# ATC/NEMA/AAPM DICOM Demonstration

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## Two-Phase Strategy for DICOM Demonstration

- Demonstrate export of ATC-Compliant DICOM RT objects (or as large a subset as possible)
- 2. Demonstrate import of RT objects exported in Phase 1.

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# DICOM Demonstration Strategy Step 1: DICOM Export

- 1. Distribute a modest test suite as a starting point for manufacturers:
  - a. CT Image Series (DICOM)
  - b. RT Structure Set (DICOM)
  - c. Instructions for creating treatment plans
- 2. Manufacturers
  - a. Import images, structures
  - **b.** Create plans
  - c. Export CT images, RT Structure Set; RT Plan, RT Dose, RT Image
- 3. Display and compare data submitted by manufacturers
  - ATC web-based review tools (RRT, NetSys)
  - Computation Environment for Radiotherapy Research (CERR)

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DICOM Demonstration Strategy Step 2: DICOM Import

- 1. Provide a library of manufacturers' DICOM objects for off-line testing:
  - a. Includes as much as possible, CT Images, RT Structure Set, RT Plan, RT Dose, RT Image from Step 1.
  - **b.** ATC web-based review tools provide displays of what data should look like.
- 2. Demonstrate exchange using
  - a. Manufacturer-to-manufacturer network exchange
  - **b.** Central Test Node (ATC, AAPM, Merge, ???)
  - c. Media exchange



# **DICOM** Test Data

- Starting point for treatment planning
- De-identified
  - Remove "hidden" identifiers
  - New instance UIDs
- Minimize obstacles for importing into treatment planning systems
  - Remove private tags
  - Structure Sets contour Z positions coincide with those of CT slices
  - Available both as LittleEndian/ImplicitVR Datasets, and as Part-10 Filesets



# DICOM "Pushset" Script

.\pushset.bat <DCMDIR> <APP\_TITLE> <HOST> <PORT\_#>

Where:

DCMDIR is the DICOM file directory one level below this scripts directory on the CD
APP\_TITLE is the AE\_TITLE of the DICOM receiver host
HOST is the host name (or IP address of the DICOM receiver)
PORT\_# is the port number (usually 104) of the DICOM receiver

Example:

.\pushset.bat ATC04PR02 STORESCP rtp1.mydomain 104



# **DICOM** Test Data Sets

### 1. ATC04PR02

- 3D-Conformal Prostate
- Conforms to RTOG protocol 0126
- 2. ATC04HN02
  - IMRT Head/Neck
  - Conforms to RTOG protocol 0225



### Case 1 (ATC04PR02) Data

- 97 CT slices
- 10 Structures:
  - PTV2 (high-dose)
  - PTV1 (low-dose)
  - Prostate/SV [CTV1]
  - CTV2
  - Bladder
  - Rcctum
  - RT Femur
  - Lt Femur
  - Penile bulb
  - Skin





### Case 1 (ATC04PR02) Plan Specifications

• 2 fraction groups (sub-plans) with 6 beams each

	ICRU Ref Point Dose (cGy)		
	Gantry	Initial	Boost
RPO	315	837	351
RLat	270	1224	514
RAO	225	837	351
LAO	135	837	351
Llat	90	1224	514
LPO	45	837	351
Total		5796	2432





### Case 1 (ATC04PR02) Target Volume Prescription

- Prescription (ICRU Ref Dose) PTV1: 57.97 Gy PTV2: 82.28 Gy (44 fractions)
- Minimum dose
   PTV1: 95% RxD = 55 Gy
   PTV2: 95% RxD = 78 Gy
- Coverage score (minimum dose)
  - 100% = no variation
  - $\geq 95\% = \text{minor variation}$
  - < 95% = major variation
- Maximum dose to PTV2
  - $\leq 107\%$  RxD = no variation
  - $\leq 110\%$  RxD = minor variation
  - > 110% RxD = major variation





### Case 1 (ATC04PR02) Normal Tissue Guidelines

#### Maximum Dose to Percent of Volume

	15%	25%	35%	50%
Bladder	80 Gy	75 Gy	70 Gy	65 Gy
Rectum	75 Gy	70 Gy	65 Gy	60 Gy





### Case 2 (ATC04HN02) Data

- 107 CT slices
- 28 structures

Brain	RT Parotid	
CTV 59.4	Rt Parotid-PTV	
CTV 70	Shoulder 2	
Cord	Skin	
INITIALREF	TMJ	
LT Parotid	avoidance	
LT Parotid-PTV	brainstem	
Larynx	cord+8mm	
Mandible	gross vol	
Optic Chiasm	inf avoidance	
Oral Cavity	lt eye	
PTV 59.4	lt optic nerve	
PTV 70	rt eye	
Pituitary	rt optic nerve	





### Case 2 (ATC04HN02) Target Volumes

- Target volumes
  - CTV70 = GTV + microscopic extensions
  - PTV70 = CTV70 + 0.5 cm
  - CTV59.4 = cervical lymph nodes at high risk
  - PTV59.4 = CTV59.4 + 0.5 cm





### Case 2 (ATC04HN02) Prescription

- 33 Fractions, 5 fractions per week, all fields treated once daily
- Rx Dose: 70Gy (PTV70), 59.4Gy (PTV59.4)
  - No variation: ≥ 95% of PTV at or above RxD or ≥ 99% of PTV at or above 93% of RxD
  - Minor variation:  $\geq$  95% of PTV at or above 93% of RxD
  - Major variation: < 95 % of PTV at or above 93% of RxD
- Dose heterogeneity
  - No variation: no more than 20% of either PTV is at or above 77Gy
  - Minor variation: nor more than 5% of either PTV is at or above 80.5 Gy



### Case 2 (ATC04HN02) Normal Tissue Constraints

Organ at Risk	Dose Limit and Criteria

Temporal lobes 60 Gy or 1% of Vol. > 65 Gy

Brainstem, optic 54 Gy or 1% of Vol. > 60 Gy nerves, chiasm

Spinal cord 45 Gy or 1 cc > 50 Gy

Mandible, TMJ 70 Gy or 1cc > 75 Gy



Parotid glandsMean dose to either gland < 26 Gy, or</th>50% of either gland receives < 30 Gy, or</td>20cc of combined glands receives < 20 Gy</td>Minor variation:40% of either gland receives < 30 Gy</td>



## DICOM Objects to Be Submitted to ATC

- CT Image
- RT Structure Set
- RT Dose (3D dose distribution) for each fraction group
- RT Plan
- RT Dose (DVH) for total dose plan
- RT Image (DRRs)













# Review Tools for Display of Data

- ITC Remote Review Tool
- RCET NetSys Client
- CERR



### **Remote Review Tool**

HEART

- CT Images (zoom, ۲ window/level)
- Structure contours ۲ (review, editing)
- Iso-dose contours ۲
- Interactive DVH ۲

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Point-dose display ۲



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## Computational Environment for Radiotherapy Research (CERR) – J. Deasy, Washington Univ.





# Submission Timeline

- Vendors submit DICOM data to ITC until June 25, 2004
  - FTP to castor.wustl.edu
  - CD-R
  - WebSys (contact Walter Bosch, <u>itc@castor.wustl.edu</u> for account)
- AAPM 46th meeting July 25-29, 2004
  - Network
  - CD-R
  - USB drive



### Email from Dwight Simon 2/14/04

Walter/et al,

I've tried to keep track of all the things that go on in WG7 over the years, but I certainly haven't done an adequate job. However, I am aware of the great amount of energy and effort that has been put into developing the RT portion of DICOM over the years and I would like to applaud your diligence and accomplishments. I believe, that you have the most complex set of DICOM objects in the standard and, this is where the difficulty has come in for getting it all to work in a fully interoperable environment.

I wish you the best of luck in this latest demonstration endeavor and would like to encourage everyone to participate to there fullest extent. I think you are on the verge of really making it happen - if you get full and in depth participation.

Best Regards, Dwight