FOB exam 3 Cheat Sheet by NoelleEvelyn via cheatography.com/168075/cs/45854/

Nucleotide

excision repair

Cheatography

| Phases of the cell cycle Mitosis | | |
|----------------------------------|--|--|
| Prophase | Chromosomes condense and spindle aparatus forms | |
| Promet- aphase | Kinetochores assembled at centromere, 2 opposite sides connected to microtubles | |
| Metaphase | Lined up on imaginary metaphase plate. Polar microt- ubles extend from each spindle, overlap in middle, pole-pole connection | |
| Anaphase | Cohesions are cleaved, daughters to opposite sides of cell. poles pulled apart | |
| Telophase | Nuclear envelope reforms, chromosomes begin to condence | |
| Cytoki- nesis | Division of cytoplasm | |

When cells divide, two gentetically identical sis

| middle, | | Helicase unwinds and removes region with damaged bases |
|---------------|----------|--|
| site sides of | | 4. DNA polymerase fills gap with undamadged strand as template |
| begin to | | 5. Nuleotide linkage (DNA ligase links the strand into esisting strand. |
| | | If sucessful continues past G1 checkpoint |
| | P53 gene | Creates CDK inhibitors if the cell is damaged so if cyclin is still present, CDK can still say no if damaged |
| ter cells are | UVRA | recgonizes DNA damage, signals to start repair, if damage cant be repaired cell wont divide anymore. |
| | recA | Facilitates DNA repair |

Mechanisms of cell cycle progression

1. Error detected in DNA by proteins

2. DNA nicking (cut at both sides of damage)

DNA synthesis in Leading strand

| Synthesized | Continously |
|------------------|----------------|
| Begins with | RNA primer |
| After RNA primer | DNA polymerase |

| DNA synthesis in lagging strand | | | |
|---------------------------------|---|--|--|
| Synthesized | in fragments (Okazaki fragments) | | |
| Initiated by | RNA polymerase | | |
| RNA polymerase | builds primers | | |
| DNA polymerase | replicates DNA off of primers | | |
| RNA primer | popped out of gaps and replaced with DNA polymerase | | |

4. Cells undamaged

G2 checkpoint

G1 checkpoint

Mitosis Mitosis

Uses

1.

2.

3.

1. No errors in replication

their products

Somatic cells

Cells big enough

Sufficient nutrients

social signals present

- 2. Activated MPF (cyclin + CDK) present
- 3. Undamaged

Metaphase checkpoint

- 1. Chromosomes attatch to spindles
- 2. Chromosomes properly segregated
- 3. MPF absent



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| Both leading and Lagging strands | | |
|---|--|--|
| Single stranded binding proteins (SSBs) | Keep stands from attatching back together | |
| Ligase | Fills in gaps or breaks in phosphodi- ester bonds of backbone | |
| Helicase | Seperares, unwinds double stranded DNA | |
| Topoisomerase | Helps with stress on wound DNA, ex. Gyrase | |

Importance of Telomeres

| Protect from | important DNA being cut out |
|---------------------------|--|
| Everytime cell divides | become shorter |
| Replication limit | prevents cancer |
| Why? | There is no 3' hydroxyl at end of lagging stand. |
| What? | G-rich series of repeats |
| Telomerase | elongates parental in 3' to 5' direction. |



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