

Big Data: How VHA Can Be Your Friend

New Orleans, LA



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Disclosure

- Vice President, Center for the Assessment of Radiological Sciences (CARS)
 - A non-profit organization dedicated to improving quality and safety of radiotherapy and radiological imaging.



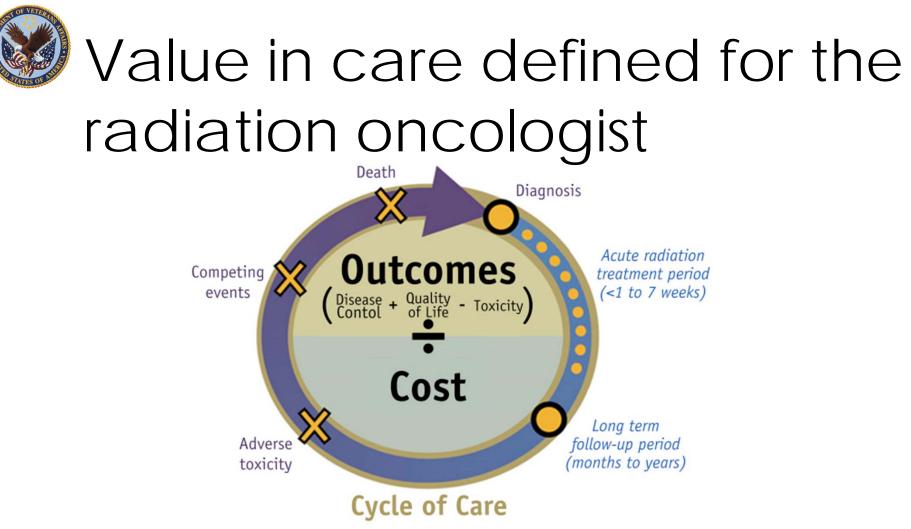


Objectives

- Discuss value proposition in radiation oncology,
- Discuss why VHA is a good test laboratory for determining value in radiation oncology,
- Describe VHAs Radiation Oncology Practice Assessment (ROPA) initiative,
- Discuss how ROPA can potentially become a model for quality and outcome assessment in radiation oncology.



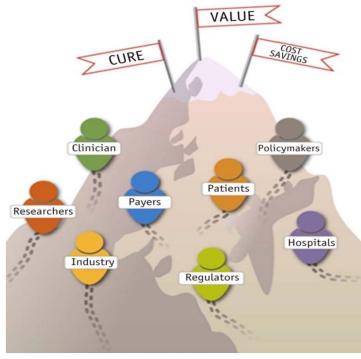




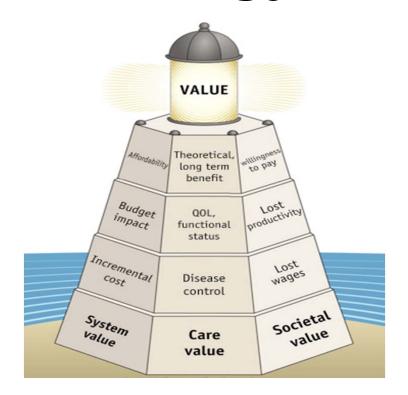
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Stakeholder in the discussion of value in oncology



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What is big data?

- Big data is a term for data sets that are so **large** or **complex** that traditional data processing application softwares are inadequate to deal with them. (Wiki)
- Big data is a term that describes the large volume of data both **structured** and **unstructured**. Big data can be analyzed for insights that lead to better decisions and strategic business moves. **(SAS)**
- Extremely large data sets that may be analyzed computationally to reveal patterns, trends, and associations.
 (Dictionary)

"5 V"; Volume, Velocity, Variety, Veracity, and Value





The Surveillance, Epidemiology, & End Results (SEER) Registries

- Includes approximately 28% of US Population
- Representative sample of all ethnicities and socio-economic backgrounds

Limitations

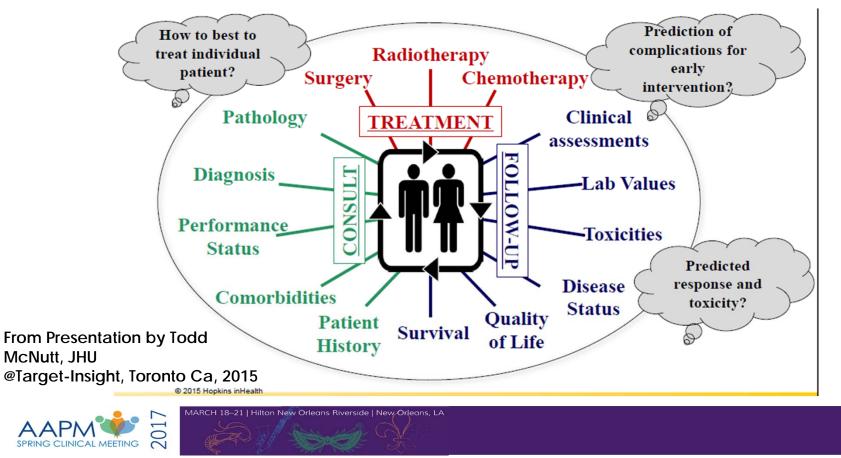
- Limited information about key health factors
- Inaccuracy; such as under ascertainment of outpatient treatments,
- Migration/loss to follow-up
- Sparse to no RT data



States included in SEER Registries; SEER 9, 13 &18

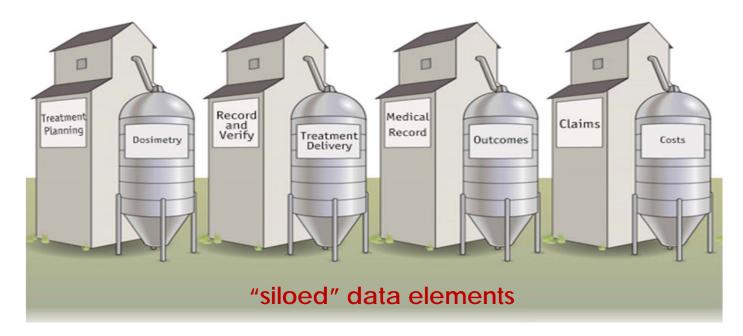


Big Data in Cancer Care





Big data Challenges in Radiation Oncology



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- Largest Integrated Health care system in United States.
 - 1,233 health care facilities
 - Incl. 168 VA Medical Centers
 - 1,053 outpatient clinics
- Serving more than 8.9 million Veteran each year.
- Annual budget: \$69 billion (2017)
- Single interconnected electronic medical record system (VISTA – CPRS) since 1983

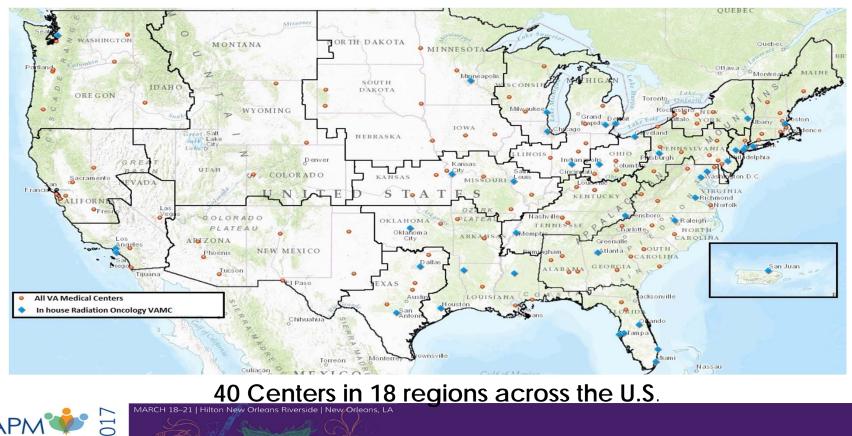






SPRING CLINICAL MEETING

VHA Radiation Oncology Centers





Radiation Oncology in VHA

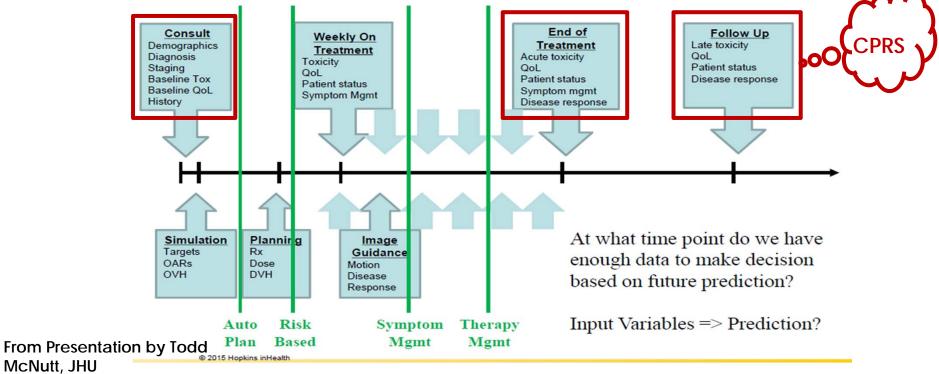
- 40 Radiation Oncology VA clinics
 - -15,000 patients treated in-house
 - -25,000 patient sent outside for RT.
 - -70+ treating radiation oncologists
 - -70+ therapeutic medical physicists
 - -72 linear accelerators

Longitudinal history of patients RT episode in Vista/CPRS.









@Target-Insight, Toronto Ca, 2015





VHA Radiation Oncology Practice Assessment(ROPA)

Purpose: Assessment of radiation delivery and cancer related outcomes for the VHA radiation oncology practices

Background: Disease-site expert panels of the American Society for Radiation Oncology (ASTRO) have identified clinical measures and associated data fields to assess the quality of radiation treatments

 These clinical measures will be used by the VHA to monitor the quality of radiation oncology and outcome assessment

 Pilot: Prostate and Lung Cancer







Scope of Data Acquisition

- Manual abstraction by visit to 40 VA Radiation Oncology Centers.
- Comprehensive evaluation of 50 cases from each center, 20-30 ASTRO vetted metrics per case
 - 20 cases prostate cancer: T1c T3, NX0M0 (Intermediate or high risk per NCCN criteria)
 - 20 cases Non-Small Cell Lung Cancer (NSCLC): Stages IIIA and IIIB
 - 10 cases Small Cell Lung Cancer (SCLC): Limited Stage.
- Most recent, serial cases in each category who have completed post-treatment follow-up examination







Data Collection for ROPA Data Sources

- Clinical data
 - Abstracted from physicians clinical note templates in CPRS used by clinicians in their routine process of care
- Radiation treatment management data
 - Abstracted from RT-EMR (e.g. ARIA, MOSAIQ)
- Treatment Planning Data
 - DICOM/DICOM-RT data abstracted from treatment planning systems (e.g. Eclipse, Pinnacle, XiO, Hi-ART...)

Data Abstraction Requirements

- No Protected Health Information (PHI) will be recorded.
- Treatment dates to be recorded as elapsed time from offset.





Clinical Measures

Defined by ASTRO Disease Site Expert Panels

- Quality Measures
 - Measures with published data that will be utilized for the practice assessment.
- Aspirational Measures
 - VA asked the panels to also provide ambitious goals or items not currently in common practice that reflect high quality.
 - Examples: Quality of life assessment prior to treatment completion, Survivorship Care Plans.

Surveillance Measures

 Measures that either do not yet have enough published data to demonstrate a link to quality (i.e. collection of molecular information) or are focused on population health (enrollment on clinical trials).





MEASURE #3: Im	aging/Staging for High Risk					
Numerator Statement	 Patients with imaging for staging, prior to the initiation of treatment, that includes: 1. CT or MRI, AND 2. Bone scan (T⁹⁹ or NaF PET). 					
Denominator Statement	All patients, regardless of age, with a diagnosis of prostate cancer, at high OR very high risk as defined by NCCN guidelines, receiving radiation therapy					
Denominator Exclusions/Exceptions	 Patients treated post prostatectomy 					
Notes	 Consensus Survey Results: 100% 					
Expected Performance Rate	 Higher = better Panel Vote: 95% CMS PQRS Measure #102 (Avoidance of Overuse of Bone Scan for Staging Low-Risk Prostate Cancer Patients). Average Performance Rate in 2011: 95.4%. in 2012: 92.9%; in 2013: 90.6% 					
Timeframe	Prior to first treatment					



DVH Metric Types

Constraint

- Metric will be used to evaluate the plan and provider's performance
- Informational
 - For the purposes of data collection
 - Not to be used to judge the appropriateness of a plan

• DVH Metric Scale

- Most DVH Constraints and DVH Informational Metrics were divided into a 3 tiered system
- Green: Pass
- Yellow: Warning
- Red: Fail







Lung Quality Measures



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Spinal Cord Dmax* Metrics

Varying Fractionation

<u>Limit</u>	<u>Green</u>	<u>Yellow</u>	<u>Red</u>	<u>Mandatory</u> <u>Constraint vs.</u> <u>Informational</u>	<u>Source</u>	<u>Fractionation</u>
50 Gy	<= 45 Gy	>45 Gy <= 50 Gy	> 50 Gy	Constraint	QUANTEC	Standard
41 Gy	<=36.9 Gy	>36.9 Gy <= 41 Gy	> 41 Gy		Turrisi, NEJM 1998, RTOG 0538	Hyper
37 Gy	<=33.3 Gy	>33.3 Gy <= 37 Gy	> 37 Gy		BED calc (aB = 3, EQD2 = 49.6 Gy)	Нуро - 10
42 6 4		>37.8 Gy <=			Timmerman / USC, confirmed w/BED (aB = 3, EQD2 = 48.7	lbung 15
42 Gy	<= 37.8 Gy	42 Gy	> 42 Gy	Constraint	Gy)	Нуро - 15
*Dose to <0.03	01.	1 Hilton New Orleans River	side New Orleans, LA			Ĩ



Various Lung Metrics

Standard Fractionation

					<u>Mandatory</u> Constraint vs.		
<u>Metric</u>	<u>Limit</u>	<u>Green</u>	<u>Yellow</u>	<u>Red</u>	Informational	<u>Source</u>	<u>Note*</u>
V20 Gy	37%	<= 33%	> 33% <= 37%	> 37%	Constraint	QUANTEC	2 lungs
V5 Gy	60%	<= 54%	> 54% <= 60%	> 60%	Informational	RTOG 1308	2 lungs
Dmean	20 Gy	<= 18 Gy	> 18 Gy <= 20 Gy	> 20 Gy	Informational	QUANTEC	2 lungs
						Rice et al,	
V20 Gy	7%	<= 6.3%	> 6.3% <= 7%	> 7%	Constraint	IJROBP 2007	1 lung
V5 Gy	60%	<= 54%	> 54% <= 60%	> 60%	Informational	Allen et al, IJROBP 2007	1 lung
v J Gy	0070	<u> </u>	> 5470 <= 0070	> 0070	mormational		Tiung
						Rice et al,	
Dmean	8.5 Gy	<= 7.7 Gy	> 7.7 Gy <= 8.5 Gy	> 8.5 Gy	Constraint	IJROBP 2007	1 lung

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*Total Lung - XXX. In order of availability, GTV>ITV>CTV>PTV



Esophagus Metrics

Standard Fractionation

Metric	<u>Limit</u>	<u>Green</u>	<u>Yellow</u>	Red	<u>Mandatory</u> <u>Constraint vs.</u> Informational	<u>Source</u>
	4 70 (4 = 0.04		4 = 0 (Palma et al,
V60 Gy	17%	<= 15.3 %	> 15.3% <= 17 %	> 17%	Informational	IJROBP 2014
Dmean	34 Gy	<= 30.6 Gy	> 30.6 Gy <= 34 Gy	> 34 Gy	Informational	QUANTEC
Dmax*	74 Gy	<= 66.6 Gy	> 66.6 Gy <= 74 Gy	> 74 Gy	Informational	RTOG 1308

* Dose to <0.035 cc





Other Metrics

Standard Fractionation

OAR	Metric	Limit	Green	Yellow	Red	<u>Mandatory</u> <u>Constraint vs.</u> Informational	Source
Brachial				<u></u>		<u></u>	<u></u>
Plexus	Dmax*	66 Gy	<= 59.4 Gy	> 59.4 Gy <= 66 Gy	> 66 Gy	Constraint	QUANTEC
Heart	V45Gy	35%	<= 31.5%	> 31.5% <= 35%	> 35 %	Informational	RTOG 1308
PTV	D95%	100% Rx	100%	>= 95% < 100%	< 95%	Constraint	RTOG 1308
PTV	Dmin*	85% Rx	>85%	>= 75% < 85%	< 75%	Informational	RTOG 1308

* Dose to <0.035 cc







Prostate Quality Measures



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Rectum Metrics

External Beam, Varying Fractionation

					Mandatory Constraint vs.		
Metric	<u>Limit</u>	<u>Green</u>	<u>Yellow</u>	Red	Informational		Fractionation
						RTOG 0126, 0415,	
V70 Gy*	25%	<=25%		>25%	Constraint	0815	Standard
V69 Gy	25%	<=25%		>25%	Informational	RTOG 0415	Нуро
V70 Gy	15%	<=15%		>15%	Informational	Michalski et al, IJROBP 2013	Standard
V75 Gy	10%	<=10%		>10%	Informational	Michalski et al, IJROBP 2013	Standard
V50 Gy	50%	<=50%		>50%	Constraint	QUANTEC	Standard





*Should be met in >= 90% of cases



Bladder, Femurs Metrics

External Beam, Standard Fractionation

<u>OAR</u>	Metric	<u>Limit</u>	<u>Green</u>	<u>Yellow</u>	Red	<u>Mandatory</u> <u>Constraint vs.</u> Informational	<u>Source</u>
Bladder*	V70 Gy	35%	<=35%		>35%	Informational	QUANTEC, RTOG 0126, 0415, 0815
Bladder	V65 Gy	50%	<=50%		>50%	Informational	QUANTEC, RTOG 0126, 0415, 0816
Femurs	V50 Gy	10%	<=10%		>10%	Informational	RTOG 0534





*Should be met in >= 90% of cases



Bowel Metrics

External Beam, Standard Fractionation

<u>OAR</u>	Metric	<u>Limit</u>	<u>Green</u>	<u>Yellow</u>	<u>Red</u>	<u>Mandatory</u> <u>Constraint vs.</u> Informational	<u>Source</u>
							QUANTEC, RTOG
Bladder	V70 Gy	35%	<=35%		>35%	Informational	0126, 0415, 0815
Bladder	V65 Gy	50%	<=50%		>50%	Informational	QUANTEC, RTOG 0126, 0415, 0816
Femurs	V50 Gy	10%	<=10%		>10%	Informational	RTOG 0534

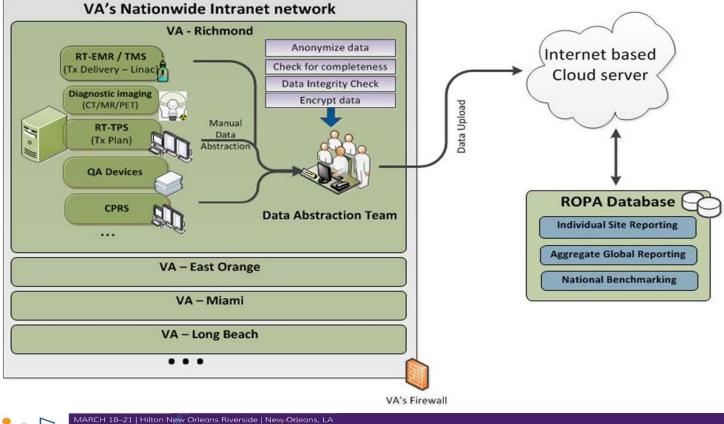




*Volume is Bladder minus CTV



VHA ROPA Workflow







VHA ROPA

Deliverables

- Facility reports: detailed radiation delivery parameters and outcomes, nationally benchmarked for 50 cases
- VHA global report: examines variability within VHA
- Benefit to the VHA enterprise: Roadmap for continuous improvement for each in-house radiation oncology practice
- Identification of metrics for future internal, remote evaluations using VA's EMR (CPRS)

Parallel Effort

Electronic abstraction of data fields for clinical measures directly from different data sources and performing periodic remote electronic re-assessment.





VHA ROPA

Data Sources

- Clinical data
 - Abstracted from disease-site specific clinical note templates in CPRS used by clinicians in their routine process of care
- Radiation treatment management
 - Abstracted from RT-EMR (e.g. ARIA, MOSAIQ)
- Treatment Planning Data
 - DICOM/DICOM-RT data abstracted from treatment planning systems
- Patient Reported Outcome data from Patient Portals

Electronic Data Abstraction

- Deployment of the data aggregation software at the local facility
 - Aggregation of data at various time points in the treatment process
 - Data integrity, completeness and validation check
- Deployment of the Enterprise Central QA Database
 - Aggregate data from all VA facilities.
 - Tools for data analysis, national benchmarking and analysis of variability within VHA







[IT Infrastructure] VA's Nationwide Intranet network VA - Richmond VA Intranet based QA Database **HINGE Software** RT-EMR / TMS Treatment related (Tx Delivery – Linac) **RO Database** Data Archival RO quality **Diagnostic imaging** RT Data assessment data (CT/MR/PET) API Data entry forms **RT-TPS** DICON (Tx Plan) Check for completeness Aggregate data Data Integrity Check Data uploaded / down securely **QA** Devices Define facility report card Encrypt data and statistics Compress Data ... **Display Statistics** Facility Report Card CPRS VA's Corporate Data Secure Web Warehouse Service Connection Data abstracted from disease site templates Abstraction of PRO data VA – East Orange ╉ 4 VA – Miami -VA – Long Beach VA's Firewall . . . **Patient Portal** Internet based **Radiation Oncology** server Patient Reported 0 Patient reported **Outcomes Module** /IARCH 18–21 | Hilton New Orleans Ri ,DB on Gateway and Exchange Applicat HINGE- Heat SPRING CLINICAL MEETING

Abstraction of Patient Specific Data elements for Practice Assessment



Disease Site Specific "Smart" Templates in Radiation Oncology

- Consensus clinical templates for all major disease sites treated with RT,
- Initial consultation, treatment planning, treatment, end of treatment, and follow up notes,
- Designed to prepopulate data from CPRS's patient chart and subsequent notes seamlessly.







Disease Site Specific "Smart" Templates in CPRS

		DSS	Oncology Suite			- 🗆 ×			
Suite Oncology Tem	plates Admin								
Consult Sir	m Directive	Pre-treatment Treat	ment Plan	eekly Treatment	End of Treatment	Follow Up			
	the Nata Des to		igy Templates			G.			
	ctive Note Pre-tre	eatment Note Treatment Pla			d Of Treament Note	Follow Up Note			
Notes			N	ote Detail					
	ViA Radiation Oncology Prostate Consult								
	TNM Sta Glesson Hirotog High Left Prostate	Score: Primary 文 Secondary 文 T sequent biopsies after initial diagnosis? Yes	Group Stage: Group Stage: Ist positive TRU No Date: Date:						

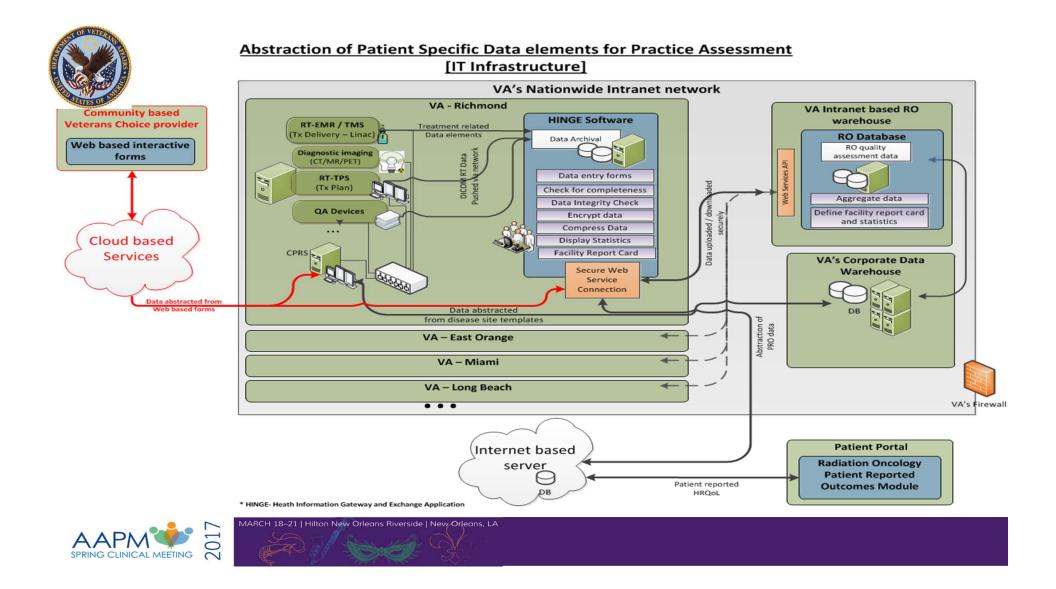
	PROSTATE TREATMENT PLAN NOTE
Uncology Prostate Consult_Rev2	
Prostate Cancer	Diagnosis: *Prostate Can
tegory: *C Low © Intermediate C High	
INM Stage:C T1 C NO MO	NCCN Risk Category: * C Low C Intermediate C High
GROUP Stage: *CICIICIII CIV *C a C b C c	STAGE: T1 NO MO
Gleason Score: Primary: 3	Gleason Score: Primary: 3
Total: 7 🕂	Secondary: +4
lst positive TRUS Biopsy Date: *Aug 10,2015 ···	Total: 7
Any subsequent biopsies after initial diagnosis? 🎷 No 🦵 Yes; date: 💴	1st positive TRUS Biopsy Date: *Aug 10,2015
Histology: *Adenocarcinoma	Any subsequent biopsies after initial diagnosis? *C Yes · No
<pre># cores positive/# cores sampled:</pre>	Non-Adendercinoma:
RIGHT 5 /9 ; LEFT 0 /0	
Prostate size on TRUS: 120 grams.	positive/# cores sampled: RIGHT 5 /9 ; LEFT 0 /0
Pre treatment PSA: 13.9 Date: Jul 13,2015	om Consult templatent PSA: 13.9
	repopulated
	FEMALE - NO DEA NECDED
✓ Diagnosis: *H€N Cancer Treatment pla	anning template
	DIAGNOSTIC TESTS REVIEWED:
TNM STAGE: C T1 C NO MO	ADDITIONAL FANCER HISTORY:
GROUP STAGE: *• I C II C III C IV *• a C b C c	Diagnosis: *H&N Cancer
Prognostic Factors:	
None	INM STAGE: II NO MO
Treatment(s):	GROUP STAGE: * CICIICINC V * a CbCc
Surgery:	
Z Radiation Therapy:	Prognostic Factors:
Anatomic Targets: Base of skull	None
RI Technique: Stereotactic	✓ Treatment(s):
Dose/Fxn: 25Gy/SFx	Surgery:
✓ Final Dose Delivered: ★C Yes C No	Chemo Completion:
Final RT Date: Oct 21,2014	Radiation Therapy:
ADDITIONAL CANCER HISTORY: Diagnosis: *Skin Cancer	
	Anatomic Targets: Base of skull
INM STAGE: c I2 c N2 M2	RT Technique: Stereotactic
GROUP STAGE: *CICIICIIICIV *CaCbCc	Dose/Fxn: 25Gy/5Fx
All None Indicates a Required Field Preview OK Cancel	Final Dose Delivered: *6 Yes 🕻 No
	Final RT Date: Oct 21 2014
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Work Flow Templates in CPRS

	VHA Radia	ation Oncology Prostate Consult -
Home	Import/Export Tools He	lp Options
Default	Page 2 Page 3	
✓ Page 4	Page 5 Page 6	Preview Finish
Page 7	Page 8 CNT Pages	
Diagnosis	CNT Pages	Template Functions
Diagnosis:		
NCCN Risk Category:	Low Interme	ediate 🥅 High
TNM Stage:		
		Group Stage:
Gleason Score: Prin	hary 🚖 Secondary	Total: 1st positive TRUS biopsy date:
Any subsequent biops	sies after initial diagnosis?	Yes No Date:
Histology:	-	
# cores positive /	# cores sampled:	
Right /		
Left /		
	grams	
Prostate size on TRU	S: Pre Treatment PSA:	Date: PSA Values:
Diagnostic Test Revie	ewed:	
🗖 🛛 Bone Scan		
CT Pelvis:		
Prostate MF	31:	
Any Prior Hormo	nes	Any Prior Radiotherapy
Prior Prostatecor	my Date:	🔽 Brachy Date:
Gleason Score:	\$	External Beam:
ECE:		Date:
🗖 SVI: 🗌		
Involved M	fargin	
VistA: Not Connec	ted USER: Not Signed In	PT: Not Signed In

Prostate End of Treatment Summary Note - 1	×
Home Tools Help Options	
✓ Default Page 2	
Page 3 Preview Finish	
CVII Pages C Tamplate Publicion	
Diagnosis Diagnosis: Age at the time of diagnosis and treatment:	
NCCN Risk Category: Low Intermediate High Gleason Score:	
TNM Stage: T N M Primary 🗘 Secondary 🗘 Total	-
1st positive TRUS biopsy date:	
Any subsequent biopsies after initial diagnosis? Yes No Date:	
Non-Adenocarcinoma	
# cores positive / # cores sampled:	
Right /	
Left /	
grams Prostate size on TRUS: Pre Treatment PSA: Date: PSA Values:	
Imaging Information	
Bone Scan:	
CT Pelvis:	
Prostate MRI:	
Any Prior Hormones Any Prior Radiotherapy	
Prior Prostatecomy	
VistA: Not Connected USER: Not Signed In PT: Not Signed In	





Summary

- Quality care is one of the dominant issues in health care today, especially in radiation oncology,
- Quality care data are most complex in radiation oncology but structured,

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- Quality of care is best assessed from the perspective of structure, process, and outcome measures.
- VHA is leading the nation in establishing an electronic infrastructure that will automatically abstract data from clinical workflow templates to assess the quality of radiotherapy and outcomes.

