

Dear HOPP SEP students,

Please read and answer the questionnaire before the class. If you have any questions about them, we will discuss them during the class.

- 1) As part of your project you are interested in modelling mutations that frequently occur in cancer patients. For this purpose, you are reviewing MSK-IMPACT clinical sequencing results using the cBioPortal platform*. One of the genes that most commonly is found to be mutated in this dataset is TP53, a known tumor suppressor gene. What are the characteristics of these types of genes?
 - a. In cancer patients, inactivating mutations are the most frequent type of mutations occurring in tumor suppressors genes.
 - b. Loss of function in tumor suppressors genes cannot occur due to non-mutational processes like promoter methylation.
 - c. Loss of tumor suppressors decreases the likelihood of malignant transformation.
 - d. Mutation in tumor suppressors genes can only be somatic, and they don't occur as germline mutations.

- 2) After acknowledging the importance of TP53 as a tumor suppressor, you are interested in understanding the effect of other tumor suppressors. Which of the following genes is not a tumor suppressor?
 - a. RB, a gene involved in controlling the pass through a cell cycle checkpoint.
 - b. APC, a gene involved in downregulating the expression of beta-catenin, a known proto-oncogene.
 - c. BRCA, a gene responsible for the reparation of DNA.
 - d. RAS, a gene involved in promoting cell growth.

Describe the differences between:

- DNA mutation vs DNA methylation:

- Gene promoter vs gene exon:

- Somatic testing vs Germline testing:

- Oncogene vs Tumor suppressor gene:

*To explore the frequency of tumor mutations using the publicly available sequencing data from MSK-IMPACT, go to cBioPortal (MSK, Nat Med 2017):
https://www.cbioportal.org/study/summary?id=msk_impact_2017