

SERVIZIO SANITARIO REGIONALE
EMILIA-ROMAGNA
Azienda Ospedaliero - Universitaria di Ferrara

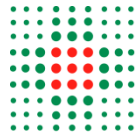
Il Percorso del Paziente con Neoplasia Neuroendocrina nella Provincia di Ferrara

Società Medico Chirurgica

12 ottobre 2019

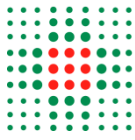
U.O.C. Radioterapia

Dr. Antonio Stefanelli



Background

- ***I tumori neuroendocrini (NET)***
*rappresentano un ampio gruppo di neoplasie rare che hanno origine dalle cellule del sistema neuroendocrino e che, per tale ragione, **sono in grado di colpire diversi organi:***
- *intestino, pancreas, polmoni, tiroide, ghiandole surrenali e timo*



NCCN Guidelines Version 1.2019 Neuroendocrine and Adrenal Tumors

CLINICAL PRESENTATIONS AND DIAGNOSIS^a

Neuroendocrine tumors of the gastrointestinal tract, lung, and thymus.

- [Multiple endocrine neoplasia, type 1 \(See MEN1-1\)](#)
- [Parathyroid](#)
- [Pancreatic neuroendocrine tumors \(PanNETs\)](#)
- [Pituitary tumor](#)
- [Bronchial/thymic](#)

Multiple endocrine neoplasia, type 2 (See MEN2-1)

- [Medullary thyroid carcinoma \(Also see NCCN Guidelines for Thyroid Carcinoma\)](#)
- [Parathyroid](#)
- [Pheochromocytoma](#)

[Merkel cell carcinoma \(See NCCN Guidelines for Merkel Cell Carcinoma\)](#)

[Neuroendocrine tumors of the gastrointestinal tract, lung, and thymus](#)^b

Clinical presentations:

- [Jejunal, ileal, colon \(See NET-1\)](#)
- [Duodenal \(See NET-1\)](#)
- [Appendix \(See NET-2\)](#)
- [Rectal \(See NET-3\)](#)
- [Gastric \(See NET-4\)](#)
- [Thymus \(See NET-5\)](#)
- [Bronchopulmonary, atypical lung carcinoid \(See NET-6\)](#)
- [Locoregional advanced disease and/or distant metastases](#)
 - ▶ [Bronchopulmonary/thymus \(See NET-7 and NET-9\)](#)
 - ▶ [GI Tract \(See NET-10\)](#)
- [Carcinoid syndrome \(See NET-11\)](#)

[Neuroendocrine tumors of the pancreas](#)^b

Clinical presentations:

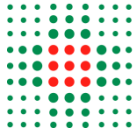
- [Nonfunctioning pancreatic tumors \(See PanNET-1\)](#)
- [Gastrinoma \(See PanNET-2\)](#)
- [Insulinoma \(See PanNET-3\)](#)
- [Glucagonoma \(See PanNET-4\)](#)
- [VIPoma \(See PanNET-5\)](#)
- [Locoregional unresectable disease and/or distant metastases \(See PanNET-7\)](#)

[Neuroendocrine tumors of unknown primary \(See NUP-1\)](#)^b

[Adrenal gland tumors \(See AGT-1\)](#)^c

[Pheochromocytoma/paraganglioma \(See PHEO-1\)](#)

[Extrapulmonary: Poorly differentiated neuroendocrine carcinoma/Large or small cell carcinoma other than lung/Unknown primary \(poorly differentiated\)](#)
([See PDNEC-1](#))



NCCN Clinical Practice Guidelines in Oncology (NCCN Guidelines®)

Neuroendocrine and Adrenal Tumors

Version 1.2019 — March 5, 2019

NCCN.org

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Cancer
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Neuroendocrine and Adrenal Tumors

[NCCN Neuroendocrine Tumors Panel Members](#)
[Summary of the Guidelines Updates](#)

[Clinical Presentations and Diagnosis \(CP-1\)](#)

[Neuroendocrine Tumors of the Gastrointestinal Tract, Lung, and Thymus \(Carcinoid Tumors\) \(NET-1\)](#)

[Neuroendocrine Tumors of the Pancreas \(PanNET-1\)](#)

[Neuroendocrine Tumors of Unknown Primary \(NUP-1\)](#)

[Adrenal Gland Tumors \(AGT-1\)](#)

[Pheochromocytoma/Paraganglioma \(PHEO-1\)](#)

[Poorly Differentiated Neuroendocrine Carcinoma/Large or Small Cell \(other than lung\) \(PDNEC-1\)](#)

[Multiple Endocrine Neoplasia, Type 1 \(MEN1-1\)](#)

[Multiple Endocrine Neoplasia, Type 2 \(MEN2-1\)](#)

[Principles of Pathology for Diagnosis and Reporting of Neuroendocrine Tumors \(NE-A\)](#)

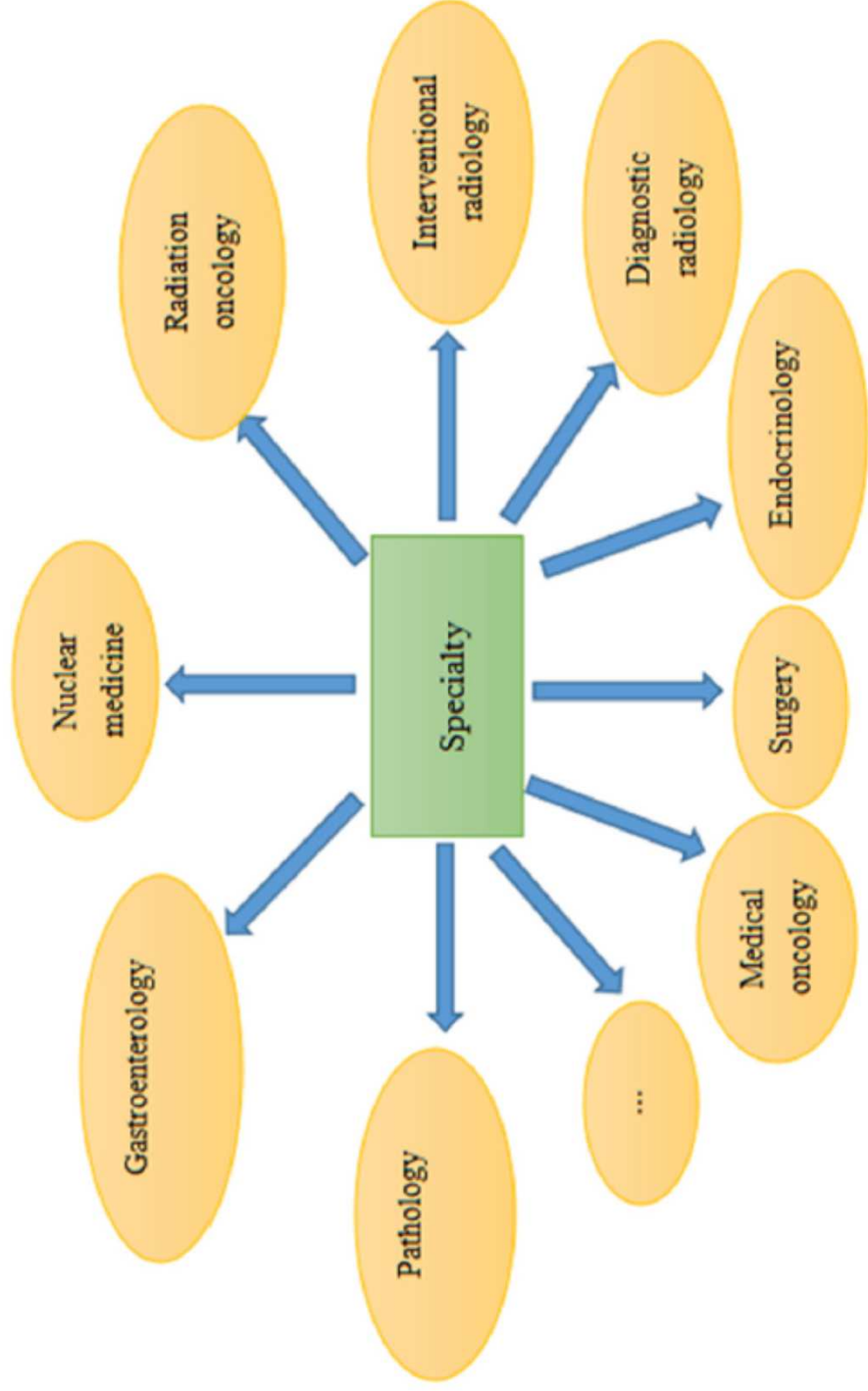
[Principles of Biochemical Testing \(NE-B\)](#)

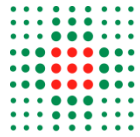
[Surgical Principles for Management of Neuroendocrine Tumors \(NE-C\)](#)

[Principles of Systemic Anti-Tumor Therapy \(NE-D\)](#)

[Principles of Peptide Receptor Radionuclide Therapy \(PRRT\) with 177Lu-dotatate \(NE-E\)](#)

[Staging \(ST-1\)](#)





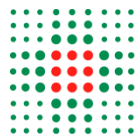
RADIOTERAPIA ESTERNA:

CONVENZIONALE (con tecniche evolute
IMRT VMAT IGRT)

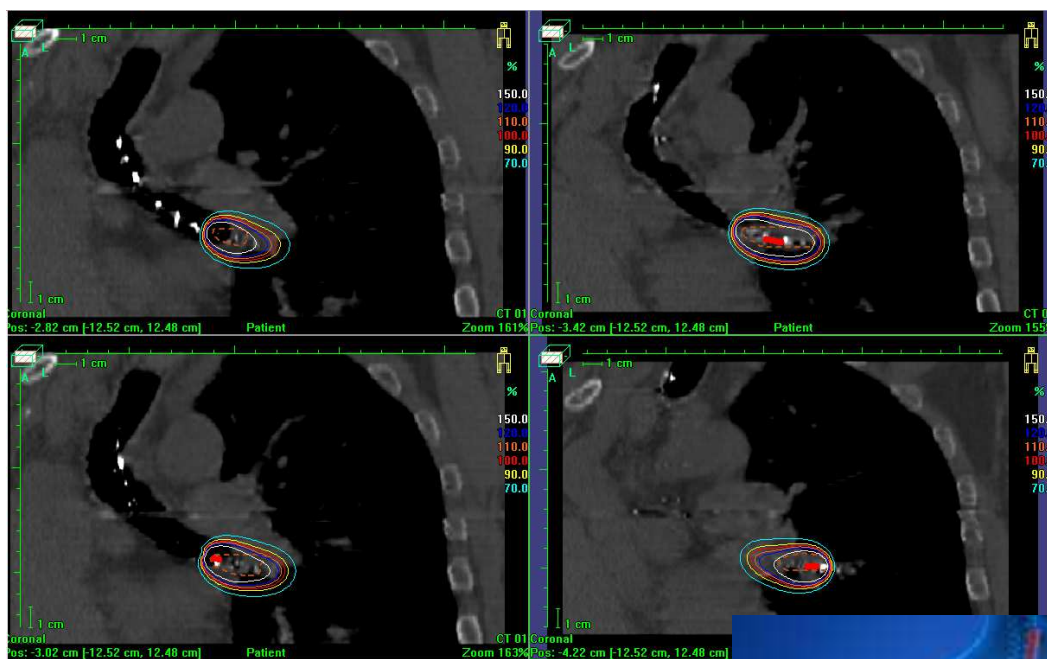
STEREOABLATIVA

IPOFRAZIONATA





BRACHITERAPIA ENDOLUMINALE





Original Article

External Beam Radiotherapy in the Treatment of Gastroenteropancreatic Neuroendocrine Tumours: A Systematic Review

Table 1
Characteristics of included studies

Reference	Primary	Grade	No. receiving RT alone/no. receiving RT with surgery	Dose/fractionation	Radiation technique (eg, CRT versus SBRT/SRS)	Radio-sensitising chemotherapy	Local RFS	Overall RFS	Overall survival	Acute toxicity (grade 3+)	Late toxicity
[4]	Pancreas	10 grade 2, 1 grade 3	11/0	50.4 Gy	NA	ChemoRT (7) – capecitabine	3/11 patients recurred	15 months	32 months	Grade 3 toxicity (1)	Grade 3 toxicity (1)
[5]	Pancreas	6 grade 1–2, 11 grade 3	0/17	50.4 Gy/28	3D–CRT	7 NA, 10 A – 14 with chemoRT (5-FU or capecitabine)	2 year RFS 85%; 3/17 patients recurred	2 year RFS 46%	56 months	NA	NA
[6]	Pancreas	6 grade 1, 1 grade 2, 2 grade 3, 7 not specified	0/16	50.4 Gy/28	2D–CRT (6), 3D–CRT (5), IMRT (4)	ChemoRT (8) – 5-FU (4) or capecitabine (4); RT (8)	1/16 patients recurred	12 months	5 year OS 28%	Grade 3 enteritis (3)	NR
[7]	Pancreas	NA	14/0	58.4 Gy	NA	NA	2.1 years		2 year	Grade 3 gastric perforation (1), grade 3 large bowel inflammation/sepsis (1), Grade 3 fatigue (1)	Grade 3 duodenal stricture (1), grade 3 gastrointestinal bleed (1), grade 5 duodenal perforation (1)
[8]	Pancreas	Well-differentiated	6/0	50.4 Gy/28 for 2 patients; unknown for other 4	3D–CRT	5-FU or capecitabine	0/6 patients progressed	0/6 patients progressed	NA	NA	NA
[9]	Pancreas	Metastatic GEPNET to bone (34) or soft tissue (11)	45/0	NA	NA	NA	RT to metastases (4)	4 months	NA	NA	NA
[10]	Pancreas	Metastatic GEPNET to brain	NA	NA	WBRT or gamma knife SRS	WBRT or gamma knife SRS	2.1 months	19 months	19 months	NA	NA
[11]	Pancreas	Metastatic GEPNET to brain	NA	NA	WBRT (n = 24), partial brain RT (n = 7), SRS (n = 6)	WBRT (n = 24), partial brain RT (n = 7), SRS (n = 6)	6.5 months	14.3 months	14.8 months	NA	NA
[12]	Pancreas	Metastatic PNET (liver and bone)	21/0	24.6 Gy	NA	NA	1.5 years	NA	NA	Nil	Nil

By contrast, the role of EBRT or chemoradiation as definitive treatment of pancreatic NETs will be difficult to define even with further evidence. As small PNETs may be



Carcinoid Tumor Arising from the Sphenoid Sinus Treated with Definitive Intensity-modulated Radiation Therapy: A Case Report

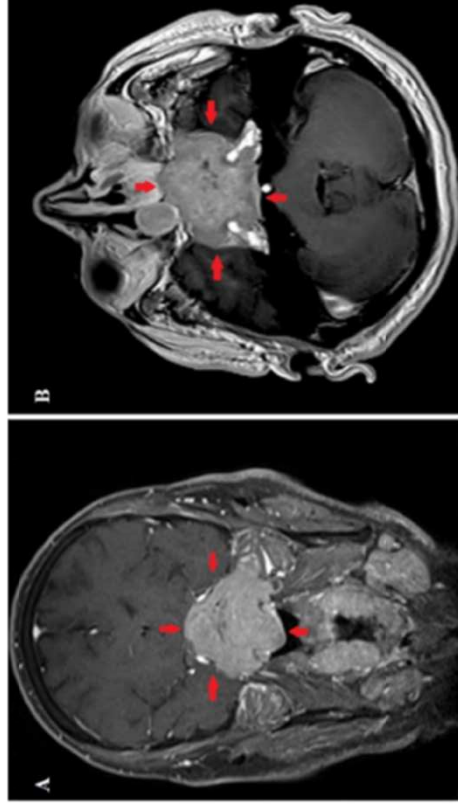


FIGURE 1: Coronal (A) and axial (B) T1-weighted magnetic resonance images with contrast showing a large unresectable sphenoid sinus tumor.

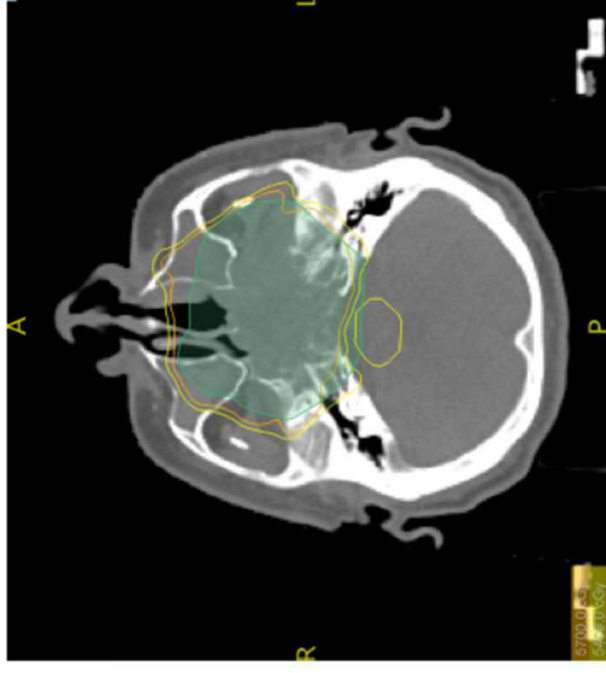
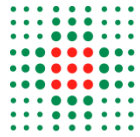


FIGURE 3: Axial image of a representative cut from the intensity-modulated radiation therapy plan showing adequate isodose lines coverage of the planning target volume.

Green: planning target volume (PTV)

Yellow: 90% isodose line

Orange: 95% isodose line



Background

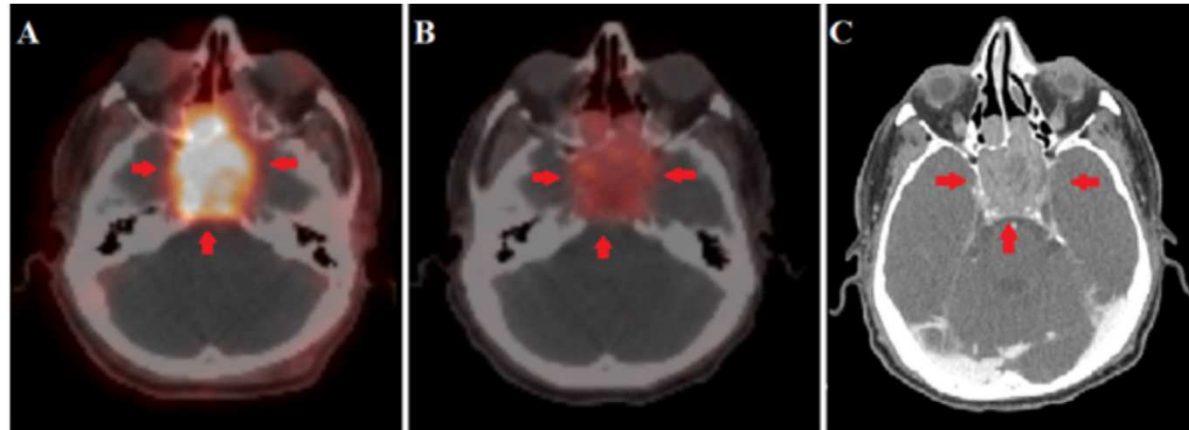
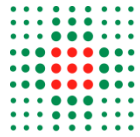


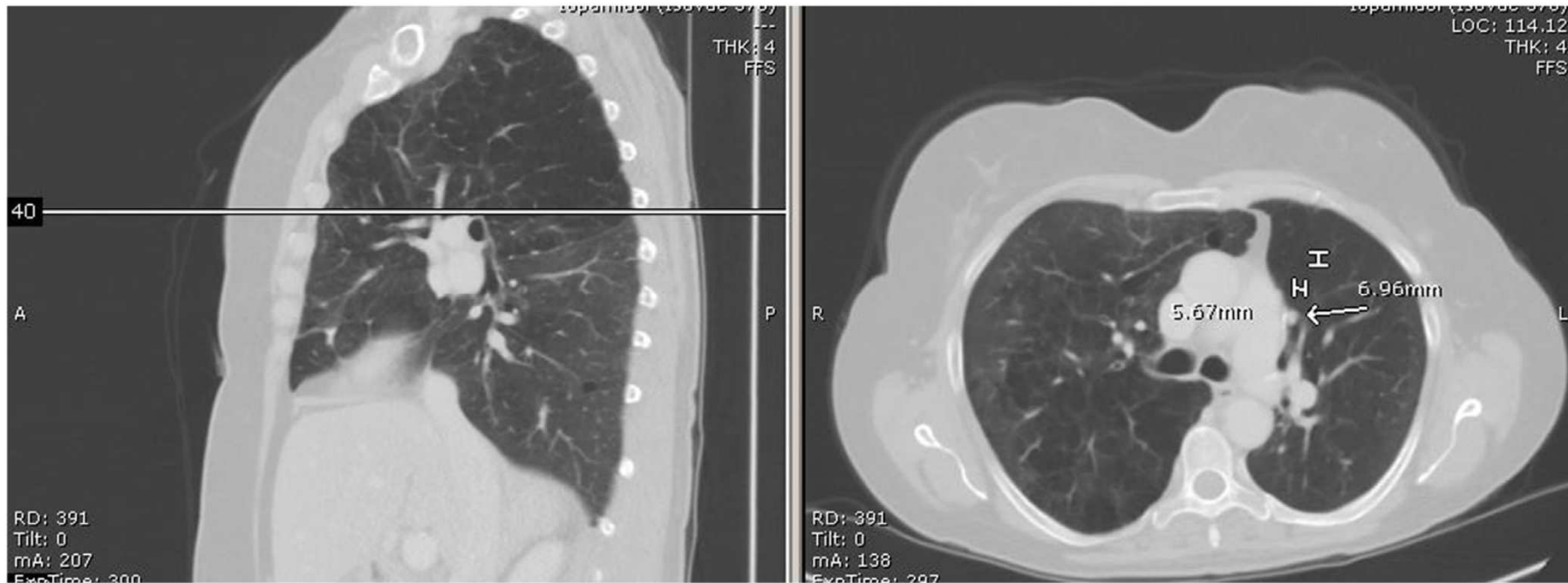
FIGURE 4: Gallium PET/CT scan (A, B); CT scan with contrast (C)

(A) Baseline gallium positron emission tomography/computed tomography (PET/CT) scan showing the large hypermetabolic sphenoid sinus tumor; (B) PET-CT scan three months after intensity-modulated radiation therapy (IMRT) showing a significant decrease in metabolic activity; (C) CT scan at the last follow-up three years after diagnosis showing stable disease.



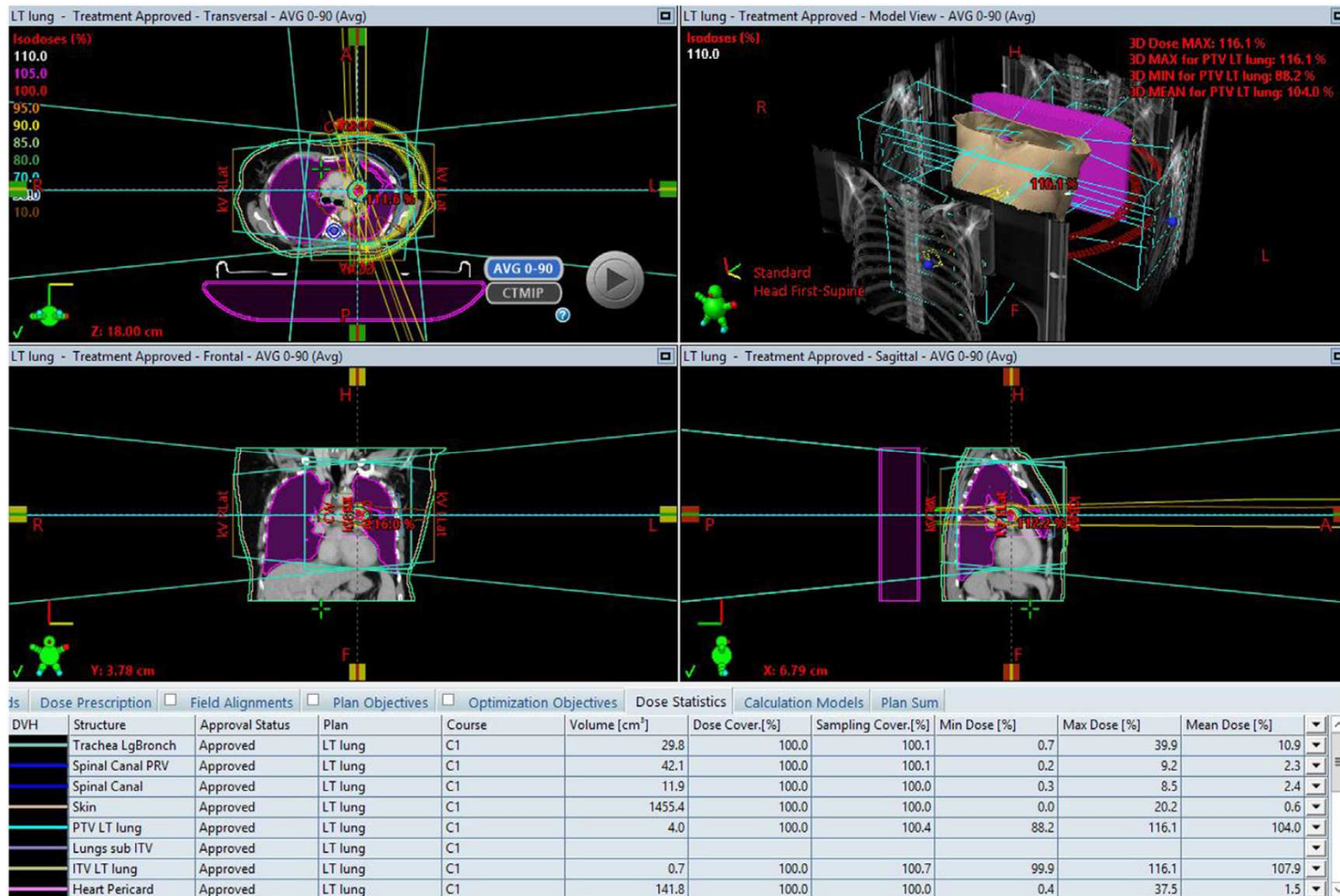
Background

**pulmonary large cell neuroendocrine carcinoma
treated with stereotactic body radiation therapy**





Background





ELSEVIER

Clinical Oncology

journal homepage: www.clinicaloncologyonline.net

Overview

Merkel Cell Carcinoma – Current Controversies and Future Directions

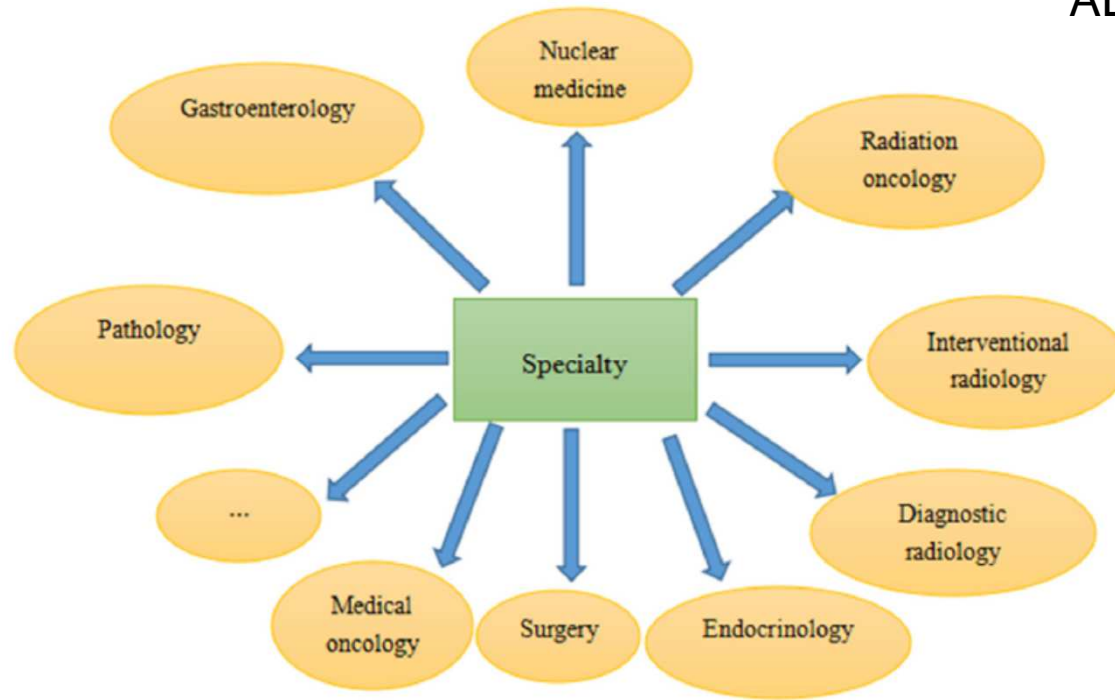
N. Steven^{*}, P. Lawton[†], M. Poulsen[‡]

Surgery and Radiotherapy for Nodal Disease

Chemotherapy for Locoregional Merkel Cell Carcinoma

Conclusions

The management of patients with MCC is built on long experience, not randomised trials. MCC requires speedy and accurate diagnosis and staging for the most suitable locoregional and systemic treatments, which are best carried out by a tertiary multidisciplinary team.



ADIUVANTE NEOADIUVANTE

CURATIVO

PALLIATIVO

EXPERTISE