Riunione Nazionale FIL Padova, 8-10 Novembre 2012

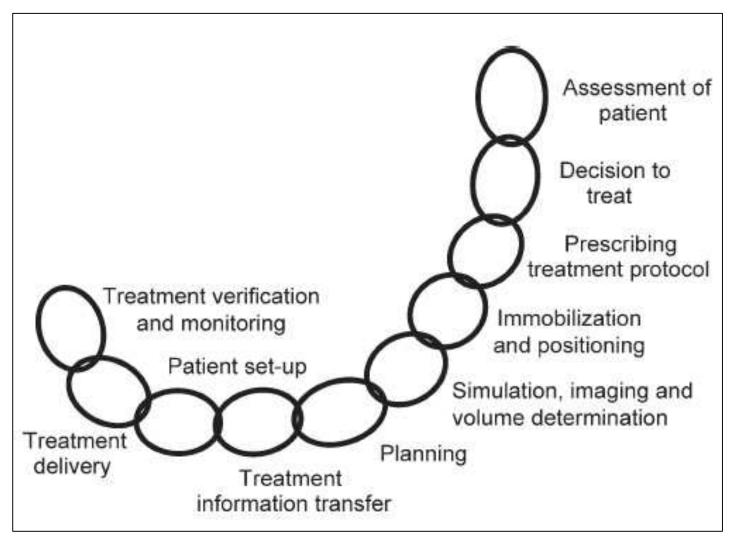
Quality Assurance in Radiotherapy



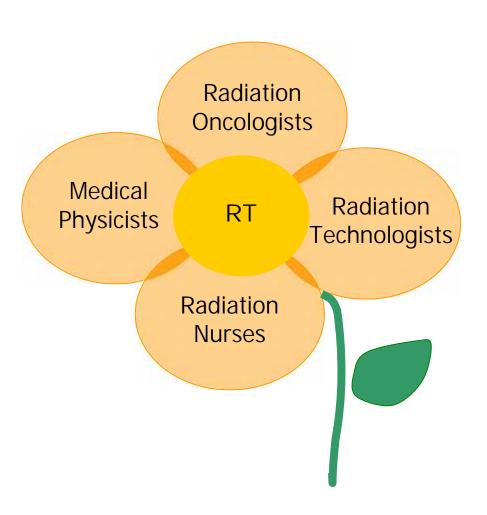
Patrizia Ciammella

Arcispedale Santa Maria Nuova Reggio Emilia

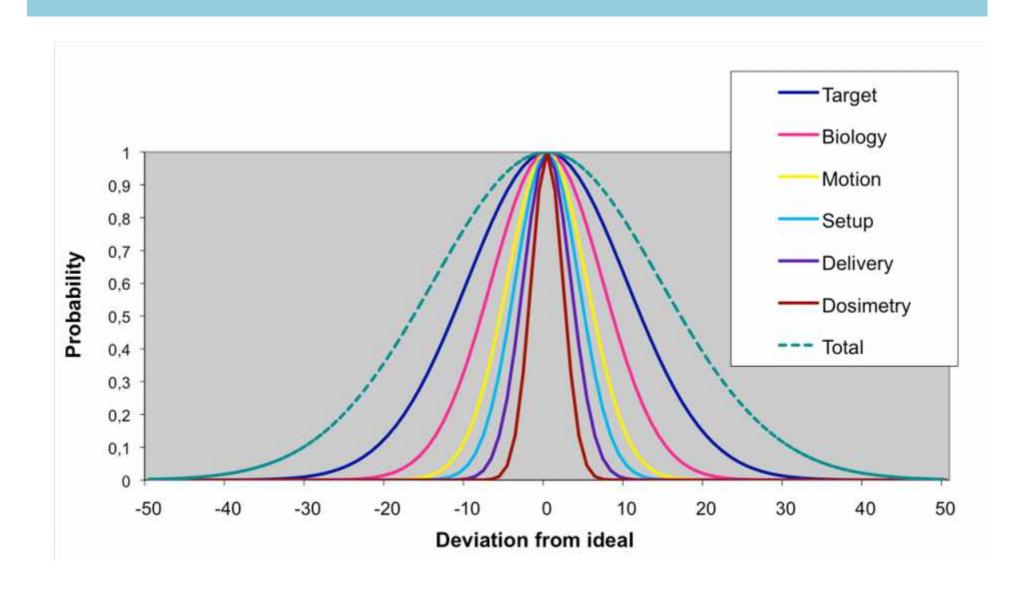
RT process



RT team



Residual treatment uncertainties



Philosophy of radiotherapy quality assurance

Scope of quality assurance protocols

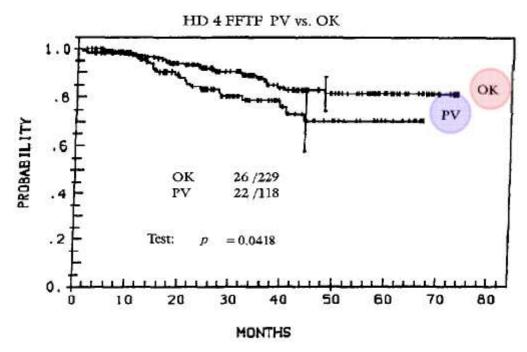


- guarantee the validity of clinical trial results
- improve the quality of clinical practice minimising the risk of errors
- increase the likelihood of desired health outcomes at the population level (equality of access, consistency, etc)

Importance of QA protocols

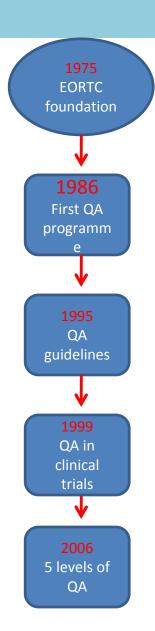
Clinical Original Contribution

RANDOMIZED TRIAL WITH EARLY-STAGE HODGKIN'S DISEASE TESTING 30 GY VS. 40 GY EXTENDED FIELD RADIOTHERAPY ALONE



5-years FFTF was significantly influenced by the quality of radiotherapeutical procedures: 70% with protocol violations (PV) vs. 82% without PV

Quality assurance in radiotherapy



QA strategy of the EORTC Radiation Oncology Group (ROG)

1982-2005

- Site visits
- Evaluation of institutions' staff and infrastructure
- Dosimetric checks of treatment units
- Cross-check of patients charts and portal images
- Radiobiological modelling of inter-institutional differences
- Mailed thermoluminescent dosimetry audits

2005-2011

- Level 1: Facility questionnaire, External reference dosimetry audit
- Level 2: Dummy run
- Level 3: Limited individual case Review
- Level 4: Extensive individual case Review
- Level 5: Complex dosimetry check

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QUALITY OF RADIOTHERAPY REPORTING IN RANDOMIZED CONTROLLED TRIALS OF HODGKIN'S LYMPHOMA AND NON-HODGKIN'S LYMPHOMA: A SYSTEMATIC REVIEW

JUSTIN E. BEKELMAN, M.D.,* AND JOACHIM YAHALOM, M.D.*

133 Publications identified in PubMed of controlled trials of Hodgkin Lymphoma and Nonreferences Hodgkin Lymphoma involving 2 in Cochrane CENTRAL. radiotherapy 3 in references Publications excluded on the basis of title and abstract 21 Not Phase III RCT 19 Secondary report of 18 Not involving EBRT 75 Publications selected for review 19 Publications excluded after obtaining full text 8 Secondary report of 7 Not Phase III RCT 3 RCT involving IBI 2 Not involving EBRT 56 Publications included from PubMed search 61 Publications included in review

Non-duplicative publications identified in Cochrane or in references

Quality Measures

Int. J. Radiation Oncology Biol. Phys., Vol. 73, No. 2, pp. 492-498, 2009

- 1. Target volume definition
- 2. Radiation dose specification
- 3. Fractionation specification
- 4. Radiation prescription point specification
- 5. QA process use for RT
- 6. QA process adherence reporting for RT

Radiotherapy reporting quality

	Adequacy of reporting	
Measures of radiotherapy reporting	n	%
1. Target volume description	23	38
2. Radiation dose specification	54	89
3. Fractionation specification	39	64
4. Radiation prescription point specification	13	21
5. Quality assurance process use	12	20
6. Quality assurance process adherence reporting*	7	11

Reporting of RT in HL and NHL RCTs is deficient. Because the interpretation, replication, and application of RCT results depend on adequate description and QA of therapeutic interventions, consensus standards for RT reporting should be developed and integrated into the peer-review process.

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Quality assurance in radiotherapy

In the last years radiation oncology is undergoing a rapid and radical transformation driving an exponential growth in the size, complexity, and quantity of data generated during the treatment process.

The increase is due, in part, to the success of image-guided oncology programs, which generate new images at each step in the treatment process — from diagnosis to verification, up to follow-up.

Modern QA → digital data exchange and timely review

Main system requirements

Task	Parameters
General	Network reliability in terms of up-time and fidelity of data transfer Avaidance of least barehvers and aeftween installation.
	 Avoidance of local hardware and software installation Web-based accessibility from any geographic location Intuitive functionally evaluation tools Uniformity of data submission procedures for all trials Adequate firewall security Central storage and archiving in a queriable database
Submission from centres to QART office	 Ability to upload operating-system independent Reads in digital data (DICOM-RT) exported from multiple TPS
Central evaluation	 Document assessment of RT plan via a web-based evaluation matrix, standardize to facilitate consensus-building

A scenario for a web-based radiotherapy treatment planning system

Patients benefit

 Accessibility of advanced radiotherapy treatment services, minimising patient's transportation

Physicians benefit

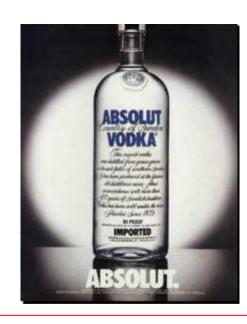
- Consultation of expert colleagues
 - Cooperation with reduce professional isolation

Organisations benefit

- Quality assurance in terms of dummy-runs and/or check of a real-patient treatment planning
- Minimisation of local-audits and the cost of on-site visit
- Evaluation and correlation of treatment outcome with radiotherapy parameters

RTQA platforms used by RTOG, EORTC or TROG:

- VodcaRT from Medical Software Solutions
- CERR software
- ITC remote review tool
- VelocityAI from Velocity Medical Solutions
- Mim software from Mimvista
- Artiview from Aquilab
- Swan
- (Widen)



- 2008: retrospective test
- 2009: prospective test
- 2010: currently utilize to QART procedures on ongoing EORTC trials

VODCA



Visualisation and Organisation of Data for Cancer Analysis

has been created to ...

- Promote in radiotherapy clinical decision-support and knowledge management technologies in patient care and clinical research;
- Disseminate methods and tools for building healthcare knowledge applications that comply with the highest quality, safety and ethical standards;
- Connect individuals, hospitals, research groups and organisations who believe in the value of knowledge management, and support the share and reuse of clinical knowledge and data.

"Prospective quality assurance program and development of an Italian network for radiotherapy in patients affected by lymphoma enrolled in the clinical trials"



Proof of Concept (POC)

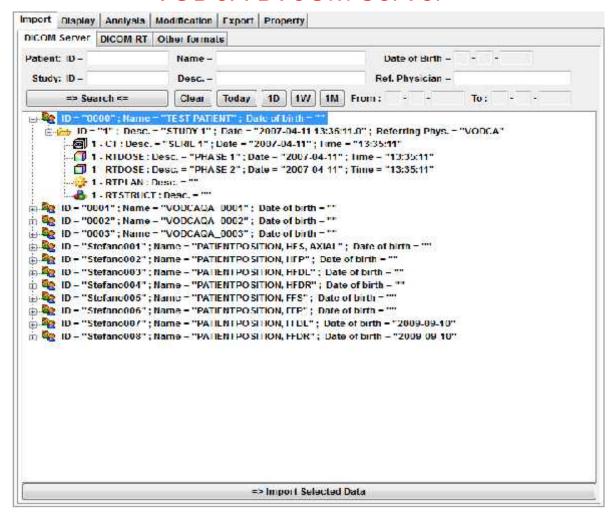
- 6 RT centres
- July-September 2012
- HL patients

VODCA

VODCA Control



VODCA DICOM Server



VODCA

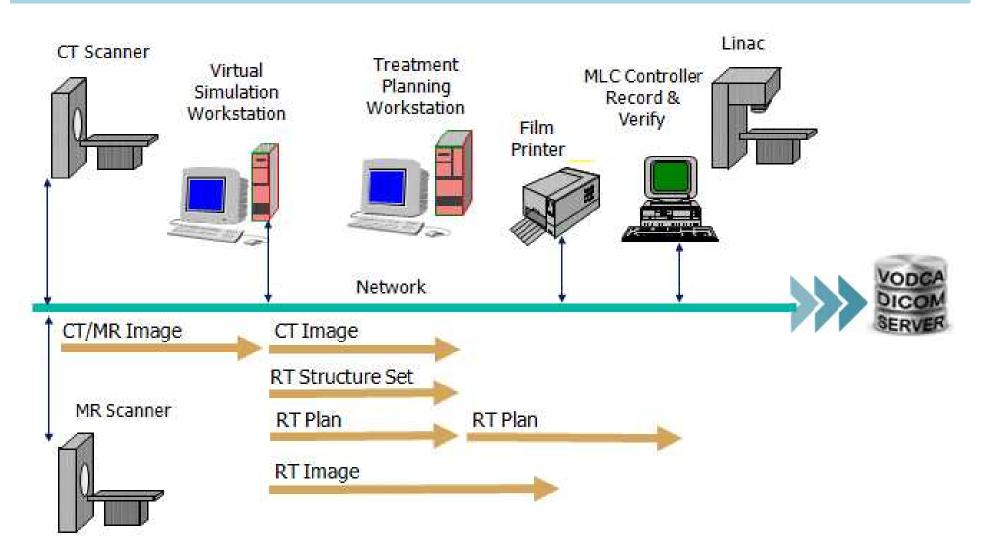
Databases

Every patient dataset has an unlimited amount of user defined fields subdivided into 3 categories:

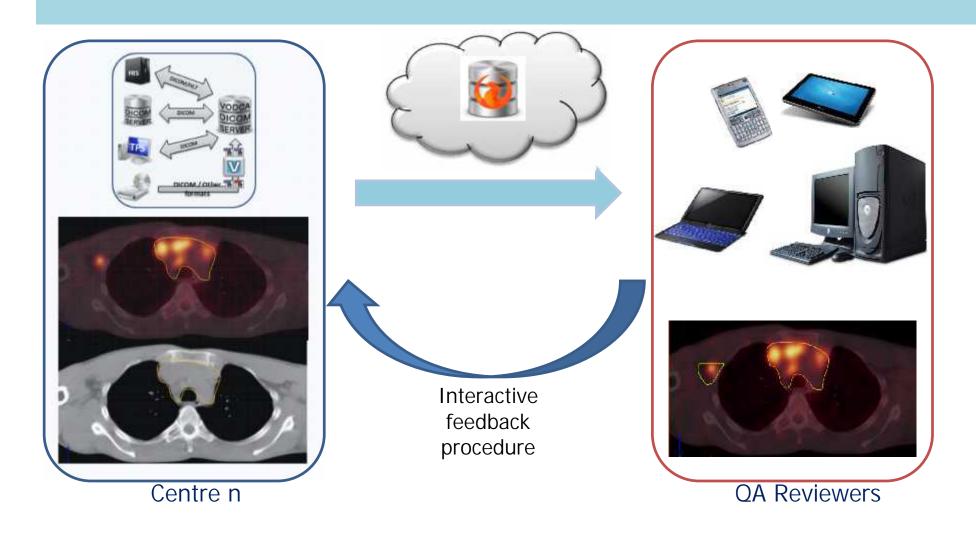
- Patient data,
- Treatment data, and
- Clinical data,

Data fields can be added or deleted at any time by the database administrator.

VODCA Network



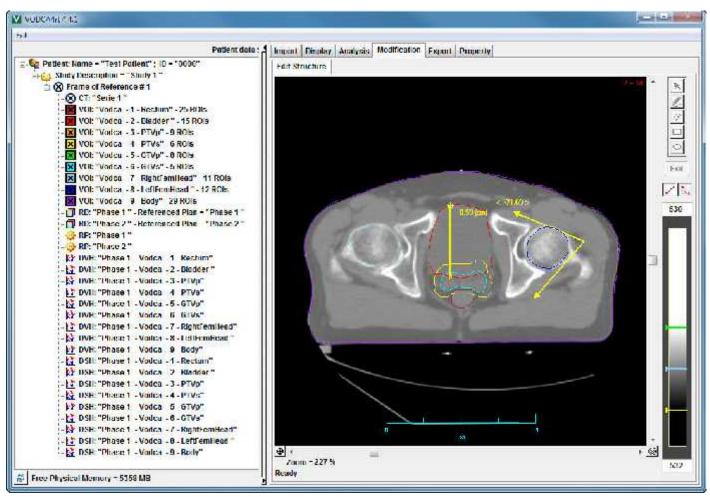
"VODCA on the cloud"



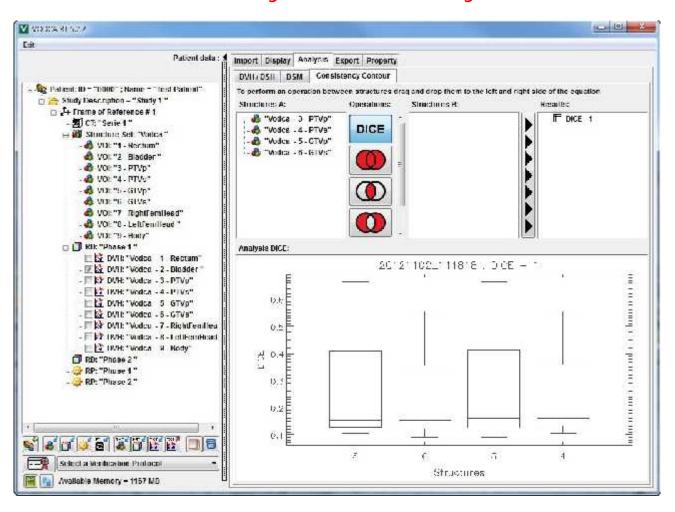
The **"VODCA Online Desktop"** is a powerful cloud-based Desktop with a customisable secure storage and it is accessible from anywhere, including tablets:

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Data Modification



Module "Analysis" – Dummy-run

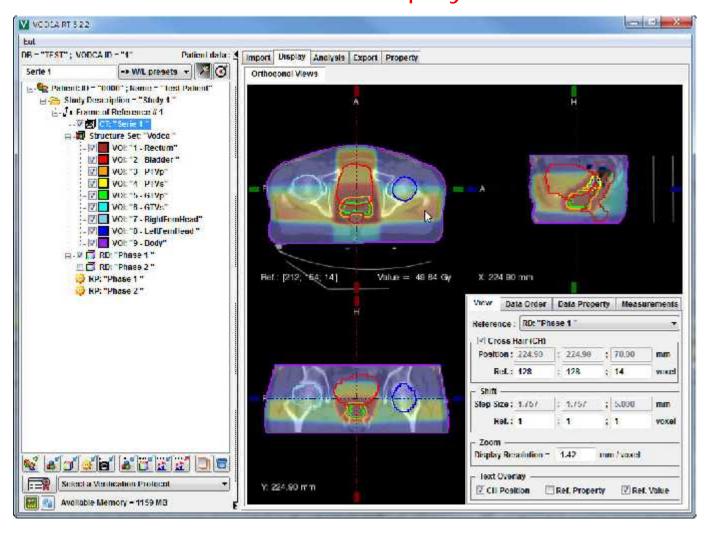


Consistency Contour

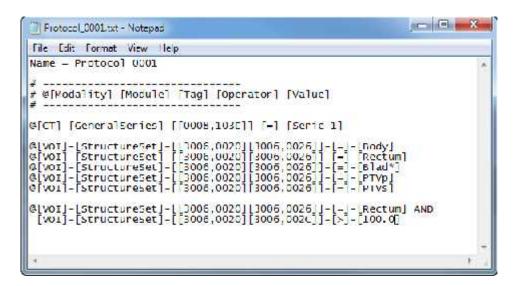
The following parameters are calculated:

- Sensibility
- Specificity
- Volume difference
- Isocentre difference

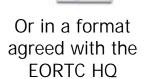
VODCA RT : Module "Display"

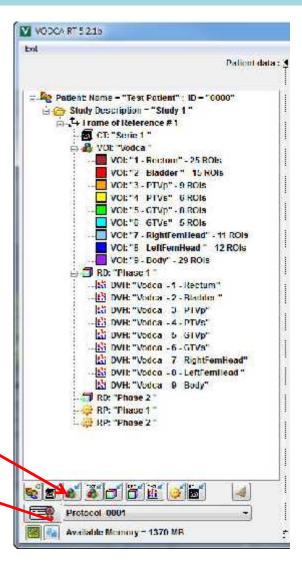


Data verification Protocol



Available [Module] and [Tag] are listed in VODCA RT "Property"





Conclusions

QART

- Widen RT-DICOM IELSG37
- Prospective trials with VODCA?

Grazie

