







## Acknowledgments

We would like to thank D-Wave Systems for providing access to their hardware and for computational assistance

In particular, Bill Macready and Mani Ranjbar

Jason Spaans

Tyler Paplham





- Cancer is one of the leading causes of morbidity and mortality worldwide
- In 2012 there were 14 million new cases of cancer
- There were 8.2 million cancer-related deaths
- Number of new cases expected to rise by 70% over next 2 decades (WHO)



#### Cancer in the European Union:

In 2012 there were 1.3 million cancerrelated deaths

This represented 25.8% of all deaths

29.2% of male deaths, 22.5% of female deaths due to cancer



#### Cancer in Germany:

In 2012 there were 222,000 cancer-related deaths

This represented 25.4% of all deaths

28.8% of male deaths, 22.4% of female deaths due to cancer

Most common types: breast, lung, prostate, colon, bladder



Cancer is treated using three methods:

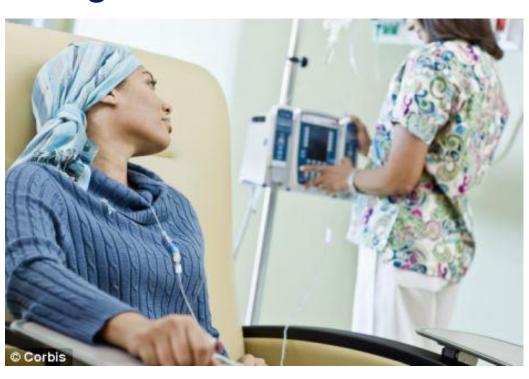
Surgery





#### Cancer is treated using three methods:

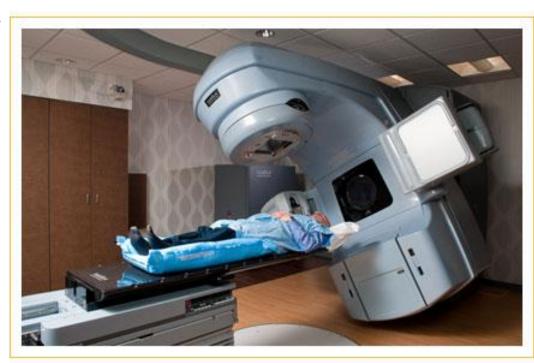
Chemotherapy





#### Cancer is treated using three methods:

Radiation Therapy

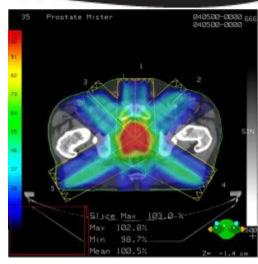


This is the subject of our work



#### **Dose Calculation**

Radiation dose distribution



Absorbed dose measured in Gy (J/kg)

Calculated from well-known physics principles

Clinical calculations use FDA-approved software



### CT Simulation



CT scan determines electron density of each voxel of patient anatomy

Allows dose calculation and anatomic structure identification (contouring)



# Medical Linear Accelerator



**Linear Accelerator Part 1** 



### Radiation Production



**Linear Accelerator Part 2** 

Energy ~ 6 MeV



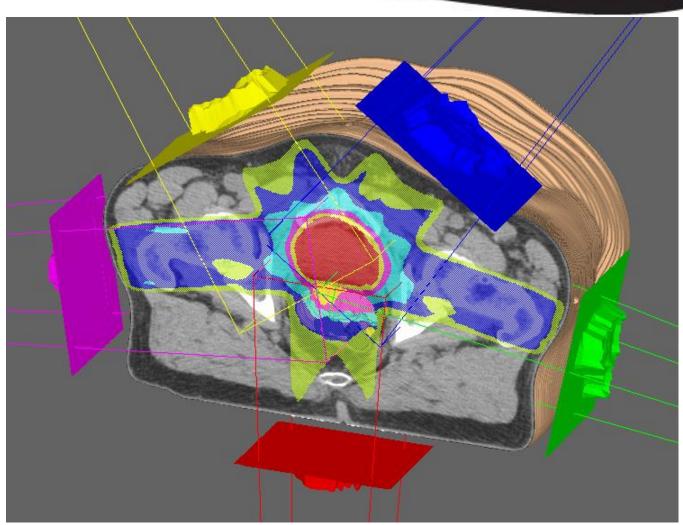
# Beam Shaping



**Linear Accelerator Part 3** 



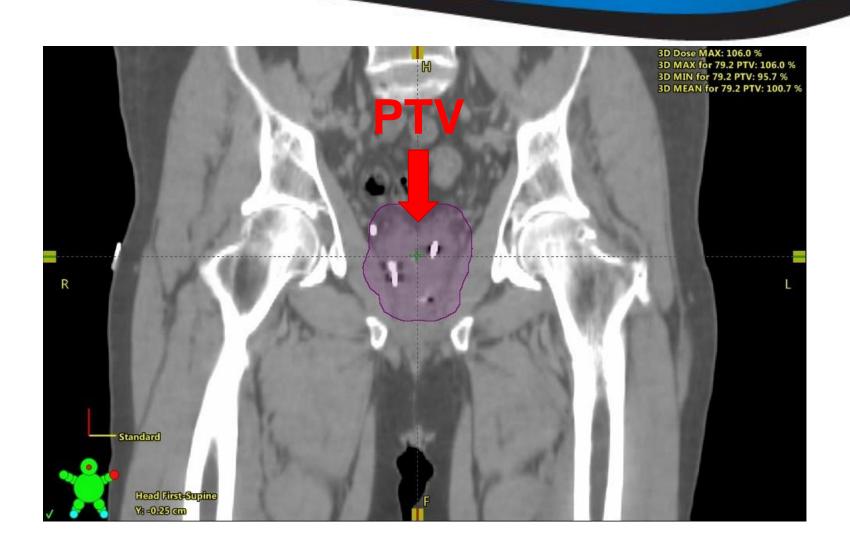
# **IMRT**



**IMRT** Treatment

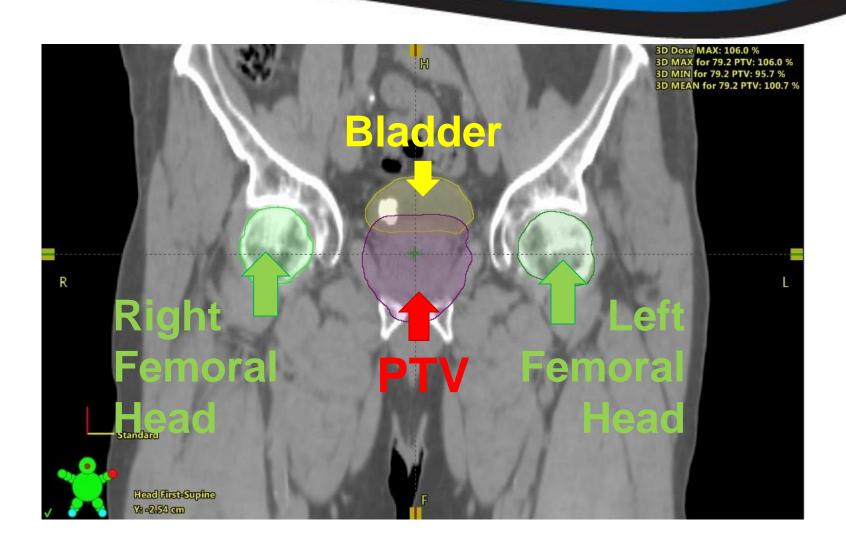


## The PTV



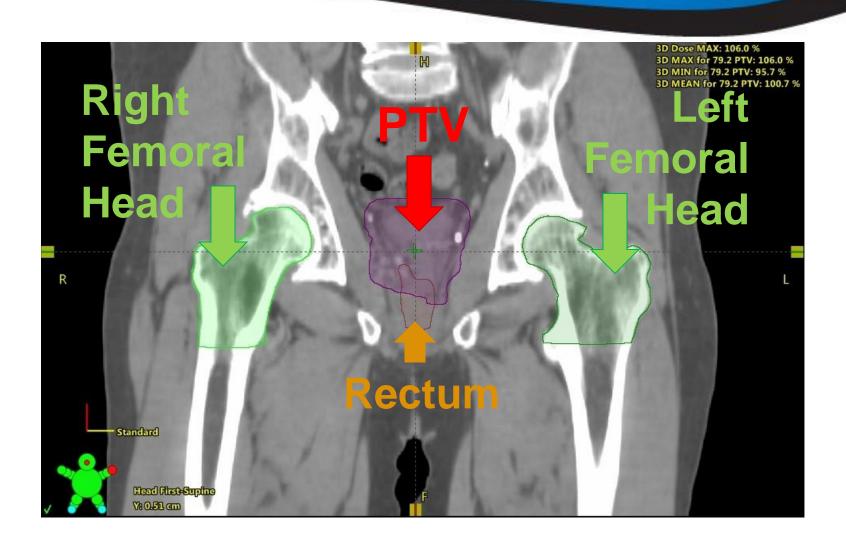


## **OARs Slice 1**



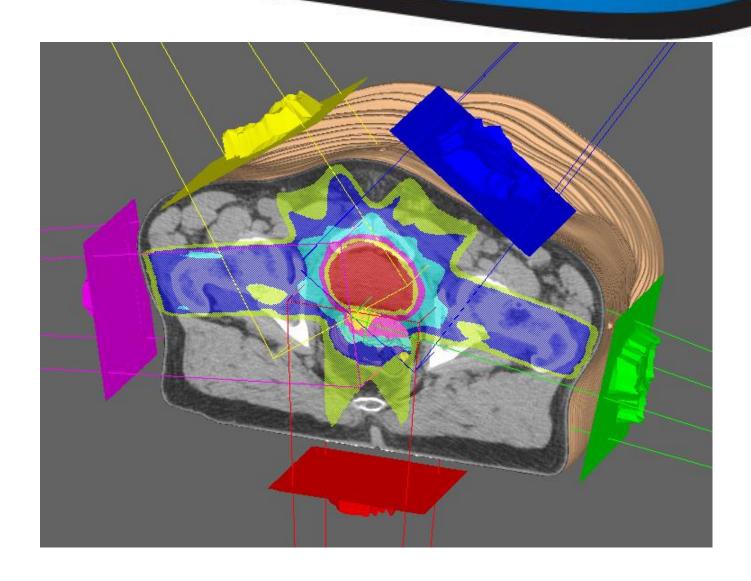


### OARs Slice 2





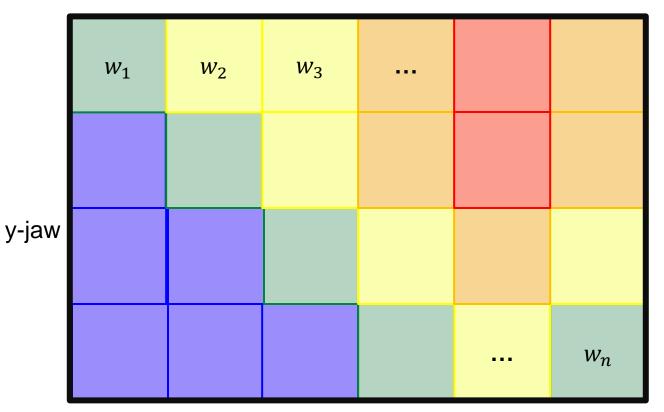
# **IMRT**





## Beamlets





y-jaw

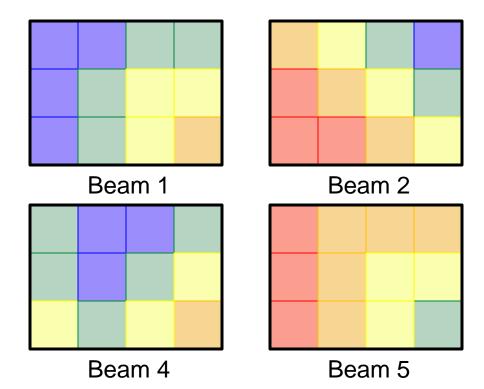
 $\mathbf{w}_{beam}$ 

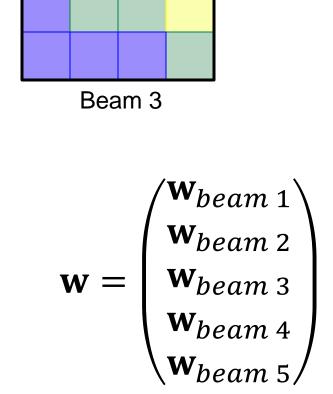
 $W_3$   $W_3$ 

x-jaw



### Beams





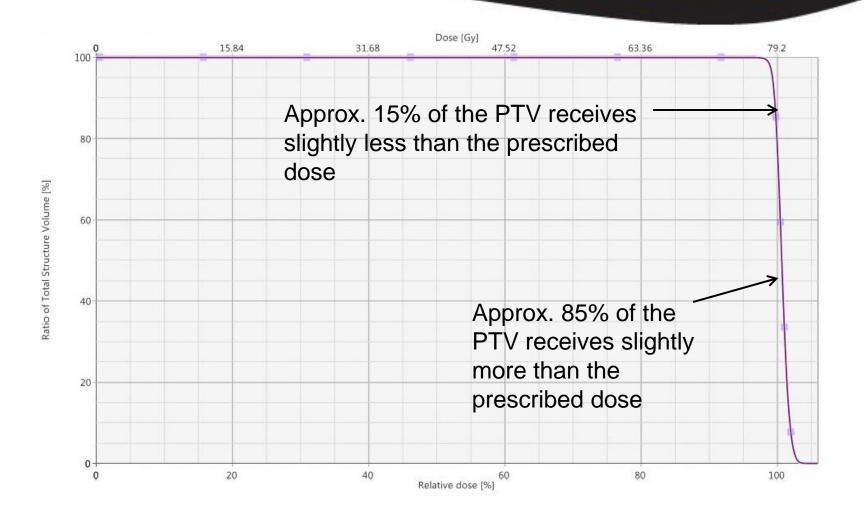


### The DVH

- A Dose-Volume Histogram (DVH) is a graphical representation of the percentage of dose received by a portion of the volume
- For a given treatment plan, the PTV and each organ has an associated DVH
- Is critical to defining the objective function

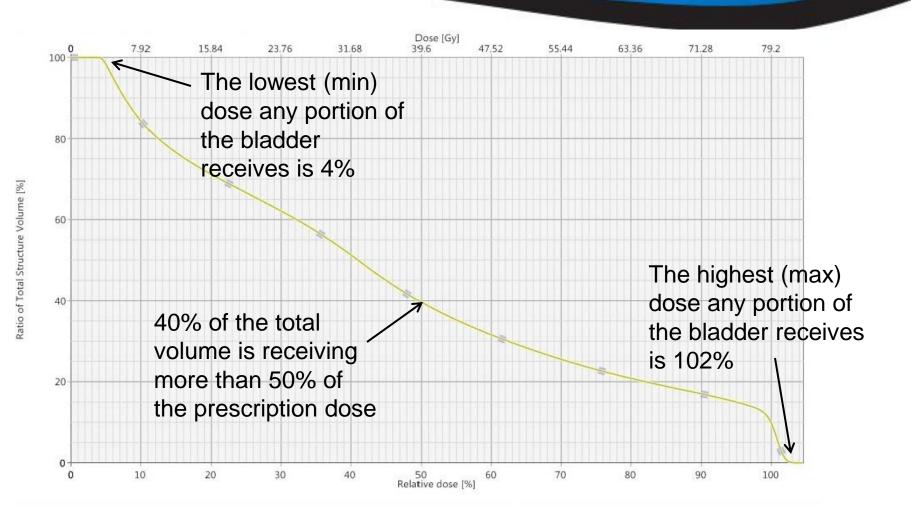


### PTV DVH



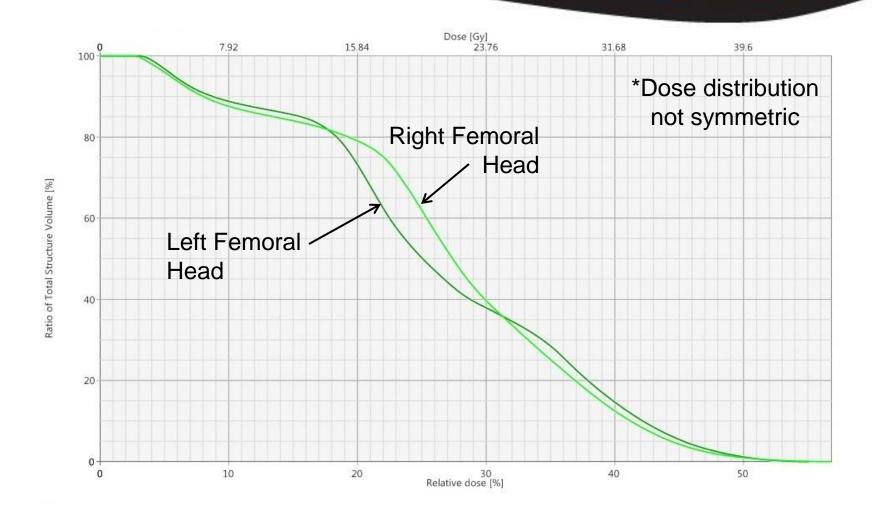


### Bladder DVH



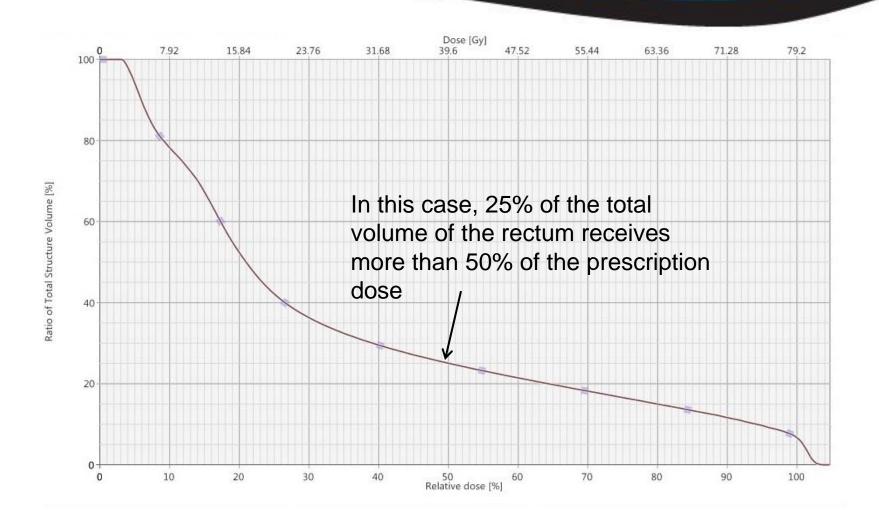


## Fem. Heads DVH





## Rectum DVH





# The Objective Function

$$F(\mathbf{w}) = \alpha (P_v - D_v(\mathbf{w}))^2 + \sum_i \sum_j \beta_i (\max[0, D_{ij}(\mathbf{w}) - C_{ij}])^2$$

 $\mathbf{w}$  is a vector of beamlet weights or intensities Minimizing  $F(\mathbf{w})$  results in optimal IMRT treatment plan



# The Target

$$F(\mathbf{w}) = \alpha (P_v - D_v(\mathbf{w}))^2 + \sum_i \sum_j \beta_i (\max[0, D_{ij}(\mathbf{w}) - C_{ij}])^2$$

 $\alpha$ =Priority of target dose (How important is it that this dose is fully administered?)

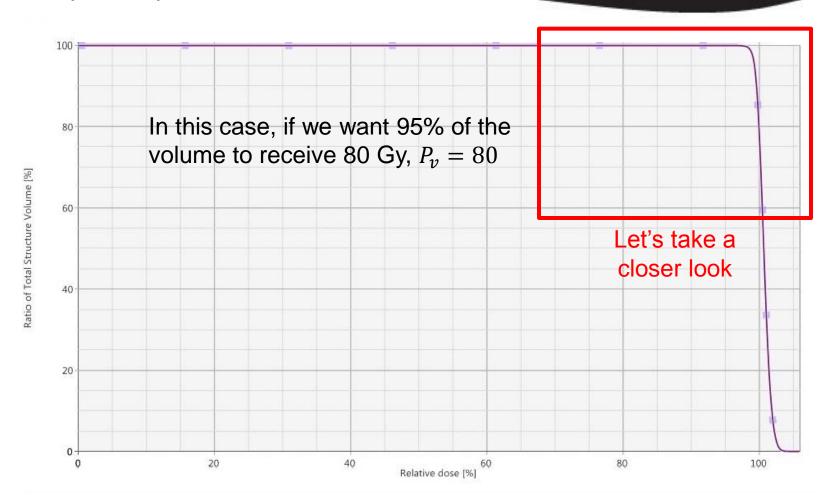
 $P_v$ =Dose prescribed to a given volume, v, of the target

 $D_v(\mathbf{w})$ =Dose actually received by volume v for weight vector  $\mathbf{w}$ 



## The Target

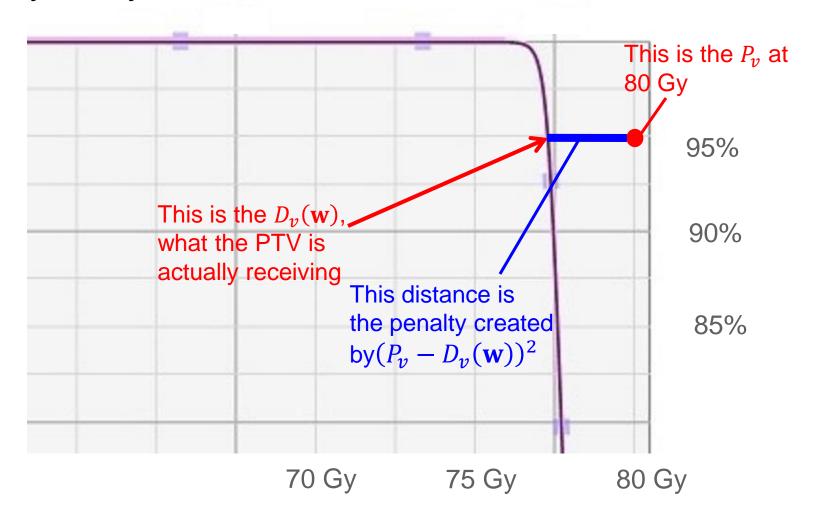
$$\alpha(P_{v}-D_{v}(\mathbf{w}))^{2}$$





# The Target

$$\alpha(P_{v}-D_{v}(\mathbf{w}))^{2}$$





### Penalties

$$\alpha(P_{v}-D_{v}(\mathbf{w}))^{2}$$

 In clinical terms, this ensures that the dose received by the target is as close as possible to the dose prescribed



# Organs at Risk (OARs)

$$F(\mathbf{w}) = \alpha (P_v - D_v(\mathbf{w}))^2 + \sum_i \sum_j \beta_i (\max[0, D_{ij}(\mathbf{w}) - C_{ij}])^2$$

 $\sum_{i}$  =Sum over each OAR, eg. bladder = 1

 $\sum_{i}$  =Sum over multiple objectives for a given OAR

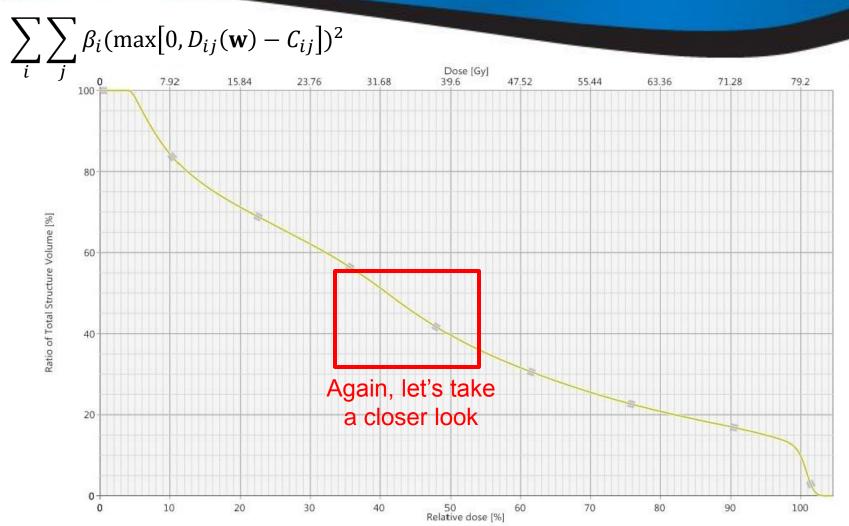
 $\beta_i$  =Priority of OAR

 $C_{i,i}$  =Objective dose

 $D_{ii}$  =Actual dose received by OAR

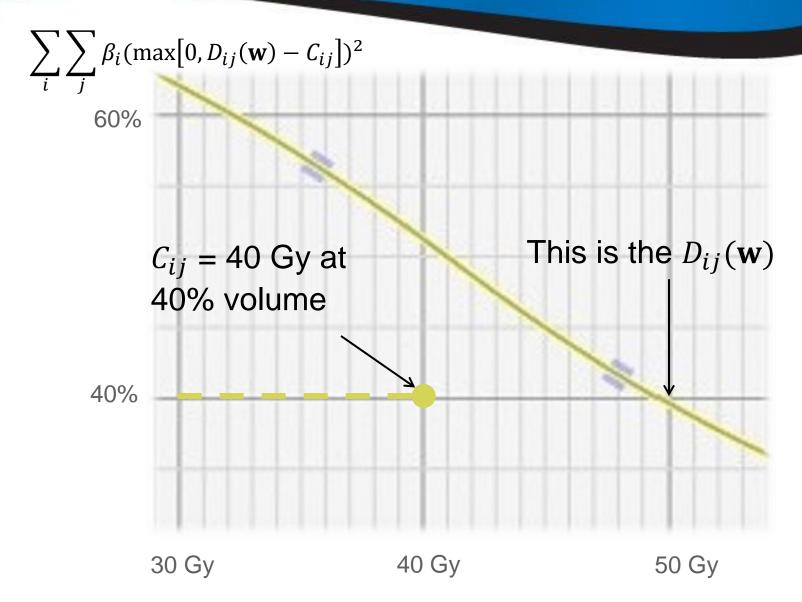


# Organs at Risk (OARs)



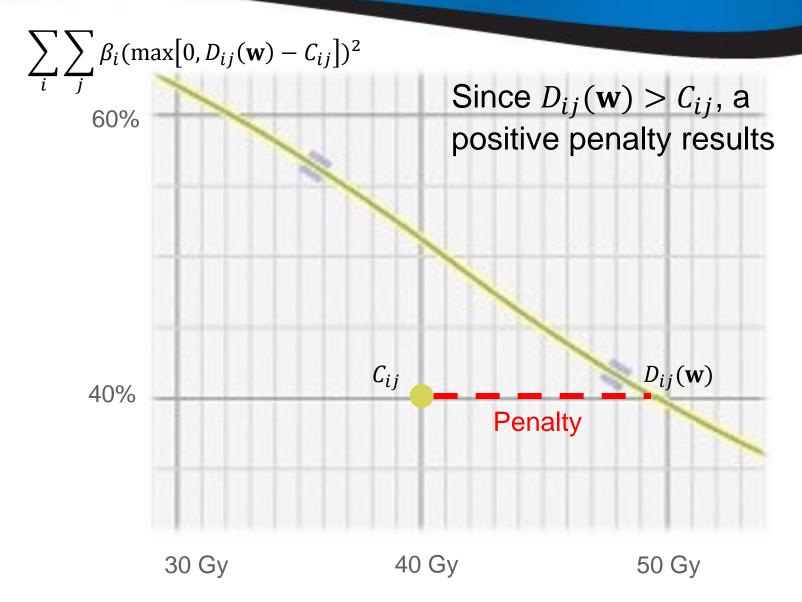


# Organs at Risk (OARs)





# Organs at Risk (OARs)





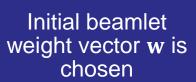
#### Penalties

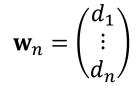
$$\sum_{i} \sum_{j} \beta_{i} (\max[0, D_{ij}(\mathbf{w}) - C_{ij}])^{2}$$

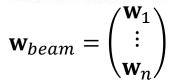
There is no reward for  $D_{ij}(\mathbf{w}) < C_{ij}$  because there is negligible clinical benefit to administering less than the objective dose to the OAR



#### The Process







Beamlet results in dose for each voxel



Beamlet dose matrices are added to create a beam dose matrix



w is modified using optimization algorithm



DVHs are created and F(w) calculated



Beam dose matrices are added to create total dose matrix

$$\mathbf{w}_{total} = \begin{pmatrix} \mathbf{w}_{beam \ 1} \\ \vdots \\ \mathbf{w}_{beam \ n} \end{pmatrix}$$



# D-Wave Systems

#### History of collaboration:

- Contacted D-Wave in 2009, put in touch with Bill
- Initially decided QA could not support
   IMRT optimization
- Visited lab in Burnaby in 2011 and revisited problem
- Worked remotely using Vesuvius chip and "Black Box" algorithm, 2012-2014



## Publication in 2015

IOP Publishing | Institute of Physics and Engineering in Medicine

Physics in Medicine & Biology

Phys. Med. Biol. 60 (2015) 4137-4148

doi:10.1088/0031-9155/60/10/4137

# First application of quantum annealing to IMRT beamlet intensity optimization

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#### Abstract

Optimization methods are critical to radiation therapy. A new technology, quantum annealing (QA), employs novel hardware and software techniques to address various discrete optimization problems in many fields. We report on the first application of quantum annealing to the process of beamlet intensity optimization for IMRT.

We apply recently-developed hardware which natively exploits quantum mechanical effects for improved optimization. The new algorithm, called



# RT Optimization

#### We defined IMRT objective function: $F(\mathbf{w})$

- w is vector of beamlet weights (intensities)
- F(w) incorporates target dose objective
- F(w) incorporates DVH objectives, with penalties if objective violated
- Objectives obtained from RTOG protocols



# Applying QA

- Vesuvius chip supported ~ 512 qubits
- Weight variables discretized to 7-digit binary variables
- Therefore, 70 beamlet weights (nonnegative, continuous) were included
- Actual clinical case would require 600-1000 beamlet weights



# SA Algorithm

#### Conventional simulated annealing (SA) features:

- Minimize function that is combo of original plus entropy
- Entropy is weighted by temp parameter T
- T is slowly reduced from large values (search space exploration) to 0 (solution)
- Can attain global minimum if cooling slow enough (but exponentially long)



#### **Evaluations**

#### Three methods compared:

- Quantum annealing
- Simulated annealing
- Tabu search: popular heuristic used in combinatorial optimization
- Methods were used to determine beamlet weights for two prostate bed cases
- Each was run for 10<sup>7</sup> function evaluations and compared for speed and score

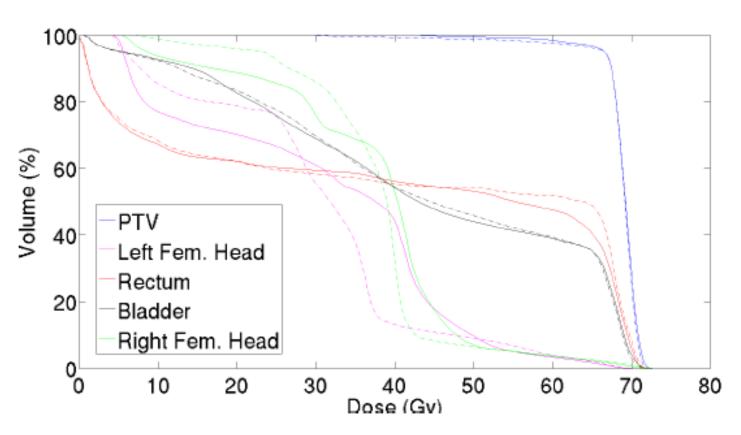


# Results

Patient	Method	Evals/sec /core	Final Score
1	QA	9.3	16.9
1	SA	9.6	6.7
1	Tabu	4.3	10.0

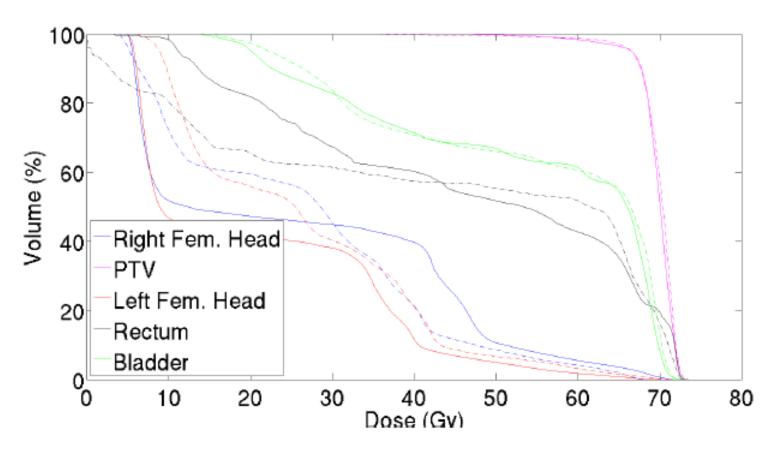
2	QA	15.4	70.7
2	SA	17.4	22.9
2	Tabu	6.3	120.0





QA (solid) and SA (dashed) for Patient 1

## **DVHs**



QA (solid) and Tabu (dashed) for Patient 2



# Wall Clock Time



<b>Patient</b>	Method	Time
1	QA	1.00
1	SA	2.89
1	Tabu	3.23

QA 1.00
SA 2.67
Tabu 3.67



# Results Summary

SA produced best score for both patients

QA was second, third

QA was fastest, by factors of 2.7 - 3.7

DVHs were compared and similar

Plans were not clinically viable due to small number of beamlets



# Future Work – VMAT



**VMAT** 

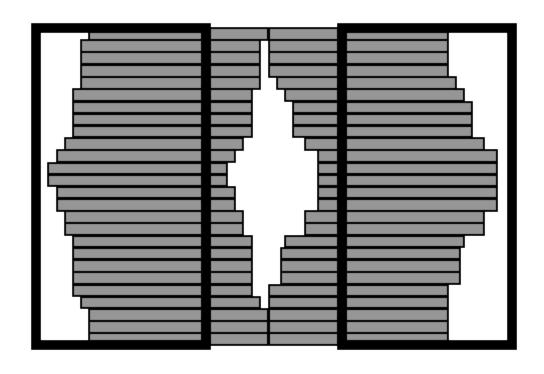




#### **VMAT Treatment**



# VMAT Optimization





#### Conclusions

This is first application of QA to IMRT optimization

Compared QA to SA and Tabu

Evaluated using clinical DVH-based objective functions

QA hardware will rapidly scale in size

Further research on application of QA to VMAT may offer promising returns



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### YouTube Embeds

Linear Accelerator

IMRT Treatment

VMAT Treatment