

# The Integumentary System

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## ◆ Skin Development

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### ◆ Overview

- Largest organ of the body
- Dual origin:

Component	Origin
Epidermis	Surface ectoderm
Dermis	Mesenchyme (mesoderm + neural crest)

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### ◆ Epidermis Development

- ◆ Early Stage
    - Initially single-layered ectoderm
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◆ 2nd Month

- Ectoderm divides → forms:
    - Periderm (epitrichium) → superficial flattened layer
    - Basal layer → proliferative layer
  - Further proliferation → intermediate layer
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◆ By 4th Month → Definitive Layers

Layer	Features
Basal (stratum germinativum)	Stem cells; forms epidermal ridges → fingerprints
Spinous (stratum spinosum)	Polyhedral cells with tonofibrils
Granular (stratum granulosum)	Keratohyalin granules
Horny (stratum corneum)	Dead keratinized cells → protective barrier

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◆ Periderm Fate

- Shed in 2nd half of intrauterine life
  - Cells present in amniotic fluid
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## ◆ Melanocytes

### ◆ Origin

- Neural crest cells

### ◆ Timeline

- Migrate into epidermis by 3rd month

### ◆ Function

- Produce melanin in melanosomes
- Transfer pigment to keratinocytes

→ Responsible for skin & hair color

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## ◆ Dermis Development

### ◆ Origin (Mesenchyme from 3 sources)

Source	Region
Paraxial mesoderm	Dermis of back
Lateral plate mesoderm	Dermis of limbs & body wall
Neural crest	Dermis of face & neck

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- ◆ 3rd-4th Month Changes

- Formation of dermal papillae:
  - Project into epidermis
  - Contain:
    - Capillary loops
    - Sensory nerve endings

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- ◆ Deep Layer

- Subcorium (hypodermis) → rich in fat

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- ◆ Special Feature at Birth

- ◆ Vernix Caseosa

- ◆ Composition

- Sebaceous gland secretions
- Degenerated epidermal cells
- Fine hair (lanugo)

- ◆ Function

- Protects skin from amniotic fluid maceration
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- ◆ Clinical Correlates

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- ◆ I. Pigmentary Disorders

- ◆ Piebaldism

- Patchy absence of melanocytes  
→ localized depigmented skin & hair
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- ◆ Waardenburg Syndrome

- Cause → defective neural crest migration

- ◆ Features

- White forelock
- Heterochromia iridis
- Depigmented patches
- Deafness

- ◆ Gene
    - PAX3 mutation (WS1 & WS3)
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- ◆ Albinism (OCA)
    - Defect in melanin synthesis/processing
      - ↓ /absent pigmentation (skin, hair, eyes)
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- ◆ Vitiligo
    - Autoimmune destruction of melanocytes
      - patchy depigmentation
  - ◆ Associations
    - Autoimmune diseases (especially thyroid disorders)
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## ◆ 2. Fingerprints (Dermatoglyphics)

- ◆ Origin
  - From epidermal ridges (basal layer)

- ◆ Sites
    - Fingertips, palms, soles
  - ◆ Features
    - Genetically determined
    - Unique to each individual
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- ◆ Clinical Importance
    - Forensics
    - Genetic studies
    - Abnormal in chromosomal disorders (e.g., Down syndrome)
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### ◆ 3. Keratinization Disorders

- ◆ Ichthyosis
  - Excess keratinization → dry, scaly skin
  - Mostly:
    - Autosomal recessive
    - Some X-linked forms

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◆ Harlequin Fetus (Severe Form)

- Thick, fissured, armor-like skin
- Often fatal shortly after birth

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🔑 High Yield Summary

- Epidermis → ectoderm
- Dermis → mesoderm + neural crest
- Melanocytes → neural crest (3rd month)
- Periderm sheds → amniotic fluid
- Fingerprints → basal layer ridges
- Vernix → protection in utero

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◆ Hair Development

◆ Origin

- From epidermis (surface ectoderm)

- Arises as solid proliferations from the basal (germinative) layer
  - Grows downward into the dermis
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## ◆ Stages of Development

### 1. Hair Bud Formation

- Solid epidermal outgrowth penetrates dermis
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### 2. Hair Papilla Formation

- Terminal end invaginates → hair papilla
  - Contains mesenchyme with:
    - Blood vessels
    - Nerve endings
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### 3. Differentiation

Component	Derivative
Central cells	Elongate → keratinize → hair shaft
Peripheral cells	Cuboidal → epithelial root sheath

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## 4. Surrounding Mesenchyme

- Forms:
    - Dermal root sheath
    - Arrector pili muscle (smooth muscle)
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### ◆ Growth

- Continuous mitosis at base → pushes hair upward

### ◆ Timeline

- End of 3rd month → first hairs appear
    - Eyebrows
    - Upper lip
  - First hair = Lanugo
    - Fine, soft, unpigmented
    - Shed before/around birth
    - Replaced by terminal hair
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### ◆ Associated Structures

- ◆ Sebaceous Glands
  - ◆ Development
    - Buds from hair follicle epithelium
  - ◆ Mechanism
    - Central cells degenerate → form sebum
  - ◆ Function
    - Sebum enters hair follicle → reaches skin surface  
→ lubricates skin & hair
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### ◆ Clinical Correlates (Hair)

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- ◆ Hypertrichosis
  - Excessive hair growth
  - Cause → ↑ number of hair follicles
- ◆ Types

- Localized → e.g., lumbosacral region (spina bifida occulta marker)
  - Generalized → whole body
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### ◆ Atrichia

- Congenital absence of hair
  - Often associated with:
    - Tooth abnormalities
    - Nail defects→ (ectodermal dysplasia spectrum)
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## ◆ Sweat Glands Development

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### ◆ I. Eccrine Sweat Glands

#### ◆ Origin

- Buds from epidermal germinative layer

#### ◆ Development

- Extend into dermis → coil → form secretory portion
  - ◆ Features
    - Mode: Merocrine (exocytosis)
    - Function: Thermoregulation
    - Distribution: Whole body (except lips, parts of genitalia)
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## ◆ 2. Apocrine Sweat Glands

- ◆ Origin
  - From epidermal buds associated with hair follicles
- ◆ Location
  - Axilla
  - Pubic region
  - Areola
  - Perianal region
  - Face
- ◆ Features
  - Start functioning at puberty (hormonal)

- Open into hair follicles
  - Secretion contains:
    - Lipids
    - Proteins
    - Pheromones
  - Odor → bacterial breakdown
  - Mode: Apocrine (part of cytoplasm lost)
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#### ◆ Clinical Correlates

- No major embryological anomalies
  - Important clinically for:
    - Type of secretion
    - Function
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## ◆ Mammary Glands

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#### ◆ Nature

- Modified sweat glands

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- ◆ Early Development

- Form as mammary lines (milk lines):
    - Extend from axilla → inguinal region
  - Most regress
    - only thoracic region persists
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- ◆ Stages

1. Sprouting Stage

- 16-24 epithelial sprouts form
  - Later canalize → lactiferous ducts
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2. Nipple Formation

- Initially:
  - Ducts open into epithelial pit
- After birth:
  - Mesenchymal proliferation → eversion → nipple formation

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- ◆ At Birth

- Only duct system present
  - No alveoli
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- ◆ At Puberty (Female)

- Under estrogen & progesterone:
    - Duct branching
    - Formation of alveoli & secretory units
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- ◆ Clinical Correlates

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- ◆ Polythelia (Most Common)

- Accessory nipples
  - Due to persistence of mammary line fragments
  - Usually along milk line (often axilla)
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## ◆ Polymastia

- Accessory breast tissue along mammary line
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## ◆ Inverted Nipple

- Failure of epithelial pit to evert
  - Clinical relevance:
    - May mimic pathological inversion
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## 🔑 Summary and High Yield Points

- Hair → ectodermal origin + mesenchymal papilla
- Lanugo → first fetal hair
- Sebaceous glands → from hair follicle epithelium
- Eccrine → merocrine, thermoregulation, all over body
- Apocrine → puberty, hair follicle, odor-producing
- Mammary gland → milk line origin
- Ducts → prenatal | Alveoli → puberty
- Lanugo = first fetal hair
- Apocrine glands → start at puberty
- Polythelia = most common mammary anomaly

- Milk line extends axilla → groin

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-> The End <-