Clinical and Experimental Radiobiology Course

Tutorial 1

Wi-Fi

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Piazza

https://piazza.com/utoronto.ca/ winter2025/mbp1301h



Temerty Medicine

Tutorial 1

- Lecture 2: Hallmarks of Cancer
 - Dr. Marianne Koritzinsky
- Lecture 3: Molecular Basis of Cell Death
 - Dr. Marianne Koritzinsky

- Lecture 4: Radiation Induced Damage & DNA Damage Response
 - Dr. Shane Harding



Tutorials: If you want CPD credits:

	Jane Smith
<u>T1</u> A ∜ C ∜ F ∜ D ∜	100%
<u>T2</u> B ♥ B ♥ B ♥ B ♥	75%

End of course: scan/photo and send to radiation.oncology@utoronto.ca





Lecture 2: Hallmarks

A genetic alteration is considered "driving" if...

- A. It occurs frequently in tumors
- B. It is somatic
- C. It represents loss of function of a tumor suppressor
- D. All of the above
- E. None of the above





Lecture 2: Hallmarks

A cancer cell typically contains approximately this number of driving mutations:

- A. 1
- B. 5
- C. 100
- D. 1000





Lecture 2: Hallmarks

Genetic alterations in cancer can influence tumor radiation response by:

- A. Disrupting DNA repair
- B. Promoting high proliferation
- C. Promoting hypoxia adaption
- D. All of the above





Lecture 3: Cell Death

What is the most common cause of cell death following irradiation?

- A. Apoptosis
- B. Autophagy
- C. Senescence
- D. Mitotic catastrophe
- E. Necrosis





Lecture 3: Cell Death

Following radiation exposure, a surviving cell that has retained reproductive integrity is said to be:

- A. Radioresistant
- B. Genomically unstable
- C. Very lucky
- D. Clonogenic





Lecture 3: Cell Death

Apoptosis is likely to influence clonogenic survival when:

- A. It occurs before the first cell division
- B. It occurs following multiple cell divisions
- C. It is characterized by membrane blebbing
- D. It is a consequence of p53 activation



From the standpoint of DNA repair, what type of insult is most lethal to cells:

- A. Low-LET
- B. Ultraviolet light
- C. Protein misfolding
- D. High LET



Which of the following genetic defects is most likely to lead to severe toxicity during radiotherapy?

- A. Mutations in the ATM gene
- B. Mutations in the p53 gene
- C. Mutations in the BRCA2 gene
- D. Mutations in a DNA glycosylase
- E. All of the above



If you could block any DNA pathway in combination with radiation, what would you target?

What limitations do you have to consider?



Reference: L4 slides 5, 37, 39, 41

What do cell cycle checkpoints accomplish:

- A. Preventing cells from mitotic entry with damage
- B. Providing time for repair
- C. Increasing radioresistance
- D. Reducing replication errors
- E. All of the above



