

ERIN C. AMERMAN FLORIDA STATE COLLEGE AT JACKSONVILLE

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MODULE 5.1 OVERVIEW OF THE INTEGUMENTARY SYSTEM

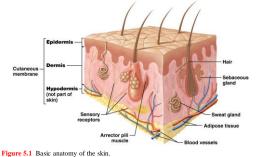
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SKIN STRUCTURE

- Skin accounts for 10–15% of an individual's total body weight making it *largest organ* in body; more than just an outer covering; complex organ with many functions important for *homeostasis* (Figure 5.1)
- Known as **cutaneous membrane**; has two main components:
 - Epidermis <u>superficial</u> layer that consists of *keratinized* stratified squamous epithelium resting on a basement membrane
 - Dermis deep to epidermis and basement membrane; consists of *loose connective tissue* and *dense irregular* connective tissue

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SKIN STRUCTURE



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SKIN STRUCTURE

- Accessory structures of skin embedded in cutaneous membrane: sweat glands, sebaceous glands, hair, and nails
- Skin contains *sensory receptors* and **arrector pili muscles** (small bands of smooth muscle associated with hair)
- Epidermis is avascular:
 - Must rely on diffusion of oxygen and nutrients from blood vessels in deeper dermis; example of Gradients Core Principle; <u>limits</u> epidermal thickness
 - About 50% of cells in epidermis are too far from adequate blood supply to sustain life; superficial layers are made up <u>entirely</u> of *dead cells*

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SKIN STRUCTURE

- Hypodermis also known as superficial fascia or subcutaneous fat, is <u>deep</u> to dermis
 - Although *not part of skin*, it does anchor skin to deeper structures like muscle and bone
 - Made of loose connective and adipose tissues; has an abundant blood supply

Figure 5.1 Basic anatomy of the skin.



CELLULITE

- Term used to describe *dimpled* or "*orange peel*" appearance of skin when collagen bands form around adipose tissue in the *hypodermis*
- Tends to develop in *thighs*, *hips*, and *gluteal area*; influenced by many factors; genetics, gender and amount and distribution of adipose tissue, and age
- Now thought to be normal condition (not disorder)
- Little evidence that any "cures" for cellulite work; only proven way to minimize appearance is a *healthy diet* and *regular exercise*; however, even diet and exercise do not generally eliminate it all together

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FUNCTIONS OF THE INTEGUMENTARY SYSTEM

Integumentary system has following functions that are critical for protecting underlying organs or for maintaining homeostasis:

- **Protection** from mechanical trauma, pathogens, and environment is most obvious function:
 - Stratified squamous, keratinized epithelium provides a *durable* but *flexible* surface; protects body from *mechanical trauma* like stretching, pressure, or abrasions
 - Provides a continuous barrier to invasion by microorganisms or pathogens that can cause disease
 - Contains cells of immune system that destroy pathogens before they invade deeper tissues

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FUNCTIONS OF THE INTEGUMENTARY SYSTEM

• **Protection** (continued):

- Glands secrete a variety of *antimicrobial substances*; sebaceous gland secretions give surface of skin a slightly acidic pH (called acid mantle); inhibits growth of many pathogens
- Provides protection from a number of environmental hazards including *absorption of ultraviolet light (UV)* <u>before</u> it damages deeper tissues
- Skin secretes hydrophobic lipid-based chemicals; repel ionic and polar covalent molecules like salt and water; critical for maintaining water and electrolyte homeostasis in a wide range of weather conditions

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FUNCTIONS OF THE INTEGUMENTARY SYSTEM

- Sensation process that enables nervous system to *perceive changes* in the body's internal or external surroundings; critical to homeostasis:
 - Skin has numerous sensory receptors or cellular structures that detect changes in internal and/or external environment
 - Receptors allow us to detect *potentially harmful* stimuli such as heat, cold, and pain; could lead to tissue damage

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FUNCTIONS OF THE INTEGUMENTARY SYSTEM

• Thermoregulation (Figure 5.2):

- Process that relies on negative feedback loops for maintenance of a *stable internal temperature*
- Example of Feedback Loops Core Principle
- Internal body temperature is determined mostly by muscle activity and many chemical reactions involved in metabolism

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FUNCTIONS OF THE INTEGUMENTARY SYSTEM

- Sequence of events that occur when body temperature rises <u>above</u> normal range; may be caused by extremes of weather or due to abnormal conditions that cause fever (Figure 5.2a):
 - Sensory receptors (thermoreceptors) in skin detect an increase in temperature in both skin itself and internal body fluids
 - Control center in hypothalamus of brain acts as a *thermostat* or thermoregulatory center; receives *input* from thermoreceptors; processes and then *responds* to sensory inputs
 - Control center stimulates sweating; sweat glands are stimulated to release a watery fluid called sweat; water carries
 - stimulated to release a watery fluid called **sweat**; water carries a great deal of heat with it when it *evaporates*; provides for an effective cooling mechanism

FUNCTIONS OF THE INTEGUMENTARY SYSTEM

• Sequence of events that occur when body temperature rises <u>above</u> normal range (continued):

- Control center stimulates cutaneous vasodilation; response triggered by hypothalamus; causes blood vessels in dermis to widen (dilate); increased blood flow through dilated vessels increases amount of heat radiated away from body into environment; cools body
- Body temperature returns to normal range and cooling mechanisms decline by negative feedback; when thermoreceptors no longer sense body temperatures <u>above</u> normal range they stop sending signals to hypothalamus; ends control center responses; sweating and vasodilation ends

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FUNCTIONS OF THE INTEGUMENTARY SYSTEM

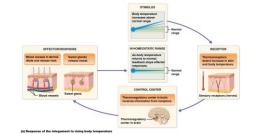


Figure 5.2a Homeostatic regulation of body temperature by integumentary system.

FUNCTIONS OF THE INTEGUMENTARY SYSTEM

- Sequence of events that occur when body temperature drops <u>below</u> normal range; usually due to cold environmental conditions (Figure 5.2b):
 - Thermoreceptors detect body temperature drop below normal range; relay information to thermoregulatory center in hypothalamus
 - Hypothalamus generates a different response than it does for an increased body temperature; blood vessels in dermis narrow (vasoconstrict) <u>reducing</u> amount of blood flow; *limits heat lost* to environment

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FUNCTIONS OF THE INTEGUMENTARY SYSTEM

- Sequence of events that occur when body temperature drops <u>below</u> normal range (continued):
 - Vasoconstriction also redirects blood flow to deeper tissues; helps to conserve heat
 - When body temperature rises back into normal range, thermoreceptors stop sending information to hypothalamus; response that hypothalamus generated for heat conservation ends; feedback loop is closed

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FUNCTIONS OF THE INTEGUMENTARY SYSTEM

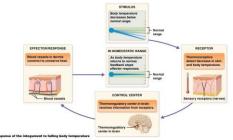


Figure 5.2b Homeostatic regulation of body temperature by integumentary system.

FUNCTIONS OF THE INTEGUMENTARY SYSTEM

• Excretion – process where *waste products* and *toxins* are eliminated from body; most occurs at <u>other</u> organs like *kidneys*; skin and its accessory structures make a small but significant contribution

FUNCTIONS OF THE INTEGUMENTARY SYSTEM

- Skin plays a critical role in vitamin D synthesis; cells found deep in epidermis convert vitamin D from an inactive form (precursor) to active form:
 - Precursor modified cholesterol molecule; converted to cholecalciferol when epidermis is exposed to UV radiation
 - Cholecalciferol is *released into blood*; modified first by liver, then by kidneys, to form calcitriol (active form of vitamin D)
 - Vitamin D is required for *calcium ion absorption* from small intestine; calcium ion is critical for *nerve function*, *muscle contraction*, *building* and *maintaining bone tissue*, and many other physiological functions

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MODULE 5.2 THE EPIDERMIS

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THE EPIDERMIS

- Epidermis most superficial layer; composed of several cell types; most numerous are keratinocytes
- Make up about 95% of epidermis; have two structural features that make epidermis *stronger* and <u>less</u> susceptible to *mechanical trauma*:
 - Manufacture keratin tough fibrous protein that makes epidermis more resistant to mechanical trauma; demonstrates Structure-Function Core Principle
 - Linked to each other by **desmosomes**; makes epidermis stronger

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THE EPIDERMIS

- Keratinocytes organized from deep to superficial into five structurally distinct strata (layers) (Figure 5.3):
 - Stratum basale (stratum germinativum) single layer of *stem cells* resting on basement membrane; closest cells to blood supply in dermis; therefore most *metabolically* and *mitotically active* cells in epidermis; involved in vitamin D synthesis and replacement of dead keratinocytes (lost from more superficial layers)
 - Stratum spinosum thickest layer, sits on top of stratum basale so still close to blood supply; also metabolically and mitotically active

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THE EPIDERMIS

- Five structurally distinct **strata** (continued):
 - Stratum granulosum
 - Three to five layers of cells with prominent cytoplasmic granules; filled with keratin bundles or a lipid-based substance; both secreted by exocytosis
 - Hydrophobic nature of lipids provides waterproofing; critical for maintaining internal fluid and electrolyte homeostasis; also leads to *isolation* and *death* of cells in this layer and in more superficial layers

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THE EPIDERMIS

- Five structurally distinct strata (continued):
 - **Stratum lucidum** narrow layer of *clear*, *dead keratinocytes*; found <u>only</u> in thick skin
 - Stratum corneum outermost layer of epidermis; consists of several layers of *dead flattened keratinocytes* with thickened plasma membranes; filled mostly with *keratin bundles* and little else; *sloughed off* or *exfoliated mechanically* as desmosomes holding neighboring cells together are lost

THE EPIDERMIS

Figure 5.3 Structure of the epidermis.

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TOPICAL MEDICATIONS

- Some medications are *toxic* if swallowed, but *safe* if used **topically** (applied to surface of skin)
- Certain topical antibiotics are fairly toxic if taken by mouth, but can be applied to skin with minimal risk of systemic absorption; they are polar molecules that <u>cannot</u> pass through epidermis to reach blood vessels in dermis; allows for *local effect only*
- Nonpolar substances cross epidermis much more easily; provides a *convenient route of administration* for certain medications such as hormones in birth control patches
- Unfortunately, many *poisons* and *toxins* (like thallium, a heavy metal) are also nonpolar; cross epidermis with same ease; therefore always good idea to wear gloves when handling chemicals

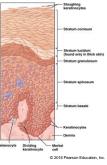
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STUDY BOOST: REMEMBERING THE STRATA OF THE EPIDERMIS

Here is a simple trick to remember **strata of epidermis**:

- "Brilliant Studying Gives Loads of Confidence"
- If you get confused as to which stratum is superficial and which is deep, think of the "B" in "basale" as standing for "bottom"; it is <u>bottom</u> layer

Figure 5.3 Structure of the epidermis.



THE EPIDERMIS

- Keratinocyte life cycle: location and functions of epidermis subjects it to both physical and environmental stress; stratum corneum is continuously shedding dead cells that must be replaced to maintain integrity of epidermis:
 - Dead keratinocytes are replaced by *mitosis* of cells in stratum basale and spinosum where blood supply is available for such activities
 - As keratinocytes in deeper strata divide they push cells above them into *more superficial layers*

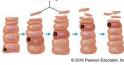
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THE EPIDERMIS

- Keratinocyte life cycle (continued):
 - Keratinocytes begin life in stratum basale or spinosum; eventually pass through <u>each</u> epidermal layer until shed from stratum corneum
 - Migration from deepest strata to stratum corneum takes a cell between 40–50 days to complete

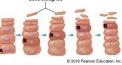
CONCEPT BOOST: UNDERSTANDING EPIDERMAL GROWTH

- Suppose for simplicity's sake that skin has *one row of cells in each epidermal stratum*, as shown:
 - Cell A undergoes mitosis, and one of its two daughter cells (cell B in diagram) is now in stratum spinosum
 - Other daughter cell of cell A divides again, producing two more daughter cells, one of which pushes cell B into stratum granulosum



CONCEPT BOOST: UNDERSTANDING EPIDERMAL GROWTH

- Suppose for simplicity's sake that skin has one row of cells in each epidermal stratum, as shown (continued):
 - **Cell B** is now quite far from blood supply; becomes *coated with lipid-based substance*; causes it to die
 - Stem cells continue to divide, pushing cell B even farther away from blood supply, into stratum lucidum and then into stratum corneum
 - **Cell B** is now a dead cell filled with keratin; will eventually be *sloughed off skin surface*



OTHER CELLS OF THE EPIDERMIS

- Dendritic (Langerhans) cells located in stratum spinosum; phagocytes of immune system; protect skin and deeper tissues from pathogens
- Merkel cells oval cells scattered throughout stratum basale; sensory receptors associated with small neurons in dermis:
 - Detect light touch and discriminate shapes and textures
 - Found in large numbers in regions that are specialized for touch; fingertips, lips, and at base of hairs
- Melanocytes located in stratum basale; produce melanin; protein skin pigment ranging from orange-red to brown-black

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THICK AND THINK SKIN

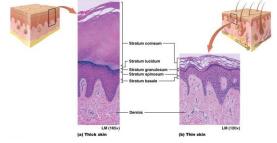
- As with all structures, form of epidermis in various parts of body differs to match its *function*, in agreement with Structure-Function Core Principle
- Palms of hand and sole of foot are subjected to a great deal of mechanical stress, so these regions of skin have adapted; remaining regions of skin are not subjected to as much stress; differences in function and exposure to stress have lead to *thick* and *thin* skin (Figure 5.4)

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THICK AND THINK SKIN

- Thick skin, about as thick as a paper towel, has all five epidermal layers and a very thick stratum corneum; does <u>not</u> have *hair follicles* but contains many *sweat glands* (Figure 5.4a)
- Areas of body not subjected to as much mechanical stress are covered with thin skin; about as thick as a sheet of printer paper, has <u>only</u> four layers; stratum lucidum is missing (Figure 5.4b)
 - Each of four layers is thinner than those found in thick skin
 - · Numerous hairs, sweat glands, and sebaceous glands present
- **Callus** additional layers of stratum corneum; form in either thick or thin skin in response to *repetitive pressure*

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THICK AND THINK SKIN

Figure 5.4 Thick and thin skin.

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MODULE 5.3 THE DERMIS

THE DERMIS

Dermis - highly vascular layer deep to epidermis

- Functions:
 - · Provides blood supply for epidermis
 - Contains sensory receptors
 - Anchors epidermis in place
- Composed of two distinct layers made up of two types of connective tissue

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THE PAPILLARY LAYER

Papillary layer – thinner most superficial of two layers; composed of *loose connective tissue* (Figure 5.5):

• Special collagen fibers are found in this layer at *dermisepidermal junction*; extends into epidermal basement membrane to *anchor epidermis to dermis*

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THE PAPILLARY LAYER

- Dermal papillae tiny projections found at surface of papillary layer where it comes into contact with epidermis:
 - Contain tiny blood vessels called capillaries arranged in loops; extend up into most *superficial part* of dermal papillae
 - Allow oxygen and nutrients to diffuse into extracellular fluid of dermis; then into cells of avascular epidermis
 - Tactile (Meissner) corpuscles also found in dermal papillae; sensory receptors that respond to light touch stimuli; more numerous in regions of body where sensation is a primary function; skin of fingertips, lips, face, and external genitalia

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THE RETICULAR LAYER

Reticular layer – <u>deep</u> thicker layer that separates dermis from hypodermis; mostly *dense irregular connective tissue* that consists largely of irregularly arranged collagen bundles:

- Collagen bundles strengthen dermis and prevent traumatic injuries from damaging deeper tissues
- Elastic fibers allow dermis to return to its original shape and size <u>after stretching</u>

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THE RETICULAR LAYER

Reticular layer (continued):

- Rich in proteoglycans that draw water into ground substance; keeps skin *firm* and *hydrated*
- Lamellated (Pacinian) corpuscles found embedded within reticular layer; sensory receptors that respond mainly to changes in *pressure* and *vibration* associated with skin
- Blood vessels, sweat glands, hairs, sebaceous glands, and adipose tissue are found in reticular layer

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THE DERMIS

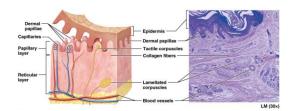


Figure 5.5 Structure of the dermis.

SKIN MARKINGS

Skin markings – small *visible lines* in epidermis created by interaction between dermis and epidermis; best seen in thick skin of palmar surfaces of hands and fingers and plantar surface of feet and toes (**Figure 5.6**)

- Dermal ridges found in areas where dermal papillae are more prominent due to presence of thick collagen bundles
- Dermal ridges *indent overlying epidermis* to create **epidermal** ridges; enhance *gripping ability* of hands and feet:
 - Epidermal ridges occur in characteristic patterns; *loops, arches,* and *whorls*; genetically determined and unique to each person
 - Sweat pores open along these ridges and leave a thin film or fingerprint on things touched with fingers

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SKIN MARKINGS

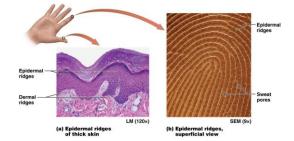


Figure 5.6 Epidermal ridges and fingerprint patterns.

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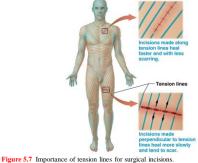
SKIN MARKINGS

Skin markings (continued):

- Reticular layer is also responsible for skin markings associated with tension or lines; *cleavage lines* and *flexure lines* (Figure 5.7):
 - Gaps found between collagen bundles in dermis create indentations in epidermis called tension or cleavage lines
 - In areas of body, such as surrounding joints, reticular layer is tightly anchored to deeper structures that create deep creases called **flexure lines**

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SKIN MARKINGS



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SKIN WRINKLES

- Hallmark of aging; due to age-related *decrease in collagen fibers, elastic fibers, proteoglycans,* and *adipose tissue* in the dermis
- <u>Reduces</u> skin's *firmness*, *hydration*, and *recoil ability* after stretching; tend to be deeper in areas of repetitive muscle movement (forehead and around eyes and mouth); *UV exposure* and *cigarette smoking* accelerate formation of wrinkles

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SKIN WRINKLES

- Appearance can be minimized by:
 - **Botox** bacterial toxin; temporarily paralyzes facial muscles; causes skin to appear smoother
 - Fillers adipose tissue, collagen, and/or proteoglycans are injected into wrinkles
 - **Topical creams** (especially nonprescription) claim to reduce appearance of wrinkles; little to no effect
- Avoidance of sun, use of sunscreens, maintenance of hydration, and avoidance of smoking can delay appearance of wrinkles

MELANIN

Skin color – mostly determined by various amounts of orange-red to black protein pigment **melanin**:

- Produced by **melanocytes** in stratum basale of epidermis (**Figure 5.8**)
- Composed of two molecules of amino acid tyrosine; chemically bonded by a series of reactions catalyzed by enzyme tyrosinase; reactions occur in a stepwise fashion within a special vesicle called a melanosome
- Protecting keratinocyte DNA from *mutations induced by UV radiation* is a primary function

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MELANIN

MODULE 5.4 SKIN PIGMENTATION

Skin color (continued):

- Melanocytes have several extensions of plasma membrane in contact with keratinocytes of stratum basale and spinosum
 - Melanosomes migrate to ends of these arms where *released* by *exocytosis*; absorbed or taken into cytoplasm of surrounding keratinocytes
 - Melanin is transported to *superficial side of nucleus* (faces exterior of body); *shields DNA* of keratinocyte like an umbrella
 - Melanin must be *made continuously* to maintain a consistent skin color as it degrades after a few days

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MELANIN

Skin color (continued):

- Melanin synthesis increases with exposure to natural or artificial UV radiation; leads to tanning or *darkening of skin pigmentation*; UV radiation has both *immediate* and *delayed* effects on skin pigmentation:
 - Immediate response to UV radiation is *oxidation of melanin* already present in keratinocytes; causes melanin to *quickly darken*
 - UV light causes DNA damage in melanocytes; stimulates melanin production leading to delayed or secondary effects of UV exposure; appear within 72 hours and last longer than melanin oxidation

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MELANIN

Skin color (continued):

- Melanin synthesis increases with exposure to natural or artificial UV radiation; leads to tanning or *darkening of skin pigmentation*; UV radiation has both *immediate* and *delayed* effects on skin pigmentation (continued):
 - Amount of UV radiation melanin can absorb is limited as is protection it provides
 - People of *all skin pigmentations* can develop sunburns and are at risk for skin cancers

MELANIN

Skin color (continued):

- Secondary function of melanin is to *reduce synthesis of vitamin D* in response to UV radiation; leads to <u>less</u> *calcium ion absorption* and maintenance of calcium ion homeostasis within a narrow range:
 - Individuals living in regions exposed to high amounts of UV radiation (such as Africa) may have developed darker skin to prevent excess vitamin D production
 - People in areas with *less UV radiation* (such as northern Europe) developed lighter skin so they could synthesize <u>enough</u> vitamin D

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MELANIN

Skin color (continued):

- Skin color depends on number of melanocytes found in a particular body region; differences lead to uneven distribution of melanin; fewer melanocytes are found on palms of hand and soles of the feet, for example
- Overall number of melanocytes is virtually identical among <u>all</u> individuals, <u>irrespective</u> of skin color; spectrum of human skin tones is due to differences in amount of tyrosinase activity and type (color) of melanin produced

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MELANIN

Skin color (continued):

- · Common variations of pigmentation:
 - Freckle small area of <u>increased</u> pigmentation; resulting from increased melanin production in local spot
 - Mole or nevus area of increased pigmentation; due to a local *proliferation* of melanocytes, <u>not</u> an increase in melanin production
 - Albinism melanocytes *fail to manufacture* tyrosinase; results in lack of skin pigmentation and greatly increased risk of keratinocyte DNA damage from UV radiation

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TANNING AND A "HEALTHY TAN"

- Tanning \$5-billion-a-year business in United States alone; number of salons has soared from 10,000 to 50,000 in last decade; salons promote notion of "healthy tan"
- THERE IS NO SUCH THING AS A HEALTHY TAN!
- UVA and UVB rays are associated with sunburning; UVA rays are linked with tanning; led salons to claim that UVA rays are safe and will not damage skin, but mechanism of increased melanin production is <u>same</u> for both types of rays; both damage DNA equally, but UVA ages skin at <u>much faster rate</u>
- ANY amount of tanning damages melanocytes and other skin elements, ages skin prematurely, and increases risk of skin cancer

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CAROTENE AND HEMOGLOBIN

Two minor pigments have an effect on skin pigmentation:

- **Carotene** *yellow-orange pigment* found in food items such as egg yolks and orange vegetables
 - *Lipid-soluble* molecule that accumulates in stratum corneum
 - Imparts a slight yellow-orange color that is particularly visible in stratum corneum of thick skin

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CAROTENE AND HEMOGLOBIN

Two minor pigments (continued):

- Hemoglobin found in red blood cells, is an ironcontaining protein that binds to and *transports oxygen* throughout body:
 - Oxygen binds to iron in hemoglobin in an oxidation reaction; same reaction that causes iron to rust; oxidized hemoglobin changes color to a bright orange-red; gives blood its characteristic color
 - Hemoglobin's effect on skin color is an <u>indirect</u> result of blood flow in dermis; color of blood in deeper dermis is visible through epidermis

SKIN COLOR AS A DIAGNOSTIC TOOL

Color changes associated with *amount of blood flow* in dermis can be useful in diagnosis of disease:

- Erythema occurs when blood flow in dermis <u>increases</u> causing a color change that makes skin *more reddish*
 - Color change is a <u>normal response to exercise</u> where blood flow in dermis has increased to maximize heat released to external environment
 - Other conditions that cause erythema include: trauma, fever, and infection

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SKIN COLOR AS A DIAGNOSTIC TOOL

Color changes associated with *amount of blood flow* in dermis (continued):

- Pallor occurs when blood flow in dermis <u>decreases</u>; results in <u>loss</u> of *normal pinkish hue*; most visible in paleskinned individuals; epidermis may take on *whitish color* of collagen in dermis
 - Normal response when body is trying to conserve heat in a cold environment
 - Can also occur when nervous and endocrine systems alter blood flow to dermis as part of a *flight or fight response*

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SKIN COLOR AS A DIAGNOSTIC TOOL

Color changes associated with *amount of blood flow* in dermis (continued):

- Cyanosis sign that someone needs immediate attention; occurs when hemoglobin has very low levels of bound oxygen; blood turns *reddish purple*; skin takes on a faint *bluish hue*; can occur when
 - Someone has difficulty breathing
 - Hemoglobin or red cell levels are <u>low</u> in blood
 - · Hemoglobin is unable to bind to oxygen

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MODULE 5.5 ACCESSORY STRUCTURES OF THE INTEGUMENT: HAIR, NAILS, AND GLANDS

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Hair

Accessory structures or appendages of integument include *hair*, *nails*, and *glands*; derived from epithelium <u>only</u>; assist in overall function of integumentary system:

• Hair (pili) – small filamentous structures that protrude from surface of skin over entire body except in regions with thick skin, lips, and parts of external genitalia (Figure 5.9)



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HAIR

- Hair too sparse in humans to play a significant role in thermoregulation, as it does in other mammals:
 - Does provide *protection* by preventing substances and organisms from external environment from *entering eyes* and *nose*
 - On head, protects underlying skin of scalp from UV radiation and mechanical trauma
 - Hairs are associated with a *small sensory neuron*; plays a role in detecting *changes in environment*

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HAIR STRUCTURE

 Hair – composed of two main parts; shaft and root; both made up of stratified squamous keratinized epithelial cells in various stages of development

Shaft

- Portion of hair that *projects* from skin's surface
- Made up of columns of dead keratinized epithelial cells that have completed keratinization process



Figure 5.9 Hair structure.

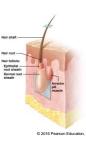


Hair structure (continued):

Root

- Segment of hair *embedded in dermis*; surrounded by a small sensory neuron
- Root is indented at its base by a projection of blood vessels from dermis called a hair papilla
- Root and hair papilla are collectively known as hair bulb
- Many epithelial cells in root are still alive; have <u>not</u> completed keratinization process

Figure 5.9 Hair structure.



HAIR STRUCTURE

- Hair structure (continued):
 - Matrix small number of keratinocytes found at base of root; actively divide
 - Root is embedded in hair follicle; an infolding of epidermis called epithelial root sheath; extends deep into dermis or even hypodermis
- Epithelial root sheath has an outer component that anchors follicle to dermis and an inner component that is anchored tightly to hair root
 Figure 5.90 Hair structure.

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HAIR STRUCTURE

- Hair structure (continued):
 - Strand of hair has three visible regions in a transverse section:
 - Inner medulla soft core <u>only</u> found in thick hair (like on head); composed of a soft keratin
 - Middle cortex highly structured and organized with several layers of keratinocytes containing *hard keratin*; provides
 - strength to strand
 Outermost cuticle consists of a single layer of overlapping keratinocytes containing hard keratin; provides mechanical strength

Figure 5.9a Hair structure.



Diale Shaft

Cuticle

HAIR STRUCTURE

- Hair structure (continued):
 - Surrounding epithelial root is a dermal root sheath; consists of connective tissue; supports follicle and keeps it separate from dermis
 - Small bands of *smooth muscle* called arrector pili muscles attach to dermal root sheath on one end and dermal papillary layer on the other
 - Contraction of these tiny muscles causes hair to stand up (piloerection); gives skin a dimpled appearance, commonly called "goosebumps"

Figure 5.9 Hair structure.



HAIR GROWTH

• Hair growth averages between 1–1.5 cm per month; varies between individuals; growth is <u>not</u> continuous but occurs in a cycle with following *two main phases*:

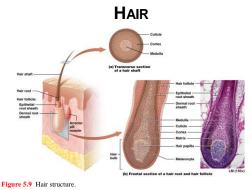
- During growth stage, *mitosis* occurs in matrix:
 - Cells divide and push cells above them farther away from blood supply where they keratinize and die
 - Stage varies in duration from a month to as long as six years;
 - depends on location of hair

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HAIR GROWTH

- Hair growth averages between 1–1.5 cm per month; *varies between individuals*; growth is <u>not</u> *continuous* but occurs in a cycle with following *two main phases* (continued):
 - During resting stage, mitosis in matrix *ends* as cells *die*:
 - Follicle shortens and hair is pushed toward surface where it remains *dormant* for a month or two
 - Falls out on its own or is pushed out by a new hair in growth stage

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HAIR PIGMENT AND TEXTURE

- Hair color and texture vary with different types of hair:
 - Lanugo thin, nonpigmented hair found covering nearly entire body of a fetus; generally fall out around birth; replaced with one of two hair types:
 - **Terminal hair** *thick*, *coarse*, and *pigmented hair*; found surrounding eyes and on scalp
 - Vellus hair thinner nonpigmented hair; found over remaining regions of body
 - Terminal hair replaces much of vellus hair <u>after</u> puberty; varies by gender with <u>more</u> hair replacement occurring in males than females

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HAIR PIGMENT AND TEXTURE

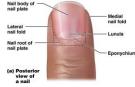
- Hair color and texture (continued):
 - Hair color is largely determined by melanin produced in matrix by melanocytes; produce a *range of colors*:
 - o Blond hair has little melanin
 - o Black hair which contains a lot of melanin
 - o Red hair has a special reddish pigment containing iron
 - Melanocytes produce less melanin with aging so hair eventually turns gray or white

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NAILS

- Nails hard accessory structures that are located at ends of digits; composed of *stratified squamous epithelium* filled with *hard keratin*
- Nail plate most visible component of nail, sits on top of an underlying epidermal nail bed; divided into:
 - Nail body visible portion of nail plate
 - Nail root portion of plate that lies under skin; where nail matrix containing actively dividing cells is found

Figure 5.10a Nail structure.



NAILS

- · Folded regions of skin surround and reinforce nail plate:
 - Proximal nail fold on proximal edge covering nail root; distal edge of this fold is called the eponychium (cuticle); consists of only stratum corneum
 - Medial and lateral nail folds on medial and lateral edges of nail plate respectively
 - Distal or free edge of nail plate – attached to underlying nail bed by an accumulation of stratum corneum called hyponychium



Figure 5.10b Nail structure.



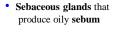
- Nail growth occurs at nail matrix; actively dividing cells push neighboring keratinocytes *distally*; die once keratinization is completed and have been cut off from blood supply; grow an average of 0.5 mm per week; toenails grow more slowly
- Nails do <u>not</u> contain melanocytes; mostly *translucent* except at region called **lunula**; half-moon shaped region of proximal nail plate that represents an *accumulation of keratin*
- Primary function of nails *protection* of underlying tissue (distal tips of the fingers and toes) from trauma; can be used as *tools*, enabling more precise gripping of items when picked up

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GLANDS

Skin contains two basic types of **glands**; both derived from epithelial cells in epidermis but located deeper in dermis

• Sweat (sudoriferous) glands that produce sweat



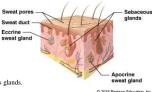


Figure 5.11 Sweat glands and sebaceous glands.

GLANDS

- Four types of sweat glands; differ structurally and in products secreted; all secrete products via exocytosis; called merocrine secretion:
 - Eccrine sweat glands (Figure 5.11a):
 - o Most prevalent type
 - o Simple coiled tubular glands found in dermis
 - Sweat, containing mostly water, waste products, and electrolytes
 - o Exits from duct through a sweat pore onto epidermal surface

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GLANDS

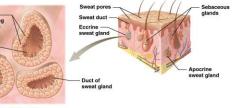
• Four types of sweat glands (continued):

• Apocrine sweat glands (Figure 5.11):

- \circ Found in specific regions of body such as axillae, anal area, and areola
- Large glands that release a *protein-rich secretion* into a *hair follicle*
- Secretions can become *odoriferous* once skin bacteria metabolize their contents
- o Influenced by sex hormones; become active after puberty

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(a) Eccrine sweat gland, frontal section

Figure 5.11a Sweat glands and sebaceous glands.

GLANDS

- Four types of sweat glands (continued):
 - · Ceruminous glands:
 - Modified apocrine glands
 - Release a thick secretion called cerumen (ear wax) into hair follicles found in ear
 - Cerumen traps incoming particles along tube leading to tympanic membrane; also lubricates
 - Mammary glands highly specialized sweat glands that produce a modified sweat product, milk

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GLANDS

- Sebaceous glands branched with clusters of secretory cells called acini surrounded by small ducts; converge to form a central duct that empties into hair follicle or small pore; makes and secretes sebum (Figure 5.11b):
 - Found everywhere on body except palms and soles; greatest number found on face and scalp
 - Secretion is influenced by sex hormones; especially male sex hormone (testosterone)
 - Dramatic increase in sebum production occurs <u>after</u> puberty; example of Cell-Cell Communication Core Principle

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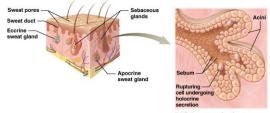
GLANDS

- Sebaceous glands (continued):
 - Sebum waxy, oily mixture of mostly lipids; released by holocrine secretion; secretory cells accumulate sebum until cell ruptures
 - Contains *cellular fragments* and *debris* in addition to lipids
 - Coats hair, providing it with a hydrophobic barrier that inhibits water loss
 - Also inhibits growth of or kills certain bacteria

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GLANDS



(b) Sebaceous gland

Figure 5.11b Sweat glands and sebaceous glands.

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ACNE

- Acne vulgaris affects 96% of adolescents and young adults to some degree
- Cause accumulation of sebum and dead cells within sebaceous glands; produces a comedone (blackhead); occasionally becomes infected by *Propionibacterium acnes*, resulting in inflammation and formation of a pustule (pimple)
- May be severe and cause *permanent scarring* in some individuals
- Male sex hormones (like testosterone) are primary cause; tends to be more pronounced in males entering puberty; decreases and may disappear by age 20–25; may persist much longer in some individuals

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MODULE 5.6 PATHOLOGY OF THE SKIN

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WOUNDS

- Wound common skin pathology; defined as any disruption in skin's integrity; include:
 - Lacerations (cuts)
 - Burns
 - Skin cancers

BURNS

Burn – wound caused by agents such as *heat*, *extreme cold*, *electricity*, *chemicals*, and *radiation*; grouped into three classes according to *extent* and *depth* of tissue damage:

- First-degree burns (superficial burns)
 - Minor wounds that <u>only</u> damage epidermis
 - Skin may develop erythema (red appearance) and some mild pain without any permanent damage

Figure 5.12a The three classes of burns.





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BURNS

Burns (continued):

- Second-degree burns (partial thickness burns)
 - Involve epidermis and part or all of dermis
 - Can result in pain, blistering, and scarring

Figure 5.12b The three classes of burns.



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BURNS

Burns (continued):

- Third-degree burns (full thickness burns)
 - <u>Most</u> damaging wounds
 - Involve epidermis, dermis, hypodermis; potentially even deeper tissue, like muscle or bone
 - <u>Not generally painful</u> at first because nerves are destroyed too

Figure 5.12c The three classes of burns.





Third-degree burn involves epidermis, dermis, hypodermis, and possibly deeper tissues.

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BURNS

Burns (continued):

• Third-degree burns (full thickness burns) (continued):

Typically result in major tissue damage and significant scarring with loss of hair follicles and diminished or absent keratin production

 Often problems with dehydration due to *massive fluid* loss from swelling; also at great risk for infection

Figure 5.12c The three classes of burns.



BURNS

Rule of nines

- Method for estimating how much of body has been affected by a burn
- Body is divided into 11 areas each representing 9% of the total body area
- Useful clinical tool for grading extent of burn; severity and extent of burn is used to direct *treatment* options

Figure 5.13 Rule of nines: estimating the extent of a burn.





SKIN CANCER

- Cancer one of most common diseases in world; caused by *mutations in DNA* that induce a cell to *lose control* of cell cycle (Figure 5.14):
 - Unchecked cell division eventually leads to formation of a large population of undifferentiated cells known as a tumor
 - Cancerous tumors are able to metastasize; tumor cells spread through blood or lymphatic vessels to other tissues and continue to divide
 - Damage caused by metastatic tumor cells *alters function* of invaded organs

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SKIN CANCER

- Three cancers affect skin; linked to UV radiation exposure; other factors that increase risk for developing cancer include exposure to:
 - Cancer-inducing chemicals, toxins, or agents called carcinogens
 - Forms of radiation

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SKIN CANCER

Basal cell carcinoma

- <u>Most</u> common of <u>all</u> cancer types, including skin cancer
- · Arises from keratinocytes in stratum basale of epidermis
- Skin that is *regularly exposed* to UV radiation is at risk for developing these tumors
- Appear as a nodule with a central crater
- · Rarely metastasize to other tissues
- Can be resolved successfully with surgical removal

Figure 5.14a The three main forms of skin cancer.



SKIN CANCER

- Squamous cell carcinoma
 - Second most common skin cancer
 - · Cancer of keratinocytes of stratum spinosum
 - Scaly plaques that may ulcerate and bleed are usually found on head and neck
 - Tumors are <u>more</u> likely to metastasize than basal cell carcinoma; *surgical removal* is still useful

Figure 5.14b The three main forms of skin cancer.



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SKIN CANCER

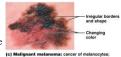
- Malignant melanoma cancer of melanocytes
 - Early detection of melanoma is critical due to its tendency to metastasize
 - "Arms" of cancerous melanocytes extend down into dermis and access dermal blood vessels; enables cells to spread to other tissues via bloodstream
 - Treated with surgical removal and possibly other options such as radiation therapy and chemotherapy
 - Prognosis depends on size of the tumor, depth to which it extends into dermis, and whether it has metastasized to other tissues

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SKIN CANCER

- Malignant melanoma can be distinguished from other skin cancers and normal moles using ABCDE rule:
 - (A): Asymmetrical shape (two sides do not match)
 - (B): Border irregularity
 - . (C): Color, usually blue-black or a variety of colors
 - (D): Diameter generally larger than 6 mm (size of a pencil eraser)
 - (E): Evolving (changing) shape and size

Figure 5.14c The three main forms of skin cancer.



blue-black color, larger diameter, and nature