



Brachytherapy - treatment of the future

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http://www.wco.pl/zb/?m=7





Derived from ancient Greek words for short distance (brachios) and treatment (therapy) and refers to the therapeutic use of encapsulated radionuclides within or close to a tumor.

It is sometimes called seed implantation and is an outpatient procedure used in the treatment of different kinds of cancer.

The efficacy of brachytherapy, as compared with the efficacy of external beam alone,

is attributable

to the ability of radioactive implants to deliver a higher concentrated radiation dose more precisely to tissues,

which contributes to improved local control, provided that the tissue is clinically delimitable and accessible.

At the same time, the surrounding healthy tissues are spared irradiation.

Kinds of brachytherapy

a. characterized by the duration of the irradiation:

there are 2 different kinds of brachytherapy: permanent, when the seeds remain inside of the body,

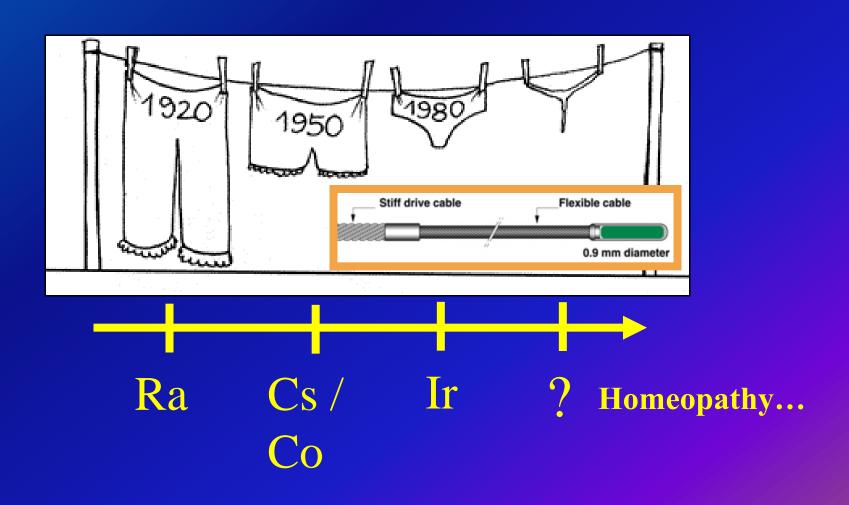
and temporary, when the seeds are inside of the body and then removed.

- b. characterized by the positioning of the radionuclides:
- interstitial brachytherapy: radioactive sources are inside the tumour,
 - contact brachytherapy or plesiobrachytherapy: radioactive sources are close to the tumour,
 - Contact brachytherapy is divided into four different kinds of brachytherapy:
 - intracavitary,
 - intraluminal,
 - endovascular,
 - surface brachytherapy

c. characterized by the dose rate (ICRU definitions):

- low dose rate (LDR) 0.4 2.0 Gy/h
- pulsed dose rate (PDR) 0.5 1.0 Gy/h
- medium dose rate (MDR) 2 12 Gy/h (history in EU,US)
 - high dose rate (HDR) > 12 Gy/h

Evolution in brachytherapy...



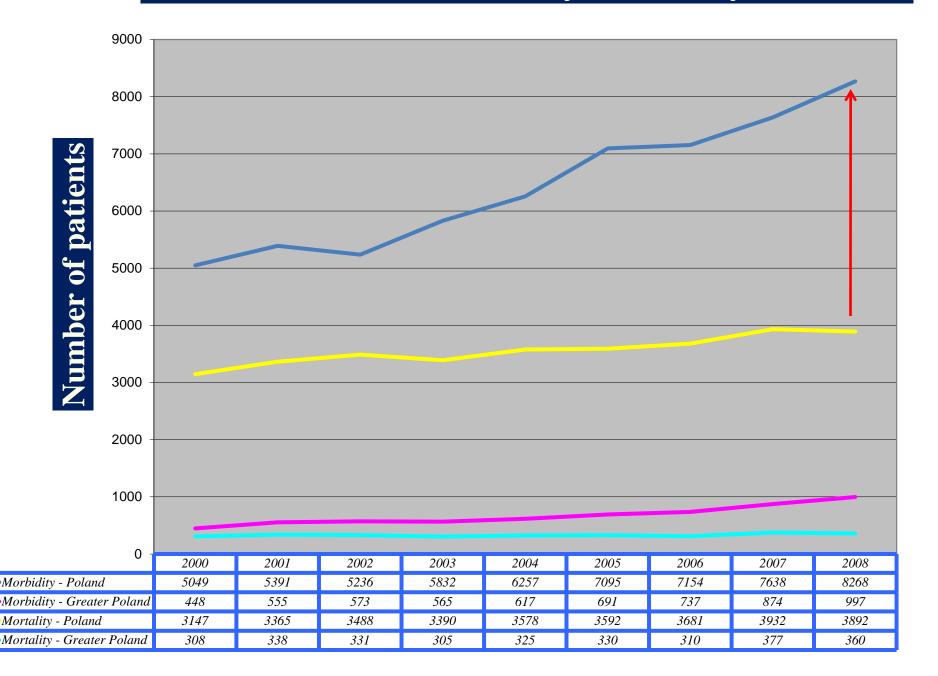
Brachytherapy curative

Prostate cancer as example

Greatest problems...



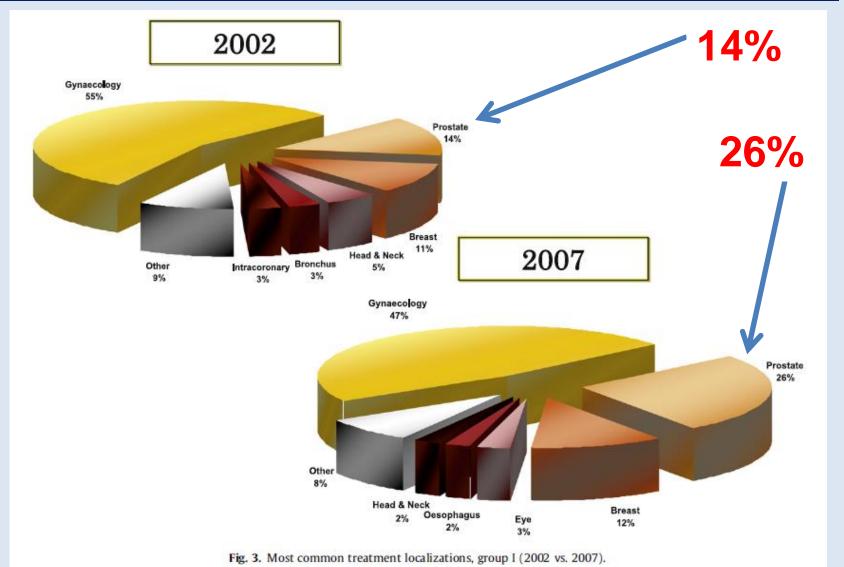
Prostate cancer - morbidity, mortality in Poland



Patterns of care for brachytherapy in Europe: Updated results

Ferran Guedea, Jack Venselaar, Peter Hoskin, Taran Paulsen Hellebust, Didier Peiffert, Bradley Londres, Montse Ventura, Jean-Jacques Mazeron, Erik Van Limbergen, Richard Pötter, Gyorgy Kovacs

Radiotherapy and Oncology 97 (2010) 514–520



> 20 YEARS AGO

Radical prostatectomy

or

External beam irradiation

10 - 20 YEARS AGO

Radical prostatectomy

or

Conventional external beam

or

Conformal external beam

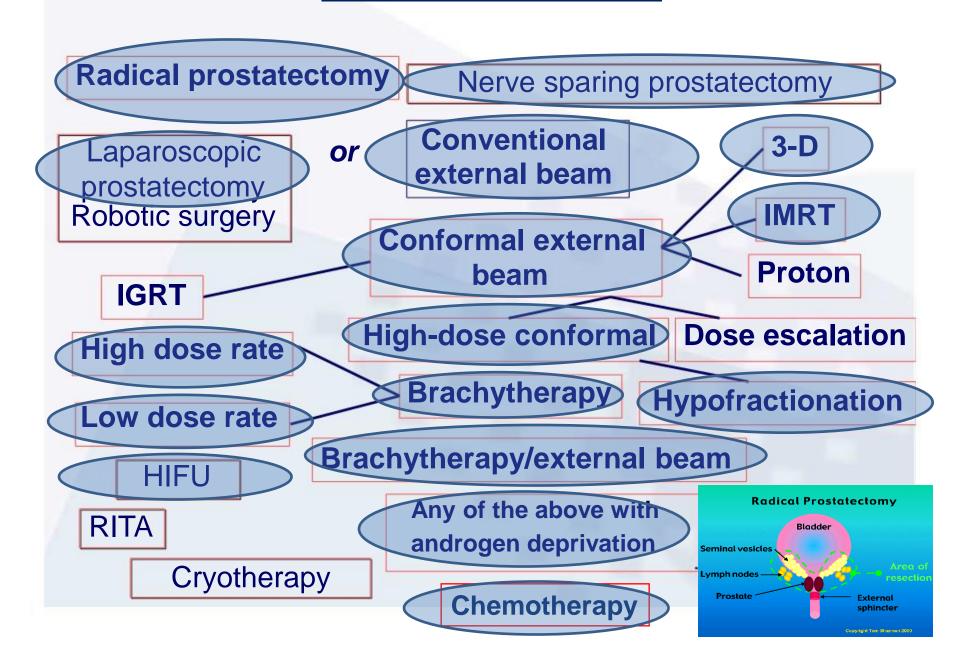
or

Brachytherapy

or

Brachytherapy/external beam

Today - Poland





Int. J. Radiation Oncology Biol. Phys., Vol. 38, No. 1, pp. 127–132, 1997 Copyright © 1997 Elsevier Science Inc. Printed in the USA, All rights reserved 0360-3016/97 \$17.00 ± .00

PII S0360-3016(97)00231-9

Clinical Investigation

AMERICAN BRACHYTHERAPY SOCIETY (ABS) CONSENSUS GUIDELINES FOR BRACHYTHERAPY OF ESOPHAGEAL CANCER

LAURIE E. GASPAR, M.D., * SUBIR NAG, M.D., † ARNOLD HERSKOVIC, M.D., †
RAO MANTRAVADI, M.D., * BURTON SPEISER, M.D. † AND THE CLINICAL
RESEARCH COMMITTEE, AMERICAN BRACHYTHERAPY SOCIETY, PHILADELPHIA, PA



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PII S0360-3016(00)01534-0

CLINICAL INVESTIGATION

Sarcoma

THE AMERICAN BRACHYTHERAPY SOCIETY RECOMMENDATIONS FOR BRACHYTHERAPY OF SOFT TISSUE SARCOMAS

Subir Nag, M.D.,* Daniel Shasha, M.D.,[†] Nora Janjan, M.D.,[‡] Ivy Petersen, M.D.,[§] and Marco Zaider, Ph.D., for the American Brachytherapy Society



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PII S0360-3016(99)00351-X

PHYSICS CONTRIBUTION

THE AMERICAN BRACHYTHERAPY SOCIETY RECOMMENDATIONS FOR PERMANENT PROSTATE BRACHYTHERAPY POSTIMPLANT DOSIMETRIC ANALYSIS

SUBIR NAG, M.D.,*[†] WILLIAM BICE, PH.D.,*[‡] KEITH DEWYNGAERT, PH.D.,*[§] BRADLEY PRESTIDGE, M.D.,*^{||} RICHARD STOCK, M.D.,*[¶] AND YAN YU, PH.D.,*[#]



BRACHYTHERAPY

Brachytherapy 6 (2007) 34-37

American Brachytherapy Society recommends no change for prostate permanent implant dose prescriptions using iodine-125 or palladium-103

Mark J. Rivard^{1,*}, Wayne M. Butler², Phillip M. Devlin³, John K. Hayes Jr.⁴, Robert A. Hearn⁵, Eugene P. Lief⁶, Ali S. Meigooni⁷, Gregory S. Merrick², Jeffrey F. Williamson⁸



Int. J. Radiation Oncology Biol. Phys., Vol. 50, No. 5, pp. 1190-1198, 2001

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0360-3016/01/5-see front matter

PII S0360-3016(01)01567-X

CLINICAL INVESTIGATION

Head and Neck

THE AMERICAN BRACHYTHERAPY SOCIETY RECOMMENDATIONS FOR HIGH-DOSE-RATE BRACHYTHERAPY FOR HEAD-AND-NECK CARCINOMA

Subir Nag, M.D.,* Elmer R. Cano, M.D., † D. Jeffrey Demanes, M.D., † Ajmel A. Puthawala, M.D., § and Bhadrasain Vikram, M.D., $^{\|}$ for the American Brachytherapy Society



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PII S0360-3016(99)00069-3

CLINICAL INVESTIGATION

Prostate

AMERICAN BRACHYTHERAPY SOCIETY (ABS) RECOMMENDATIONS FOR TRANSPERINEAL PERMANENT BRACHYTHERAPY OF PROSTATE CANCER

Subir Nag, M.D.,*[†] David Beyer, M.D.,*[‡] Jay Friedland, M.D.,*[§] Peter Grimm, D.O.,*^{||} and Ravinder Nath. Ph.D.*[¶]



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PII S0360-3016(00)00555-1

RAPID COMMUNICATION

AMERICAN BRACHYTHERAPY SOCIETY RECOMMENDATIONS FOR CLINICAL IMPLEMENTATION OF NIST-1999 STANDARDS FOR 103PALLADIUM BRACHYTHERAPY

DAVID BEYER, M.D.,* RAVINDER NATH, Ph.D.,[†] WAYNE BUTLER, Ph.D.,[‡] Gregory Merrick, M.D.,[‡] John Blasko, M.D.,[§] Subir Nag, M.D.,[‡] and Colin Orton, Ph.D.,[†] For the American Brachytherapy Society, Oakbrook, IL



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PII S0360-3016(00)00497-1

CLINICAL INVESTIGATION

Cervix

THE AMERICAN BRACHYTHERAPY SOCIETY RECOMMENDATIONS FOR HIGH-DOSE-RATE BRACHYTHERAPY FOR CARCINOMA OF THE CERVIX

Subir Nag, M.D.,* Beth Erickson, M.D.,[†] Bruce Thomadsen, M.D.,[‡] Colin Orton, Ph.D.,[§]
Jeffrey D. Demanes, M.D.,^{||} and Daniel Petereit, M.D.,[‡] for the American Brachytherapy Society

Int. J. Radiation Oncology Biol. Phys., Vol. 48, No. 3, pp. 779-790, 2000 Copyright © 2000 Elsevier Science Inc. Printed in the USA. All rights reserved 0360-3016/00/\$-see front matter

PII S0360-3016(00)00689-1

CLINICAL INVESTIGATION

Endometrium

THE AMERICAN BRACHYTHERAPY SOCIETY RECOMMENDATIONS FOR HIGH-DOSE-RATE BRACHYTHERAPY FOR CARCINOMA OF THE **ENDOMETRIUM**

Subir Nag, M.D.,* B NILENDU GUPTA, PI



Contents lists available at ScienceDirect

Radiotherapy and Oncology 91 (2009) 150-156

Radiotherapy and Oncology

journal homepage: www.thegreenjournal.com



GEC-ESTRO recommendations

GEC-ESTRO recommendations for brachytherapy for head and neck squamous cell carcinomas

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Int. J. Radiation Oncology Biol. Phys., Vol. 76, No. 4, pp. 963-972, 2010 Copyright © 2010 Elsevier Inc. Printed in the USA. All rights reserved 0360-3016/10/\$-see front matter



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Vol. 74, No. 4, pp. 987–1001, 2009 ncology. Published by Elsevier Inc. Printed in the USA. 0360-3016/09/\$-see front matter

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HOLZ, M.D.,

GH, M.D.,



ESTRO/EAU/EORTC

REPORT

Daniel Ash^{a,*}, Ant

AMERICAN SOCIETY FOR THERAPEUTIC RADIOLOGY AND ONCOLOGY (ASTRO) EMERGING TECHNOLOGY COMMITTEE REPORT ON ELECTRONIC BRACHYTHERAPY

doi:10.1016/j.ijrobp.2009.10.068

ASTRO GUIDELINE

AMERICAN SOCIETY COLLEGE OF RADIOLOG OF I CATHERINE C. PARK, M.D., * SUE S. YOM, M.D., Ph.D., * MATTHEW B. PODGORSAK, Ph.D., † ELEANOR HARRIS, M.D., ROBERT A. PRICE, JR., Ph.D., ALISON BEVAN, M.D., Ph.D., ELEANOR HARRIS, M.D., ALISON BEVAN, M.D., Ph.D., JEAN POULIOT, Ph.D.,* ANDRE A. KONSKI, M.D., M.B.A.,§ AND PAUL E. WALLNER, D.O. ELECTRONIC BRACHYTHERAPY WORKING GROUP.

BETH A. ERICKSON, M.

JOHN K. HAYES, M.D., M.S., I-CHOW J. HSU, M.D., DAVID E. MORRIS, M.D., RACHEL A. RABINOVITCH, M.D., # JONATHAN D. TWARD, M.D., Ph.D., ** AND SETH A. ROSENTHAL, M.D. ††

R. Рн.D.,[‡] TRANK A. VICINI, IVI.D., TIMOTHY J. WHELAN, IVI.D., JULIA WHITE, IVI.D., JENNIFER Y. WO, M.D., AND JAY R. HARRIS, M.D. ¶¶

Practical Radiation Oncology (2011) 1, 60-71

Int. J. Radiation Oncology Biol. Phys., Vol. 79, No. 2, pp. 335-341, 2011

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ASTRO GUIDELINE

AMERICAN SOCIETY FOR RADIATION ONCOLOGY (ASTRO) AND AMERICAN COLLEGE OF RADIOLOGY (ACR) PRACTICE GUIDELINE FOR THE TRANSPERINEAL PERMANENT BRACHYTHERAPY OF PROSTATE CANCER

Seth A. Rosenthal, M.D.,* Nathan H. J. Bittner, M.D., M.S.,† David C. Beyer, M.D.,‡ D. Jeffrey Demanes, M.D., BRIAN J. GOLDSMITH, M.D., ERIC M. HORWITZ, M.D., Geoffrey S. Ibbott, Ph.D., W. Robert Lee, M.D., Subir Nag, M.D., W. Warren Suh, M.D., AND LOUIS POTTERS, M.D.^{‡‡}

Special Article

Palliative thoracic radiotherapy in lung cancer: An American Society for Radiation Oncology evidence-based clinical practice quideline

George Rodrigues MD, MSca,*, Gregory M.M. Videtic MD, CM, FRCPCb, Ranjan Sur MD, PhDc, Andrea Bezjak MD, FRCPCd, Jeffrey Bradley MDe, Carol A. Hahn MD^f, Corey Langer MD^g, Keith L. Miller MD^h, Benjamin J. Moeller MD, PhD¹, Kenneth Rosenzweig MD¹, Benjamin Movsas MD^k





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ESTRO/EAU/EORTC recommendations on permanent seed implantation for localized prostate cancer

Daniel Ash^{a,*}, Anthony Flynn^a, Jan Battermann^b, Theodorous de Reijke^c, Paulo Lavagnini^d, Leo Blank^e

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Received 18 September 2000; accepted 27 September 2000

	Recommended Do well	Optional Fair	Investigational Do poorly			
PSA (ng/ml)	< 10	10-20	>20			
Gleason score	5-6	7	8-10			
Stage	T1c-T2a	T2b-T2c	Т3			
IPSS	0-8	9-19	>20			
Prostate volume (g)	<40	40-60	>60			
Q _{max} ml/s	>15	15-10	<10			
Residual volume cm ³			>200			
TURP <u>+</u>			+			

National Comprehensive Cancer Network 1.2009 www.nccn.org Risk categories (N0 M0) – Clinically localized

	Low	Intermediate	High	Very high, locally advanced
Т	1-2a	2b, 2c	3a	3b, 4
PSA	< 10	10-20	>20	Any
Gleason	< 7	7	>7	any

Metastatic: any T, N1 M0 any T, any N, M1

Contraindications

ESTRO/EAU/EORTC RECOMMENDATIONS ON PERMANENT SEEDIMPLANTATION FOR LOCALISED PROSTATE CANCER

- expected survival time < 5 years,
 - 2. metastasis,
- 3. TURP in the past with considerable damage to the prostate,
 - persistent hematuria, the regular use of aspirin or anticoagulants,
 - 5. prostate volume > 60 cm cc.

Temporary - HDR





Radiotherapy and Oncology 74 (2005) 137-148

www.elsevier.com/locate/radonline

GEC/ESTRO-EAU recommendations on temporary brachytherapy using stepping sources for localised prostate cancer

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^bUniversity Clinic for Radiotherapy and Radiobiology, Vienna General Hospital, Vienna, Austria

^cDepartment of Urology, University of the Saarland, Homburg, Germany

^dDepartment of Radiotherapy, Barmherzige Schwestern Hospital and St Vincenc Clinic, Linz, Austria

^eDivision of Clinical Physics, Department of Radiation Oncology, Erasmus MC, Daniel Den Hoed Cancer Center, Rotterdam, The Netherlands

^fDepartment of Urology, AMC, Amsterdam, The Netherlands

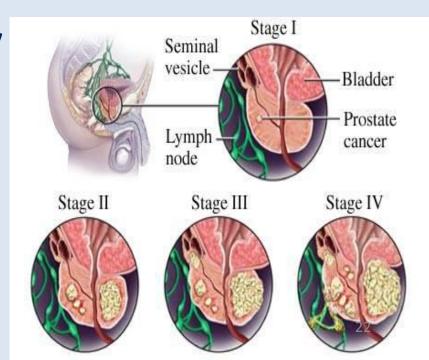
^gDepartment of Urology, City Hospital, Kiel, Germany

Received 3 August 2004; accepted 2 September 2004 Available online 22 October 2004

Monotherapy (ABS, GEC-ESTRO)

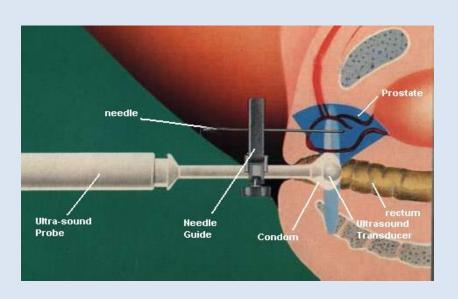
- 1. T1 i T2,
- 2. NO,
- 3. M0,
- **4.** PSA <10, Gleason ≤6,
- 5. > 5 years life expectancy





HDR brachytherapy: diagnostic, equipment, team

similar to seeds



Team



Experience in:

- TRUS (done by radiotherapist),
- dosimetry, treatment planning (physicist and radiotherapist),
- needles (seeds) implantation (radiotherapist and physicist),
- radiotherapy knowledge.

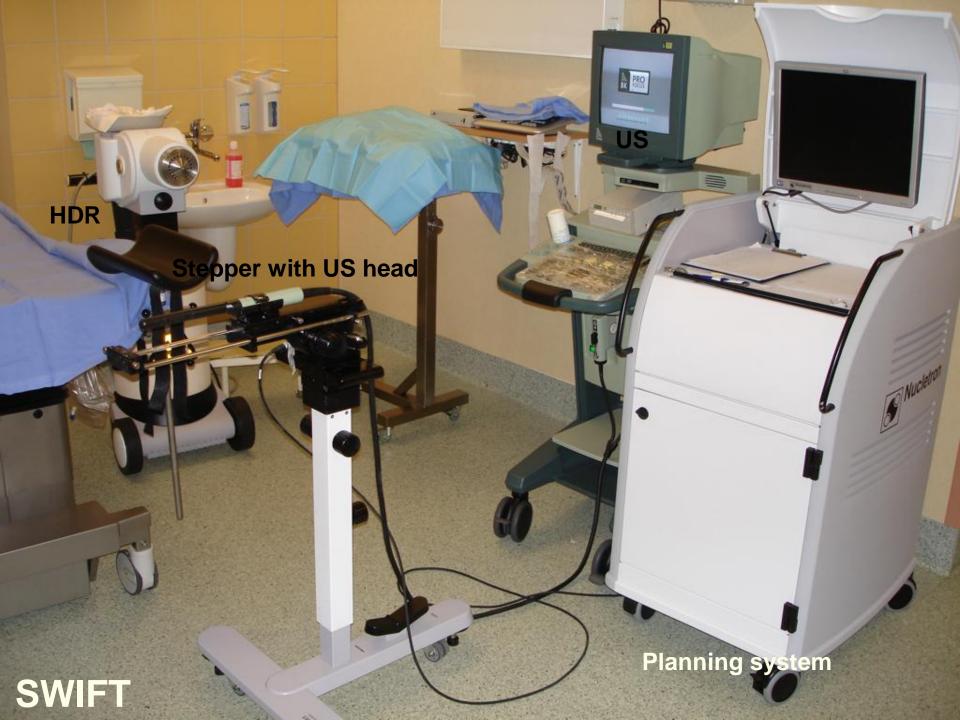
Team:

- radiotherapist,
- urologist, radiologist or radiotherapist with ultrasound skills,
- physicist,
- 2-3 nurses,
- anesthetist,
- nurse anesthetic,
- X-ray technician.

Brachytherapy - equipment

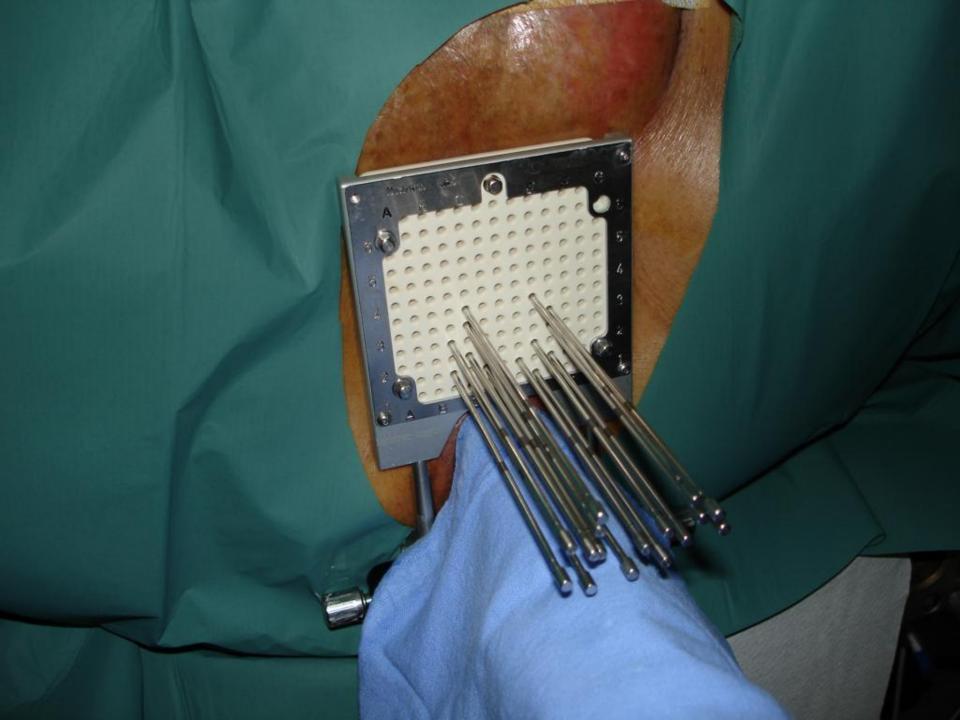
High quality - image guided source placement - SWIFT

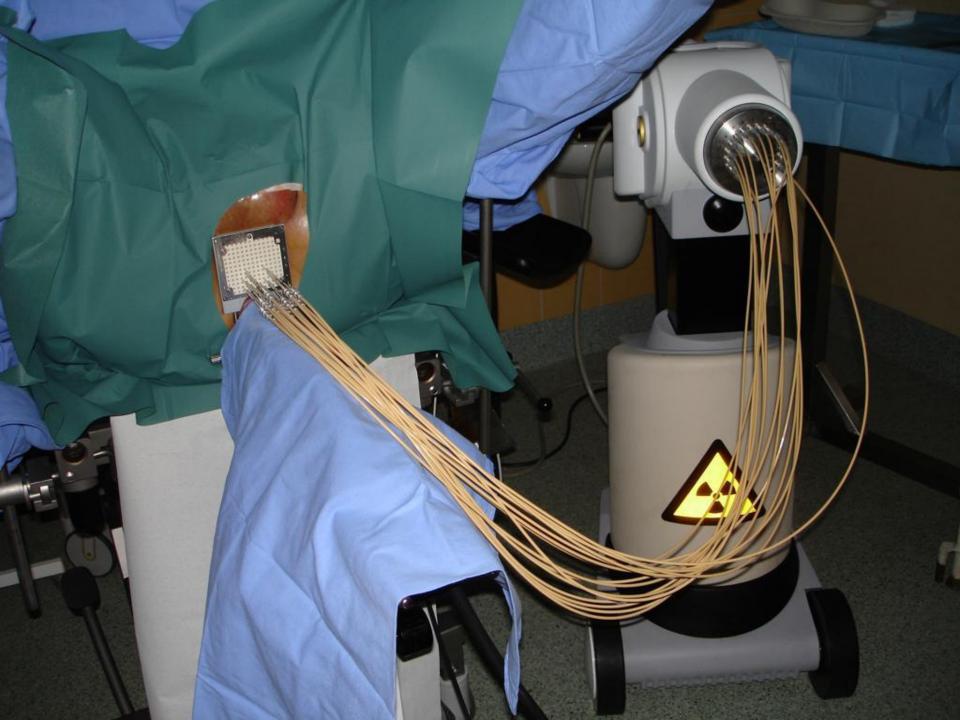
- High quality digital TRUS with template software,
- X-ray/CT for post-implant dosimetry,
- Stepper, stepping unit,
- Treatment planning system.











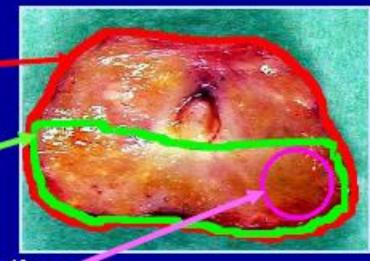


Dose Prescription: IMBRT

Different Target and Treatment Philosophies:

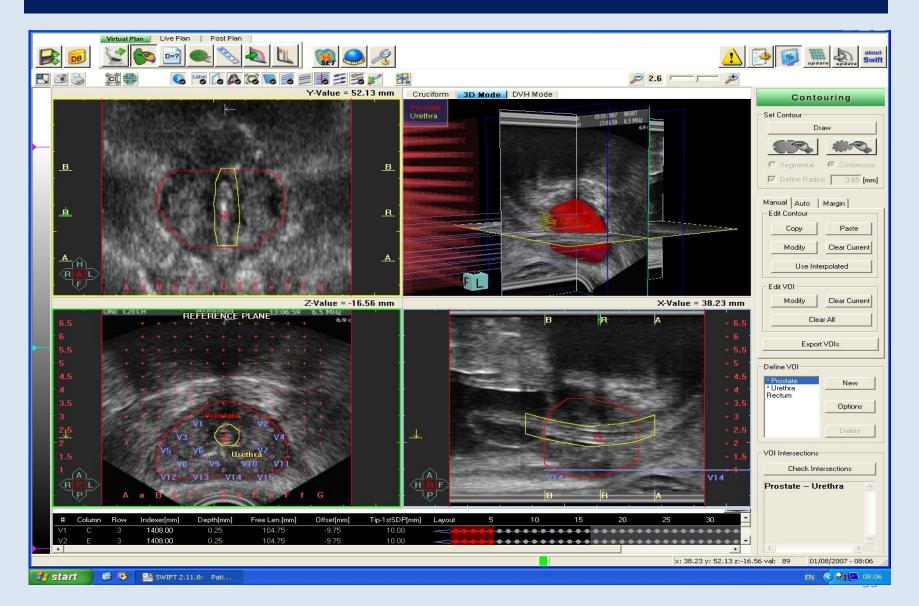
CTV 1 → Prostate Capsule

CTV 2 → Peripheral Zone

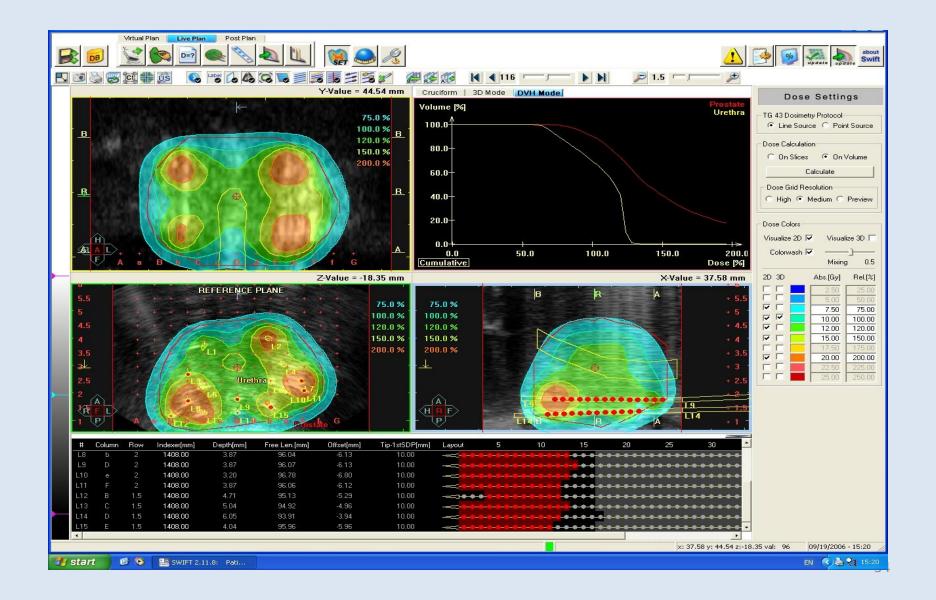


CTV 3 → Visible Tumor Infiltration

"Virtual planning"



"Real-time planning"



Conclusions

Remote temporary afterloading brachytherapy has several advantages:

- accurate positioning of the source by first implanting non-active guide needles,
- possibility to choose the source positions over the length of the needle,
 - no target movement during radiation,
- stepping source technology allowing for dose and volume adaptation due to adjustment of source dwell locations and times according to 3D imaging based individual dose prescription before irradiation,
 - shortening the time and costs.



PII S0360-3016(99)00069-3

CLINICAL INVESTIGATION

Prostate

AMERICAN BRACHYTHERAPY SOCIETY (ABS) RECOMMENDATIONS FOR TRANSPERINEAL PERMANENT BRACHYTHERAPY OF PROSTATE CANCER

Subir Nag, M.D.,* † David Beyer, M.D.,* ‡ Jay Friedland, M.D.,* $^{\$}$ Peter Grimm, D.O.,* $^{\parallel}$ and Ravinder Nath, Ph.D.* †

*Prostate Brachytherapy Quality Assurance Group, Clinical Research Committee, American Brachytherapy Society, Reston, VA;

†The Ohio State University, Columbus, OH; ‡Arizona Oncology Services, Phoenix, AZ; 5Moffitt Cancer Center, Tampa, FL;

|Swedish Medical Center, Seattle, WA; and †Yale University, New Haven, CT

Brachytherapy as a Boost to EBRT:

- T2b, T2c or
- Gleason 8-10 or
- PSA > 20 ng/ml

Other possible indications for Brachytherapy as a Boost to EBRT:

- Perineural invasion,
- Multiple positive biopsies, bilateral positive biopsies,
- MRI positive for capsular penetration.

Brachytherapy (including Boosting EBRT) in Conjunction with Androgen Deprivation:

Patients with initially large prostate (>60 cc) that have downsized sufficiently

Brachytherapy

Permanent



Int. J. Radiation Oncology Biol. Phys., Vol. 44, No. 4, pp. 789-799, 1999

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0360-3016/99/\$-see front matter

PII S0360-3016(99)00069-3

CLINICAL INVESTIGATION

Prostate

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†The Ohio State University, Columbus, OH; [‡]Arizona Oncology Services, Phoenix, AZ; [§]Moffitt Cancer Center, Tampa, FL;

|Swedish Medical Center, Seattle, WA; and [¶]Yale University, New Haven, CT





Radiotherapy and Oncology 57 (2000) 315–321

www.elsevier.com/locate/radonline

ESTRO/EAU/EORTC recommendations on permanent seed implantation for localized prostate cancer

Daniel Ash^{a,*}, Anthony Flynn^a, Jan Battermann^b, Theodorous de Reijke^c, Paulo Lavagnini^d, Leo Blank^e

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^bDepartment of Radiotherapy, Academisch Ziekenhuis, Utrecht, Germany

^cDepartment of Urology, Academisch Medisch Centrum, Amsterdam, The Netherlands

^dInstituto Tumori, Genoa, Italy

^eDepartment of Radiotherapy, Free University, Amsterdam, The Netherlands

Received 18 September 2000; accepted 27 September 2000





Seeds - producers

Greatest:

1.McMaster University's Reactor (Canada),

2.National Research
Universal (NRU) Reactor
(Canada)

- 1. IsoRay Medical™, Cesium-131
- 2. Best Medical International, Inc.

Best Industries Iridium (Ir 192)

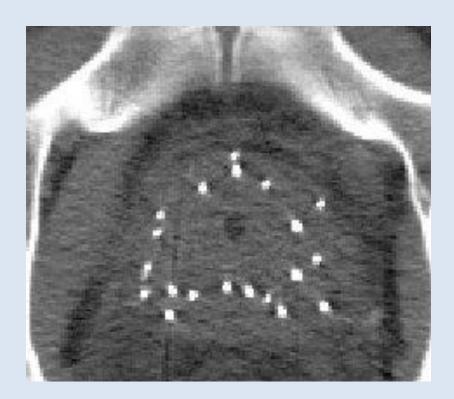
- 3. Theragenics TheraSeeds Iodine-125, Palladium-103
- 4. Amersham Healthcare Iodine (I-125)

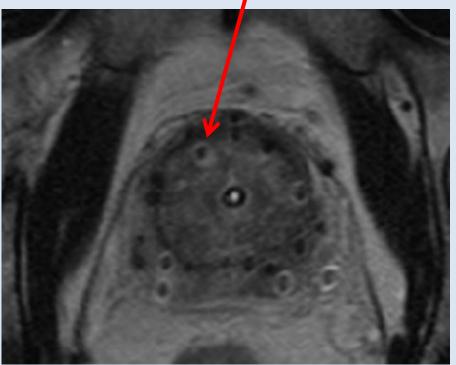
- 1. IBt Bebig
- 2. Nucletron
- 3. Varian VariSeed
- 4. Oncura RAPID Strand
 - 5. BARD ProSeed
 - 6. IsoAid

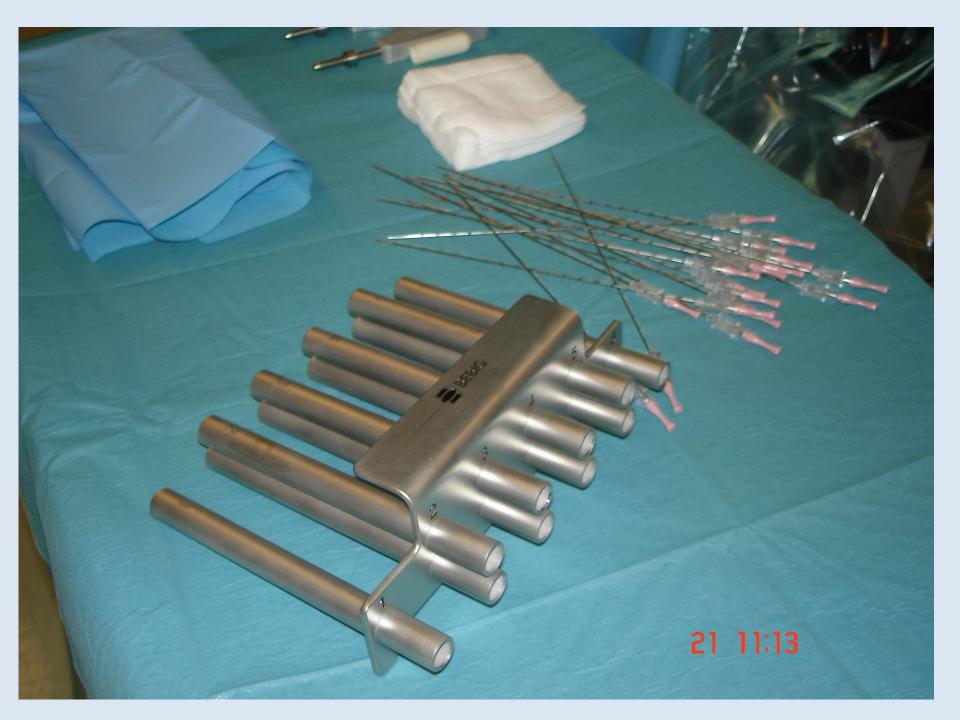


In CT imaging the high contrasts leads to absorption artefacts which can be handled using a soft tissue filter for image reconstruction on the CT device.

125.S17 seeds also provide a unique quality in MRI imaging. While normal seeds only appear as dark dots in the image, similar to a blood vessel, S17 seeds show a bright halo around the dark dot, this allows you to clearly identify the seeds.





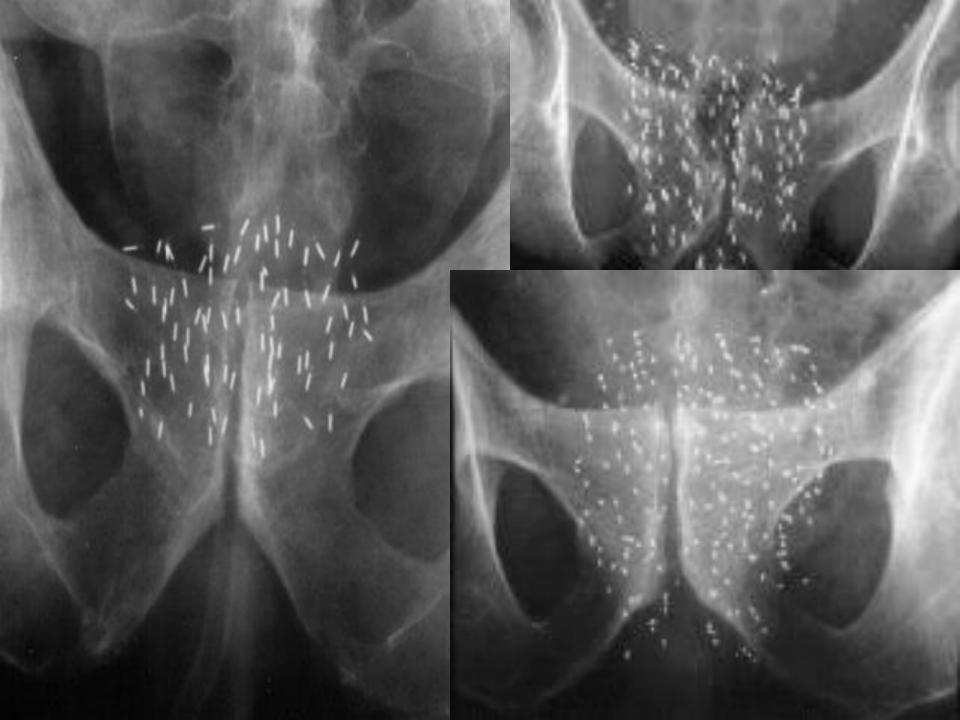


Doses:

I-125 monotherapy 140-160 Gy (144-145 Gy)
I-125 + 40-50 Gy EBRT 100-120 Gy

Pd-103 monotherapy 110-120 Gy (125 Gy) Pd-103 + 50 Gy EBRT 60-90 Gy

Cs-131 monotherapy 115 Gy





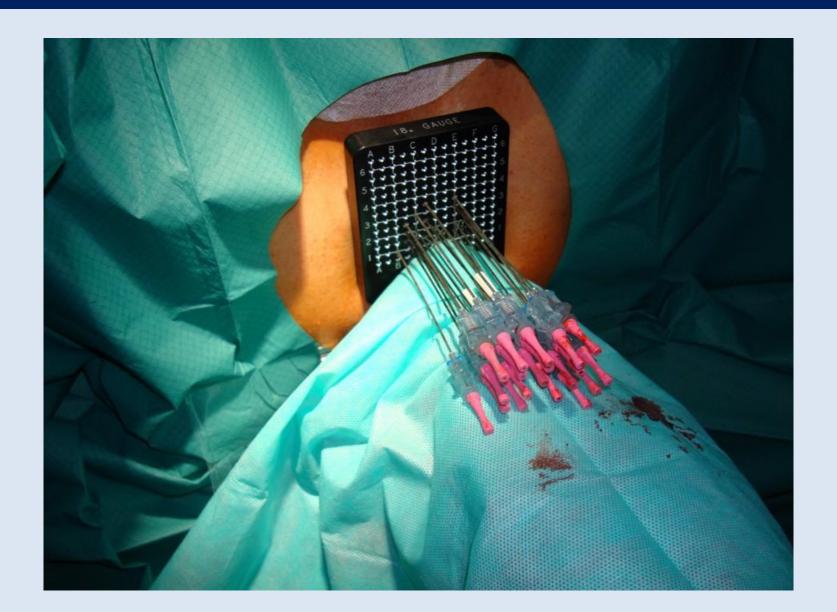
Post implant dosimetry

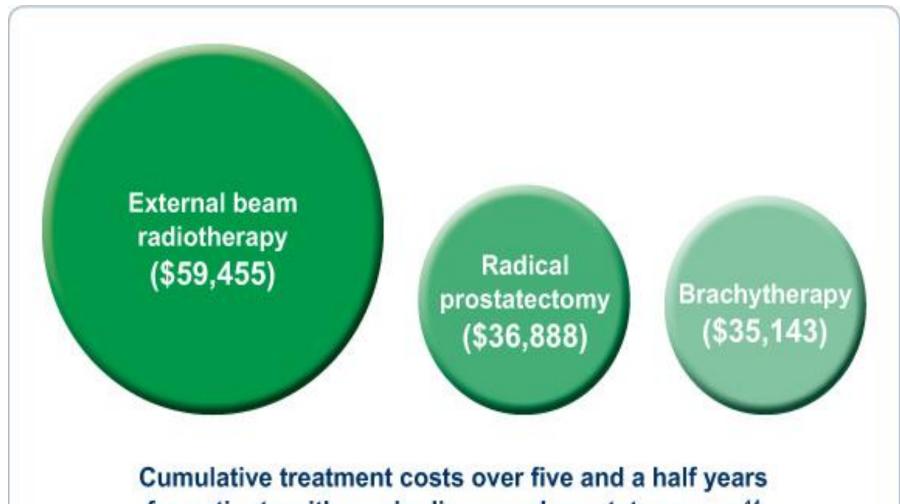
It is usual to perform the CT scan 4±6 weeks after implantation when oedema has settled.

It is recommended that the following indices are recorded for all patients:

- 1. The volume implanted.
- 2. The number of seeds.
- 3. The number of needles used.
- 4. The total activity implanted.
- 5. The prescribed dose.
- 6. The D90, that is the dose that covers 90% of the prostate volume as defined from post implant imaging.
- 7. The V100, that is the percentage of the prostate volume that has received the prescribed dose.
- 8. V150, the volume that has received 50% more than the prescribed dose.

Utrecht technique – used in 2008-2009





for patients with newly-diagnosed prostate cancer.14

Seeds

Sole LDR brachytherapy:

Advantages

- good treatment results (similar to surgery) 1,2
 - relatively small rate of complications
 - short treatment time (1-3 days)

Disadvantage

- in the past seed migration possibility
 - small risk of relatives irradiation
 - costs

¹⁾ Sylvester, J. E., P. D. Grimm, et al. (2010). "Fifteen-Year Biochemical Relapse-Free Survival, Cause-Specific Survival, and Overall Survival following I(125) Prostate Brachytherapy in Clinically Localized Prostate Cancer: Seattle Experience." Int J Radiat Oncol Biol Phys

²⁾ Ragde, H., L. J. Korb, et al. (2000). "Modern prostate brachytherapy. Prostate specific antigen results in 219 patients with up to 12 years of observed follow-up." Cancer 89(1): 135-41.

HDR

Advantages

- good treatment results (similar to seeds) 1,2
 - possibility of dose verifacion
 - complications similar to ¹⁾
 - positive radiobiology
 - no staff exposure to radiation

Disadvantage

- different fractionation schemas
- in monotherapy small trial's number

¹⁾ Demanes, D. J., A. A. Martinez, et al. (2011). "High-Dose-Rate Monotherapy: Safe and Effective Brachytherapy for Patients with Localized Prostate Cancer." Int J Radiat Oncol Biol Phys.

²⁾ Mark, R. J., P. J. Anderson, et al. (2010). "Interstitial High-Dose-Rate Brachytherapy as Monotherapy for Early Stage Prostate Cancer: Median 8-Year Results in 301 Patients "Brachytherapy 9(Supplement 1): S76.

Brachytherapy palliative

Lung cancer as example

Radical treatment (<5 -10% of BT patients): Indications

- 1. Curative intent as a "boost" to EBRT T1-2 No-1 Mo
 - **1** LC
 - before EBRT- remission of atelectasis, reclassification.
- 2. <u>Alone definitive brachytherapy for small tumors</u> T1-2 No Mo
 - in patients with occult carcinoma or tumors potentially resectable, with diameter <
 2 cm, disqualified for surgery or EBRT (Japan, USA).
- 3. <u>Postoperative brachytherapy of the bronchial stump</u> <u>after resection with positive resection margins (R2).</u>
- 4. As a boost for minor residual disease within a combined non-surgical radical approach.

Palliative treatment (>90 % of BT patients): Indications

1. The main indication is treatment of lifethreatening complications such a dyspnea,

obstructive pneumonia or atelectasis, cough or haemoptisis resulting from endobronchial or endotracheal tumour growth.

2. Treatment of endobronchial or endotracheal recurrent tumour growth in previously irradiated areas or in combination with EBRT for metastatic lung cancers.

Brachytherapy treatment schemas - indications, doses

Indications fo brachytherap Radical combine treatment: schema I;	Poznań I phase otal dose 4	4.6	II phase	III phase	IV phase
	otal dose 4	40.			
treatment: schema I:			1 fr. x 6 Gy, ref. point	EBRT 16 Gy in 8	1 fr. x 6 Gy,
,	11. aa 2 Gy (2 a-p fields)		0.5 - 1 cm	fr. (changed fields)	ref. point 0.5 - 1
clinical stage T1-3 N1-3 M0					cm
Radical combined	EBRT: total dose 4	4 Gy in	EBRT 16 Gy in 8 fr.	HDR-BT - in 1, 3	
treatment: schema II;	22 fr. aa 2 Gy (2 a-p fields)		(changed fields)	and 5 weeks of	
clinical stage T1-3 N1-3				$EBRT - 3 \times 10$	
M 0				Gy.	
Radical sole	Total dose 36 - 42 Gy in 6 - 7 fr. with interval of 4 - 7 days between fractions				
<u>reatment</u> , radiologically					
occult cancer T1-2N0					
Radical treatment	After EBRT with total dose of 50 - 60 Gy		To consider increasing		
after surgery, R2			the total use using HDR- BT HDR. Fr. dose from 1		
			x 6 Gy till 3 fr. x 6 Gy (18		
			Gy), depending on EBRT		
			dose		
Radical treatment:	Sole brachytherapy				
stump infiltration	75 10 Cy with interval of 4 - 7				
Cy in 3 fr. of 6 Cy with interval of 4.7 days, in notionic treated earlier with ERDI					
<u>Palliative</u>	Poznań dose > 50 Gy				
treatment	se 22,5 Gy in 3 fr. of 7,5 Gy Gy with interval of 4 -7 days – in patients not irradiated or				
	treated earlier with EBRT – dose < 50 Gy				
	1 x 10 Gy in case of Sometimes dose can be repeated after few weeks, in cases with clinical				ses with clinical
<u>Palliative</u> <u>treatment</u>	se 22,5 Gy in 3 fr. of 7,5 Gy Gy with interval of 4 -7 days – in patients not irradiated or treated earlier with EBRT – dose < 50 Gy				

WHO scale > 2

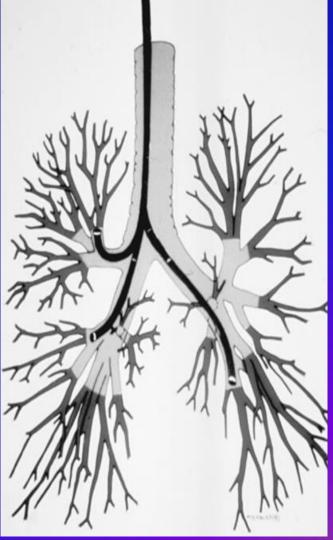
remission or visible during bronchoscopy

Contraindications:

- 1. peripheral location of the tumor,
 - 2. Pancoast tumor (?),
- 3. pressure location outside of bronchii,
- 4. contraindication to bronchoscopy (relative).

Bronchofiberoscope, endobronchial catheter French 5-6, applicator adapter





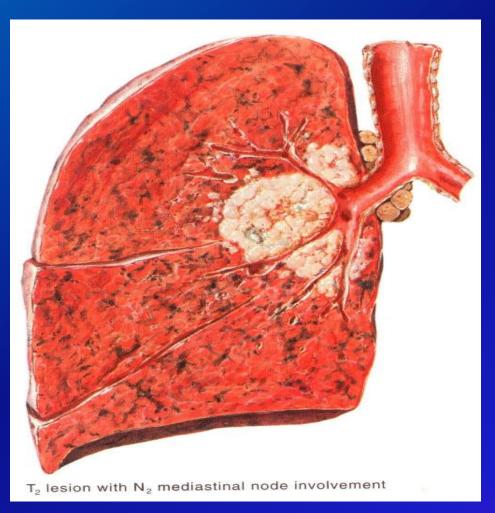


Lumencath Applicator Set

Fritz Adjustable Intralumenal Applicator Set

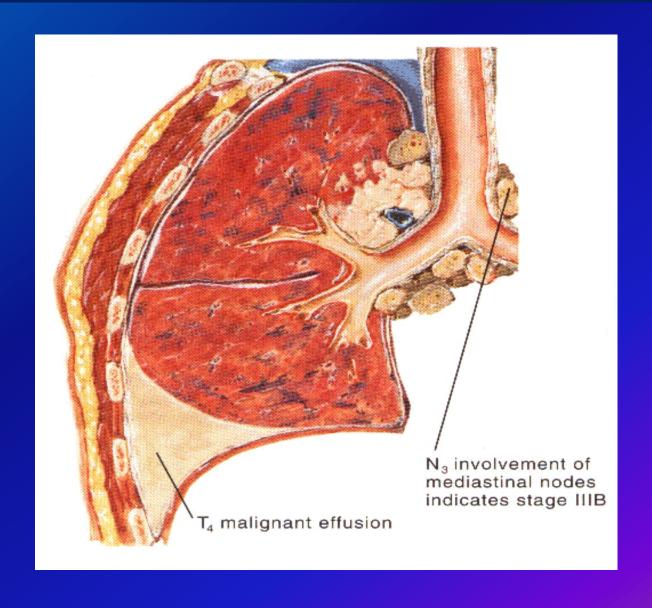


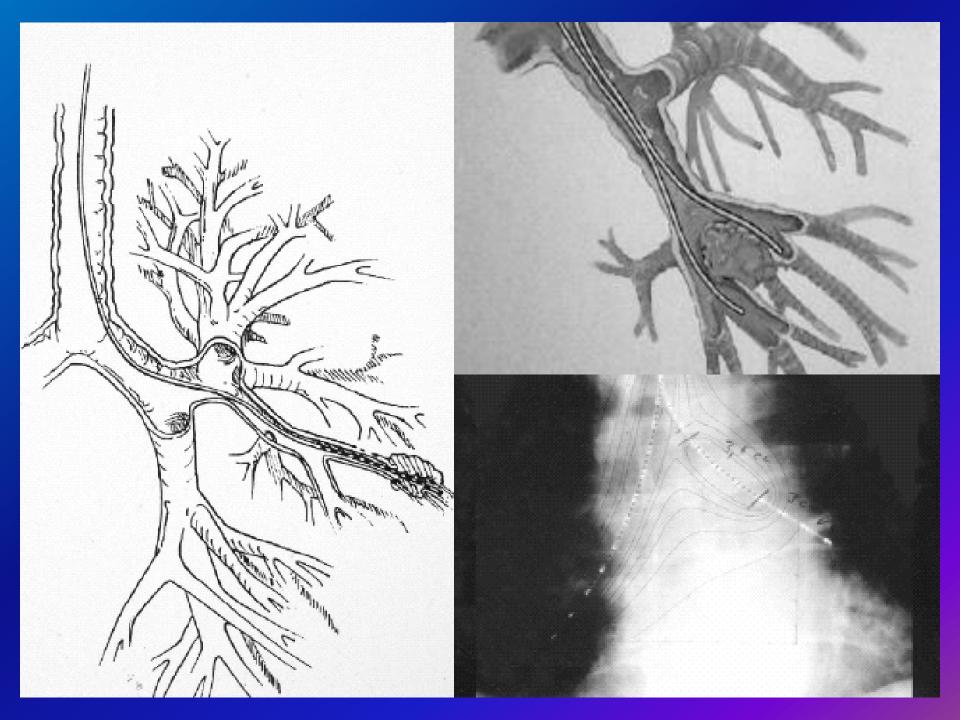
Location of the tumor for brachytherapy





Lack of indications for brachytherapy





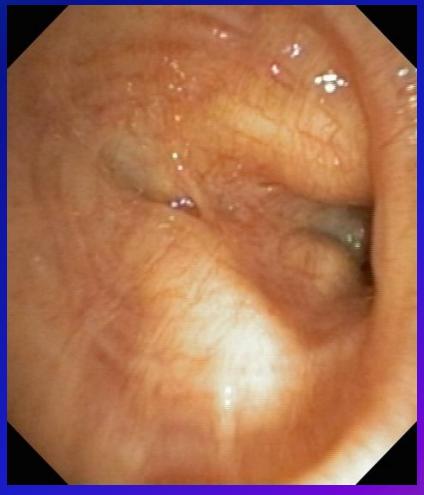
Stump



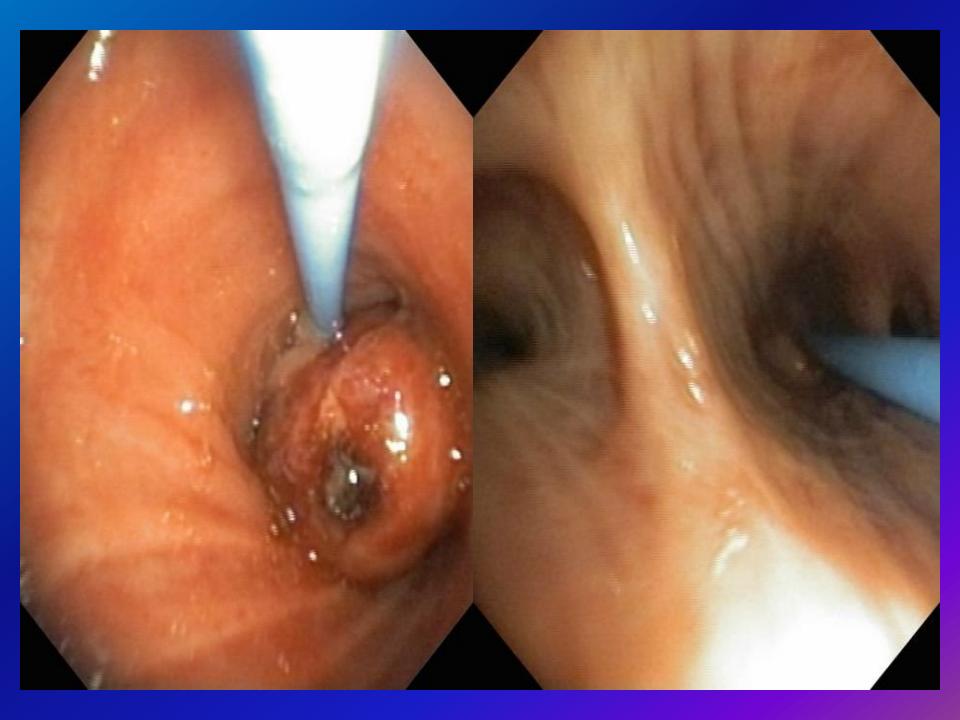


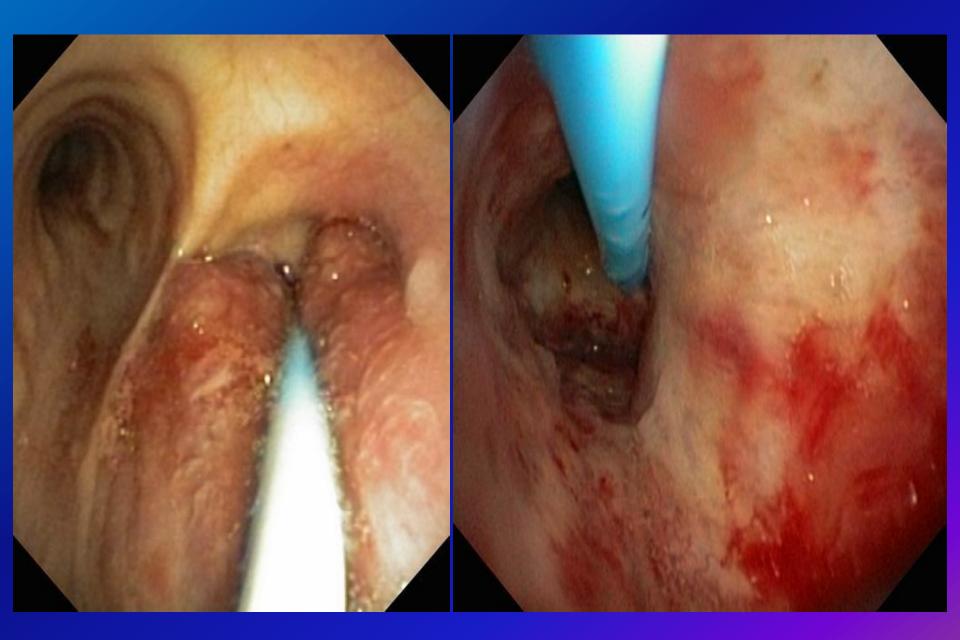
Stump











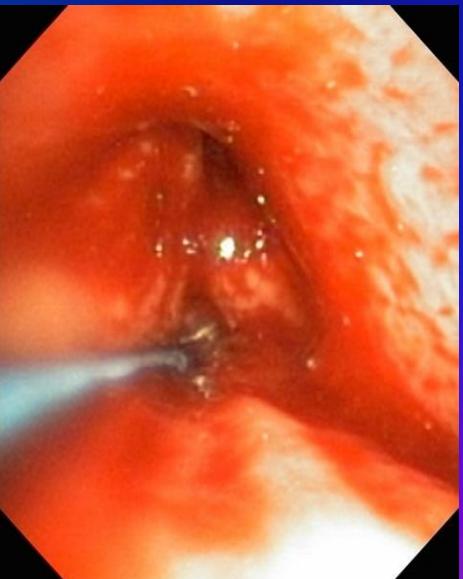


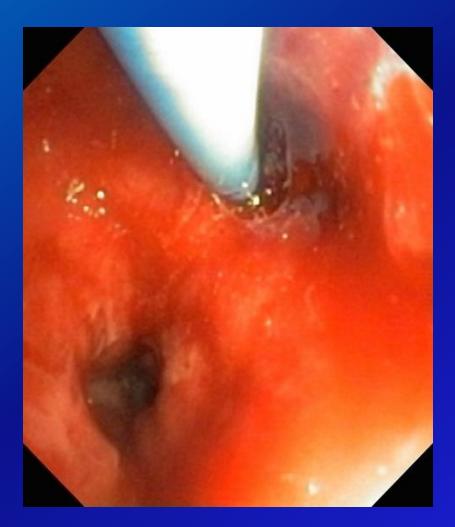
















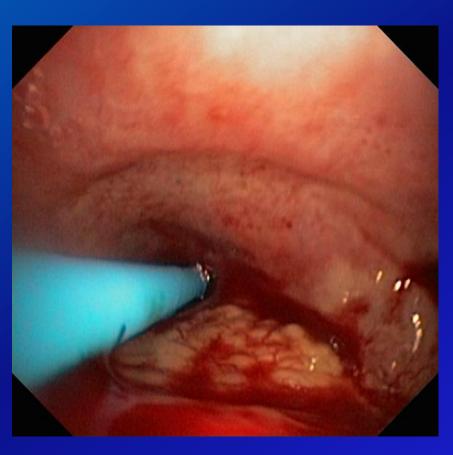




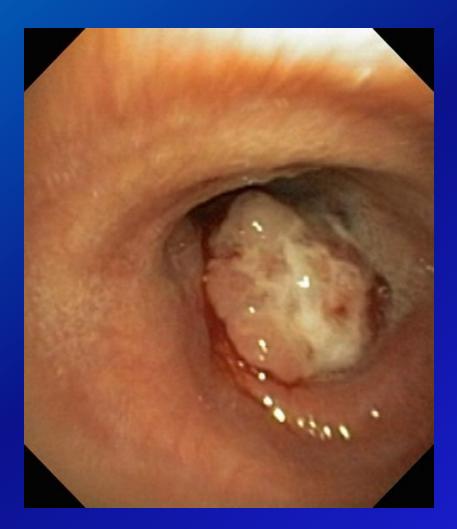










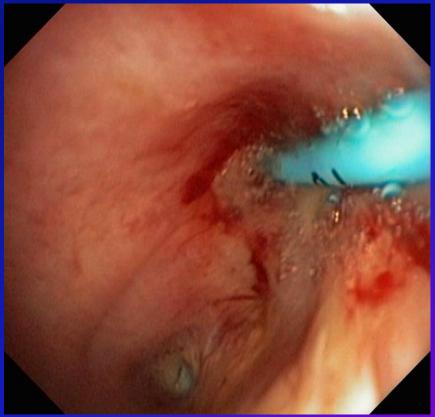






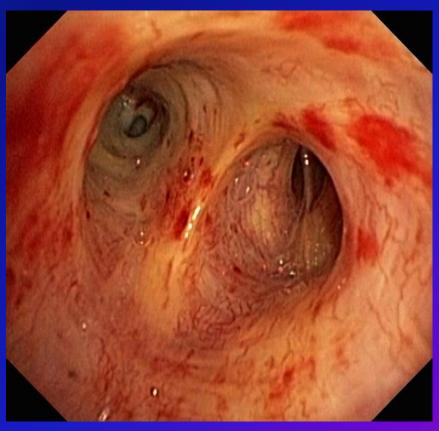






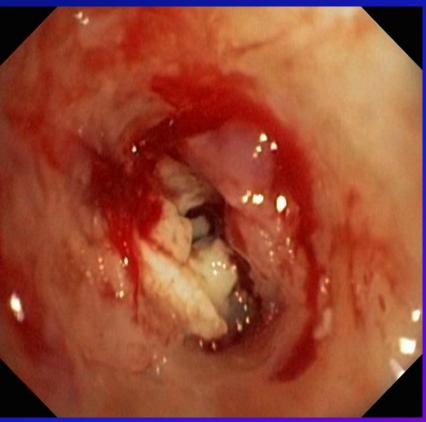
Late radiation injury





Late radiation injury





Late radiation injury

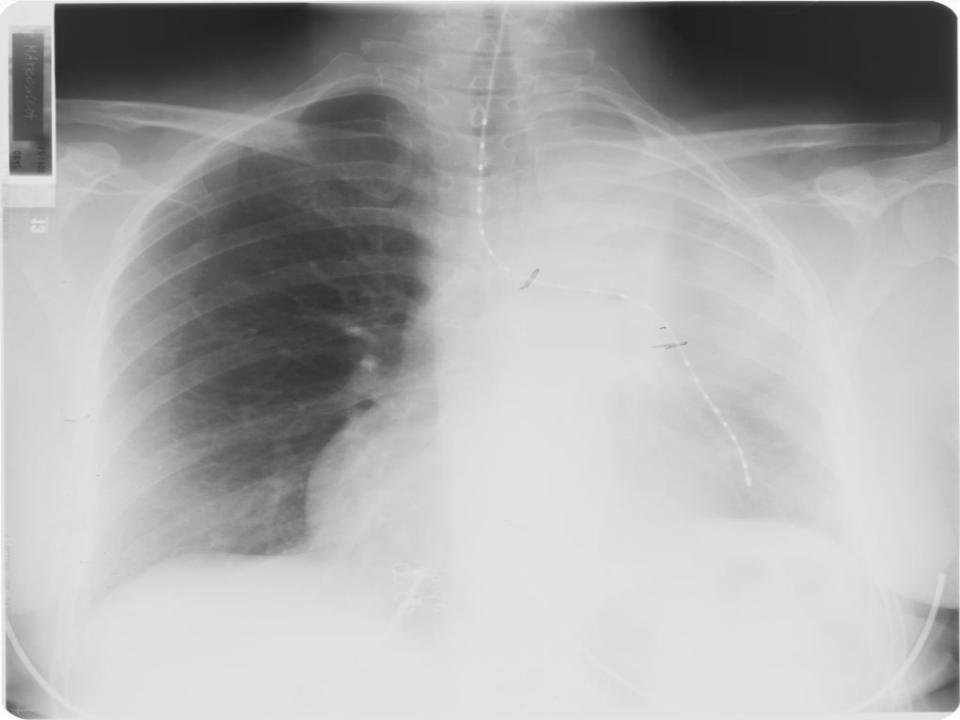


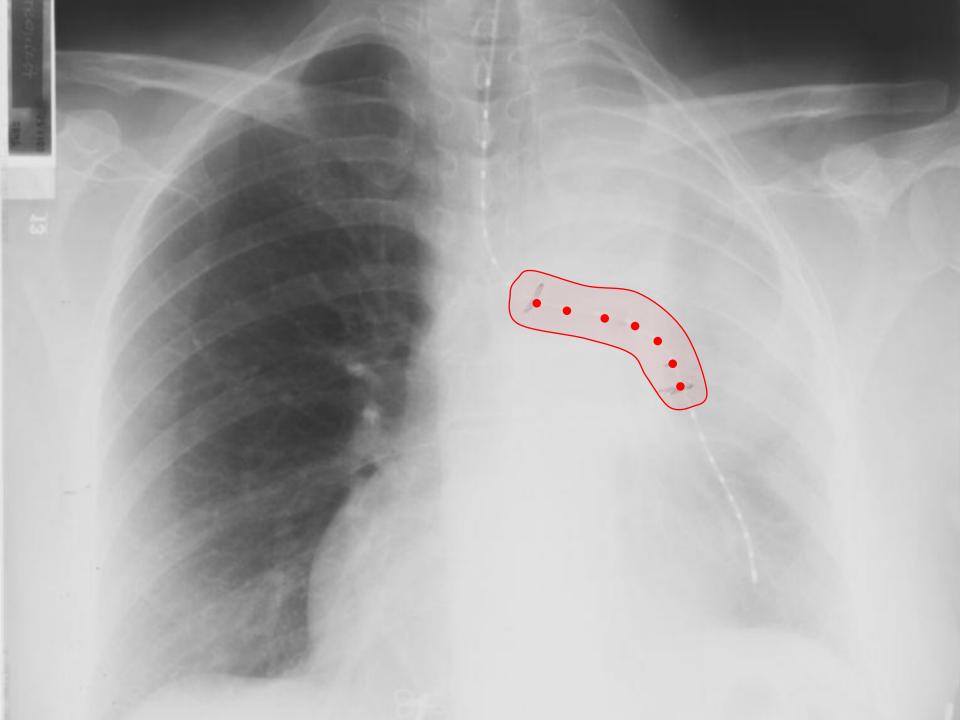


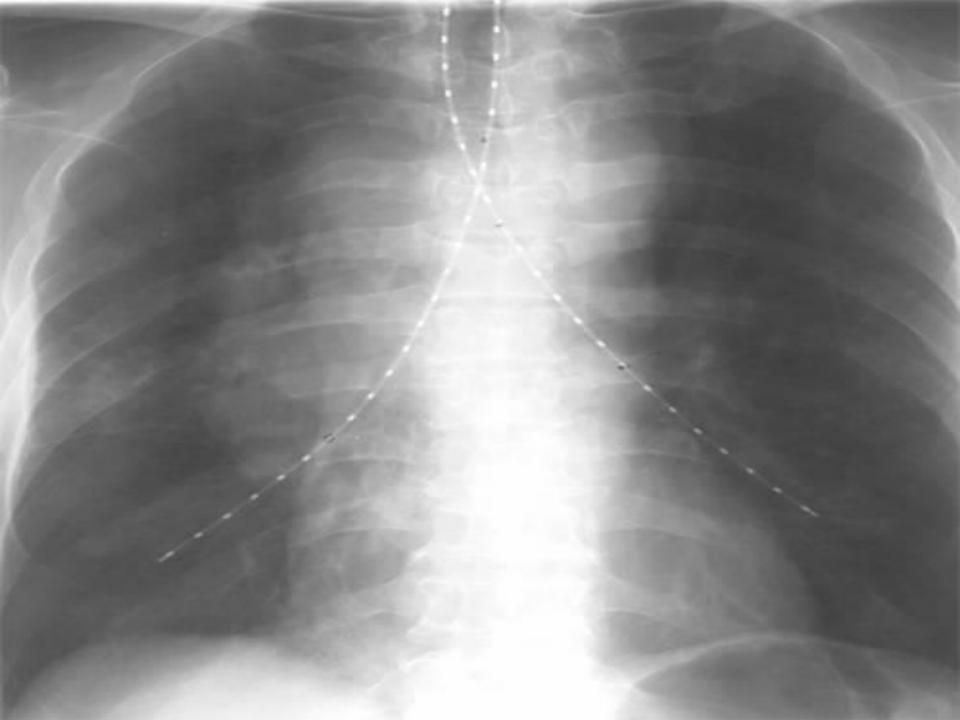


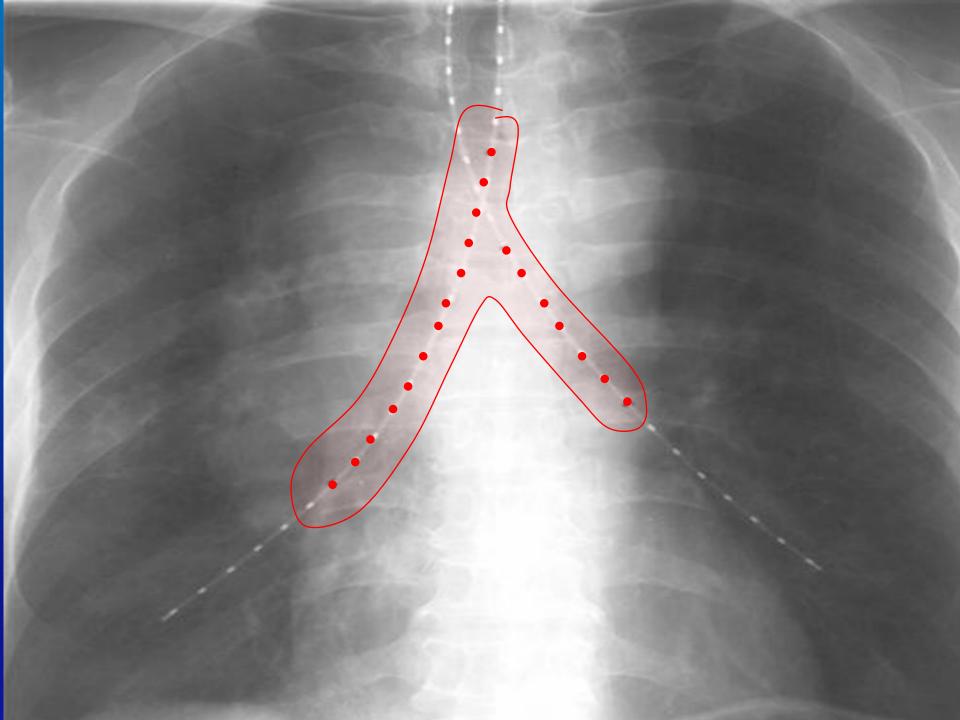


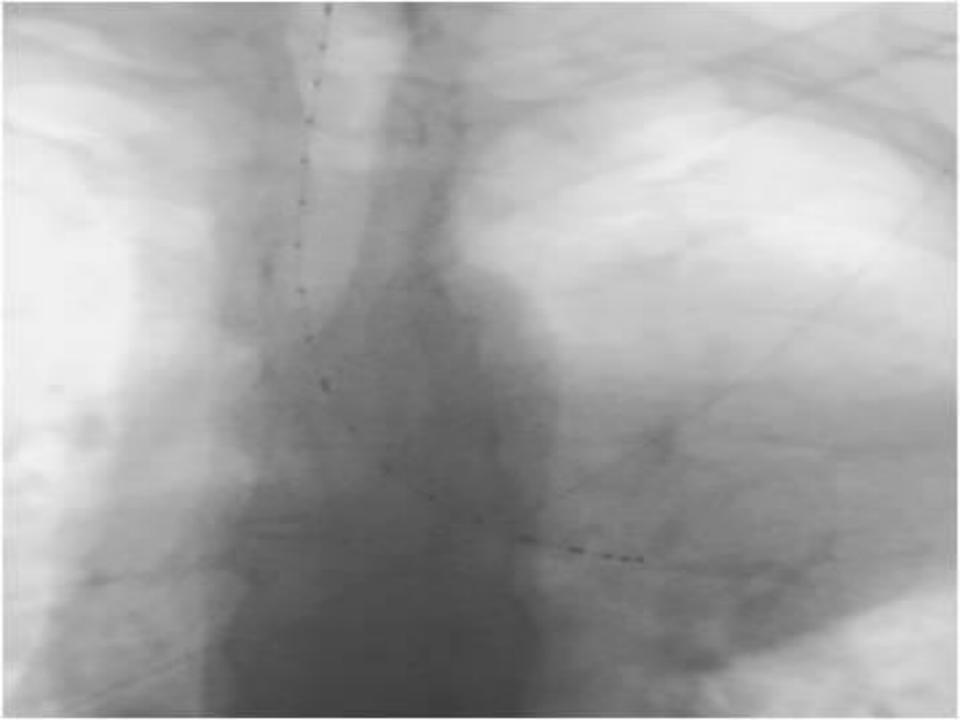




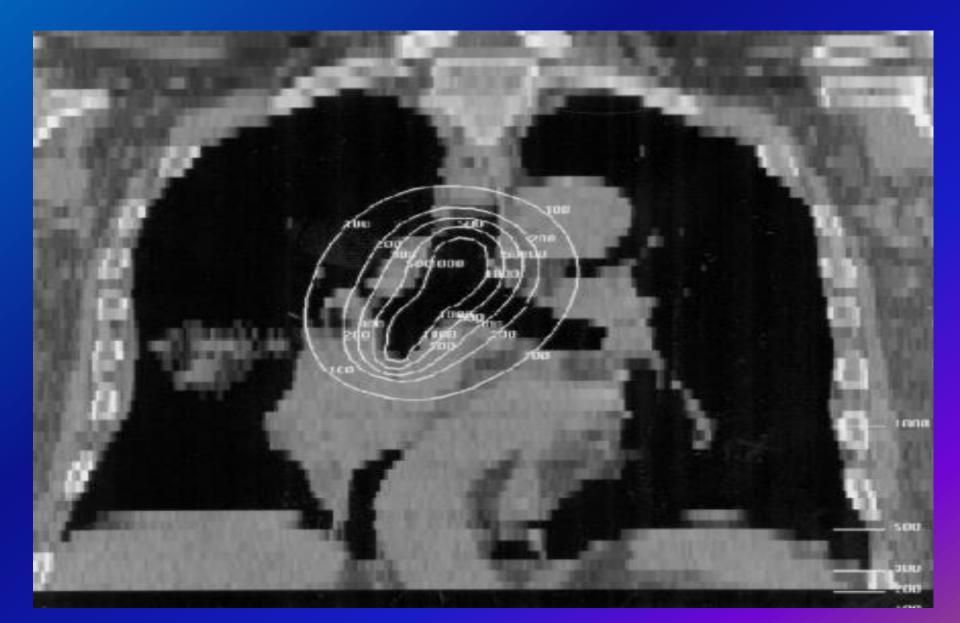






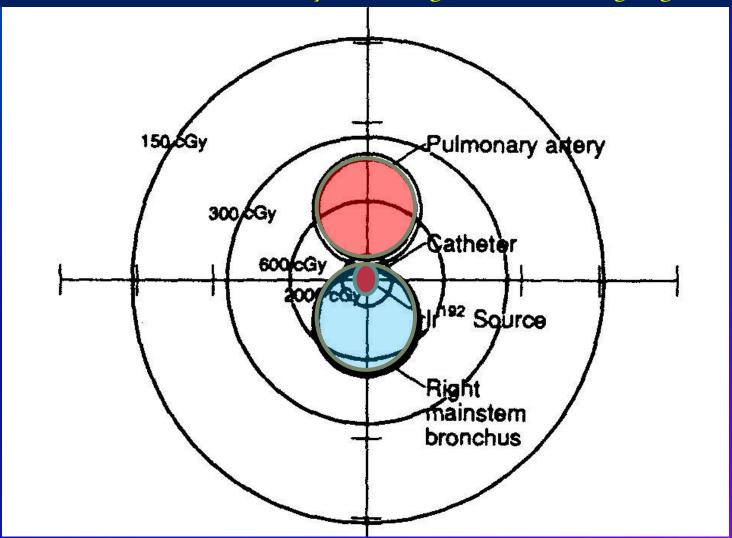


Coronal reconstruction plane with superimposed dose distribution achieved with dose prescription relative to the mucosa. Rapid dose-fall visible



Isodoses placed on schematically situated right main bronchus and pulmonary artery. Catheter with inserted isotope Ir192 is located nearby artery wall. In this case irradiation dose, growing constantly with shortening of distance to source, is very high and greater in artery wall then in tumor.

The risk of bronchus and artery wall damage and haemorrhage is great



Permanent implants (seeds)

Requirements

1. adequate performance status,

2. T1-2, No Mo,

3. easy postero - lateral access to the tumor.

Brachytherapy of inoperable tumors

- 1. Inoperable tumors, peripheral location,
- 2. Steel or elastic applicators used for seeds,
- 3. Primary or single metastasis with diameter < 2-3 cm,
- 4. Total dose 100 160 Gy, 0,5 1,0 cm outside target,

Brachytherapy after surgery

after resection, non radical, using elastic needles seeds are implanted

Convenient location

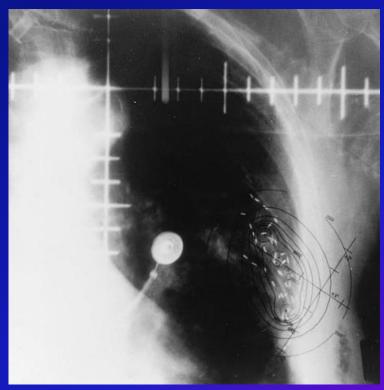


Clinical appearance of T1N0M0 lesion

Intraoperative brachytherapy for Stage I lung cancer c A. CHEN et al. Int. J. Radiation Oncology Biol. Phys., Vol. 44, No. 5, pp. 1057–1063, 1999



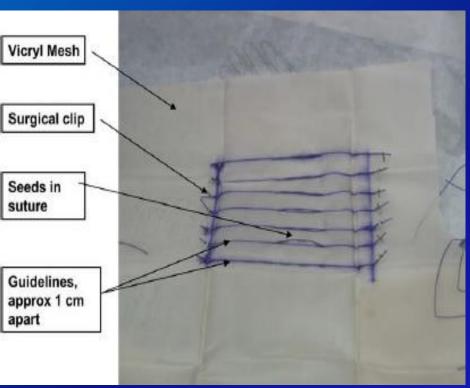
Insertion of the 125I seeds through the thoracoscopy port with video assistance.



Postoperative orthogonal simulation film with isodose curves.

A.J. Stewart et al.

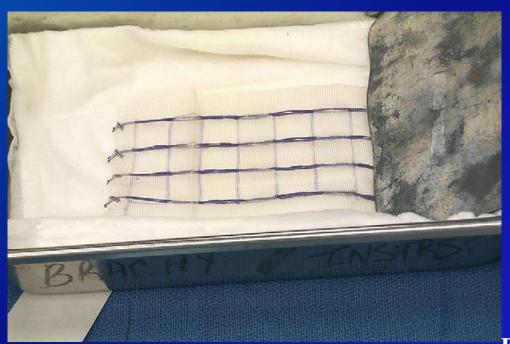
Brachytherapy 8 (2009) 63-69



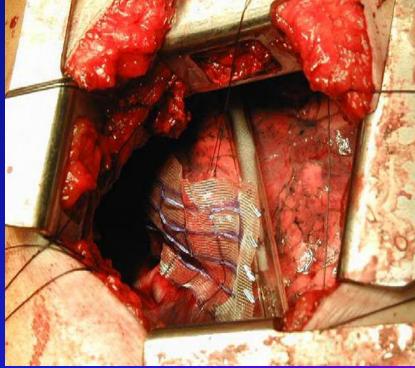
• (A) Planar implant made using 125I seeds in suture within a Vicryl mesh. (B) Completed implant being placed into mediastinum using long-handled tools. The seeds will be straightened to give optimal dosimetry before lung reinflation.



Tolerance of the aorta using intraoperative iodine-125 interstitial brachytherapy in cancer of the lung
Mark G. Trombetta, Athanasios Colonias, Daryl Makishi, Robert Keenan, E. Day Werts, Rodney
Landreneau, David S. Parda;
Brachytherapy 7 (2008) 50-54



125-I Vicryl mesh implant

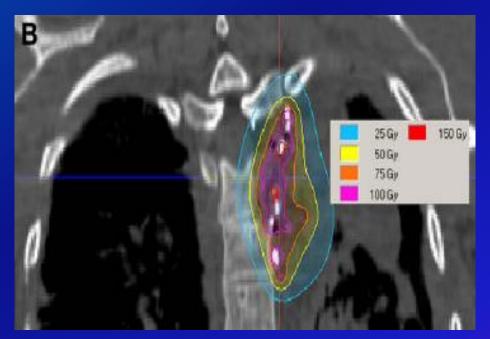


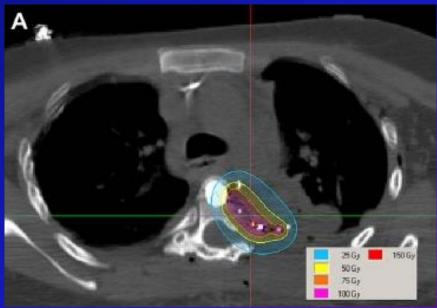
Placement of the 125I implant along the sublobar resection staple line.

Intraoperative seed placement for thoracic malignancydA review of technique, indications, and published literature Alexandra J. Stewart, Subhakar Mutyala, Caroline L. Holloway, Yolonda L. Colson, Phillip M. Devlin Brachytherapy 8 (2009) 63-69

Intraoperative 125I seed placement has been used in conjunction with sublobar resection in patients with lung cancer who are medically unfit for lobar resection . This technique is currently being evaluated in the USA in a multi-institution randomized prospective trial

by the American College of Surgeons Oncology Group (ACOSOG) Z4032





Axial (A) and sagittal (B) isodose distributions for a planar permanent 125I paraspinal implant.



Thank you for your attention