Decision support for adaptive radiotherapy: in Vivo Dosimetry

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Patient may change during the treatment course



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At what point should we adapt

- What is understood as "adaptation"?
- When should the treatment be "replaned"?

At what point should we adapt

- What is understood as "adaptation"?
 - Any change made during the delivery so that the patient conforms to our patient model used for treatment planning.
 - Can it be adapting the position, can be adapting to motion (breath hold/gatting)
- When should the treatment be "replaned"?
 - When the differences cannot be compenasted by repositionning/table shifts and rotations, motion control strategies
 - When the immobilisation devices do not fit to the patient and lose their purpose
 - When the there can be "relevant" differences in the resulting dose distribution

Some points to consider

- Imaging can be used for real time treatment monitoring:
 - Patient position and tumor/organ motion.
- Adaptive radiotherapy: Adapting to tumor size and position.



Advances in dose delivery need more accurate patient positioning and monitoring



Beaton, L., Bandula, S., Gaze, M.N. *et al.* How rapid advances in imaging are defining the future of precision radiation oncology. *Br J Cancer* **120**, 779–790 (2019).

We are already adapting to changes with IGRT



We are already adapting to changes with IGRT



Adapt each fraction to the changing shape of the tumor



From Uulke van der Heide EMA 2024

Available data



Syntetic CT from CBCT



Deformable registration + structure set in the CBCT



Dose recalculation (original RT plan)



Comparison of actual delivered dose with the planned dose



Discussion with RO



There exist some commercial solutions

In vivo dosimetry as a suport tool for adaptive RT



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4 linacs







1 x Clinac 2100

3 x Truebeam





iVD for every patient: 1800 patients/year



TPS:



Secondary calculation and transit In-Vivo dosimetry: PerFraction (Sun Nuclear)

IOS:

ARIA

Transit in-Vivo Dosimetry: Clinical Use

Patient-specific quality assurance



IVD experience at Sant Pau Hospital





Patient platform

 Upload to Queue New P 	2171 Patient									SUN corp	NUC oratio	
Patient Name or ID O Pla	an Status ▼ Phase ▼	Treatment Site -	Machine •	Event Status -	Date Range -							
Patient List 0												Clear Filters and Sort
NAME		~ ID		PLAN		PRE-T	REATME	AD TH	IN-VIVO	MONITORING	REC	ENT ACTIVITY
	_			1PROST-V				\bigcirc	0	19 🚺 9	0	24 JUL. 2020
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Patient-specific quality assurance



PerFRACTION™ platform



Workflow



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EPID-based transit IVD



Limitations of EPID-based transit iVD:

- Large fields (size of EPID)
- Lateral fields (collision risk)
- Fields with couch rotation (collision risk)



checking the field size to avoid irradiating EPID electronics!!

Analysis

Analysis 1D Points	RT TX record , logfiles in all sessions	 Point Dose Abs. Dose Difference (cGy) 20 Enable Search Radius, using Distance (mm) From General 	✓ 2D Analysis
Analysis 2D Beams (2D)	EPID integrated images	General Editable templates Difference (%) Distance (mm) 5	Uses General Settings Baseline Predicted Dose Fraction 2 (17 FEBR. 2021 1:27) Fraction 12 (03 MARÇ 2021 11:05) Fraction 17 (10 MARÇ 2021 11:20) Fraction 19 (12 MARÇ 2021 10:57)
Analysis 3D Targets OAR	RT TX record Logfiles CBCT	Normalization Local	Image Source Calculated On Structure Tolerances MEAN D90% D95% MAX CBCT (17 FEBR. 2021 12:49) ▼ Image Source Image Source Image Source Image Source Image Source Plan CT CBCT (17 FEBR. 2021 12:49) Image Source Image Source Image Source Image Source When calculating on CBCT CBCT (17 FEBR. 2021 12:49) Image Source Image Source Image Source Image Source Image Source CBCT (17 FEBR. 2021 12:49) Image Source Image Source Image Source Image Source Image Source When calculating on CBCT Expanded Distance (cm) Image Source Image Source </td

2D analysis: transit IVD

<u>2D image analysis</u>: integrated image

🥑 Beams (2D)



Expected image:

- Relative: 1st fraction EPID image
- Absolute: predicted dose by the platform using RT plan and CT simulation

Predicted Dose

Fraction 2 (17 FEBR. 2021 1:27) Fraction 12 (03 MARÇ 2021 11:05) Fraction 17 (10 MARÇ 2021 11:20) Fraction 19 (12 MARÇ 2021 10:57)

2D analysis: transit IVD

<u>2D image analysis</u>: integrated image







Breast, extremities, palliatives

Tolerancia 90%

Breast or palliative

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3D analysis

<u>3D image analysis</u>: log-files + CBCT



In-Vivo and log-files based analysis



Flag alerts assesing



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Results: failed fractions

10% of daily sessions flag alerts

EPID-based iVD analysis

By technique

Counts/frequency: 3DCRT (150, 18.2%), IMRT (173, 21.0%), VMAT (431, 52.3%), SBRT (59, 7.2%), SRS (11, 1.3%)

By location

Counts/frequency: General unespecified (3, 0.4%), Brain (34, 4.1%), Head and Neck (264, 32.0%), Esophagus (3, 0.4%), Lung (84, 10.2%), Breast (98, 11.9%), Mediastinum (15, 1.8%), Gastrointestinal (19, 2.3%), Abdomen (67, 8.1%), Pelvis (81, 9.8%), Prostate (63, 7.6%), Genitourinary (4, 0.5%), GYN (0, 0.0%), Colorectal (2, 0.2%), Kidney (2, 0.2%), Extremities (61, 7.4%), Bone (24, 2.9%)





By treatment unit

ounts/frequency: Clinac 2 (153, 18.6%), TB1 (128, 15.5%), TB0 (322, 39.1%), TB3 (221, 26.8%)



Results: failed fractions

10% of daily sessions flag alerts

By location

Counts/frequency: General unespecified (3, 0.4%), Brain (34, 4.1%), Head and Neck (264, 32.0%), Esophagus (3, 0.4%), Lung (84, 10.2%), Breast (98, 11.9%), Mediastinum (15, 1.8%), Gastrointestinal (19, 2.3%), Abdomen (67, 8.1%), Pelvis (81, 9.8%), Prostate (63, 7.6%), Genitourinary (4, 0.5%), GYN (0, 0.0%), Colorectal (2, 0.2%), Kidney (2, 0.2%), Extremities (61, 7.4%), Bone (24, 2.9%)



Cause of alerts

Counts/frequency: Patient position (297, 36.0%), Anatomical changes body contour (121, 14.7%), Anatomical changes internal structures/tumor (97, 11.8%), Intrafraction motion (74, 9.0%), Positioning devices (26, 3.2%), EPID adcquisition (71, 8.6%), Preditcted dose calculation (29, 3.5%), MLC motion/position (9, 1.1%), Beam stability (1, 0.1%), Beam interrupted (88, 10.7%), Unknown (137, 16.6%), Other (40, 4.8%)



Results: failed fractions



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Clinical examples



Ex 1: Patient position



0 0 CTV 5280 CTV 5940 CTV_6996 0 86,56 % Gamma 99,90 % Gamma 99,91 % Gamma METRIC TPS QA Δ% METRIC TPS QA Δ% METRIC TPS QA Δ% 1,91 1,87 2.13 2,11 2.04 2.01 Mean -1,82 Mean -1,27 Mean -1,12 D95 1,60 1,51 -5,55 1,80 1,77 D95 D95 2,08 2,06 -1,00 -1,61



Cold Hot





Shoulders position

Ex 1: Patient position







Ex 2: Patient anatomy



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Ex 3: Patient position







Ex 3: Patient position change of practice

On line adaptive ??

IMPLEMENTATION OF IGRT



Conclusion

Real-time monitoring of patient surface and beam control during the whole treatment session accounts for higher in-vivo dosimetry gamma rates obtained using SGRT. This highlights an increase in treatment accuracy.

Ex 4: Intrafraction motion



Ex 5: Respiratory movement

Fraction 1 Approve

intrafraction motion: breath motion



Action : transit iVD next

												1-181-179-0	6 MV		87,47 %	61.325	7.102	584			
Beams (2	20)											- SID: 150 c	D	elivered		_	Ga	imma	SID: 150 cm	Expected	
BEAM NA	ME		ENERGY			PER	CENT	POINT	S F/	AILED HIGH	FAILE	ED L 180									
0 2-179	-181-0		6 MV				89,09 %	68	3.082	7.428		100 - 80 - 60 -		2						-	
1-181	-179-0		6 MV				87,47 %	61	1.325	7.102		40 - 20 - 0 - -20 -				_		2			
												-40 - -60 - -80 -	-2	1			4			4	
Targets												-120- -140- -160-		7				13			
CTV_504 61,59 % (0 Gamma	۲	CTV_4500 71,31 % G	amma	۲	CTVp_450 70,67 % Ga	0 amma	۲	CTVn_45 67,25 % 0	00 Gamma	۲	F Points Average	50 - 130 - 110 - 90 - 70 - 50 - 3	30 -10 10 30 50 3	0 90 110 130 150 170 0,43 0.49	190-190-170-150-130-110 Q Zoom	.90'.70'.50'.30'.1	0 10 30 50 70 90 110 130 150 17 0 2 Reset V OAbsolute D	o 190-170-150-130-11 ose(Gy) 0,00) -90 -70 -50 -30 -10 10 3r	0 50 70 90 110 130 150 170 190 0,33 0,33
Mean	1 98 2 07	Δ%	METRIC	194 201	2.33	METRIC	194 200	3 50	METRIC	1.98 2.06	Δ% 4 12	Mean	1.98 2.06	4 18	Mean	190 196	3.09	×	Gamma N/A <mark>0</mark>		1 21
D95	1,95 1,95	-0,10	D95	1,79 1,78	-0,50	D95	1,34 1,78	-0,55	D95	1,94 1,94	0,00	D95	1,92 1,92	-0,05	D95	1,76 1,75	-0,28				
OARs													A					0			
SpinalCo	ord	۲	Lungs		۲	Liver		0			1	-			A			a successful	1		

SpinalCo	ď		U	Lungs			0	Liver			
100,00 %	Gamma			99,23 % Ga	amma			99,89 % Ga	imma		
METRIC	TPS	QA	Δ%	METRIC	TPS	QA	Δ%	METRIC	TPS	QA	Δ9
Max	0,60	0,59	-1,48	Max	2,06	2,24	8,70	Мах	1,98	2,08	5,0





Ex 5: Respiratory movement

Approve 👻 r	espiratory motion. Demanat	notion. Demanat repetir en expiració forçada.								
ENERGY	PERCENT	POINTS	FAILED HIGH	FAILED LOW						
6 MV	68,70 %	63.694	19.939	0						
6 MV	71,90 %	69.161	19.436	0						
	Approve re ENERGY 6 MV 6 MV	Approve respiratory motion. Demanat ENERGY PERCENT 6 MV 68,70 % 6 MV 71,90 %	Approve respiratory motion. Demanat repetir en exp ENERGY PERCENT POINTS 6 MV 68,70 % 63.694 6 MV 71,90 % 69.161	Approve respiratory motion. Demanat repetir en expiració forçada. ENERGY PERCENT POINT S FAILED HIGH 6 MV 68,70 % 63.694 19.939 6 MV 71,90 % 69.161 19.436						

Action : plan with breath

 \rightarrow

Δ%



Targets

0 0 0 0 CTV 5040 CTV 4500 CTVp 4500 CTVn 4500 PTV 5040 PTV 4500 19,50 % Gamma 35,97 % Gamma 32,93 % Gamma 34,96 % Gamma 28,61 % Gamma 45,74 % Gamma METRIC TPS QA Δ% METRIC TPS QA Mean 1,98 2,25 13,17 Mean 1,94 2,16 11,04 Mean 1,94 2,16 11,74 Mean 1.98 2.19 1,98 2,21 Mean 1,90 2,08 9,48 10,26 Mean 11.85 D95 1,95 2,01 3,12 D95 1,79 1,84 2,79 D95 1,78 1,83 2,63 D95 1,94 1,94 0,15 D95 1,92 1,96 1,91 D95 1,76 1,79 1,70

OARs

Lungs I Liver I Samma			0	Heart 90,16 % Gamma			SpinalCord 🥑 100,00 % Gamma			Kidney_R			Kidney_L 100,00 % Gamma				
METRIC	TPS QA	Δ%	METRIC	TPS QA	Δ%	METRIC	TPS QA	۵%	METRIC	TPS QA	Δ%	METRIC	TPS QA	Δ%	METRIC	TPS QA	۵%
Мах	2,06 2,36	14,56	Max	1,98 2,35	18,69	Мах	2,04 <mark>2,38</mark>	16,99	Max	0,60 0,60	-0,16	Max	0,36 0,34	-3,86	Мах	1,42 1,42	-0,07



Pau

Ex 5: Respiratory movement

Following fractions...

Plan with breath control



Beams (2D)						
BEAM NAME		ENERGY	PERCENT	POINTS	FAILED HIGH	FAILED LOW
1-181-179-0	A	6 MV	94,65 %	61.088	2.650	618
2-179-181-0	▲	6 MV	97,76 %	67.612	1.515	0



Q Zoom 🔵

🕂 Pan 🕥

2 Reset

Absolute Dose(Gv

10 130 150 170 190 191

0,31 0,38

190,170,150,130

Points Average Points Standard Deviation

Targets

CTV_5040 📀		0	CTV_4500			0	CTVp_4500)	0	CTVn_4500	0				
99,06 % Ga	mma			99,43 % Ga	imma			99,42 % Ga	imma			98,92 % Ga	amma		
METRIC	TPS	QA	Δ%	METRIC	TPS	QA	Δ%	METRIC	TPS	QA	Δ%	METRIC	TPS	QA	Δ%
Mean	1,98	2,02	1,60	Mean	1,94	1,97	1,18	Mean	1,94	1,96	1,33	Mean	1,98	2,00	1,10
D95	1,95	1,96	0,30	D95	1,79	1,79	0,16	D95	1,78	1,78	0,11	D95	1,94	1,93	-0,56

BODY SpinalCord				Lungs				Liver							
99,80 % Ga	mma			100,00 % G	amma			99,63 % Ga	imma			99,94 % Ga	mma		
METRIC	TPS	QA	Δ%	METRIC	TPS	QA	Δ%	METRIC	TPS	QA	Δ%	METRIC	TPS	QA	۵%
Mean	0,28	0,28	0,35	Mean	0,37	0,37	-0,52	Mean	0,47	0,47	0,00	Mean	0,47	0,48	1,04
Мах	2.07	2.10	1.15	Мах	0.60	0.60	-0.65	Max	2.06	2.05	-0.67	Max	1.98	2.05	3.3



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Ex 6: Patient anatomy



Atelectasis



Conclusions



Take-home messages

We have the data to assess whether a patient would benefit from replanning/adapting

BUT, the information has to be processed to be appropriate for those assessments (deformable registration / structure deformation/ syntetic CT generator/ Dose calculation/ Dose accumulation)

Automation is needed to be able to detect patients that would benefit from replanning/adapting.

Take-home messages

In vivo dosimetry allows to flag treatments that could benefit from an adaptive strategy without adding any extra-dose due to CBCT

Our in vivo procedures are a safety network as we detect patient variations before the RO goes through off line review.

Experience with EPID-based in vivo verification of advanced treatments shows that medical physicists are much more involved in assuring the quality of the actual patient treatment than when only performing a pre-treatment dose verification measurement



In Memoria to Ben Mijnheer

He had a pivotal role in medical physics in Europe and a true pioneer in in vivo dosimetry and EPID-based IVD in particular