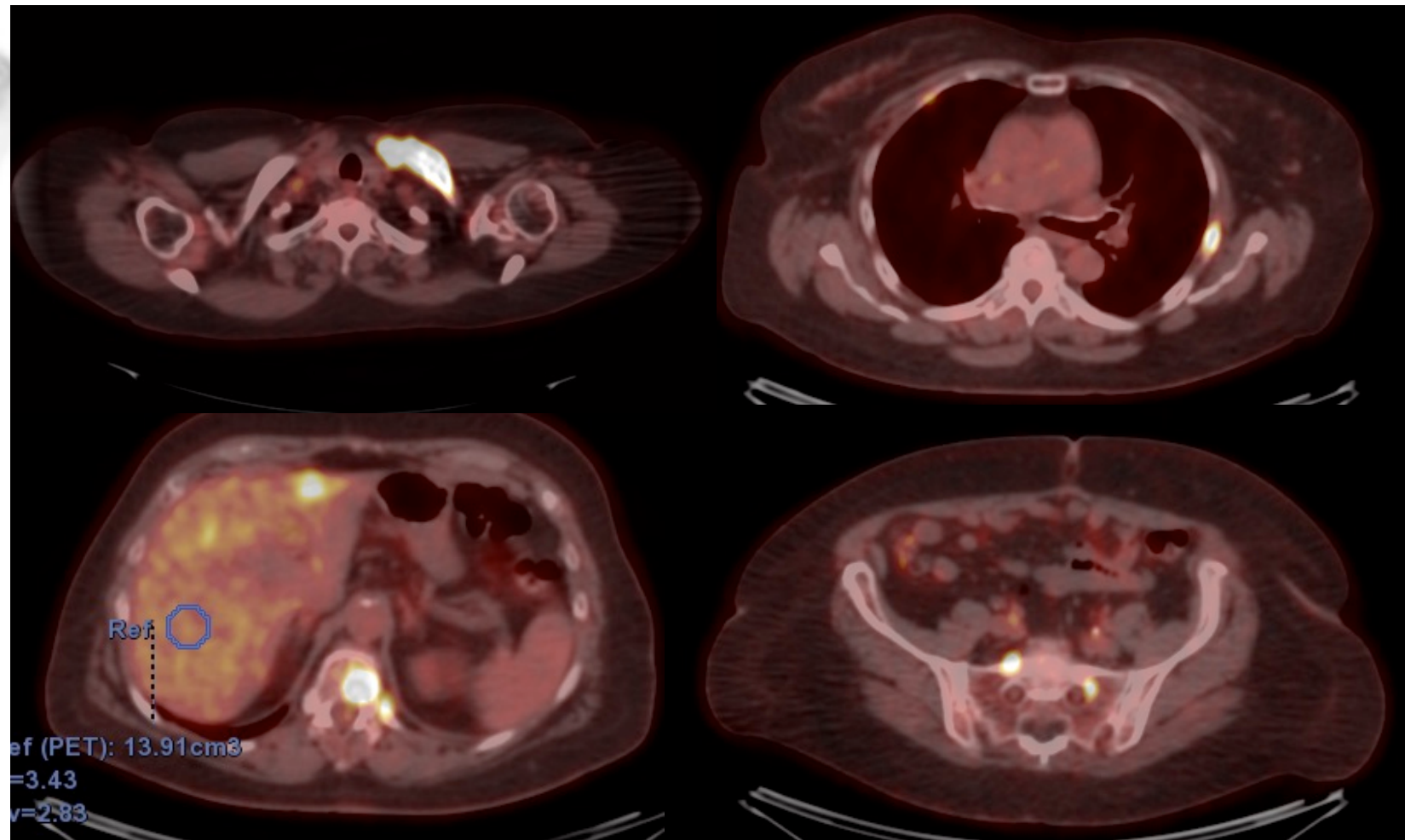
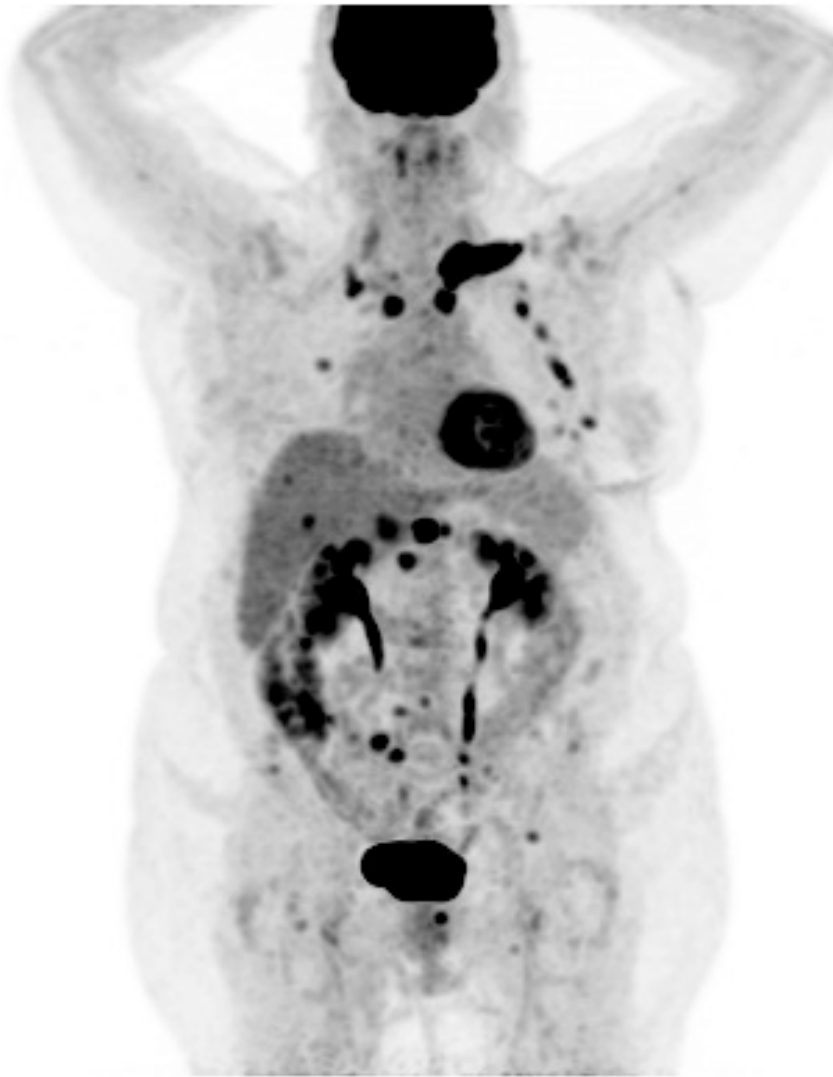
Traitement par FASLODEX – PALBOCICLIB

Adressée par Oncologue Médical pour radiothérapie stéréotaxique



Que faites vous?

1. Stéréotaxie des deux lésions osseuses
2. Traitement antalgique par 30Gy/10
3. Traitement antalgique par 8Gy/1
4. Absence de traitement par RTH car terrain irradié
5. Traitement médical seul
6. Autre



4 cancers primitifs

Breast
Prostate
Lung
Colorectal

1,597 patients
enregistrés

1468 patients évalués

Severe adverse event by concomitant systemic therapy.

System Organ Class + Preferred term

Comcomitant systemic treatment

	(SBRT pop: N=941)				(SBRT pop: N=527)			
	no				yes			
	Grade 3 N (%)	Grade 4 N (%)	Grade 5 N (%)	Grade ≥3 N (%)	Grade 3 N (%)	Grade 4 N (%)	Grade 5 N (%)	Grade ≥3 N (%)
PATIENTS' WORST GRADE	3 (0.3)		1 (0.1)	4 (0.4)	3 (0.6)		1 (0.2)	4 (0.8)
INFECTIONS AND INFESTATIONS								
Empyema	1 (0.1)			1 (0.1)				1 (0.2)
Pneumonia					1 (0.2)			
INJURY, POISONING AND PROCEDURAL COMPLICATIONS								
Radiation Pneumonitis					1 (0.2)			1 (0.2)
Radiation Skin Injury	1 (0.1)			1 (0.1)				
METABOLISM AND NUTRITION DISORDERS								
Decreased Appetite	1 (0.1)			1 (0.1)				
MUSCULOSKELETAL AND CONNECTIVE TISSUE DISORDERS								
Bone Pain					1 (0.2)			1 (0.2)
NERVOUS SYSTEM DISORDERS								
Brain Oedema	1 (0.1)			1 (0.1)				
Cerebral Haemorrhage	1 (0.1)			1 (0.1)				
RESPIRATORY, THORACIC AND MEDIASTINAL DISORDERS								
Pneumonitis					1 (0.2)			1 (0.2)

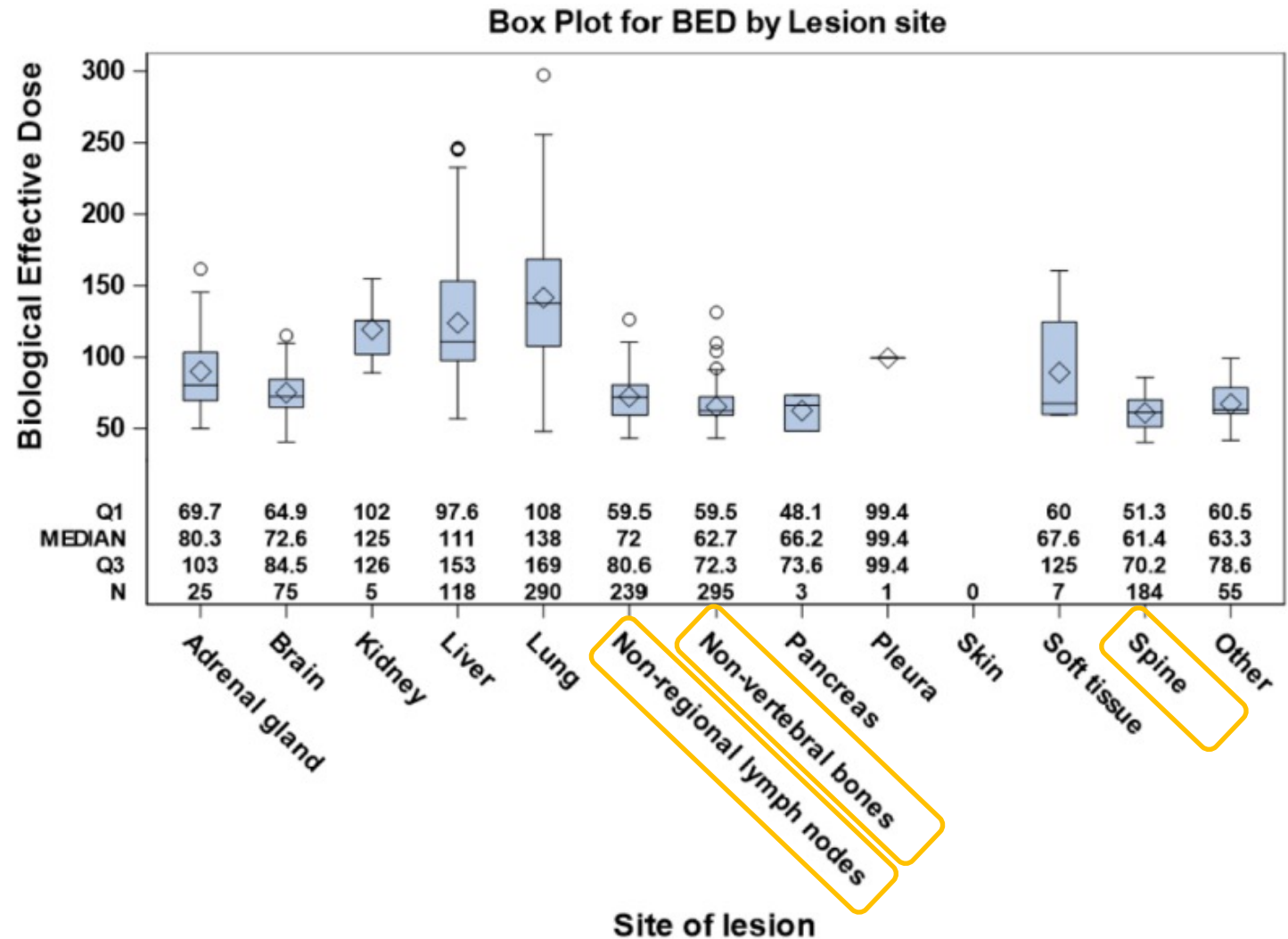
EORTC-ESTRO E2
RADlatE
OligoCare study

Traitements réalisés

Cohorte
prospective

Pan-
tumeur

- ✓ α/B à 10 pour lung cancer et CRC
- ✓ α/B à 2.5 pour breast cancer
- ✓ α/B à 1.5 pour prostate cancer
- ✓ 1004 patients inclus

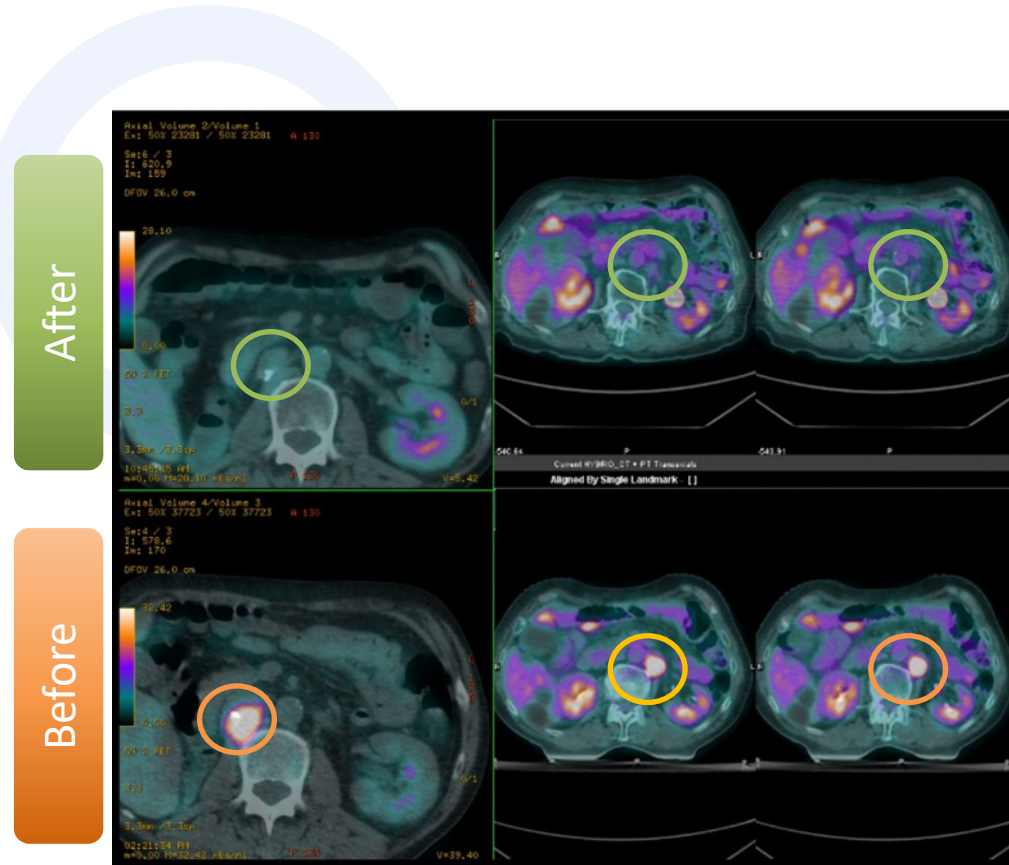
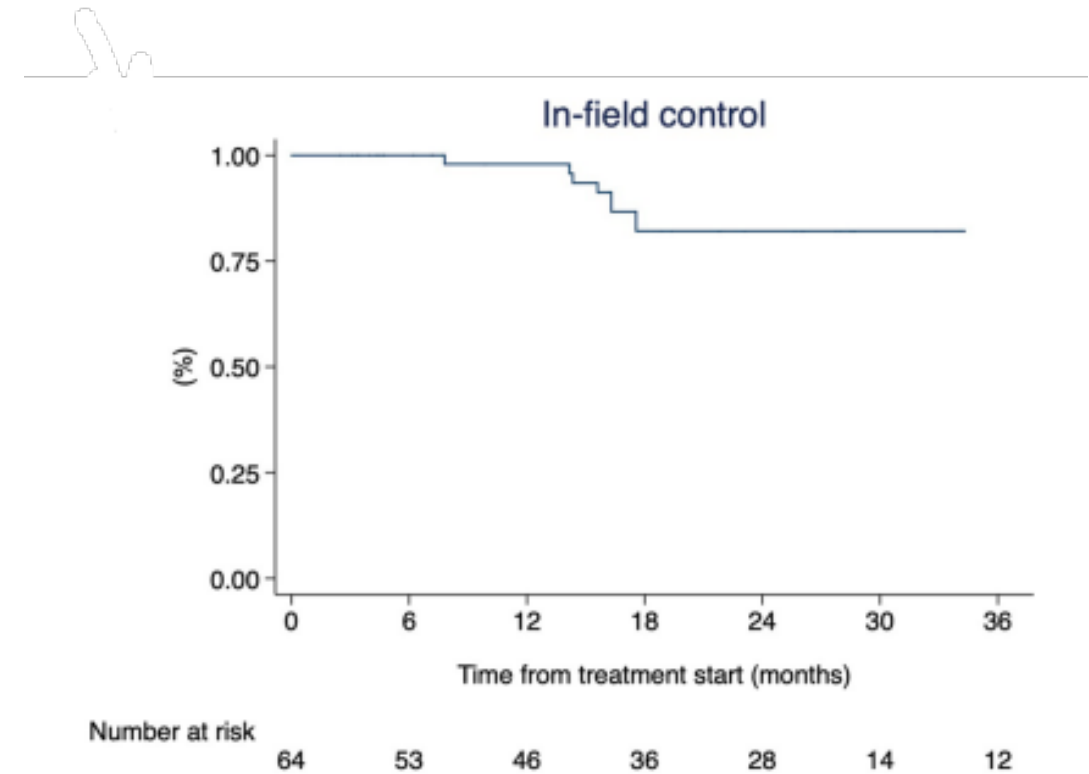


Indications

Author	Study design	N	N with A-P LN	Primary tumor	Median FU (months)	Median total dose (range)	Median number of fractions (range)	Median d/f (range)	Toxicity classification	QA score	Study	N pts A-P LN	Acute toxicity in grade:				Late toxicity in grade:			
													1	2	3	4	1	2	3	4
Alsuhaibani, 2019 [19]	Retrospective	21	11	GI	17	n.a. (30–60)	n.a. (3–5)	n.a. (n.a.)	CTCAE V4.0	3	Alsuhaibani (2019) [19]	11	n.a.	2	0	0	n.a.	0	0	0
Barney, 2012 [20]	Retrospective	47	13	Various	12	45 (20–60)	5 (1–5)	10 (n.a.)	CTCAE V3.0	4	Barney (2012) [20]	13	n.a.	n.a.	0	0	n.a.	n.a.	0	0
Bignardi, 2011 [21]	Retrospective	19	19	Various	12	45 (36–45)	6 (6)	n.a. (6–7.5)	CTCAE V3.0	3	Bignardi (2011) [21]	19	4	0	0	0	1	0	1	0
Bouman, 2017 [22]	Retrospective	43	34	Prostate	31	n.a. (30–35)	n.a. (3–5)	n.a. (7–10)	n.a.	3	Bouman (2017) [22]	34	1	2	0	0	0	0	0	0
Burkon, 2020 [12]	Retrospective	90	57	Various	35	n.a. (27–45)	n.a. (3–8)	n.a. (5–15)	CTCAE (version n.a.)	3	Burkon (2020) [12]	57	n.a.	n.a.	0	0	0	0	0	0
Caivano, 2023 [23]	Retrospective	174	82	Various	NA	36 (14–76)	n.a. (1–8)	n.a. (4–23)	CTCAE V4.4	3	Caivano, (2023) [23]	82	n.a.	n.a.	0	0	0	0	0	0
Corvò, 2013 [24]	Retrospective	36	36	Various	28	35 (12–50)	5 (2–10)	n.a. (4–9)	CTCAE V4.0	3	Corvò (2013) [24]	36	23	0	0	0	0	0	0	0
Cozzi, 2022 [25]	Retrospective	74	74	Prostate	31	40 (33–40)	5 (3–5)	8 (8–11)	CTCAE V4.0	3	Cozzi (2022) [25]	74	0	0	0	0	0	0	0	0
Cuccia, 2023 [26]	Retrospective	69	66	Prostate	16	35 (30–40)	5 (3–6)	n.a. (n.a.)	CTCAE V4.0	3	Cuccia (2023) [26]	66	n.a.	n.a.	0	0	n.a.	n.a.	0	0
Detti, 2015 [27]	Retrospective	30	30	Prostate	12	n.a. (24–36)	n.a. (1–5)	n.a. (6–24)	CTCAE V4.0	3	Detti (2015) [27]	30	0	1	0	0	1	0	0	0
Franzese, 2017 [28]	Retrospective	26	26	Prostate	29	40 (25–45)	6 (4–6)	n.a. (n.a.)	CTCAE V4.0	3	Franzese (2017) [29]	35	2	3	0	0	n.a.	n.a.	n.a.	n.a.
Franzese, 2017 [29]	Retrospective	35	35	CRC	15	n.a. (30–45)	n.a. (6–13)	n.a. (3–7.5)	CTCAE V3.0	3	Franzese (2017) [28]	26	5	0	0	0	0	0	0	0
Franzese, 2016 [30]	Retrospective	71	71	Various	18	45 (45)	6 (6)	7.5 (7.5)	CTCAE V4.0	3	Franzese (2016) [30]	71	10	2	0	0	0	0	0	0
Franzese, 2020 [31]	Prospective	52	52	Various	24	48 (48)	4 (4)	12 (12)	CTCAE V4.0 & RTOG/EORTC	5	Franzese (2020) [31]	52	4	0	0	0	0	0	0	0
Gawish, 2023 [32]	Retrospective	17	17*	Prostate	16.6	48 (30–60)	12 (5–20)	4 (3–8)	n.a.	3	Gawish (2023) [32]	17	n.a.	n.a.	0	0	n.a.	n.a.	0	0
Ingrosso, 2017 [33]	Retrospective	40	39	Prostate	24	n.a. (12–50)	n.a. (1–5)	n.a. (5–12)	RTOG/EORTC criteria	4	Ingrosso (2017) [33]	39	n.a.	1	0	0	n.a.	0	1	0
Kang, 2010 [34]	Retrospective	59	30	CRC	32	42 (35–51)	3 (3)	n.a. (12–17)	CTCAE V2.0	4	Kang (2010) [34]	30	n.a.	n.a.	0	2	n.a.	n.a.	n.a.	n.a.
Kneebone, 2018 [35]	Prospective	57	39	Prostate	16	n.a. (30–50)	n.a. (1–5)	10 (10)	CTCAE V4.0	5	Kneebone (2018) [35]	39	n.a.	n.a.	0	0	n.a.	n.a.	0	0
Kutuk, 2022 [36]	Retrospective	96	52	Various	10	48.5 (30–60)	5 (3–15)	n.a. (n.a.)	CTCAE V4.0	2	Kutuk (2022) [36]	52	n.a.	n.a.	0	0	n.a.	n.a.	0	0
Lepinoy, 2019 [37]	Retrospective	62	35	Prostate	42	36 (30–66)	n.a. (n.a.)	7.5 (2–15)	CTCAE V4.0	4	Lepinoy (2019) [37]	35	0	2	1	0	2	12	2	0
Loi, 2018 [38]	Retrospective	23	23	Prostate	22	24 (24)	1 (1)	24 (24)	CTCAE V4.0	3	Loi (2018) [38]	23	2	0	0	0	0	0	0	0
Loi, 2018 [39]	Retrospective	91	89	Various	23	n.a. (40–48)	n.a. (5–6)	n.a. (7–9)	CTCAE V4.0	3	Loi (2018) [39]	89	26	13	0	0	5	5	0	0
Matoba, 2020 [40]	Retrospective	15	15	HCC	18	n.a. (45–49.5)	n.a. (6–9)	n.a. (5.5–7.5)	CTCAE V4.0	4	Matoba (2020) [40]	15	8	1	0	0	0	0	0	0
Nicosia, 2022 [41]	Prospective	63	63	Prostate	17	35 (14–40)	n.a. (n.a.)	n.a. (5–21)	CTCAE V5.0	6	Nicosia (2022) [41]	63	0	0	0	0	0	0	0	0
Ost, 2016 [42]	Retrospective	72	72	Prostate	36	n.a. (24–50)	n.a. (3–10)	n.a. (5–10)	CTCAE V4.0	3	Ost (2016) [42]	72	n.a.	n.a.	n.a.	n.a.	12	3	0	0
Park, 2015 [43]	Retrospective	85	83	Cervix	20	39 (27–51)	n.a. (3–10)	13 (n.a.)	CTCAE V4.0	4	Park (2015) [43]	83	n.a.	n.a.	1	0	0	9	2	2
Pasqualetti, 2016 [44]	Prospective	29	17	Prostate	12	n.a. (24–27)	n.a. (1–3)	n.a. (9–24)	CTCAE V4.0	5	Pasqualetti (2016) [44]	17	n.a.	0	0	0	n.a.	0	0	0
Pezzulla, 2021 [45]	Prospective	38	38	Prostate	27	n.a. (20–50)	n.a. (1–5)	n.a. (9–24)	CTCAE V4.0	6	Pezzulla (2021) [45]	38	n.a.	n.a.	0	0	n.a.	n.a.	0	0
Regnery, 2022 [46]	Prospective	26	26	Various	10	n.a. (25–40)	n.a. (3–7)	n.a. (5–9)	CTCAE V5.0	5	Regnery (2022) [46]	26	9	2	0	0	n.a.	n.a.	n.a.	n.a.
Shahi, 2020 [11]	Retrospective	51	48	Various	22	35 (25–40)	5 (5)	7 (5–8)	CTCAE V4.0	4	Shahi (2018) [11]	48	n.a.	n.a.	1	0	1	0	0	0
Siva, 2018 [47]	Prospective	33	13	Prostate	NA	20 (20)	1 (1)	20 (20)	CTCAE V4.0	6	Siva (2018) [47]	13	n.a.	n.a.	0	0	n.a.	n.a.	0	0
Wang, 2016 [48]	Retrospective	22	22	Various	33	39 (21–51)	5 (3–8)	8 (5–13)	CTCAE V4.0	4	Wang (2016) [48]	22	n.a.	n.a.	0	0	0	0	0	0
Werensteijn, 2021 [49]	Prospective/retrospective	90	90	Prostate	21	n.a. (30–35)	n.a. (3–6)	n.a. (6–10)	RTOG/EORTC criteria	5	Werensteijn (2021) [49]	90	48	3	0	0	18	8	0	0
Yang, 2022 [50]	Prospective/retrospective	101	101	Various	11	40 (25–50)	5 (n.a.)	8 (5–10)	CTCAE V5.0	5	Yang (2022) [50]	101	n.a.	n.a.	n.a.	n.a.	53	4	3	0
Yeung, 2017 [51]	Retrospective	18	11	Various	34	n.a. (30–60)	n.a. (4–8)	n.a. (5–8)	CTCAE V4.0	3	Yeung (2017) [51]	11	n.a.	3	0	0	n.a.	0	0	0
Median:		47	37		21					3	Total toxicity		142	35	3	2	93	41	9	2
											Reported in studies (n)*		16	20	33	33	21	25	32	32
											Total patients in studies		718	796	1356	1356	1122	1200	1438	1438
											% patients with toxicity		19.8	4.4	0.2	0.1	8.3	3.4	0.6	0.1

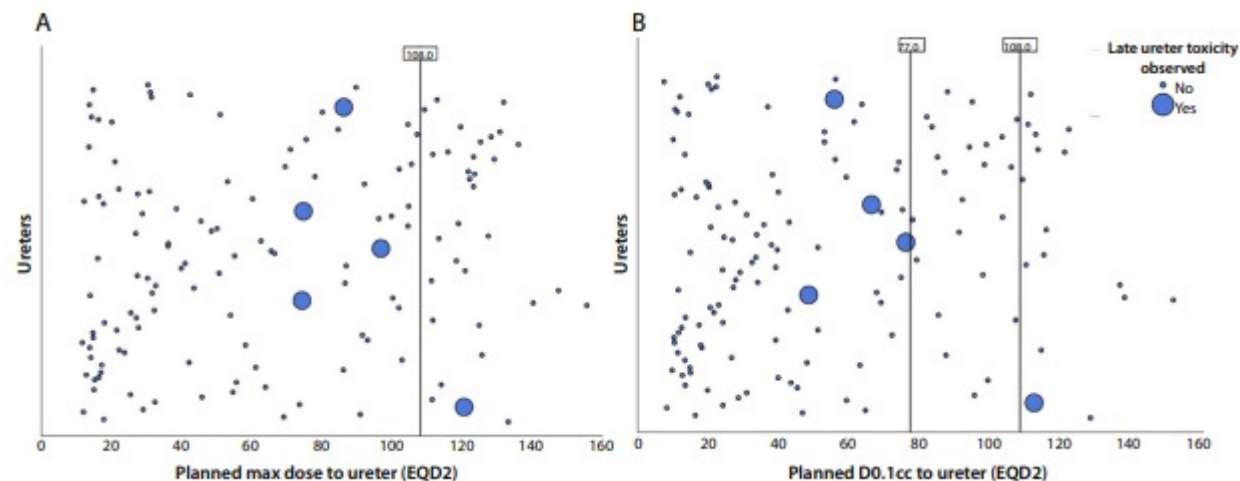
Résultats

- ✓ 52 patients
- ✓ 64 lésions
- ✓ LC à 1, 2 et 3 ans : 97.9%, 82.1% and 82.1%



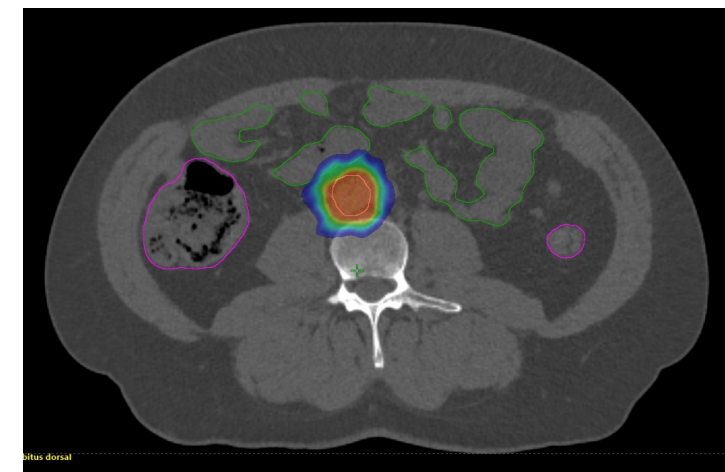
Quels schémas

- ✓ 48 Gy en 6 fractions
 - ✓ 45 Gy en 5 fractions
 - ✓ 35 Gy en 5 fractions
 - ✓ 40 Gy en 5 fractions
 - ✓ 36Gy en 6 fractions
- ✓ Difficulté à prédire la toxicité
 - ✓ Fractionnement
 - ✓ Mobilités OAR



Dosimetric parameters of the gastro-intestinal organs in EQD₂¹⁰.

		D _{max} (EQD ₂ ¹⁰)	D _{0.2cc} (EQD ₂ ¹⁰)	D _{0.5cc} (EQD ₂ ¹⁰)	D _{1cc} (EQD ₂ ¹⁰)	D _{2cc} (EQD ₂ ¹⁰)	D _{5cc} (EQD ₂ ¹⁰)	D _{10cc} (EQD ₂ ¹⁰)
All treatments (n = 55)	Median	61.5	48.6	43.7	38.6	29.6	22.6	15.3
	IQR	43.9–72.2	30.1–59.6	25.9–54.9	21.7–50.5	17.2–42.0	12.1–30.2	9.2–24.4
	Range	7.3–90.2	5.9–72.9	5.0–61.9	5–61.9	4.5–55.4	3.7–45.7	3.0–38.6
Patients with toxicity (n = 20)	Median	67.4	55.9	50.4	44.5	38.1	27.1	20.9
	IQR	54.1–67.4	41.8–66.0	35.3–60.8	29.3–52.3	23.4–45.1	16.2–36.5	12.6–29.4
	Range	12.7–84.3	10.1–72.9	9.1–67.1	8.1–61.9	7.3–55.4	6.5–45.7	5.9–38.6
Patients without toxicity (n = 35)	Median	60.0	43.7	37.8	30.2	22.9	15.6	11.7
	IQR	33.3–71.3	23.9–57.9	21.6–53.9	19.3–47.3	15.4–39.2	11.4–29.0	8.9–21.0
	Range	7.3–90.2	5.9–72.7	5.4–67.3	5.0–61.6	4.5–54.7	3.7–45.3	3.0–38.0
P-value*		0.090	0.037	0.036	0.031	0.033	0.025	0.031



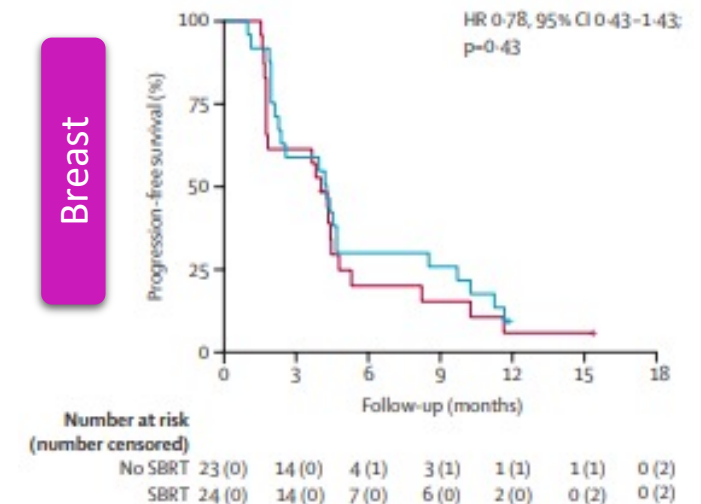
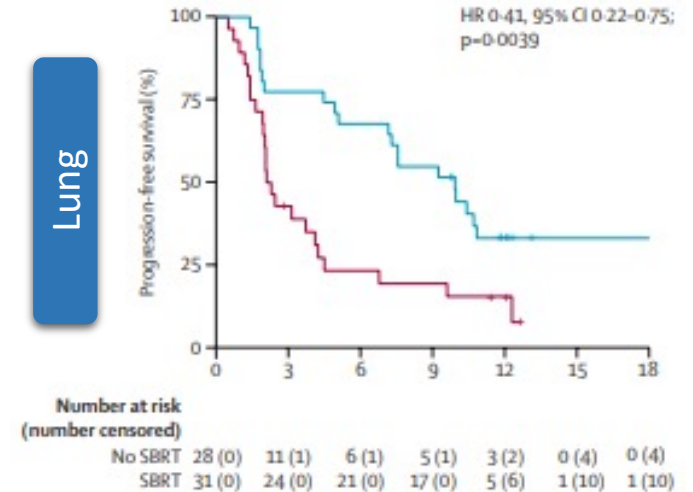
Indications

Bone

- Ré-irradiation
 - Protection de la dose à la moelle
 - Fractionnement potentiellement classique
- Cancer radiorésistant
 - Mélanome, Cancer du rein
- Patients oligométastatiques
 - 1 à 3 (5) métastases
 - 1 à 3 organes
- *Toutes lésions douloureuses?*
- *Radiothérapie préventive?*

?

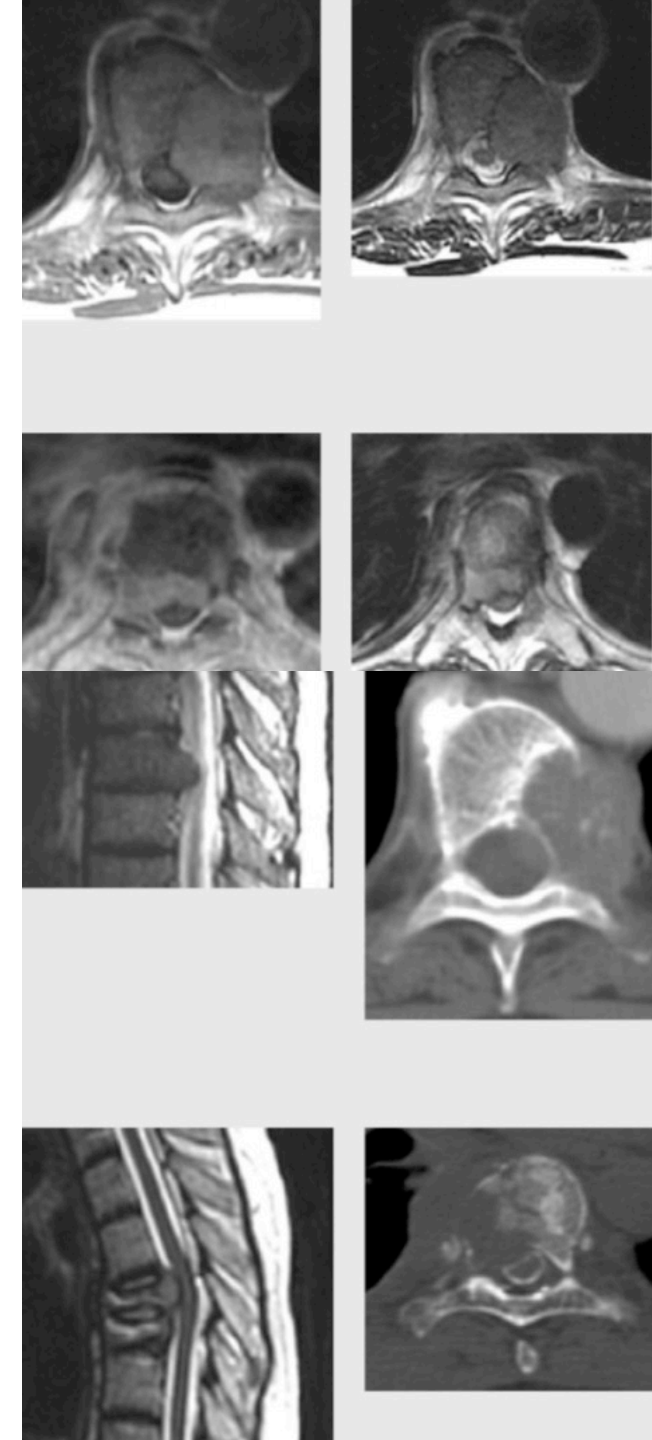
Cancer du sein?
Cancer de la prostate?
Cancer pulmonaire?



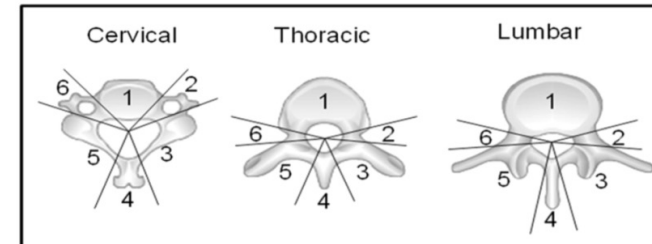
Définitions des volumes

Bone

Rachis



- Os périphériques
 - Intérêt IRM +/- ou TEP
 - CTV = GTV + 5mm (respect corticale)
- Rachis
 - Recommandations RTOG
 - IRM millimétrique si possible en position de traitement
 - T1 + Gadolinium pour le GTV
 - T2 pour la Moelle épinière / Queue de cheval



GTV involvement	ISRC GTV anatomic classification	ISRC bony CTV recommendation	CTV description
Any portion of the vertebral body	1	1	Include the entire vertebral body
Lateralized within the vertebral body	1	1, 2	Include the entire vertebral body and the ipsilateral pedicle/transverse process
Diffusely involves the vertebral body	1	1, 2, 6	Include the entire vertebral body and the bilateral pedicles/transverse processes
GTV involves vertebral body and unilateral pedicle	1, 2	1, 2, 3	Include entire vertebral body, pedicle, ipsilateral transverse process, and ipsilateral lamina
GTV involves vertebral body and bilateral pedicles/transverse processes	3	2, 3, 4	Include entire vertebral body, bilateral pedicles/transverse processes, and bilateral laminae
GTV involves unilateral pedicle	2	2, 3 ± 1	Include pedicle, ipsilateral transverse process, and ipsilateral lamina, ± vertebral body
GTV involves unilateral lamina	3	2, 3, 4	Include lamina, ipsilateral pedicle/transverse process, and spinous process
GTV involves spinous process	4	3, 4, 5	Include entire spinous process and bilateral laminae

Abbreviations: CTV = clinical target volume; GTV = gross tumor volume; ISRC = International Spine Radiosurgery Consortium.

E.M. Dunne et al. / Radiotherapy and Oncology 145 (2020) 21–29

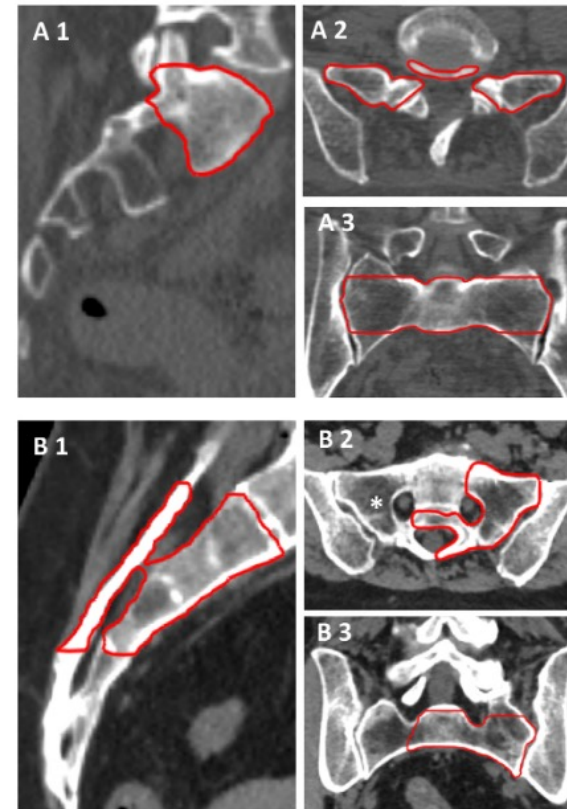
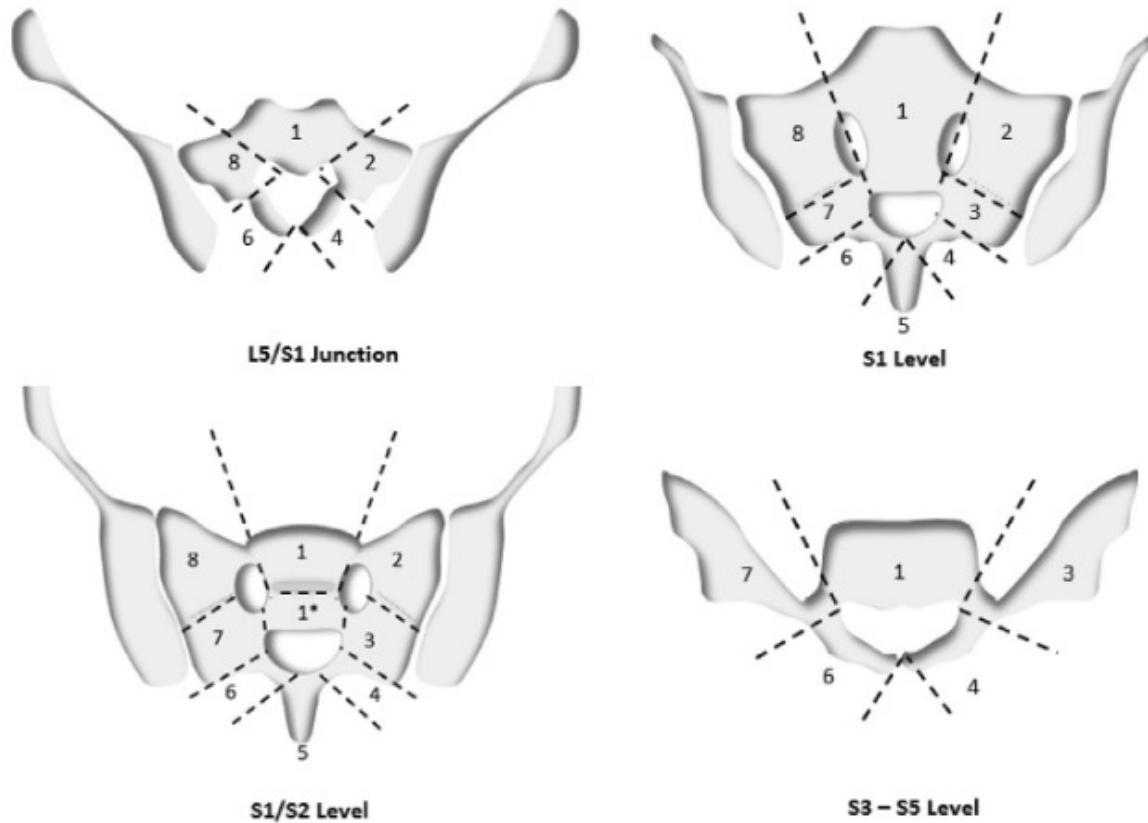
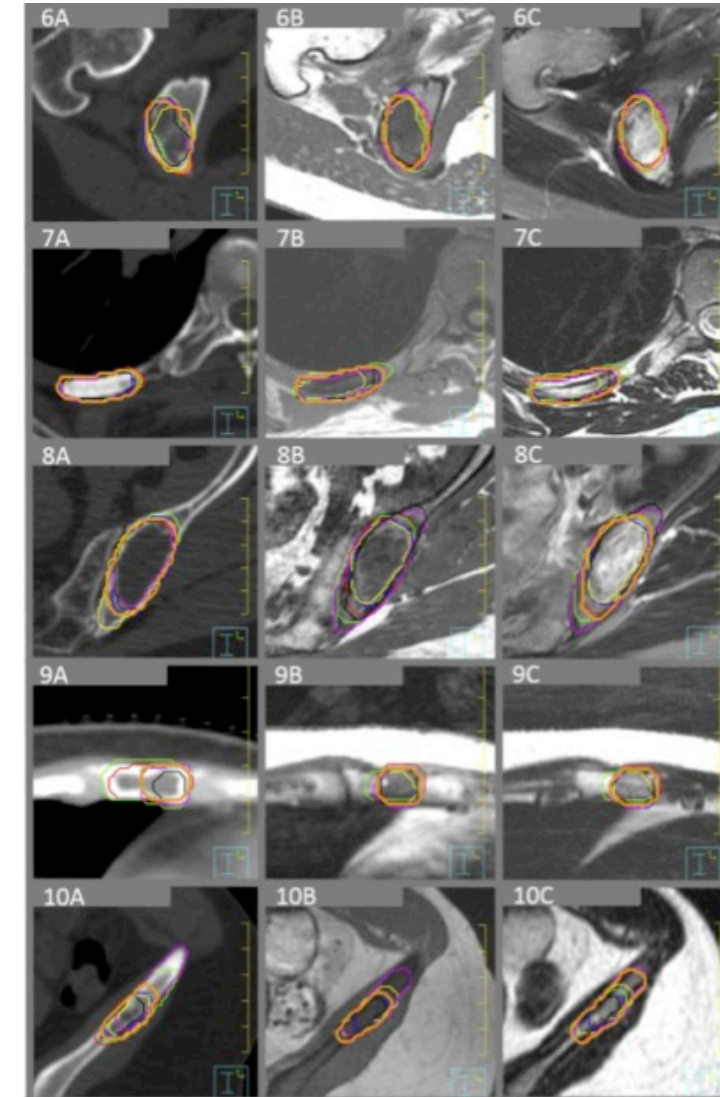
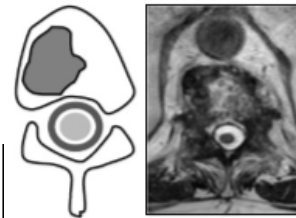
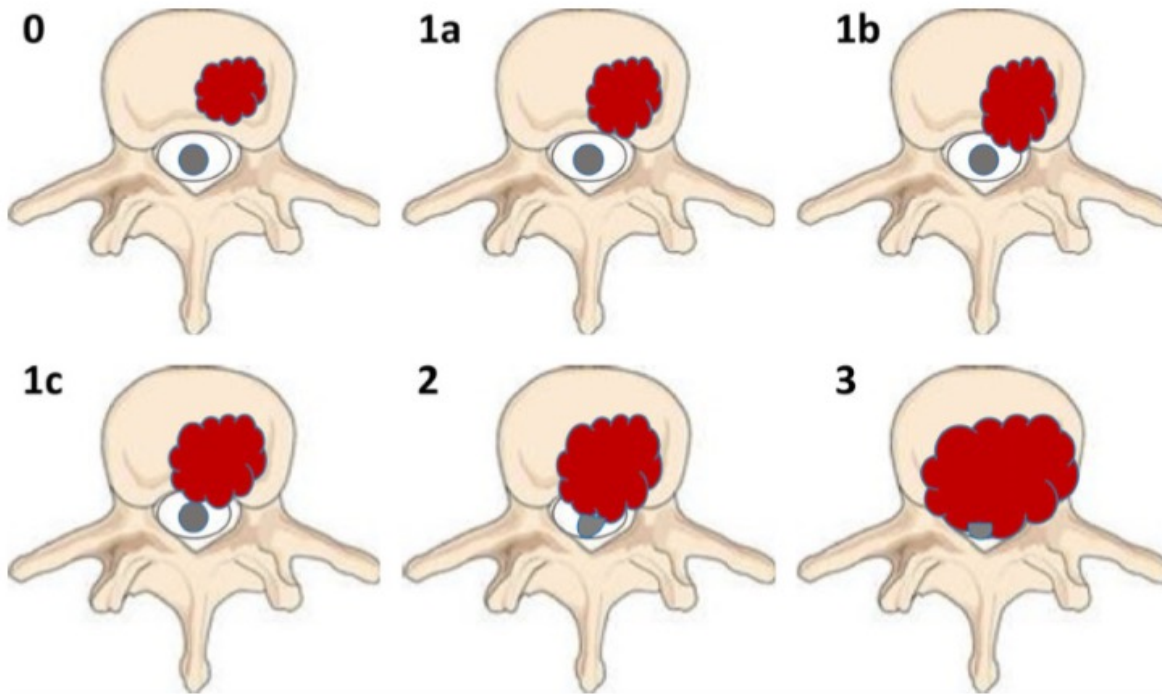


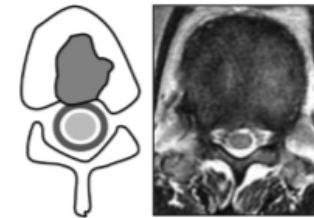
Fig. 3. Sagittal, axial and coronal CT images at the level of S1 (A1–A3, Case 8) and S2 (B1–3, Case 4). * Ossification line separating the right anterior and posterior ala (B2). Abbreviations: CT, computed tomography.



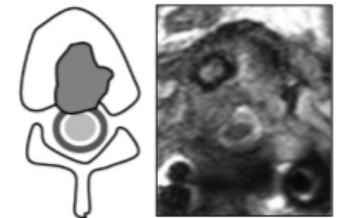
Contre indication



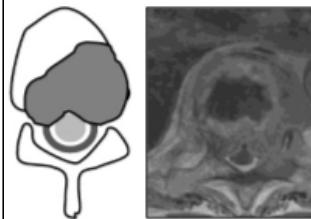
Grade 0
Bone involvement only
No canal compromise



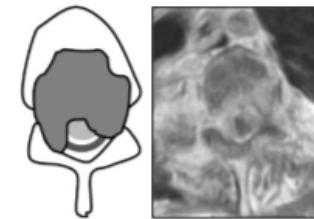
Grade I
Involvement of epidural fat



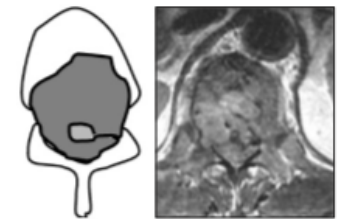
Grade II
Impingement of thecal sac



Grade III
Impingement of spinal cord

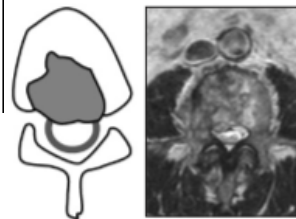


Grade IV
Compression and/or
displacement of spinal cord
Partial block of CSF

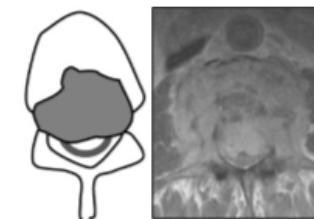


Grade V
Spinal cord compression and
Complete block of CSF

At cauda level

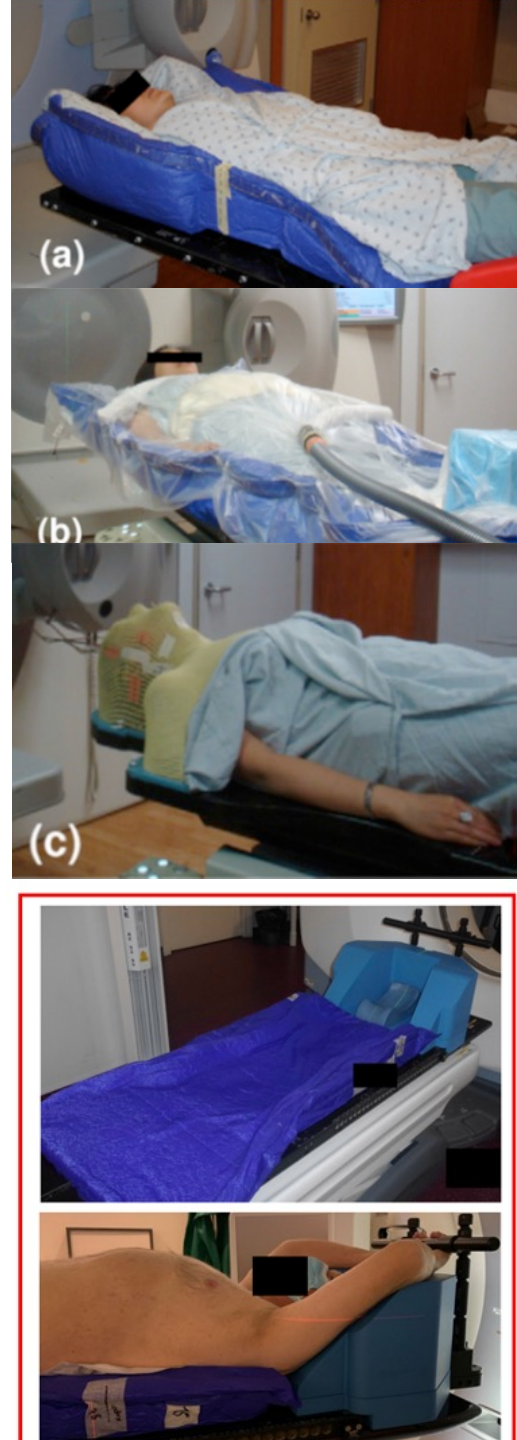
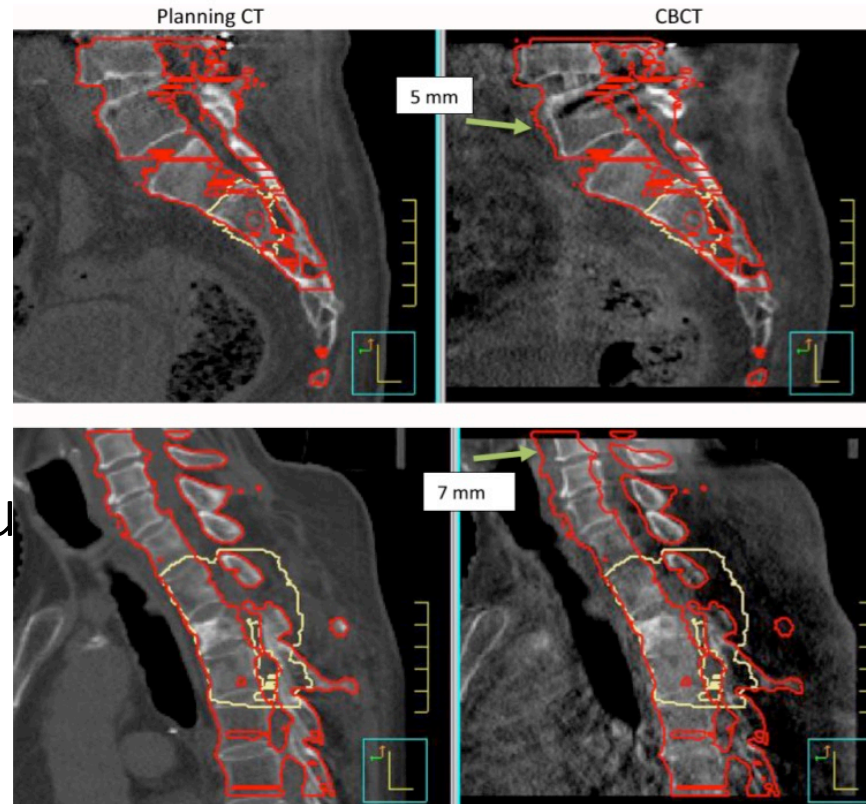


Grade II
≤50% canal



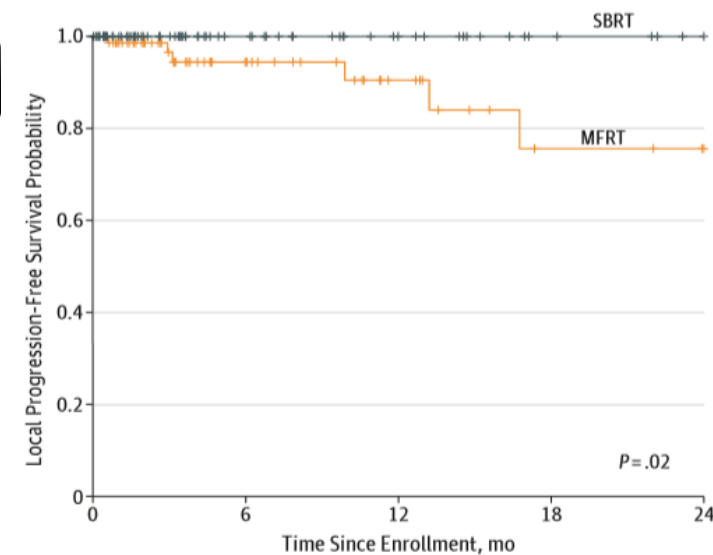
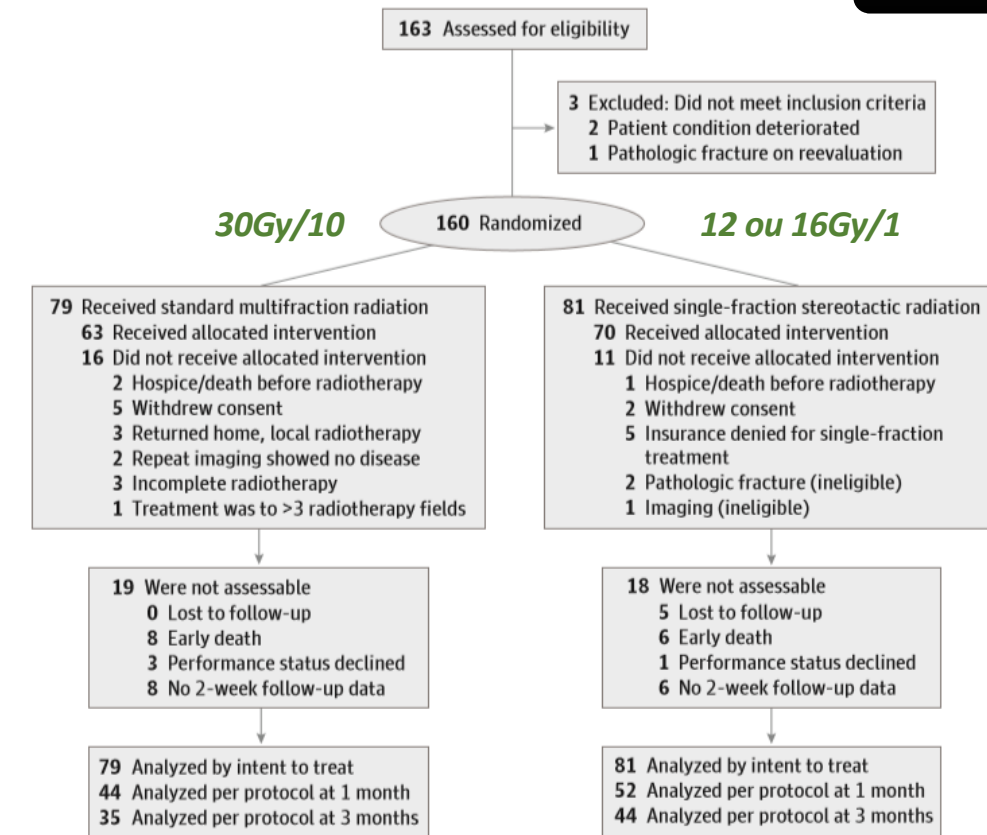
Grade IV
>50% canal compromise

- Dépend:
 - De la localisation
 - De la précision attendue
 - Du mode de traitement
 - Type de contrôle
 - Durée du traitement
- Attention aux volumes contigus
- Importance IRM/ TDM

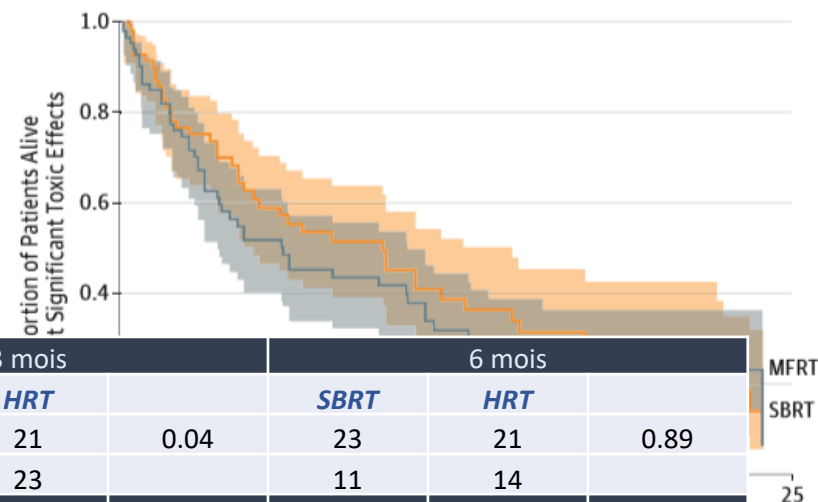


Quelle efficacité?

Rachis



No. at risk	0	6	12	18	24
SBRT	81	32	17	8	4
MFRT	79	31	17	8	6

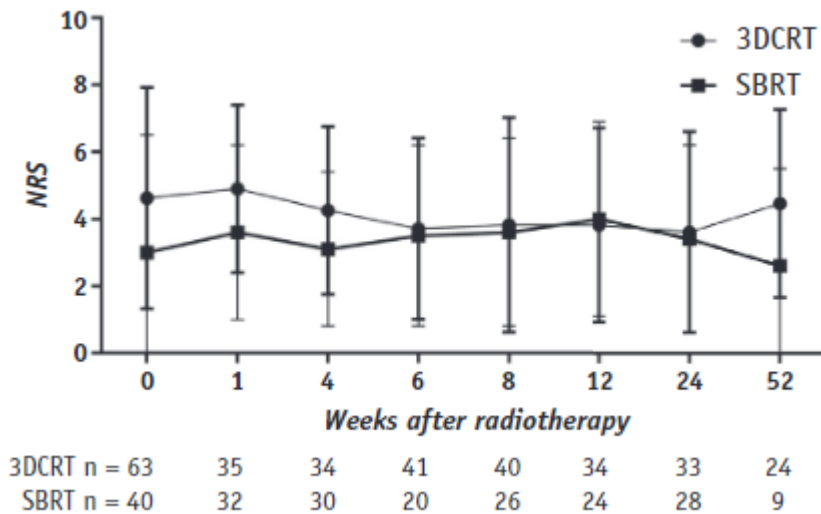


ITT		1 mois			3 mois			6 mois		
		SBRT	HRT		SBRT	HRT		SBRT	HRT	
	Respondeurs	44	30	0.19	39	21	0.04	23	21	0.89
	Non respondeurs	20	25		15	23		11	14	
PP										
	Respondeurs	69	55	0.15	73	49	0.04	68	61	0.78
	Non respondeurs	31	45		27	51		32	39	

Quelle efficacité?

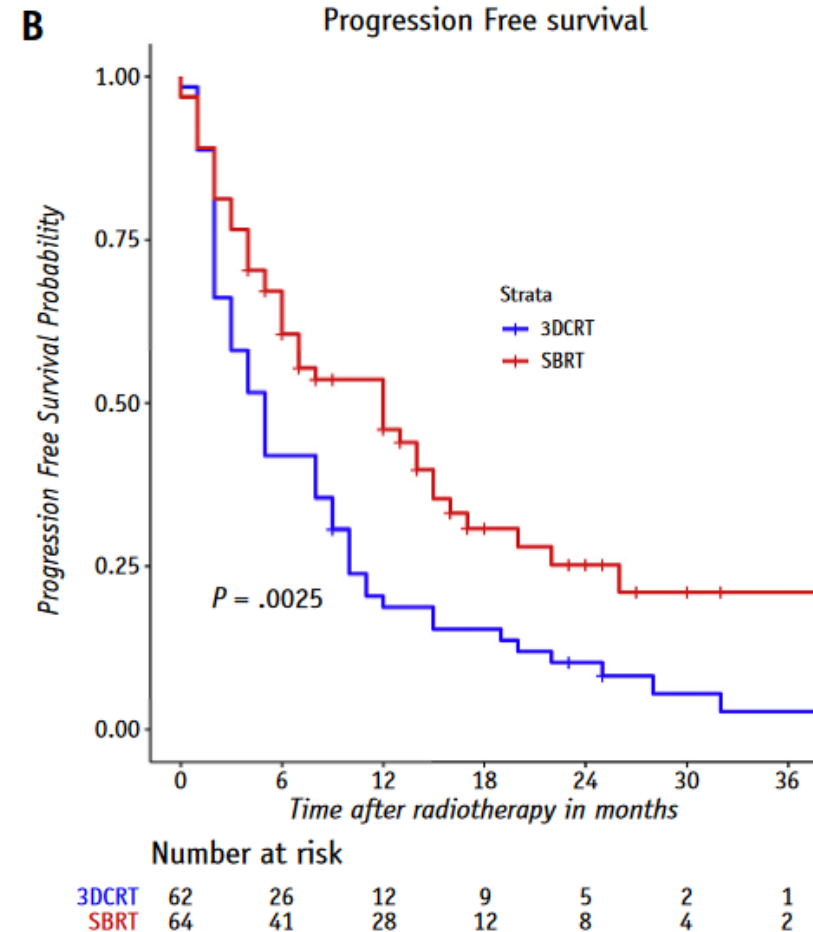
Bone

douleur

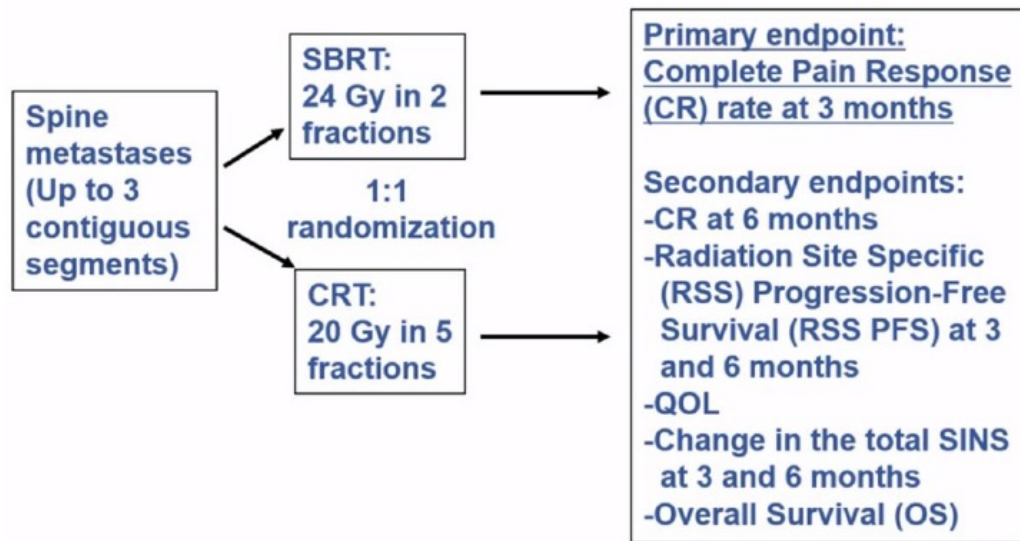


PRESENT Cohorte

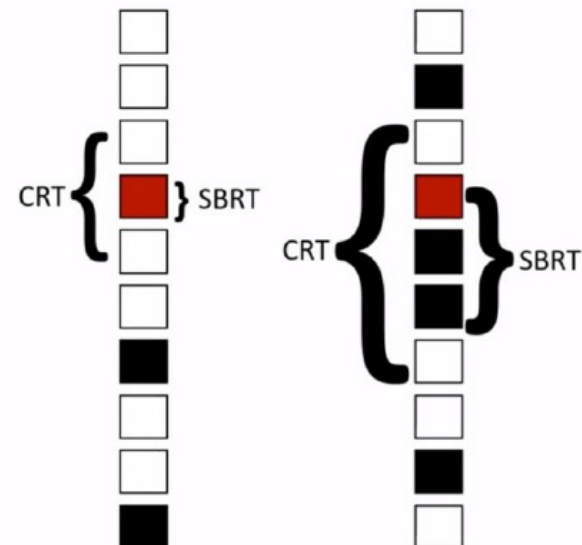
PFS



Schema

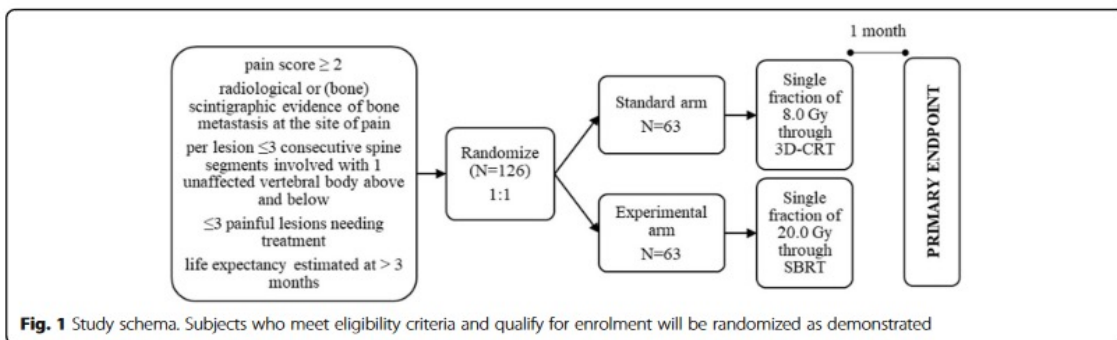


- Stratification factors: radioresistant (GI, RCC, melanoma, sarcoma) vs. radiosensitive, and presence vs. absence of extra-osseous extension (Mass type)



Red: Target spinal segment volume
Red +/- Black: Treatment Volume
{: Included segments in a SBRT treatment volume and segments encompassed in a CRT treatment field adjusting for margins

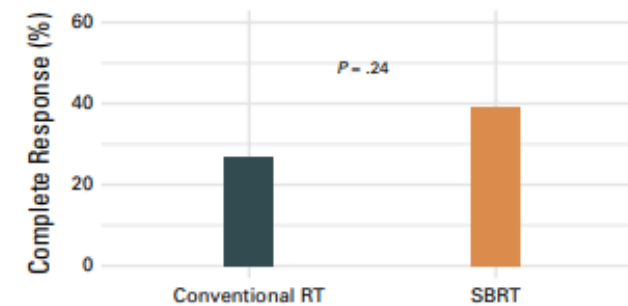
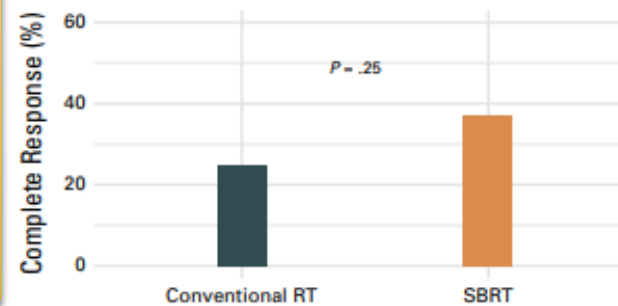
Response	CRT (N=115)	SBRT (N=114)
3 month assessment		
Complete response	14 %	35%
Partial response	25%	18%
Stable disease	30%	24%
Progressive disease	12%	6%
Indeterminant	19%	18%
Mean change in total SINS (standard deviation)	-0.49 (1.61)	-0.94 (1.69)
Response	CRT (N=115)	SBRT (N=114)
6 month assessment		
Complete response	16%	32%
Partial response	16%	9%
Stable disease	27%	23%
Progressive disease	7%	4%
Indeterminant	34%	32%
Mean change in total SINS (standard deviation)	-0.74 (1.99)	-0.73 (1.86)



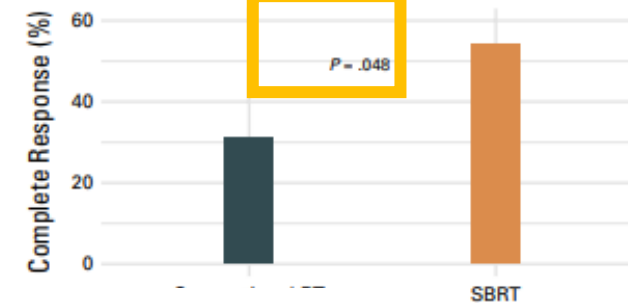
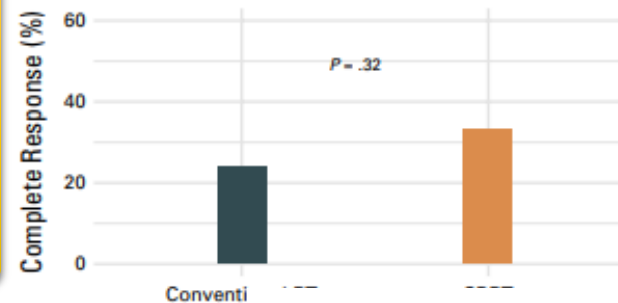
1 mois

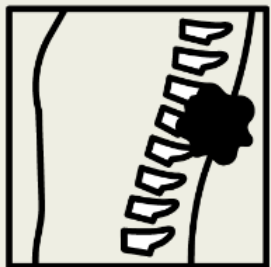
ITT

PP



3 mois



POPULATION**184 Men, 155 Women**

Adults with 1 to 3 newly diagnosed vertebral metastases

Mean age, 62.6 y
(range, 23-93 y)**SETTINGS / LOCATIONS****59 Institutions in**
the US, Canada,
and Israel**INTERVENTION****214** Participants randomized and analyzed at 3 mo**138 Stereotactic radiosurgery (SRS)**

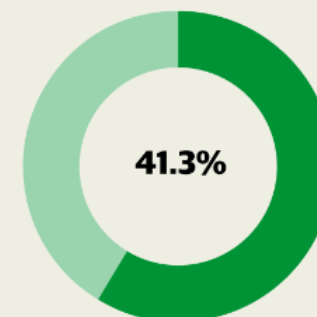
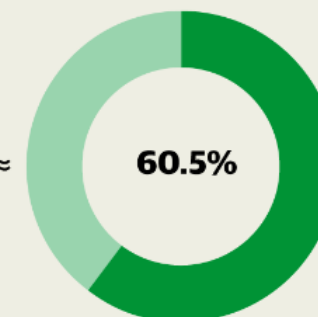
16-Gy or 18-Gy single-dose SRS delivered to the involved spine only

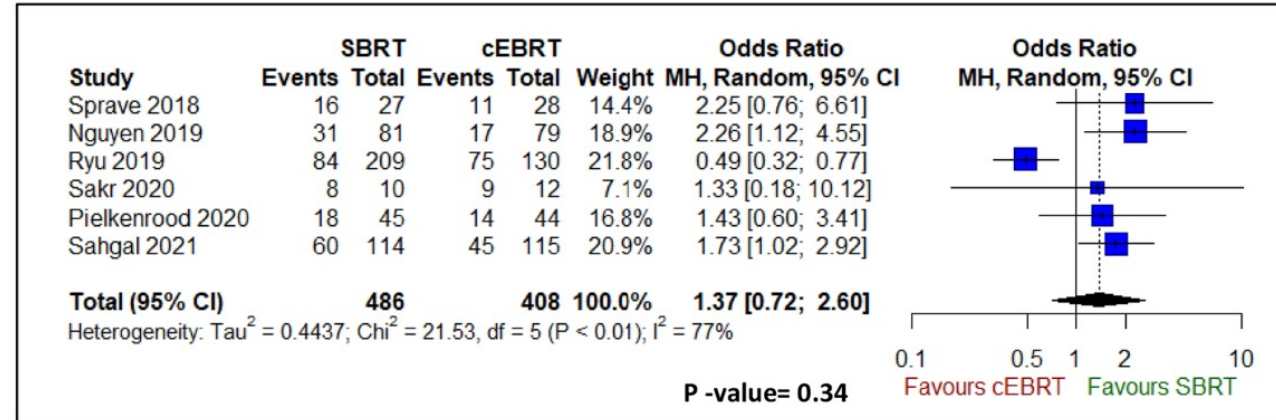
76 Conventional external beam radiotherapy (cEBRT)

8-Gy single-dose cEBRT administered to the involved spine plus 1 spine segment above and below

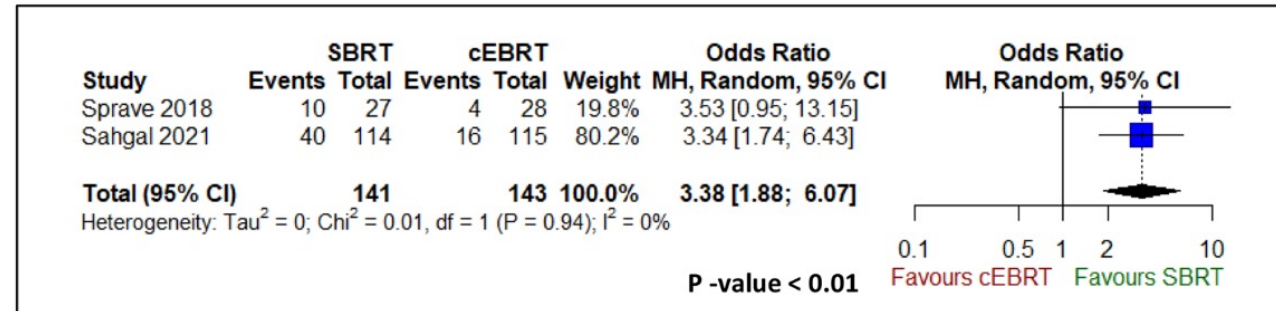
PRIMARY OUTCOMEPain response at 3-mo posttreatment, defined as pain score of 0 (or ≥ 3 point decline) at the index site and no increases in narcotic pain medication or secondary site pain score; pain measured using the Numerical Rating Pain Scale (NRPS; range, 0 [no pain] to 10 [extreme pain])**FINDINGS**

There was no significant difference between SRS and cEBRT in the proportion of participants with pain response

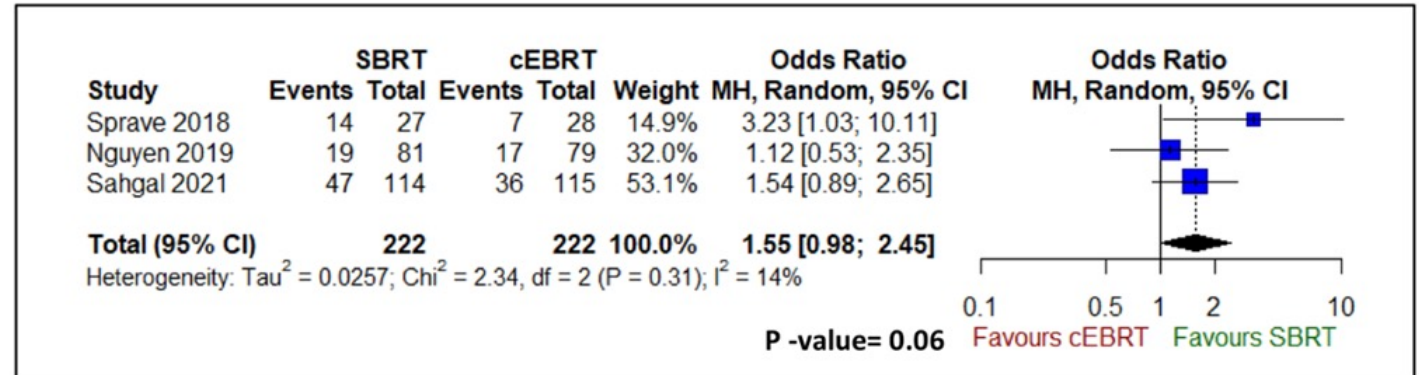
Proportion of participants with pain response at 3 mo**SRS****57 of 138 Participants****cEBRT****46 of 76 Participants****Between-group difference, SRS vs cEBRT:** -19 percentage points (95% CI, -32.9 to -5.5); 1-sided *P* value = .99



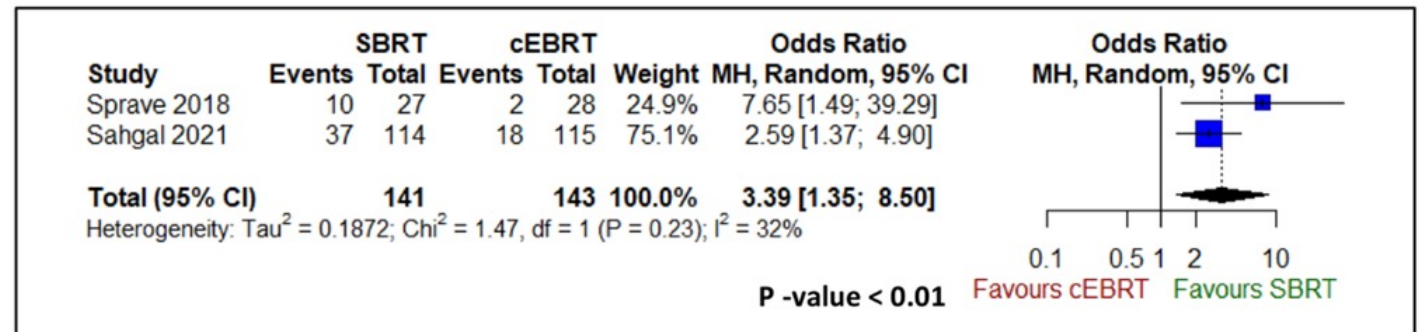
A: Overall pain response at 3 months



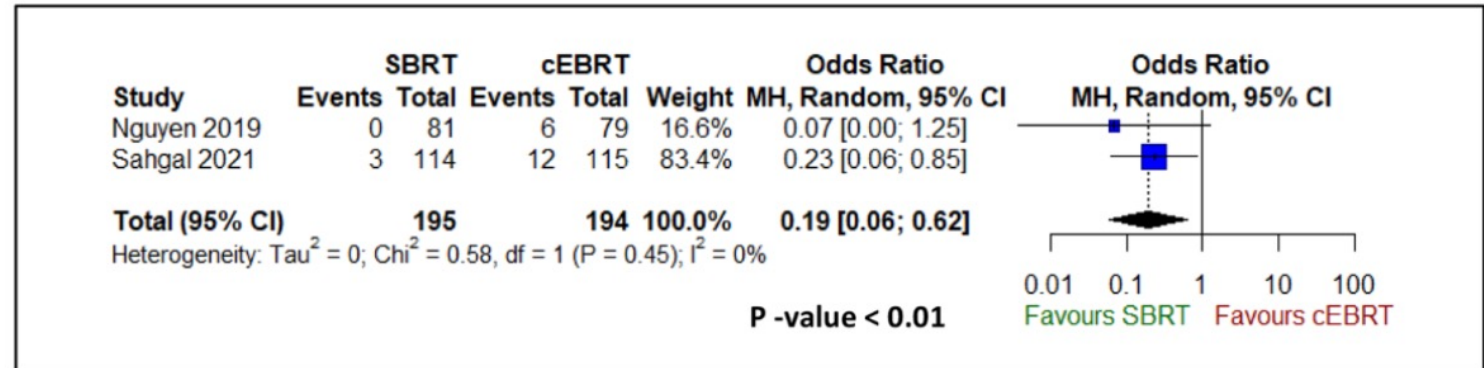
B: Complete pain response at 3 months



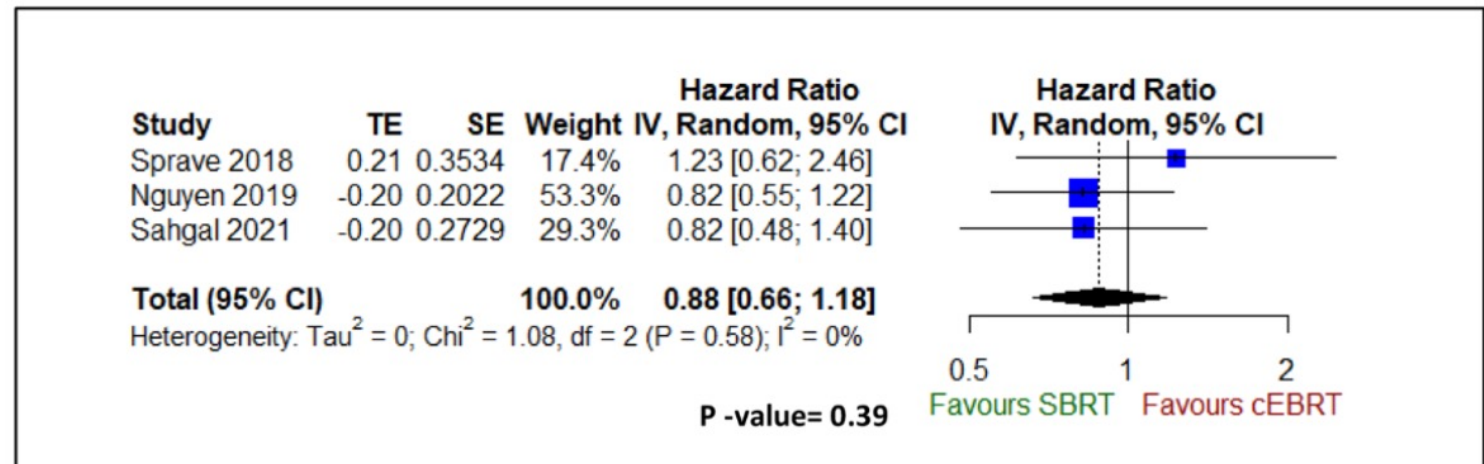
C: Overall pain response at 6 months



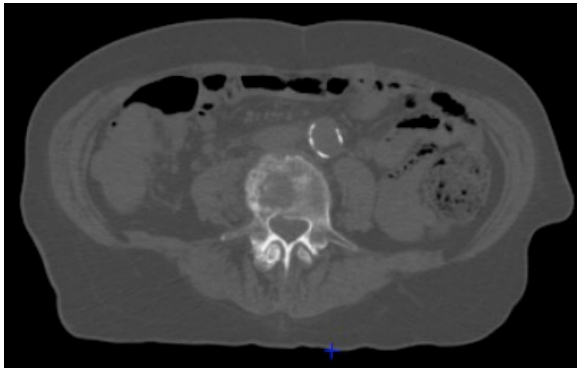
D: Complete pain response at 6 months



A: Local progression



B: Overall survival

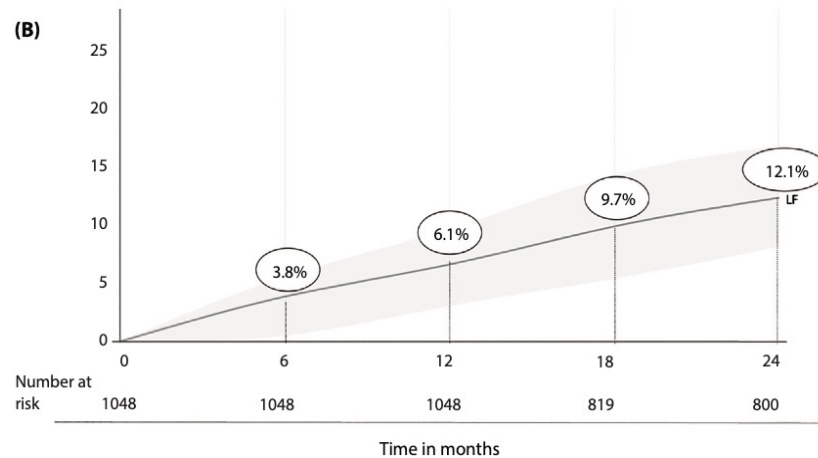
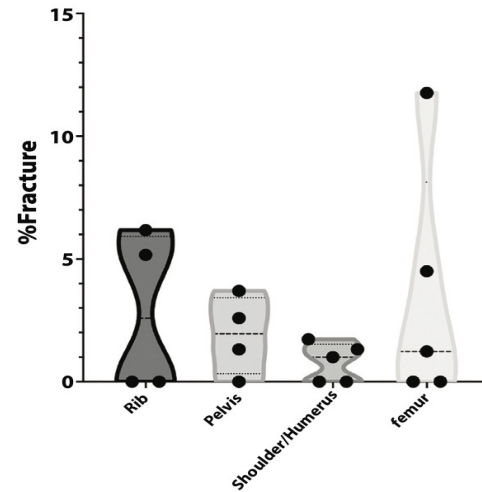
**Table 3.** Significant Predictors of VCF on Univariate and Multivariate Analysis

Factor	Univariate <i>P</i>	Multivariable Fine and Grey Model		
		<i>P</i>	HR	95% CI
Vertebral body collapse	< .001	Global, < .001		
≥ 50% VCF		.0189	6.92	1.38 to 34.77
< 50% VCF		< .001	8.98	4.48 to 18.00
No VCF but > 50% of vertebral body involved		< .001	4.46	2.08 to 9.57
Dose/fraction, Gy	< .001	Global, < .001		
≥ 24		< .001	5.25	2.29 to 12.01
20-23		< .001	4.91	1.96 to 12.28
Alignment	.0027	< .001	2.99	1.57 to 5.70
Bone lesion type	< .001	.0022	3.53	1.58 to 7.93
Paraspinal/epidural extension	.0036	NS		

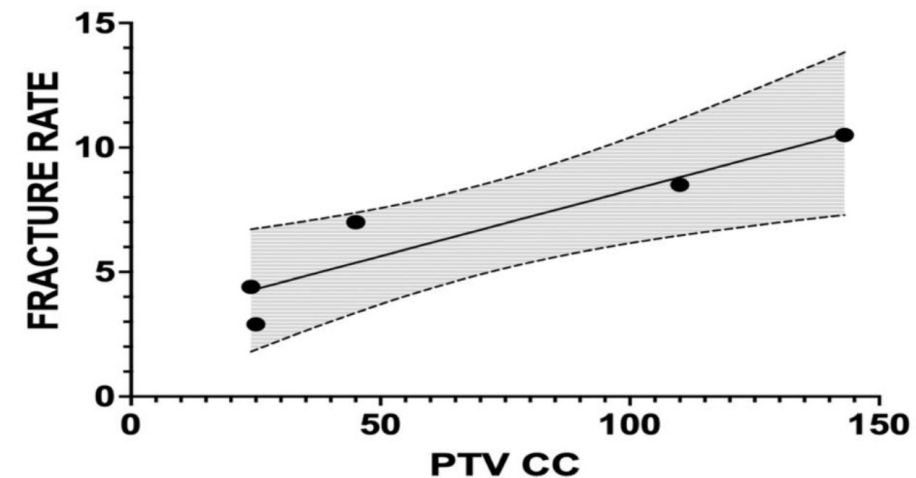
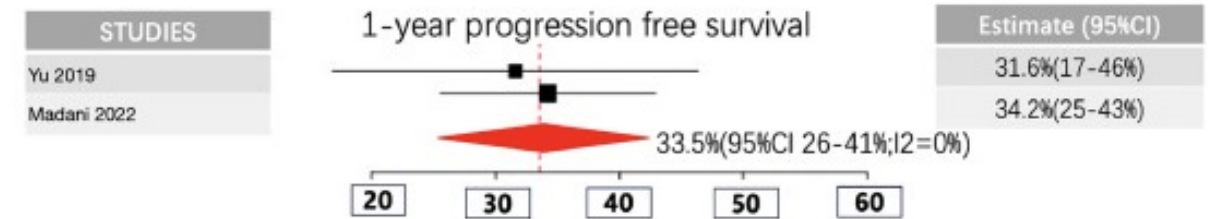
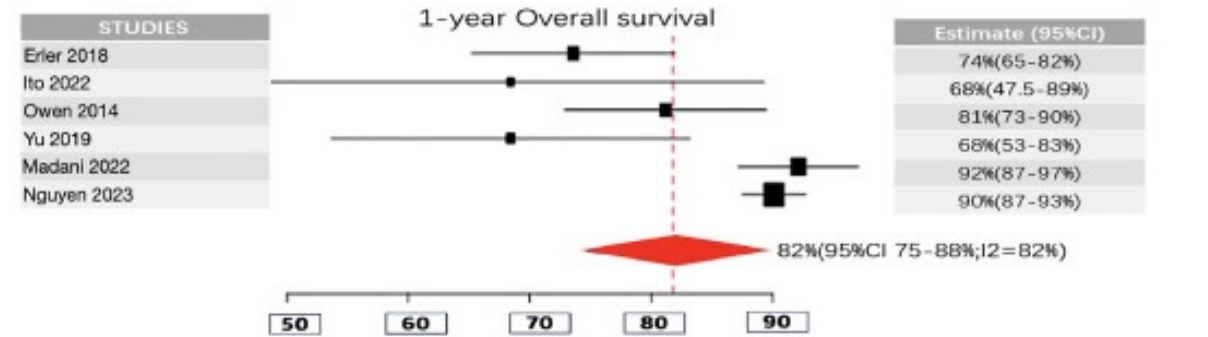
NOTE. For vertebral body collapse, the reference is no VCF and less than 50% vertebral body involvement; for dose/fraction, the reference is ≤ 19 Gy/fraction; the reference for alignment was normal, and yphosis/scoliosis and subluxation/translation were grouped as only one patient had subluxation; and the reference for bone lesion was grouped according to mixed and osteoblastic tumor versus osteolytic, given that the majority of VCFs occurred in lytic tumors.

Abbreviations: HR, hazard ratio; NS, not significant; VCF, vertebral compression fracture.

- Fracture: cotes, fémur
- Flaire-up: 7.5%



Moraes, IJROBP, 2024, Singh, IJROBP, 2024



Quelle efficacité?

Bone

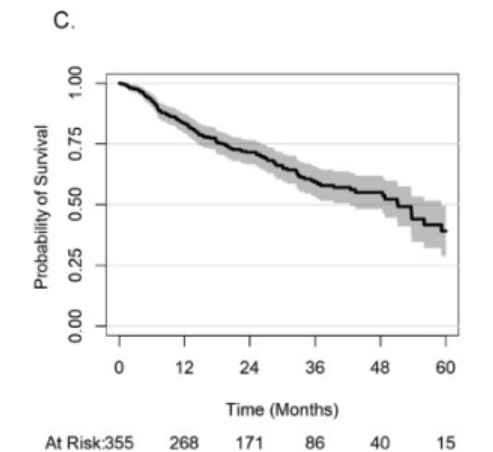
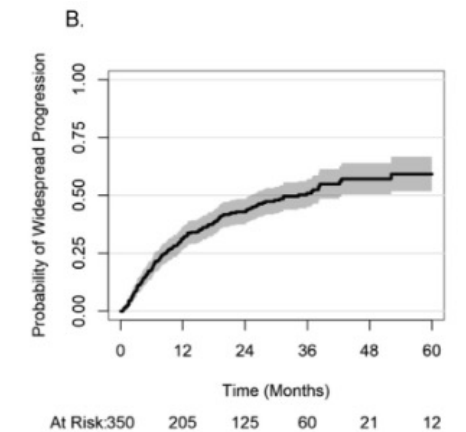
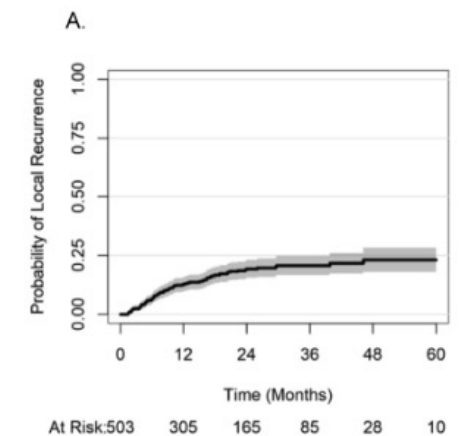
Table 3

Final multivariable regression models for local recurrence.

Covariate for All Bone Lesions	Sub-HR (95% CI)	p value
Radioresistant Histology	2.49 (1.61–3.87)	<0.001
Treatment at initial oligometastatic presentation to SBRT	0.58 (0.34–0.97)	0.038
PTV size \geq median	2.11 (1.28–3.46)	0.0033
PTV Dmin (BED10) \geq median	0.53 (0.33–0.87)	0.011

Covariate for Non-Spine Bone Lesions	Sub-HR (95% CI)	p value
Primary Histology		
Prostate (ref)	1	
Renal cell	10.8 (3.21–36.1)	<0.001
NSCLC	6.48 (2.05–20.5)	0.0015
Other	2.60 (0.75–9.0)	0.13
PTV size \geq median	5.02 (1.39–18.2)	0.014

Covariate for Spine Lesions	Sub-HR (95% CI)	p Value
Radioresistant Histology	2.11 (1.25–3.57)	0.0051
PTV Dmin (BED10) \geq median	0.46 (0.26–0.82)	0.0085
Epidural Disease	1.99 (1.13–3.49)	0.016



Cohorte
prospective

Quelle efficacité?

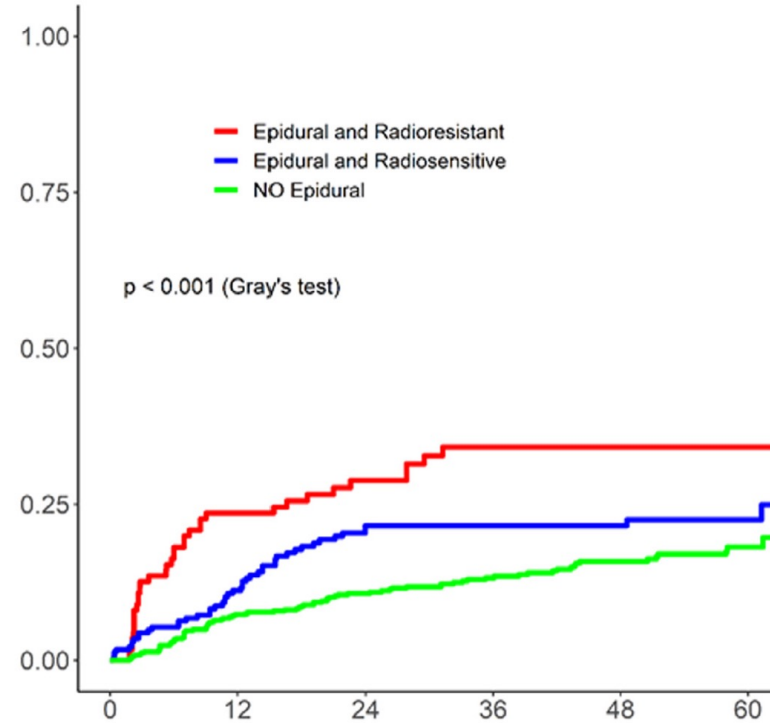
Rachis

Bone

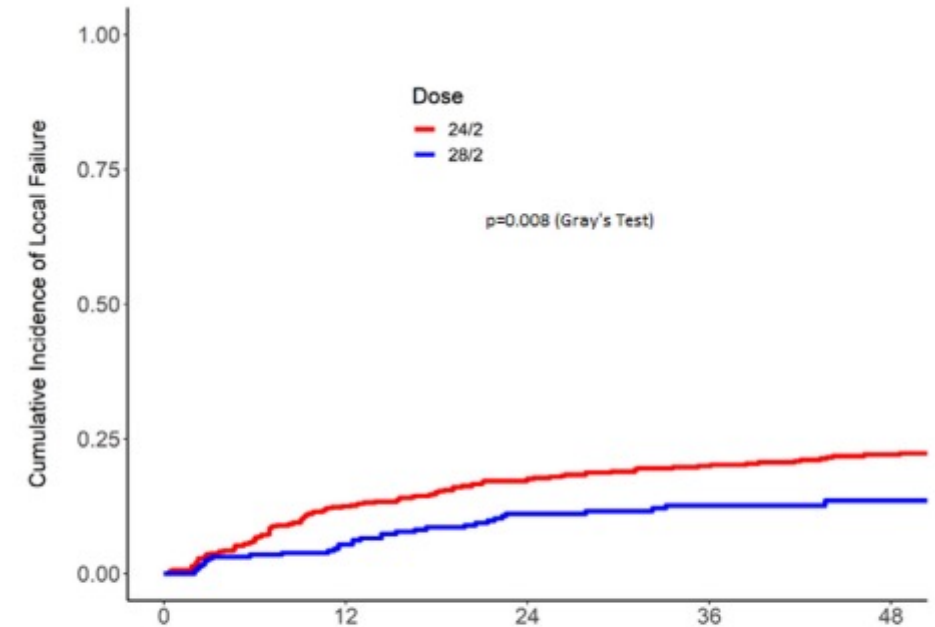
28Gy/2 (159)

24Gy/2 (343)

Cumulative Incidence of Local Failure



	Months since start of treatment					
	0	12	24	36	48	60
	Numbers at risk					
Epidural and Radioresistant	114	44	24	12	4	3
Epidural and Radiosensitive	230	129	79	57	34	12
NO Epidural	594	376	269	184	112	48



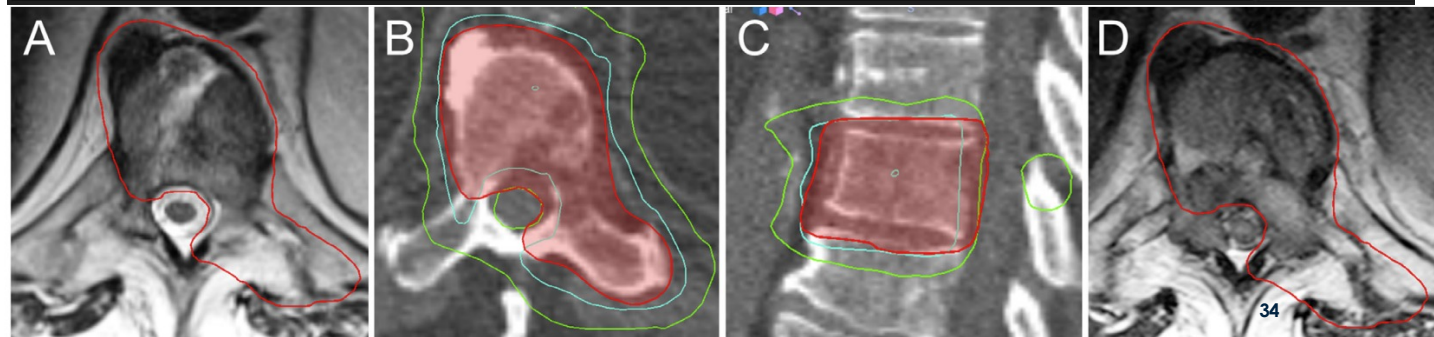
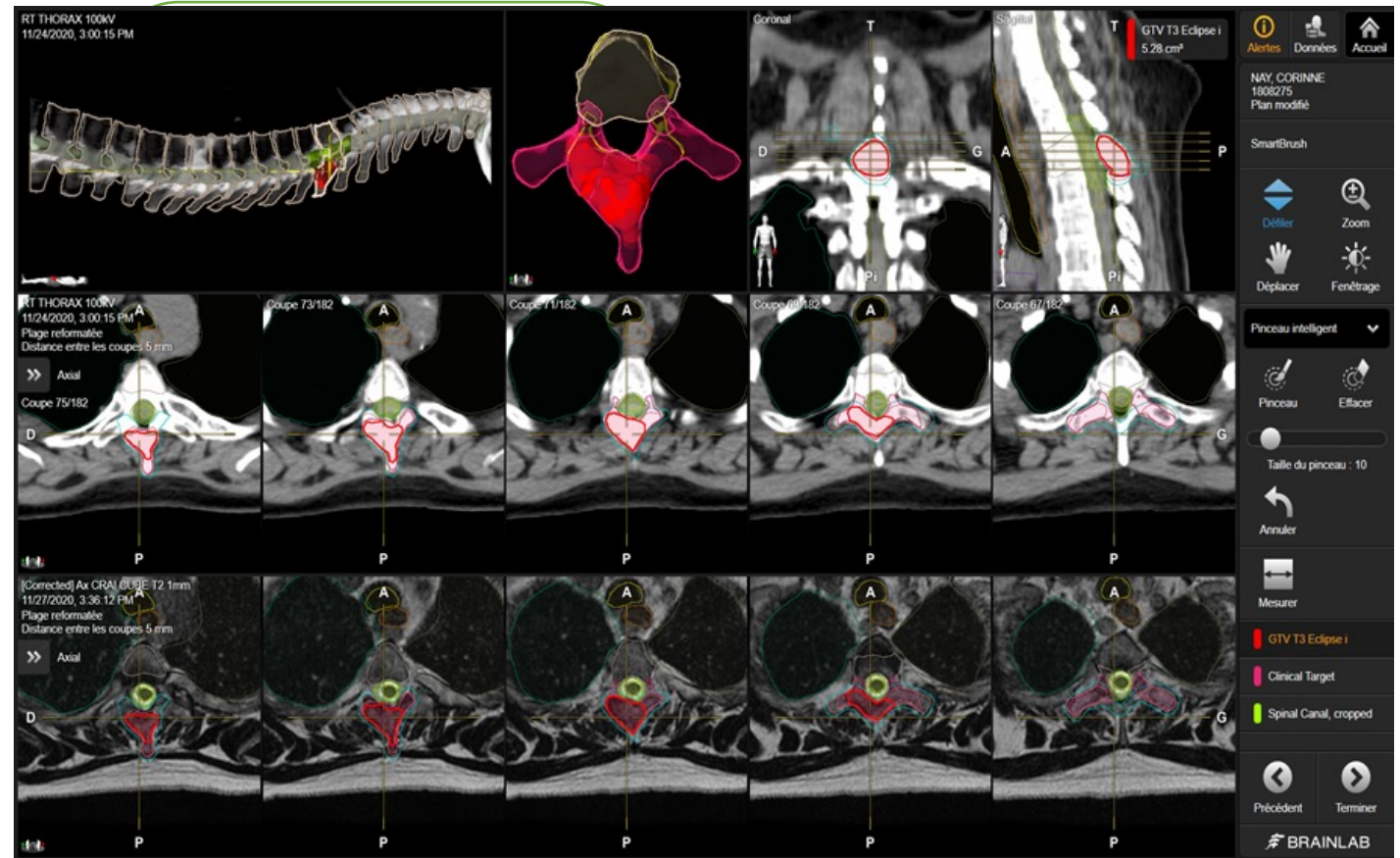
Dose	Months since start of treatment				
	0	12	24	36	48
24/2	646	359	238	163	107
28/2	301	192	135	91	43
Numbers at risk					

Quality control

Rachis

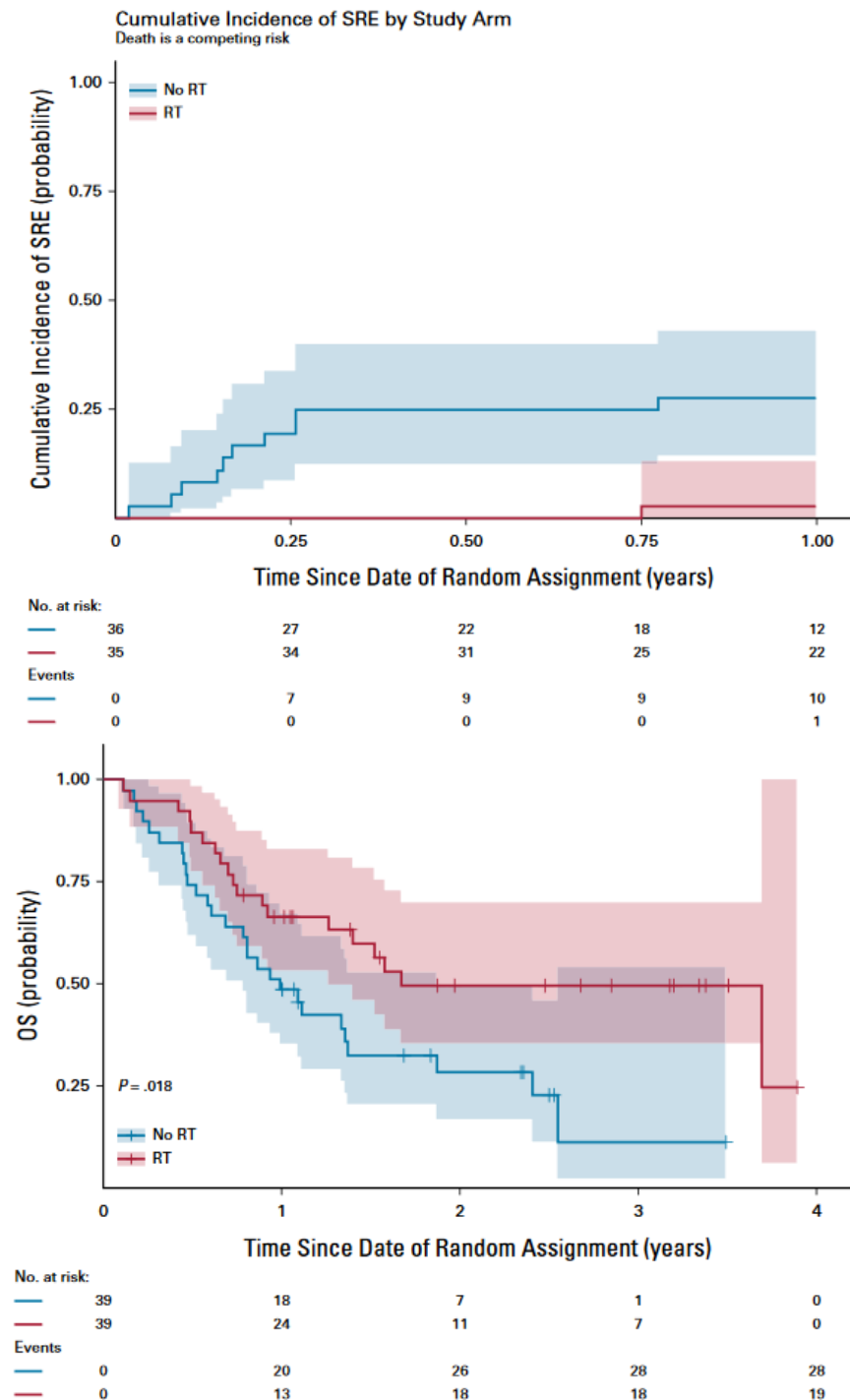
Bone

- ✓ 283 patients
- ✓ 360 lesions
- ✓ 17% of deviations



Radiothérapie préventive

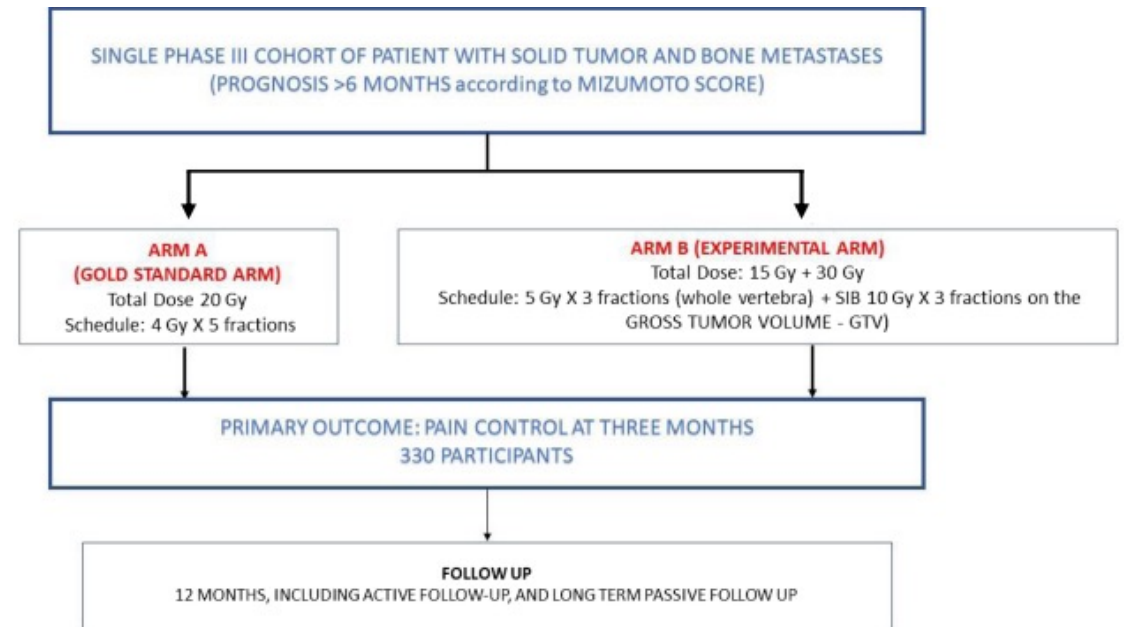
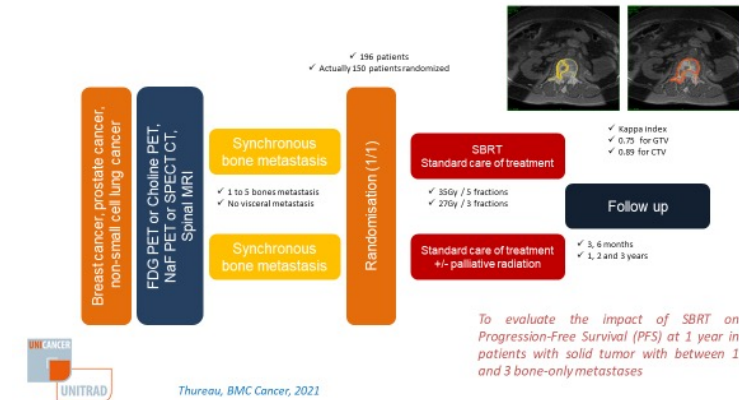
- Radiothérapie prophylactique
- Phase II
- 78 patients randomisés
- 37 lésions rachidiennes
- Facteurs d'OS en multivarié
 - RT
 - Type de primitif
- Type de RT
 - 27/3 19 lésions
 - 20/5, 30/10 ou 8/1



Conclusion

- Pour quels patients?
 - Oligométastatique
 - Oligoprogressif
- Quels fractionnements?
 - Importance de la dose
- Quels objectifs cliniques?
 - Douleur
 - Control local
 - PFS, OS....

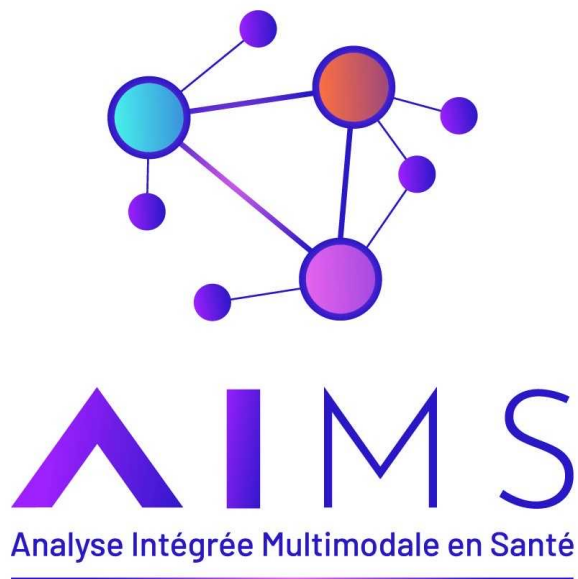
STEREO OS - NCT0314332



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Pr F. Guisier
Pr C. Collet Savoye
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Mr O. Rastelli, Mme L. Burel
Mr LF. Pepin
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Dr JC. Faivre,
Dr A. Arnaud**



Radiotherapy for bone metastases

RECORAD 2025

N Martz

