



# Sovradosaggio con radioterapia stereotassica dopo Radioterapia ad Intensità Modulata (IMRT) in pazienti con carcinoma del rinofaringe.

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**CLINICAL INVESTIGATION**

**Head and Neck**

**EXPERIENCE IN FRACTIONATED STEREOTACTIC BODY  
RADIATION THERAPY BOOST FOR NEWLY DIAGNOSED  
NASOPHARYNGEAL CARCINOMA**

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**Boost con CK-SBRT frazionata 12-15 Gy (3 Gy/fr) dopo RT  
convenzionale (64.8-68.4 Gy - 1.8 Gy frazione)**

**Controllo locale a 3 anni 93.1%**

**Conclusioni:**

**CK-SBRT come boost è un trattamento fattibile, ottiene un buon  
controllo locale senza complicanze severe.**

**(64 Pz 2003 – 2006)**

**CLINICAL INVESTIGATION**

**Head and Neck**

**EXCELLENT LOCAL CONTROL WITH STEREOTACTIC RADIOTHERAPY BOOST  
AFTER EXTERNAL BEAM RADIOTHERAPY IN PATIENTS WITH  
NASOPHARYNGEAL CARCINOMA**

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**Boost CK-SBRT dopo IMRT (66 Gy 2-2.2 Gy fr) :**

**7-15 Gy in unica frazione**

**Freedom From Local Relapse a 5 aa : 98%; OS: 69%**

**Conclusioni:**

**SBRT buon controllo locale**

**Bisogna migliorare la delineazione del target per evitare  
complicanze a lungo termine.**

**Necessità di nuovi farmaci.**

**82 Pz**

## Casistica (2010-2014)

Tot 5 pz

- CT neoadiuvante e/o concomitante
- Trattamento IMRT precedente
- Stadio di malattia cT2-cT4 all'esordio
- Massimo volume malattia post IMRT 24 cc

## **CASISTICA (2010-2014)**

<b>Maschi</b>	<b>3</b>	
<b>Femmine</b>	<b>2</b>	
<b>Ca. indiff. tipo RNF</b>	<b>3 Pz</b>	
<b>Ca. squamoso</b>	<b>2 Pz</b>	
<b>Età media</b>	<b>53,2 aa</b>	<b>Range 39 - 72</b>
<b>Follow up medio</b>	<b>35.5 mesi</b>	<b>Range 26 - 45</b>

	<b>TNM</b>	<b>STADIO</b>
<b>Pz 1</b>	cT2 N1	II
<b>Pz 2</b>	cT2 N2	III
<b>Pz 3</b>	cT4 N1	IV a
<b>Pz 4</b>	cT4 N0	IV a
<b>Pz 5</b>	cT2 N2	III

## CHEMIOTERAPIA

<b>Pz 1</b>	Cisplatino sett. concomitante + CHT sec. schema Al Sarraf a dose ridotta di consolidamento.
<b>Pz 2</b>	PEE x 2 neoadiuvante (Cispl. +Etoposide+Epirubicina) , Carbopl. concomitante e PEE x2 successivo.
<b>Pz 3</b>	Al Sarraf x 2 neadiuvante e Cisplatino concomitante
<b>Pz 4</b>	PEE x2 neoadiuvante, Cisplatino concomitante e poi PEE x2
<b>Pz 5</b>	Cispl + FU neoadiuvante ; Cisplatino sett. concomitante

# TRATTAMENTO RT

4 Pz    PTV 1 60 Gy    PTV 2 54 Gy    SIB-IMRT (30 fr)

1 Pz    PTV 1 63.90 Gy    PTV 2 60 Gy    SIB-IMRT (30 fr)



## Tossicità RTOG

	Grado 1	Grado 2	Grado 3	Grado 4
<b>Cute</b>	0	5	0	0
<b>Mucose</b>	0	4	1	0
<b>Disfagia Odinofagia</b>	2	2	1	0

## PET – TAC di ristadiatione dopo almeno 40 gg dal termine RT/CT concomitante

	Pre RT	Post RT	
Pz 1	cT2 N1	cT1 N0	Sensibile diminuzione del SUV
Pz 2	cT4 N1	cT2 N0	Sensibile diminuzione del SUV
Pz 3	cT4 N1	cT2 N0	Sensibile diminuzione del SUV
Pz 4	cT4 N2	cT3 N0	Sensibile diminuzione del SUV
Pz 5	cT2 N2	cT1N0	Sensibile diminuzione del SUV

**Intervallo termine IMRT inizio boost CK-SBRT da 1.5 a 3 mesi e 12 mesi per il paziente recidivato (rispettivamente di 1.5, 2.5, 3, 3 e 12 mesi)**

**Trattamento CK-SRBT :**

- **confezionamento nuovo mould**
- **TAC di centratura: spirale, slides 1 mm senza intervalli, dal vertex a C7**

**Pianificazione**

**contouring PTV: lesione visibile alla PET post EBRT + 2/3 mm di margine**

<b>Dose (Gy)</b>	<b>N° fr</b>	<b>Isodose di prescrizione</b>	<b>N° Pz</b>
<b>12</b>	<b>2</b>	<b>80%</b>	<b>1</b>
<b>16</b>	<b>2</b>	<b>77%</b>	<b>2</b>
<b>16</b>	<b>2</b>	<b>80%</b>	<b>3</b>
<b>20</b>	<b>2</b>	<b>80%</b>	<b>4</b>
<b>24</b>	<b>3</b>	<b>80%</b>	<b>5</b>

## SUMMARY OF SUGGESTED DOSE CONSTRAINTS FOR CRITICAL ORGANS

Stereotactic body radiation therapy : The report of AAPM Task Group 101

SERIAL TISSUE	MAX CRITICAL VOLUME ABOVE THRESHOLD	ONE FRACTION		THREE FRACTIONS		FIVE FRACTIONS		ENDPOINT (≥GRADE 3)
		THRESHOLD DOSE (Gy)	MAX POINT DOSE (Gy)**	THRESHOLD DOSE (Gy)	MAX POINT DOSE (Gy)**	THRESHOLD DOSE (Gy)	MAX POINT DOSE (Gy)**	
Optic pathway	<0.2 cc	8 Gy	10 Gy	15.3 Gy (5.1 Gy/fx)	17.4 Gy (5.8 Gy/fx)	23 Gy (4.6 Gy/fx)	25 Gy (5 Gy/fx)	Neuritis
Cochlea			9 Gy		17.1 Gy (5.7 Gy/fx)		25 Gy (5 Gy/fx)	Hearing loss
Brainstem (not medulla)	<0.5 cc	10 Gy	15 Gy	18 Gy (6 Gy/fx)	23.1 Gy (7.7 Gy/fx)	23 Gy (4.6 Gy/fx)	31 Gy (6.2 Gy/fx)	Cranial neuropathy
Spinal cord and medulla	<0.35 cc <1.2 cc	10 Gy 7 Gy	14 Gy	18 Gy (6 Gy/fx) 12.3 Gy (4.1 Gy/fx)	21.9 Gy (7.3 Gy/fx)	23 Gy (4.6 Gy/fx) 14.5 Gy (2.9 Gy/fx)	30 Gy (6 Gy/fx)	Myelitis
Spinal cord subvolume (5–6 mm above and below level treated per Ryu)	<10% of subvolume	10 Gy	14 Gy	18 Gy (6 Gy/fx)	21.9 Gy (7.3 Gy/fx)	23 Gy (4.6 Gy/fx)	30 Gy (6 Gy/fx)	Myelitis

I limiti di dose sono stati calcolati in relazione alle tabelle di riferimento ed alla dose totale già ricevuta con IMRT.

<b>Constraints (Gy)</b>	<b>Pz 1</b>	<b>Pz 2</b>	<b>Pz 3</b>	<b>Pz 4</b>	<b>Pz 5</b>
<b>N° frazioni</b>	<b>3</b>	<b>2</b>	<b>2</b>	<b>2</b>	<b>2</b>
<b>Midollo allungato</b>	<b>10 Gy</b>	<b>8 Gy</b>	<b>6 Gy</b>	<b>7.5 Gy</b>	<b>8 Gy</b>
<b>Chiasma</b>	<b>4 Gy</b>	<b>4 Gy</b>	<b>5 Gy</b>	<b>5 Gy</b>	<b>4 Gy</b>
<b>Midollo</b>	<b>14 Gy</b>	<b>10 Gy</b>	<b>4 Gy</b>	<b>5 Gy</b>	<b>5 Gy</b>
<b>nn. ottici</b>	<b>14 Gy</b>	<b>4 Gy</b>	<b>10 Gy</b>	<b>10 Gy</b>	<b>2 Gy</b>

<b>Volume PTV</b>	
<b>Pz 1</b>	<b>25.5 cc</b>
<b>Pz 2</b>	<b>13.4 cc</b>
<b>Pz 3</b>	<b>7.6 cc</b>
<b>Pz 4</b>	<b>24.4 cc</b>
<b>Pz 5</b>	<b>14.3 cc</b>

Setup Isocentric Conformal Evaluate Finetune

Dose Calculation

Algorithm: Monte Carlo

Resolution: High

Uncertainty %: 1

Calculate

Prescription

Prescription

Reference Point

Use max dose point

Dose (cGy): 1500.00

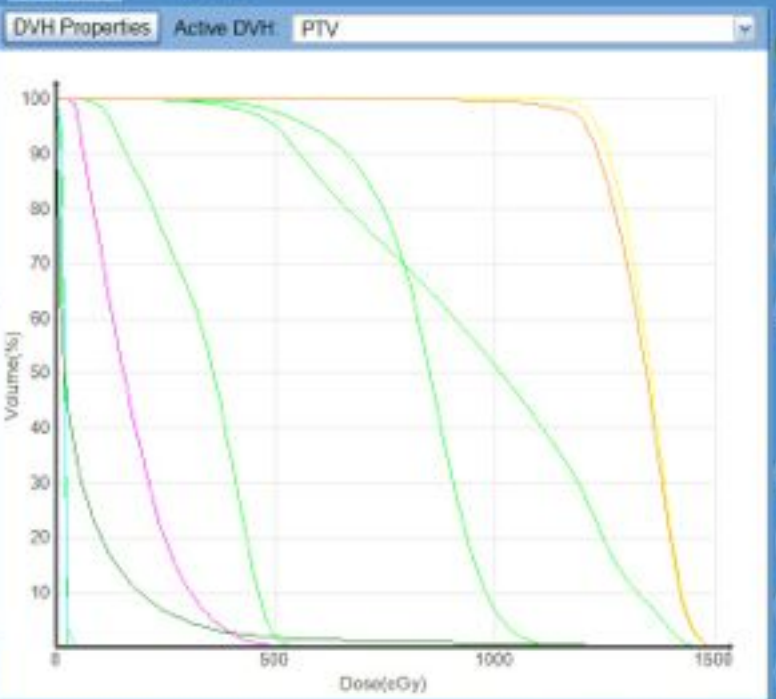
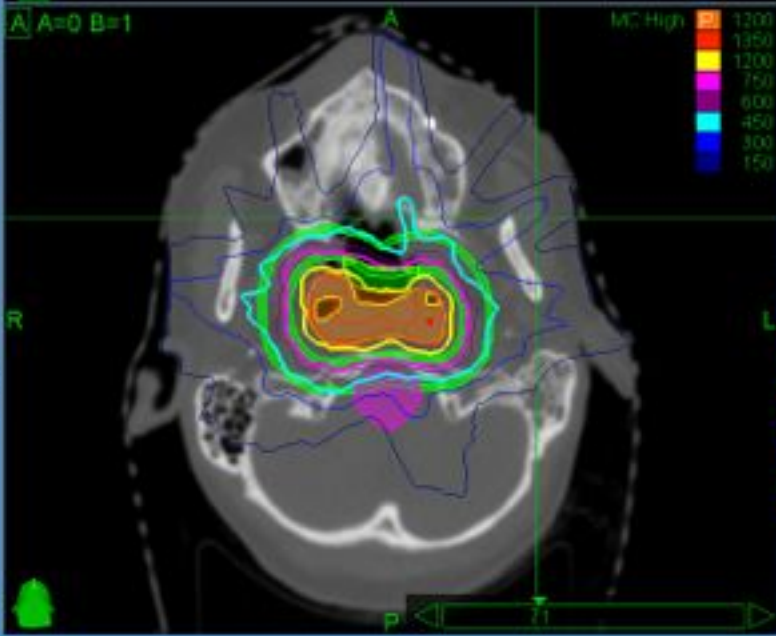
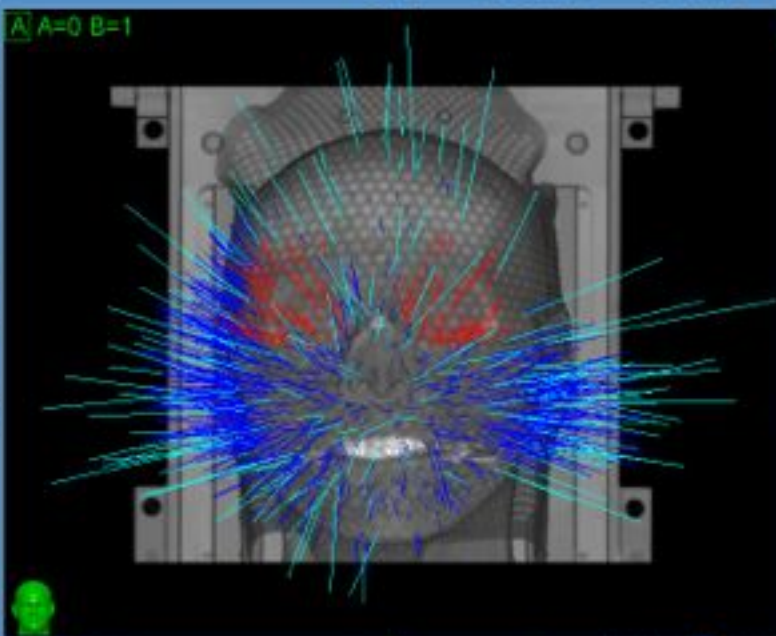
Point: Go to >>

15.23, 48.11, 162.50

Set to Cross-hair Point

Save Plan

Save Plan



Nodes	90	Total MU	13345.54
Beams	167	Min MU	23.21
Max Dose (cGy)	1500.00	Max MU	263.33

Dose Statistics Table

VOI	Min (cGy)	Mean (cGy)	Max (cGy)	CI	nCI	HI	Coverage
PTV	920.46	1335.07	1500.00	1.53	1.59	1.25	96.26%
Left Eye	11.79	19.58	37.56				
Right Eye	12.78	21.16	49.05				
midollo	37.78	174.14	728.88				
Tune 1	304.50	836.20	1183.42				
Tune 2	89.75	335.57	582.71				
plv_carne	1160.23	1347.96	1500.00	1.06	1.08	1.25	99.03%
Critical 9	920.46	1335.07	1500.00				
buchi	242.09	973.76	1489.51				

Standard Display

Plan

fix15\_12GyAt80\_2tz

2012-05-10 12:21:38

Rx

80%, 1200.00 cGy



**Non rilevata Tossicità Acuta o Tardiva dopo CK-SBRT  
per il follow up fino ad ora eseguito**

<b>Risposta</b>		<b>Mesi</b>
RC	NED	44
RC	NED	31
RP 50%	Exitus per pro	7
RC	Viva con M+	35
RC <b>Pz</b>	NED	26

## Conclusioni

**CK-SBRT come sovradosaggio in pazienti trattati con SIB-IMRT per carcinoma del Rinofaringe è un trattamento fattibile e ben tollerato che dà un buon controllo locale di malattia.**



Original article

Evolution of treatment for nasopharyngeal cancer – Success and setback in the intensity-modulated radiotherapy era

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- Miglioramento tecniche di RT (IMRT dose escalation/IGRT)
- Miglioramento Imaging pre-trattamento
- Nuovi farmaci CHT
- Risultati nei T4 soddisfacenti?



Revue générale

Boost et réirradiation stéréotaxiques des cancers des voies aérodigestives supérieures

Stereotactic irradiation in head and neck cancers

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SBRT come boost o come reirradiazione per migliorare il controllo locale nei tumori della testa collo.



**Grazie per l'attenzione**

<b>Tossicità Acuta dopo IMRT</b>	<b>(RTOG)</b>
<b>Pz 1</b>	Gr. 2 skin; Gr. 4 mucous membrane; Gr. 3/4 Dysphagia or odynophagia
<b>Pz 2</b>	Gr. 2 skin and mucous membrane; Gr. 1 Dysphagia or odynophagia
<b>Pz 3</b>	Gr. 2 skin mucous membrane; Gr. 1 Dysphagia or odynophagia
<b>Pz 4</b>	Gr. 2 skin and mucous membrane; Gr. 2 Dysphagia or odynophagia
<b>Pz 5</b>	Gr. 2 skin and mucous membrane Gr. 2 dysphagia or odynophagia