Learning Objectives

- Define teratogenesis, mutagenesis, and carcinogenesis.
- Describe the relevance of replication, transcription, and translation to teratogenesis, mutagenesis, and carcinogenesis.
- Summarize the mechanism of action for teratogenesis, mutagenesis, and carcinogenesis.
- Discuss examples of known teratogens, mutagens, and carcinogens.

Molecules of Life

- Toxicants can react with or modify DNA or RNA.
- Replication → perpetuate genetic information.
- Transcription and translation → express genetic information.

The Cell

- Replication
- Transcription
- Translation

Protein Functions

- Antibodies.
- Receptors.
- Enzymes
- Neurotransmitters, hormones
- Channels and pores.

Endpoints

- Teratogenesis.
- Mutagenesis.
- Carcinogenesis.
### DNA Replication
- Structure implies replication
- Occurs via multiple enzyme action
- Helix unravels, strands part, DNA replicates
- Mitosis, meiosis
- Not always perfect
  - Repair enzymes

### Replication
- Duplicates cell DNA.
- Mitosis - one somatic cell with 2n chromosomes divides to create two cells with 2n chromosomes (humans, n = 23).
- Number, quality and quantity of chromosomes per cell is conserved.
  - Triggers for mitosis (receptors + proteins).
    - External signals.
    - Hormones.
    - Internal factors.
    - Growth factors.

### Replication, 2
- Meiosis - germ cells are cells that divide into gametes.
  - 2 cell divisions.
  - Four daughter cells.
    - Each with a different set of chromosomes.
    - Each with 1 set that will be joined by another in fertilization.

### DNA Transcription
- DNA is copied via expendable mRNA
- mRNA codes for specific proteins
- Occurs in nucleus of cell

### DNA Translation
- Occurs in the cytosol
- Interaction of mRNA, tRNA, amino acids and enzymes
- tRNA has three-base codons which correspond to different aa
- AA are added one at a time to form chain - polypeptide
- Polypeptide corresponds to protein with a specific aa sequence

### Transcription and Translation
- DNA: double strand of nucleotides.
  - Nucleotide = nucleic acid, sugar and phosphate.
  - Cytosine, Thymine, Uracil; Adenine, Guanine.
  - Base pairing = A-T, G-C.
  - Gene: sequence of bases that code for a specific sequence of amino acids (protein).
  - Codon: sequence of 3 bases that code for a single amino acid, i.e.
    - AGC → Serine.
    - AAA → Lysine.
Transcription and Translation, 2

- Transcription = copying.
  - DNA unzips and enzymes make RNA "copy".
  - Differences:
    • T → U (UA not TA).
    • Deoxyribose → ribose.
  - mRNA formation; transport to cytoplasm.
- Translation = protein formation.
  - mRNA (blueprint).
  - rRNA (support).
  - tRNA (a.a. transport).

DNA Structure - Function

- Nucleotides form chains
- 3 nucleotides form a codon
- Multiple codons form genes
- Multiple genes form chromosomes
- Multiple chromosomes form DNA

DNA/RNA Complex

DNA/RNA Complex, 2

Errors in DNA Replication, Transcription and Translation

- Base pairing
- Repair enzymes and other enzymes
- Regulatory genes, operons, termination sequences
- Methylation patterns
- Post transcriptional/translation processing

DNA/Chemical Interactions

- Alkylation - covalent adduct between DNA and chemical
- Intercalation - noncovalent binding of chemical between two adjacent base pairs
- Cross-linkage - Inter or intrastrand covalent binding of chemical
- Breakage - scission of one or both strands of DNA
Principles of Environmental Toxicology

Aflatoxin B1 – DNA Adduct

Principles of Environmental Toxicology

Teratogenesis

- Teratology: the study of the frequency, causation, and development of congenital malformations.
- Complex mechanisms and timing of disruptive interaction during embryogenesis.
- Some natural “bad path” spontaneous abortion.
  - Humans: critical 1st 8 wks gestation.
    • Embryonic stage.
      • Morphological defects in specialized tissues and organs.
  - Fetal stage exposure.
    • Developmental or neoplastic endpoints.
- Known human teratogens.

Principles of Environmental Toxicology

Example: Teratogenesis

Five Legged Frog

Principles of Environmental Toxicology

Example: Teratogenesis, 2

Ovine Cyclopia
Veratrum Californicum
W. False Helebore

Principles of Environmental Toxicology

Example: Teratogenesis, 3

Crooked Calf Disease
Lupine Quinolizidine alkaloids
Anagyrine

Principles of Environmental Toxicology

Case Study: Lupine Alkaloid Birth Defects

- In September 1980, a baby boy born in the mountainous back-country of northwestern California (Trinity County) was brought to the UC Medical Center in Sacramento with severe bone deformities in his arms and hands, including a partial absence of forearm bones (radial aplasia) and absent thumbs.
- Extensive medical histories and genetic analyses of his parents indicated that the probable cause was environmental rather than hereditary.
- His mother feared that somehow exposure to herbicide spraying was responsible. Association of forest spraying and a reportedly high incidence of birth defects in northwestern California and southern Oregon has been highly publicized in recent years and has become controversial. Indeed, it appears likely that this herbicide had been applied to a forested ridge several miles distant from the mother’s home more than a year before the child’s conception.
Case Study: Lupine Alkaloid Birth Defects

- The mother provided the evidence that her goats also gave birth to kids stillborn or with deformed legs during and after the period of her pregnancy, and that puppies born to a dog fed the goat's milk during pregnancy were likewise deformed.
- Local goat's milk has become a common food item in the area, and the child's mother drank it regularly herself throughout pregnancy.
- A thorough survey of nearby areas where the goats had regularly browsed at the time of the mother's early pregnancy showed that a perennial lupine, identified as the widely distributed Lupinus latifolius, often formed the principal low-growing forage as well as wild tobacco (Nicotiana), poison hemlock (Conium), and skunk cabbage (Veratrum).
- Circumstantial evidence.

Mutagenesis

- Somatic cell mutations → metabolic dysfunction; carcinogenesis.
- Germ cell mutation → heritable change.
- Point mutation.
  - Base substitution (including analogues).
  - Frame shift.
- Chromosomal aberration.
  - Structural anomaly.
  - Numerical anomaly.

Karotypes

Patterns photographed during metaphase help examine for chromosomal defects.

Ames Test for Chemical Mutagenicity

- Salmonella bacteria strain with histidine coding defect.
- Mutagenic chemicals can change the defect to allow cell division and growth.
- Add salmonella + test chemical + rat hepatocytes (for biotransformation).
  - Growth indicates mutagenic effect.

Carcinogenesis

Multi-step, multi-factorial disease
### Cancer Definitions

- **Cancer.**
  - A malignant tumor that has the ability to metastasize or invade into surrounding tissues.

- **Tumor (Neoplasm).**
  - A general term for the uncontrolled growth of cells that becomes progressively worse with time.

- **Neoplasia.**
  - The growth of new tissue with abnormal and unregulated cell proliferation.

### Cancer Definitions, 2

- **Benign tumor.**
  - A tumor that does not metastasize.

- **Metastasis.**
  - Ability to establish secondary tumor growth at a new location.

### Cancer Definitions, 3

- **Carcinoma.**
  - Malignant tumor arising in the epithelium.
  - Most common form of cancer.
  - Usually spreads in the lymphatic system.

- **Sarcoma.**
  - Malignant tumor in muscle or connective tissue.
  - Usually spread in the blood stream.
  - Frequently metastasizes to the lung.

### Multistage Carcinogenesis: Initiation

- Chemical-virus-spontaneous causes DNA lesion
- Cell division perpetuates DNA lesion
- No outcome if not promoted
  - Some chemicals can initiate and promote
  - May remain indefinitely if not promoted
- One hit
  - No threshold; irreversible

### Properties of Initiated Cells

- No phenotypic differences
- Excess/deficiency of enzymes
  - e.g. δ-GT, G-6-P, Fe exclusion, ATPase
- Resistance to cytotoxic chemicals
  - Faster or slower metabolism
- Impaired cellular communication
- Enhanced response to growth factors
- Resistance to terminal differentiation

### Multistage Carcinogenesis: Promotion

- Change in micro-environment of cells
- Chemical, viral, spontaneous-induced clonal proliferation of initiated cells
- Growth control factors; receptors; immune function; endocrine control; communication; metabolic; apoptosis
- Multi-hit, high dose
  - Reversible
  - Threshold
**Multistage Carcinogenesis: Progression**

- Complete loss of growth control
- Karyotype instability
- Loss/gain of chromosomal fragments
- DNA demethylation/deregulation
- Gene amplification
- Error prone DNA repair
- Irreversible
- Same mechanisms as promotion

**Classification of Carcinogens**

- **Genotoxic.**
  - Act directly on DNA or expression of DNA during translation.
  - DNA replication errors.
  - Point mutations.
  - Chromosomal aberration.

- **Epigenetic.**
  - Non-DNA reactive.
  - Potentiators.
  - Cell, hormone, immune function modifiers.

**Genotoxic Carcinogens**

- Chemical capable of producing cancer by directly altering the genetic material of target cells.
- Direct carcinogens (no metabolic activation).
  - Alkylating agents.
- Indirect carcinogens (metabolic activation).
  - Polycyclic aromatic hydrocarbons.
  - Aromatic amines.
  - Nitrosamines.
  - Natural substances.
    - Mycotoxins.
    - Inorganic carcinogens.
  - Ni, Cr, Cd, As.

**Epigenetic Carcinogens**

- Cytotoxic carcinogens.
  - Nitrilotriacetate, BHA, BHT.
- Tumor promoters.
  - DDT, Dioxin
- Hormones.
  - Estradiol, DES
- Immunosuppressants.
  - Cyclosporin A
- Particulates.
  - Asbestos.

**PAH Carcinogenic Activation**

- 7,8-Epoxide
- 7,8-Diol

**Proven Human Carcinogens**

- Chemicals.
  - Aflatoxins, 4-aminobiphenyl, As, benzene, benzidine, Be, bis-chloroethylether, Cd, Cr(VI), soot, mineral oils, mustard gas, 2-naphthylamine, Ni, vinyl chloride.
- Substance abuse.
  - Alcohol, betel nuts, cigarettes.
- Dust and fiber.
  - Asbestos, silica, soots, talcum, wood dust.
- Chronic infection.
  - H pylori, Hepatitis B/C, HIV, liverfluke, papilloma virus, schistosomes.
**Initiator Chemicals in Food**

- Most genotoxic chemicals
- PAHs
- Aromatic amines
- Heterocyclic amines
- Mycotoxins
- Nitrosamines
- Nitrosamides

**Promoting Agents in Food**

- Butylated hydroxy toluene (BHT)
- Saccharin
- Cholic acid
- Tetrachloro-dibenzo-dioxin (TCDD)
- Alcohol

**Chemical Cancer Assessment**

<table>
<thead>
<tr>
<th>Group</th>
<th>Class</th>
<th>Human carcinogen</th>
<th>Sufficient human evidence</th>
<th>Probable human</th>
<th>Limited human evidence</th>
<th>Possible human</th>
<th>Possible animal evidence</th>
<th>Probable human</th>
<th>Limited animal evidence</th>
<th>Possible human</th>
<th>Limited animal evidence</th>
<th>Group F</th>
<th>No evidence</th>
<th>2 animal tests or epidemiological evidence</th>
</tr>
</thead>
<tbody>
<tr>
<td>Group C</td>
<td>Possible human</td>
<td>Limited animal evidence</td>
<td>Possible human</td>
<td>Inadequate human evidence</td>
<td>Possible human</td>
<td>Limited animal evidence</td>
<td>Possible human</td>
<td>Limited animal evidence</td>
<td>Probable human</td>
<td>Limited animal evidence</td>
<td>Possible human</td>
<td>Limited animal evidence</td>
<td>Group F</td>
<td>No evidence</td>
</tr>
<tr>
<td>Group D</td>
<td>Not classifiable</td>
<td>Inadequate animal evidence</td>
<td>Possible human</td>
<td>Inadequate human evidence</td>
<td>Possible human</td>
<td>Limited animal evidence</td>
<td>Possible human</td>
<td>Limited animal evidence</td>
<td>Probable human</td>
<td>Limited animal evidence</td>
<td>Possible human</td>
<td>Limited animal evidence</td>
<td>Group F</td>
<td>No evidence</td>
</tr>
</tbody>
</table>

**Causes of Cancer**

- Diet, 35%
- Tobacco, 30%
- Sexual behavior, 7%
- Alcohol, 3%
- Infection, 10%
- Occupational exposure, 4%
- UV/radiation, 3%
- Pollution, 2%

**2006 Estimated US Cancer Cases***

<table>
<thead>
<tr>
<th>Site</th>
<th>Men</th>
<th>Women</th>
</tr>
</thead>
<tbody>
<tr>
<td>Prostate</td>
<td>720,280</td>
<td>679,510</td>
</tr>
<tr>
<td>Lung &amp; bronchus</td>
<td>13%</td>
<td>12%</td>
</tr>
<tr>
<td>Colon &amp; rectum</td>
<td>10%</td>
<td>11%</td>
</tr>
<tr>
<td>Urinary bladder</td>
<td>6%</td>
<td>6%</td>
</tr>
<tr>
<td>Melanoma of skin</td>
<td>5%</td>
<td>4%</td>
</tr>
<tr>
<td>Non-Hodgkin lymphoma</td>
<td>4%</td>
<td>4%</td>
</tr>
<tr>
<td>Kidney</td>
<td>3%</td>
<td>3%</td>
</tr>
<tr>
<td>Oral cavity</td>
<td>3%</td>
<td>3%</td>
</tr>
<tr>
<td>Leukemia</td>
<td>3%</td>
<td>2%</td>
</tr>
<tr>
<td>Pancreas</td>
<td>2%</td>
<td>2%</td>
</tr>
<tr>
<td>All Other Sites</td>
<td>18%</td>
<td>22%</td>
</tr>
</tbody>
</table>

**2006 Estimated US Cancer Deaths***

<table>
<thead>
<tr>
<th>Site</th>
<th>Men</th>
<th>Women</th>
</tr>
</thead>
<tbody>
<tr>
<td>Lung &amp; bronchus</td>
<td>26%</td>
<td>22%</td>
</tr>
<tr>
<td>Colon &amp; rectum</td>
<td>10%</td>
<td>6%</td>
</tr>
<tr>
<td>Prostate</td>
<td>9%</td>
<td>6%</td>
</tr>
<tr>
<td>Pancreas</td>
<td>6%</td>
<td>6%</td>
</tr>
<tr>
<td>Leukemia</td>
<td>4%</td>
<td>4%</td>
</tr>
<tr>
<td>Liver &amp; intrahepatic bile duct</td>
<td>4%</td>
<td>3%</td>
</tr>
<tr>
<td>Esophagus</td>
<td>4%</td>
<td>3%</td>
</tr>
<tr>
<td>Non-Hodgkin lymphoma</td>
<td>3%</td>
<td>3%</td>
</tr>
<tr>
<td>Kidney</td>
<td>3%</td>
<td>2%</td>
</tr>
<tr>
<td>All other sites</td>
<td>23%</td>
<td>23%</td>
</tr>
</tbody>
</table>
### Lifetime Probability of Developing Cancer, by Site, Men, 2000-2002*

<table>
<thead>
<tr>
<th>Site</th>
<th>Risk</th>
</tr>
</thead>
<tbody>
<tr>
<td>All sites†</td>
<td>1 in 2</td>
</tr>
<tr>
<td>Prostate</td>
<td>1 in 6</td>
</tr>
<tr>
<td>Lung and bronchus</td>
<td>1 in 13</td>
</tr>
<tr>
<td>Colon and rectum</td>
<td>1 in 17</td>
</tr>
<tr>
<td>Urinary bladder</td>
<td>1 in 28</td>
</tr>
<tr>
<td>Non-Hodgkin lymphoma</td>
<td>1 in 46</td>
</tr>
<tr>
<td>Melanoma</td>
<td>1 in 52</td>
</tr>
<tr>
<td>Kidney</td>
<td>1 in 64</td>
</tr>
<tr>
<td>Leukemia</td>
<td>1 in 67</td>
</tr>
<tr>
<td>Oral Cavity</td>
<td>1 in 73</td>
</tr>
<tr>
<td>Stomach</td>
<td>1 in 82</td>
</tr>
</tbody>
</table>

*For those free of cancer at beginning of age interval. Based on cancer cases diagnosed during 2000 to 2002.†All sites include basal and squamous cell skin cancers except urinary bladder.


### Lifetime Probability of Developing Cancer, by Site, Women, US, 2000-2002*

<table>
<thead>
<tr>
<th>Site</th>
<th>Risk</th>
</tr>
</thead>
<tbody>
<tr>
<td>All sites†</td>
<td>1 in 3</td>
</tr>
<tr>
<td>Breast</td>
<td>1 in 8</td>
</tr>
<tr>
<td>Lung &amp; bronchus</td>
<td>1 in 17</td>
</tr>
<tr>
<td>Colon &amp; rectum</td>
<td>1 in 18</td>
</tr>
<tr>
<td>Uterine corpus</td>
<td>1 in 38</td>
</tr>
<tr>
<td>Non-Hodgkin lymphoma</td>
<td>1 in 55</td>
</tr>
<tr>
<td>Ovary</td>
<td>1 in 68</td>
</tr>
<tr>
<td>Melanoma</td>
<td>1 in 77</td>
</tr>
<tr>
<td>Pancreas</td>
<td>1 in 79</td>
</tr>
<tr>
<td>Uterine bladder‡</td>
<td>1 in 88</td>
</tr>
<tr>
<td>Uterine cervix</td>
<td>1 in 135</td>
</tr>
</tbody>
</table>

‡Includes invasive and in situ cancer cases.

### Colorectal Cancer

**Risk Factors**
- Over 50
- Previous occurrence
- Family history
- High fat/low fiber diet
- Smoking
- Alcohol consumption
- Sedentary
- Overweight

### Kidney Cancer
Liver Cancer

Basal Cell Carcinoma

Tobacco Use in the US, 1900-2002

Trends in Consumption of Five or More Recommended Vegetable and Fruit Servings for Cancer Prevention, Adults 18 and Older, US, 1994-2003

The American Cancer Society recommends that individuals eat five or more servings of vegetables and fruits a day for cancer prevention.