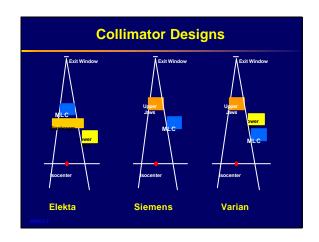
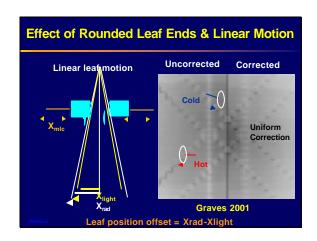


### Outline Acceptance testing Detectors for commissioning Phantoms Dosimetry analysis tools Commissioning tests Varying complexity and geometry Potential Pitfalls Summary

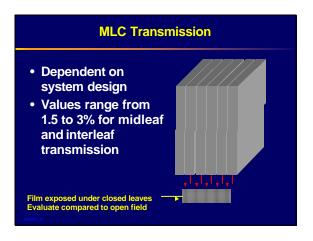
### Prior to purchase and installation, review manufacturer's acceptance tests If needed, adapt tests and tolerances with manufacturer in purchase order Test basic functionality of equipment Tests may be dependent on the MLC design

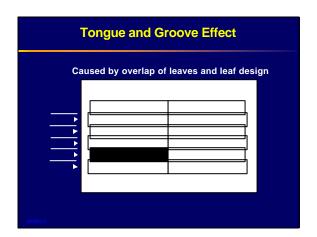


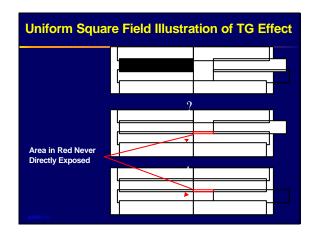
Physical MLC Characteristics					
	Elekta	Siemens	Varian		
Field Width	40x40 cm <sup>2</sup>	40x40 cm <sup>2</sup> (40x27 cm <sup>2</sup> )	40x40 cm <sup>2</sup>		
Leaf ends	Rounded	Divergent	Rounded		
Leaf width	1 cm	1 and 6.5 cm	0.5 and 1 cm		
Length	32.5 cm	31 cm	16 cm (14.5		
Inter-	No	No	cm Carriage) Yes		
digitation	=1 cm gap				

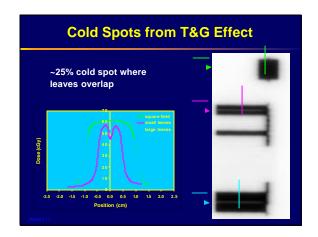












## DMLC and/or SMLC Mode Tests Leaf speed Dose rate evaluation Leaf position tolerance and reproducibility Leaf acceleration and deceleration Rounded leaf tip transmission Beam stability (output, flatness, symmetry, linearity) Interrupted treatments

# Multiple incorrect leaf positions All leaves move across field and deliver dose to small field gap (0.1 or 0.2 cm) Sensitive to errors of 1 mm in leaf position AMXXX13

### 

water phantom for IMRT delivery

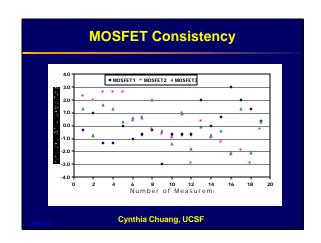
### 

### **Detector Characterization** Linearity **Energy dependence** Dose Stem and cable effects (cGy) Angular response 60 Calibrated if for absolute measurements 40 Small field detectors 30 required for small field characterization Sensitive to position Detector should be smaller than homogeneous region of dose to be measured

1-D Detector Characteristics					
Measurement Volume (cm <sup>3</sup> )	Sensitive Area (cm²)	Diameter (cm)	Thickness (cm)	Effective Point of Measurement (cm)	
0.009	0.24	0.6	NA	0.2	
0.3	0.49	0.4	0.06	0.6	
NA	0.011	0.45	0.006	0.07	
0.015	0.010	0.2	NA	0.06	
NA	0.04	NA	0.1	NA NA	
0.0019	0.056/0.073	0.73	0.026	0.1	
	0.009 0.3 NA 0.015	0.009 0.24  0.3 0.49  NA 0.011  0.015 0.010	(cm²) Area (cm²)  0.009	Volume (cm²)         Area (cm²)         (cm)         (cm)           0.009         0.24         0.6         NA           0.3         0.49         0.4         0.06           NA         0.011         0.45         0.006           0.015         0.010         0.2         NA           NA         0.04         NA         0.1	

1-D Detectors				
DETECTOR	DISADVANTAGES			
Micro-chamber	Poorer resolution than diodes			
Stereotactic diode	Over-respond to low energy photons Martens et al. 2000			
MOSFET  Diamond  JMM03 18	Non-linear dose response for <30 cGyChuang et al 2002 < resolution than diodes, expensive, Rustgi et al, Laub et al			

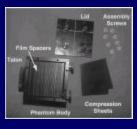
### **MOSFET System** Excellent spatial resolution Automatic and immediate readout · Can be re-used immediately · Linear dose response Response OSFET independent of depth C. Chuang, UCSF

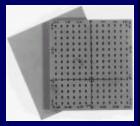


### **TLDs**

- TLDs must be characterized
  - Time consuming
- Reusable
- Achievable accuracy: 2-3%
- Automatic reader required for multiple **TLD** measurements

**TLD Holder for Phantom Measurements** 





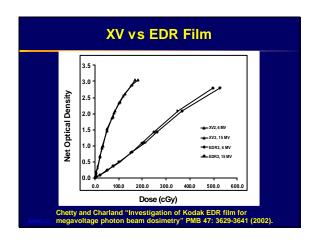
D.A. Low et al. "Phantoms for IMRT Dose Distribution Measurement and Treatment Verification, Int J Radiat Oncol Biol Phys 40: 1231-1235 (1998).

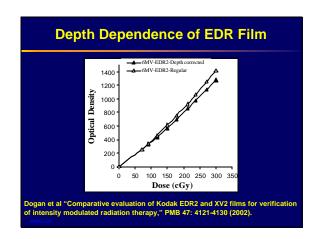
### **2-D Detectors**

- Film
  - Radiographic: XV and EDR
  - Radiochromic
- Beam imaging system, CCD, SLIC, **AMFPI** 
  - EPID systems attached to gantry
  - Investigated more for pre-treatment QA currently

### Radiographic Film

- Advantages
  - Excellent spatial resolution
  - Readily available
  - Less expensive than other 2-D systems
- Disadvantages
  - Over-response to low energies
  - Dependent on QA of film batch
  - Dependent on processor and digitizer QA
  - Sensitive to storage conditions
  - Need to measure the response to dose for each experiment





### **Radiochromic Film: Advantages**

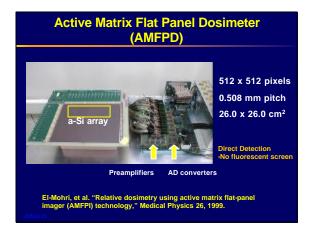
- Decreased sensitivity to low energy photons compared to radiographic film
- No processing
  - Film changes color with irradiation
- Insensitivity to visible light
- High spatial resolution

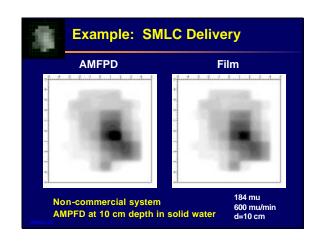
Niroomand-Rad et al. "Radiochromic film dosimetry: recommendations of AAPM Radiation Therapy Committee Task Group 55," Med Phys 25: 2093-2115.

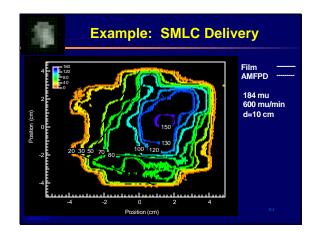
### **Radiochromic Film: Disadvantages**

- Non-uniform film response
  - Can be minimized by using double-exposure technique
- Response dependent on time and temperature
- · More expensive than radiographic film
- Digitizer response is dependent on the light source of the digitizer and may need to be modified

Niroomand-Rad et al. "Radiochromic film dosimetry: recommendations of AAPM Radiation Therapy Committee Task Group 55," Med Phys 25: 2093-2115.







### **Gel Dosimetry: Advantages**

- Obtain 3-D information in one irradiation
- Gels can be prepared with different density therefore ideal for heterogeneous measurements
- · Gel can be used in containers of different shapes
- Ideal for anthropomorphic phantoms

### **Gel Dosimetry: Disadvantages**

- · Sensitive to many factors such as time, preparation, temperature
- Optical reader requires cylindrical container for gel
- MR time is often limited and expensive (unless dedicated scanner)
  - Long scan times are required to increase accuracy of readout
  - E.g. 5% accuracy over 10 hr scan time (Gum et al
- Interface of gel and container results in less accurate readout at edges of the gel
- Not ready for routine use in the community

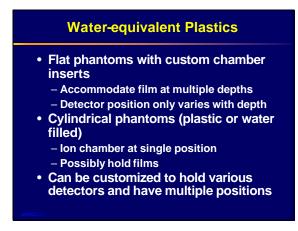
### **Gel Dosimetry** NR Dose Maps IMRT Plan Slice 1 Slice 2 Slice 3 0 20 40 60 80 100 120 leviation -60 -40 -20 0 30 40 60

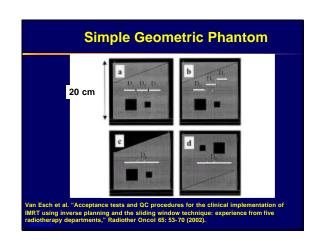
### **Considerations for Phantoms**

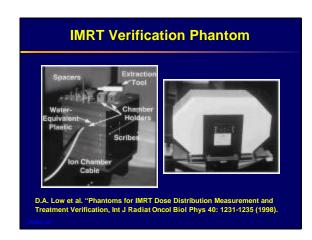
- · Fiducials for reproducible setup of phantom and detectors
- Flexibility to accommodate different detectors
- Simple vs. anthropomorphic
- Homogeneous or heterogeneous

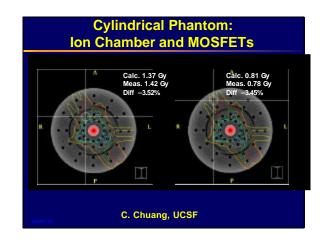
### **Water Tank**

- Restricted to gantry at 0 degrees - Unless mylar window for 90 degrees
- · Flexibility in chamber position
- Important for basic depth dose and profile measurements
- Output, flatness, symmetry, and linearity assessment

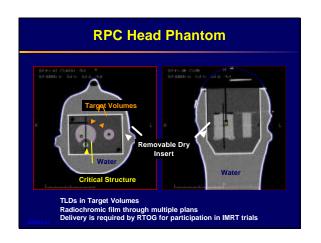


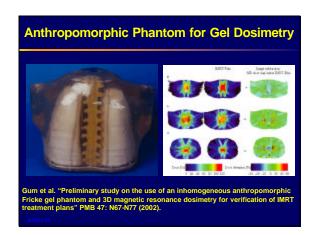


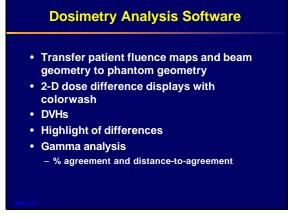


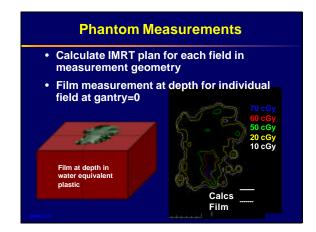


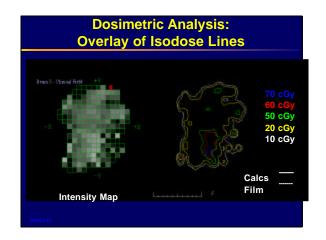


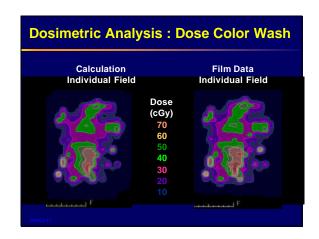


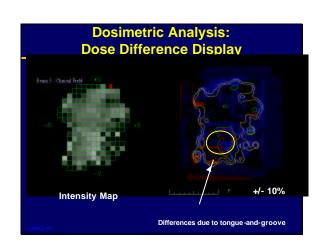


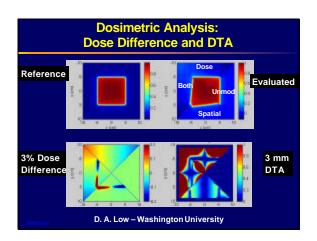


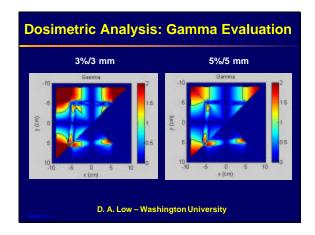












### **Simple Geometry Tests**

- Leaf position reproducibility in dynamic or step-and-shoot mode
- Effect of gravity on leaf position accuracy and reproducibility
- Sweeping gap test
- Fence test

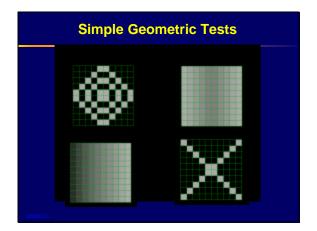
JMM03

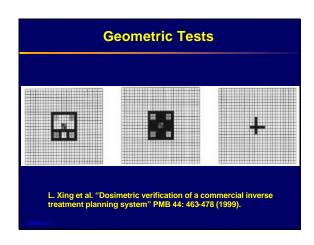
### **Simple Geometry Tests**

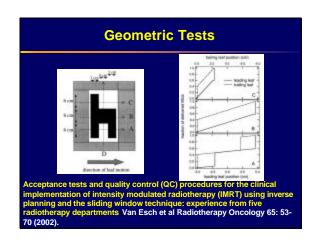
- Leaf speed stability
- Leaf acceleration/deceleration
- Output checks for small to large fields

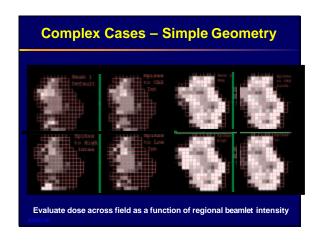
   Including smallest field size 1 x 1 cm<sup>2</sup>
- MLC limits (field size restrictions)
- Depth dose curve measurements

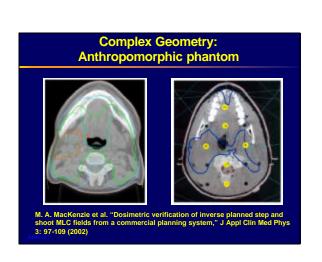
JMMC

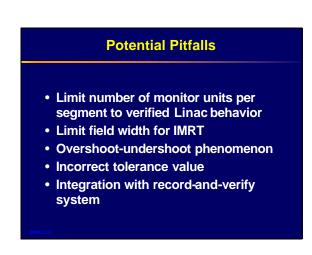




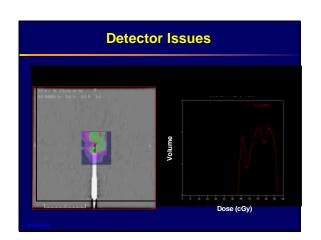




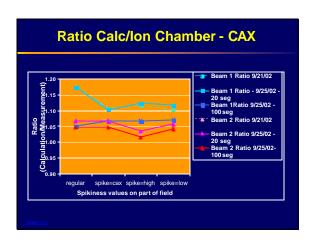


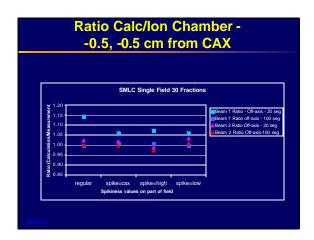


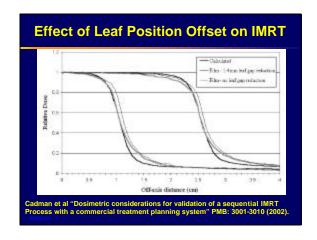
## Verify all equipment is functioning properly Film processor, digitizer Detectors, cables, electrometers (automatic leakage correction) TLD reader, ovens Input/output to treatment planning system Standardize measurement setup when possible Monitor software and hardware changes and QA

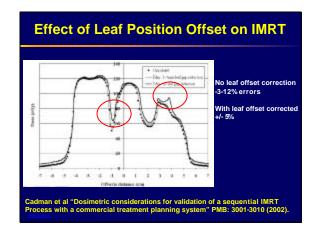


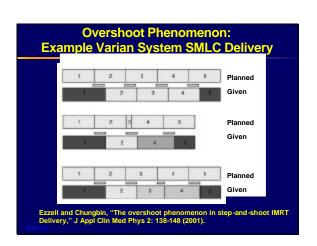
## Small ion chambers are very sensitive to position Position should be considered with respect to MLC design Example: CAX of Varian MLC is a junction of four 0.5 cm wide leaves

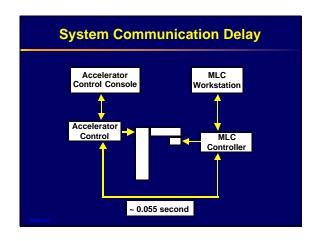






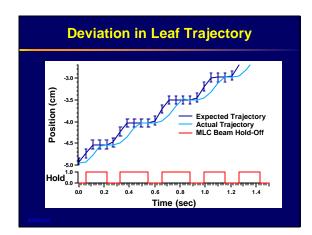


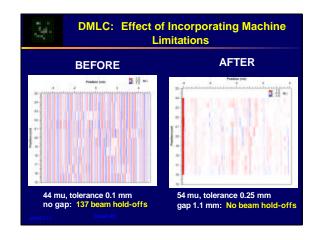




### Varian 21 EX Expected and actual leaf positions Beam hold-off events Position tolerance setting Recorded approximately every 55 ms Information can be compared to imported

2-D information





### **Summary**

- Basic MLC characteristics should be tested in static mode prior to IMRT testing
- IMRT tests should be specific to delivery mode and device
- Be aware of potential issues with delivery systems that may need further investigation
- Multiple detectors and phantoms are typically required for IMRT commissioning
- Quantitative dose analysis tools are required for proper evaluation of delivery

### **Summary**

- Measurements may show dosimetric mechanical differences that planning systems may not model at this time
- Need to know the limits of the mechanical systems and combination of software + delivery
- Continued need to improve software for delivery system, measurement devices, phantoms, and dose analysis tools

### Acknowledgements

University of Michigan Dale Litzenberg Jeff Radawski Tim Nurushev Benedick Fraass

Dan Low – Washington University Cynthia Chuang - UCSF