

Apparato Tegumentario

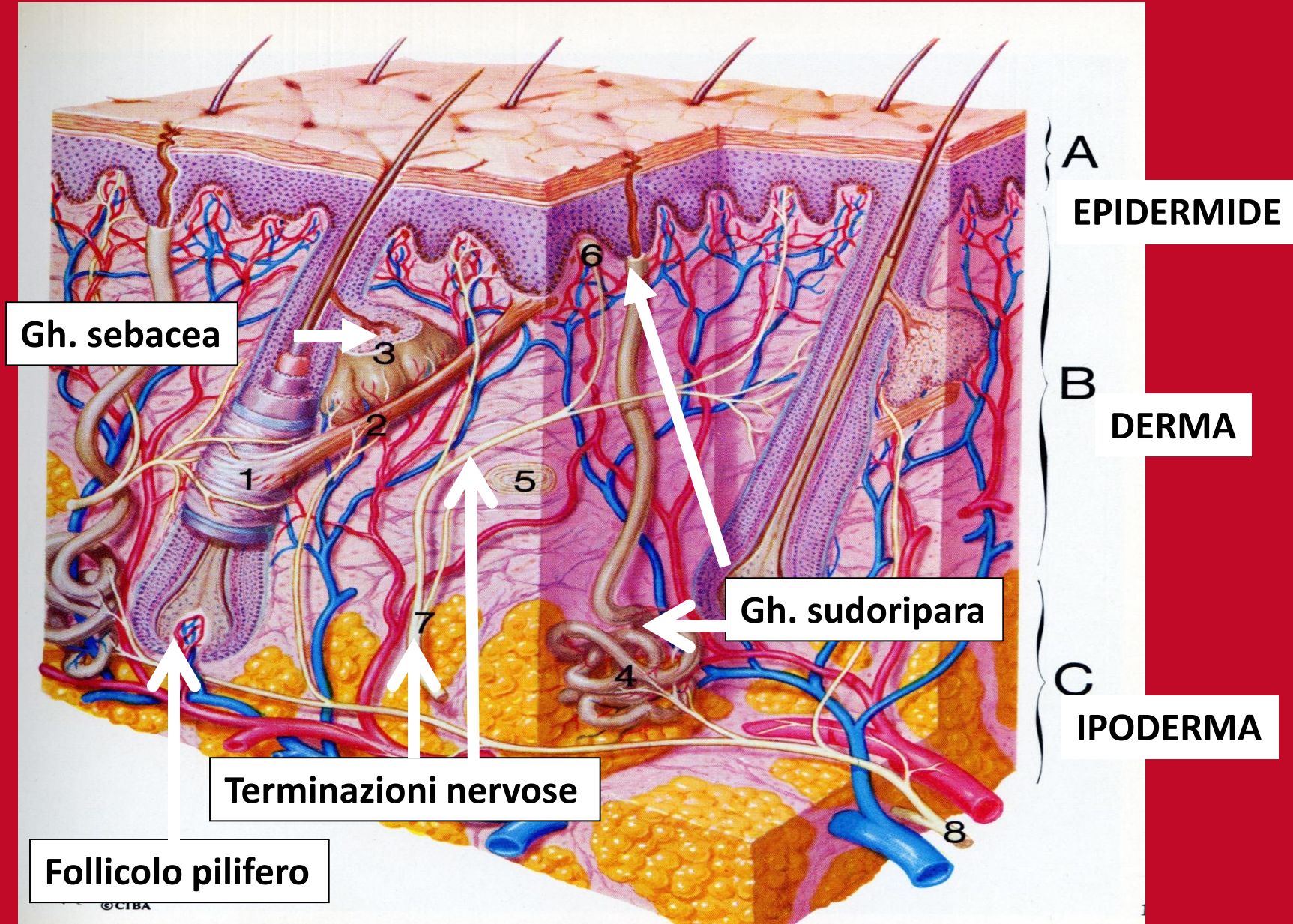
Cute

Epidermide, derma e tela sottocutanea

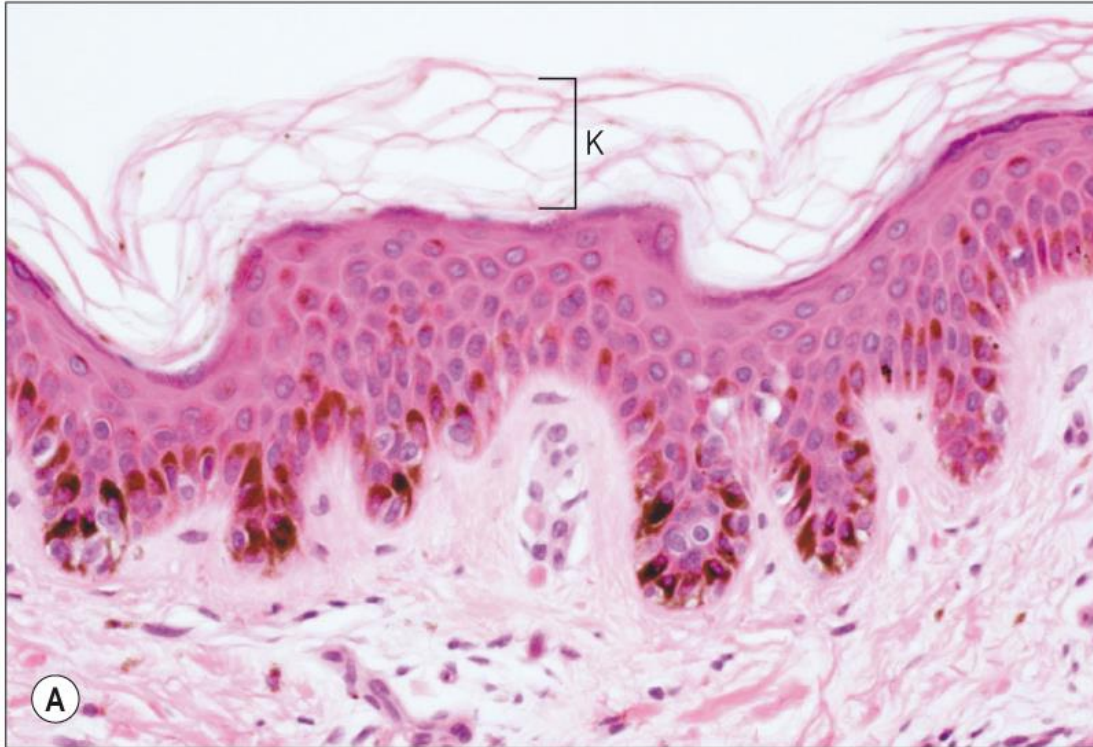
Annessi cutanei

Formazioni cornee (peli e unghie) e annessi ghiandolari (ghiandole sudoripare e sebacee, ghiandola mammaria)

Struttura cutanea



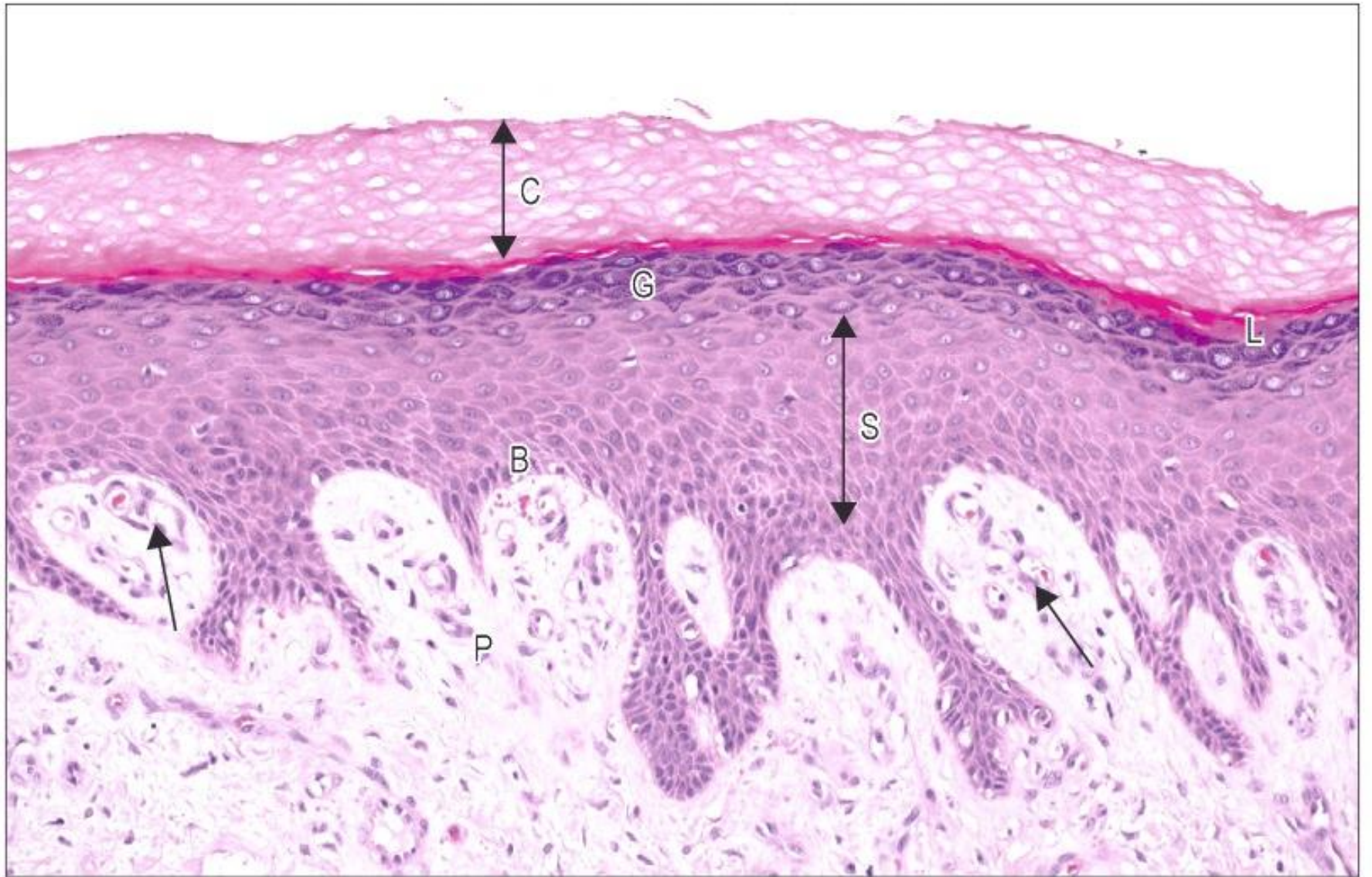
**Strato
corneo**
↑
**Strato
lucido**
↑
**Strato
granuloso**
↑
**Strato
spinoso**
↑
**Strato
basale**

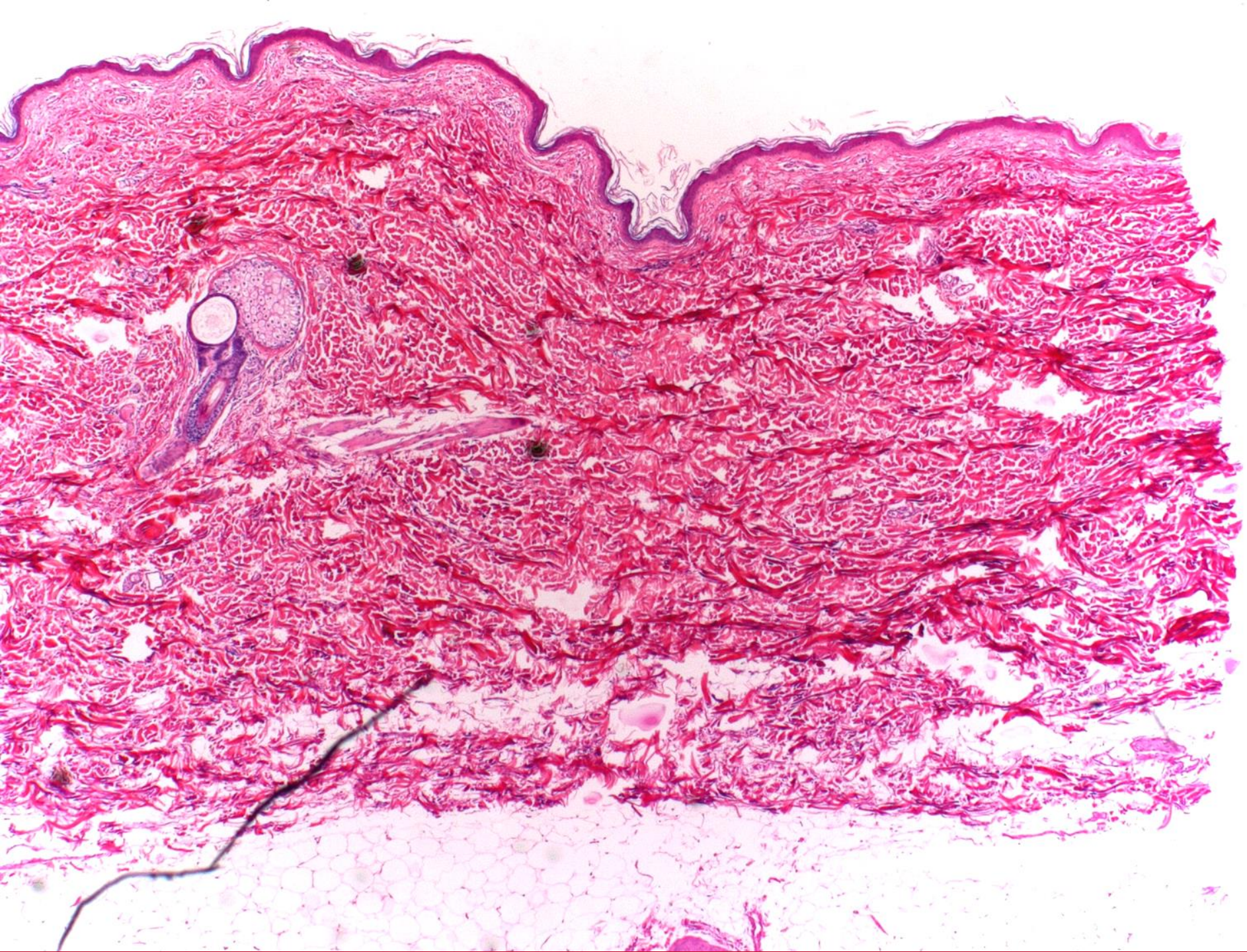


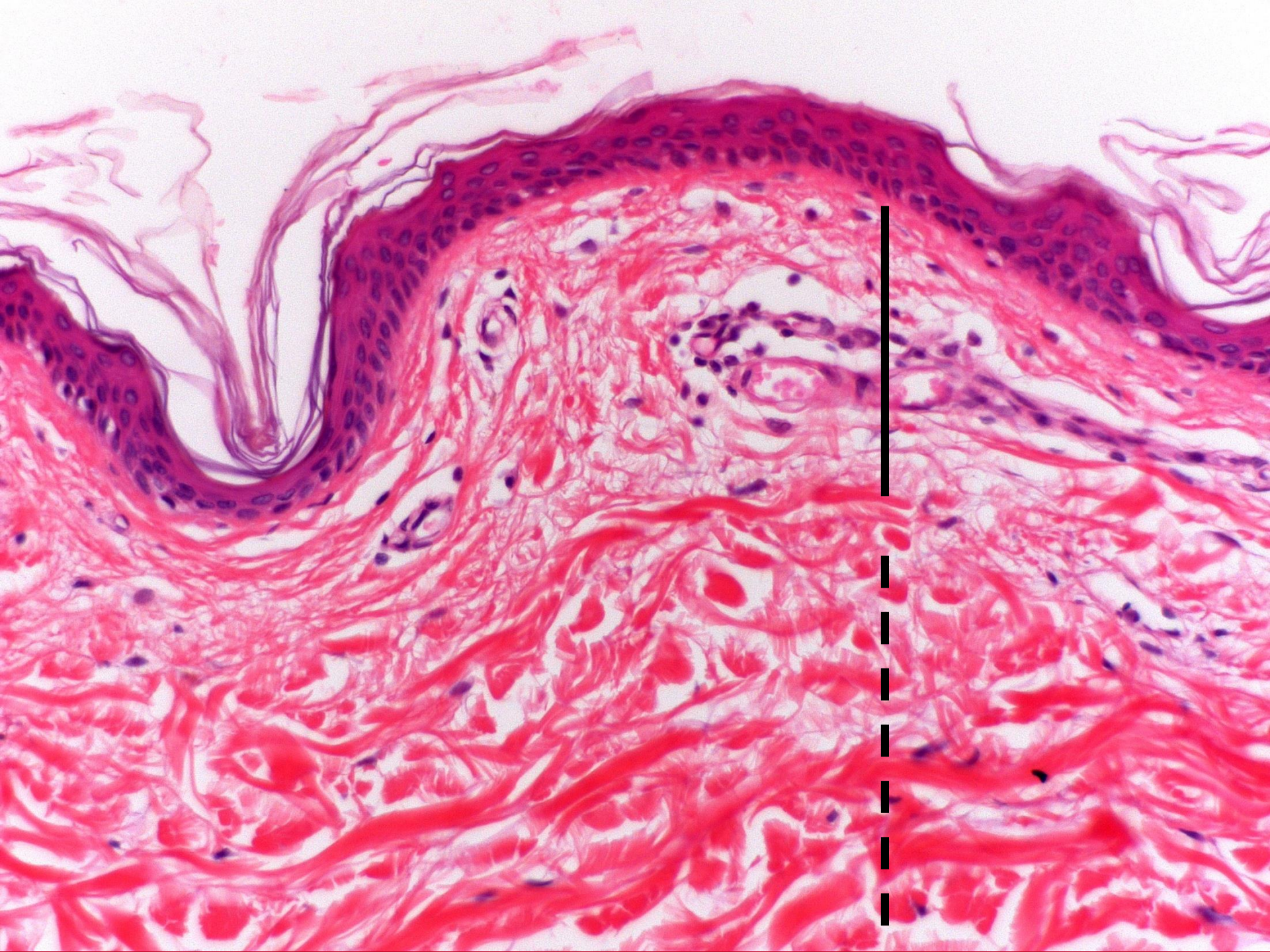
Nell'epidermide si trova inoltre un particolare tipo di cellule:

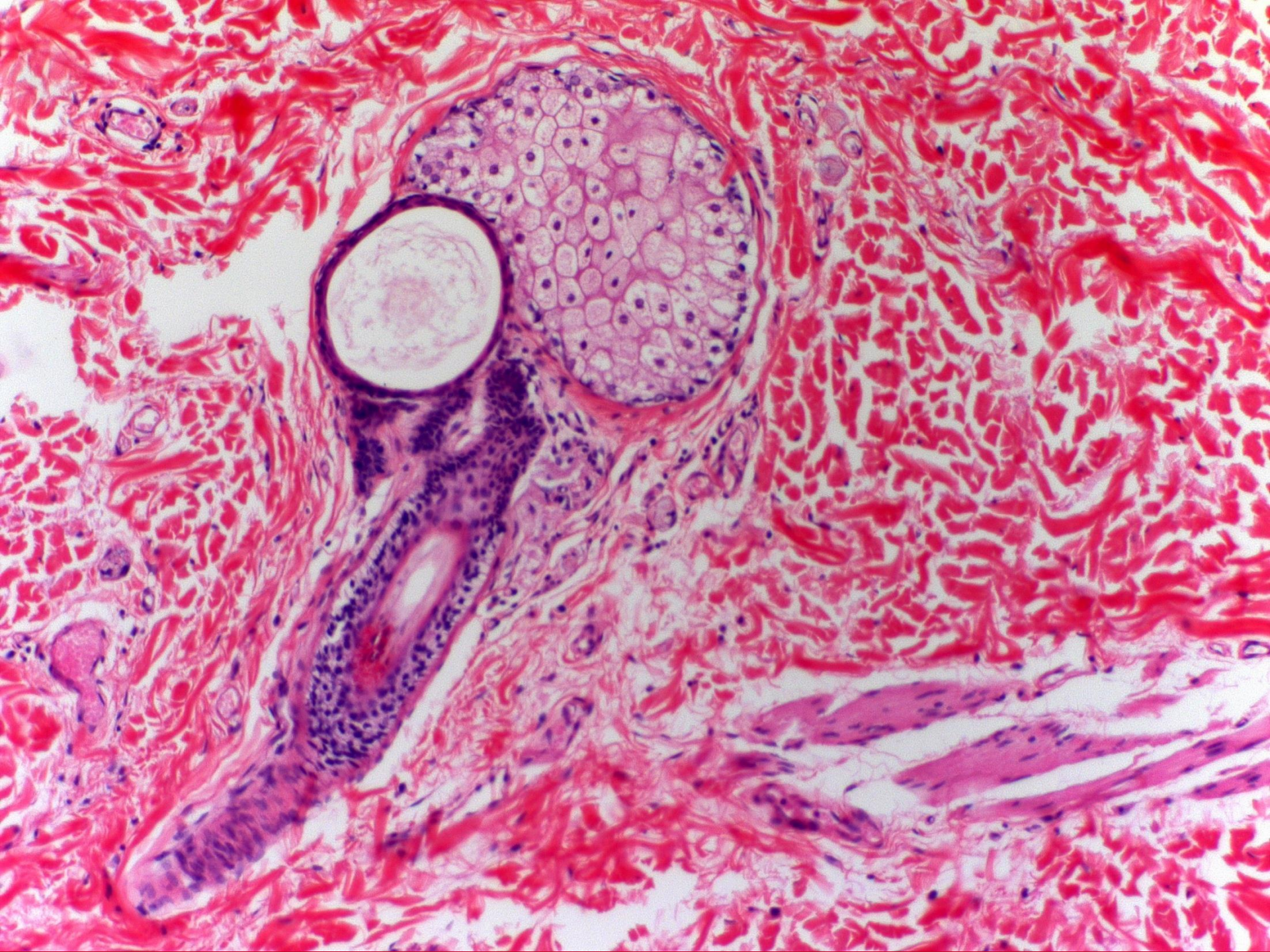
i melanociti

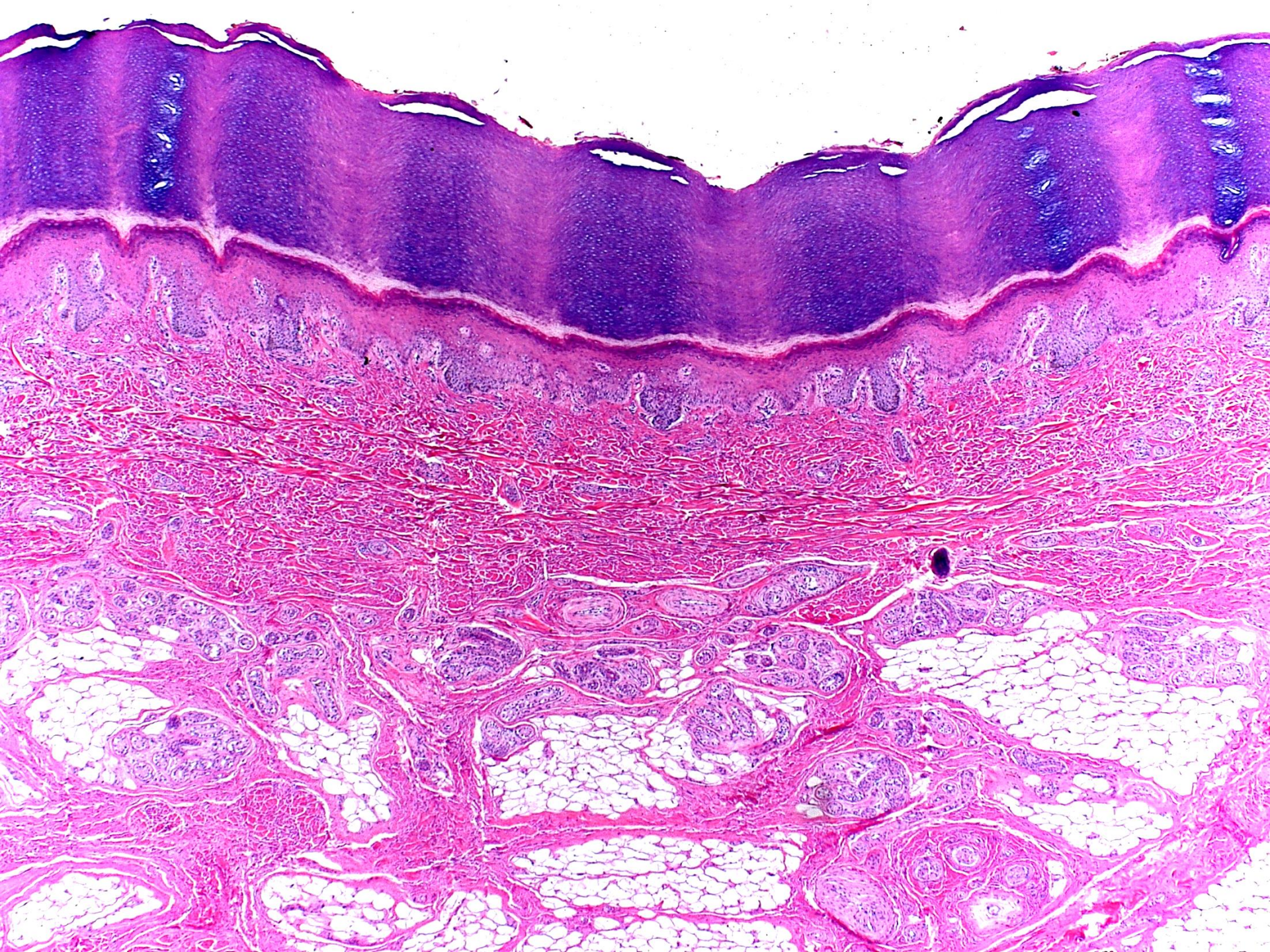
che elaborano un particolare pigmento, la melanina.

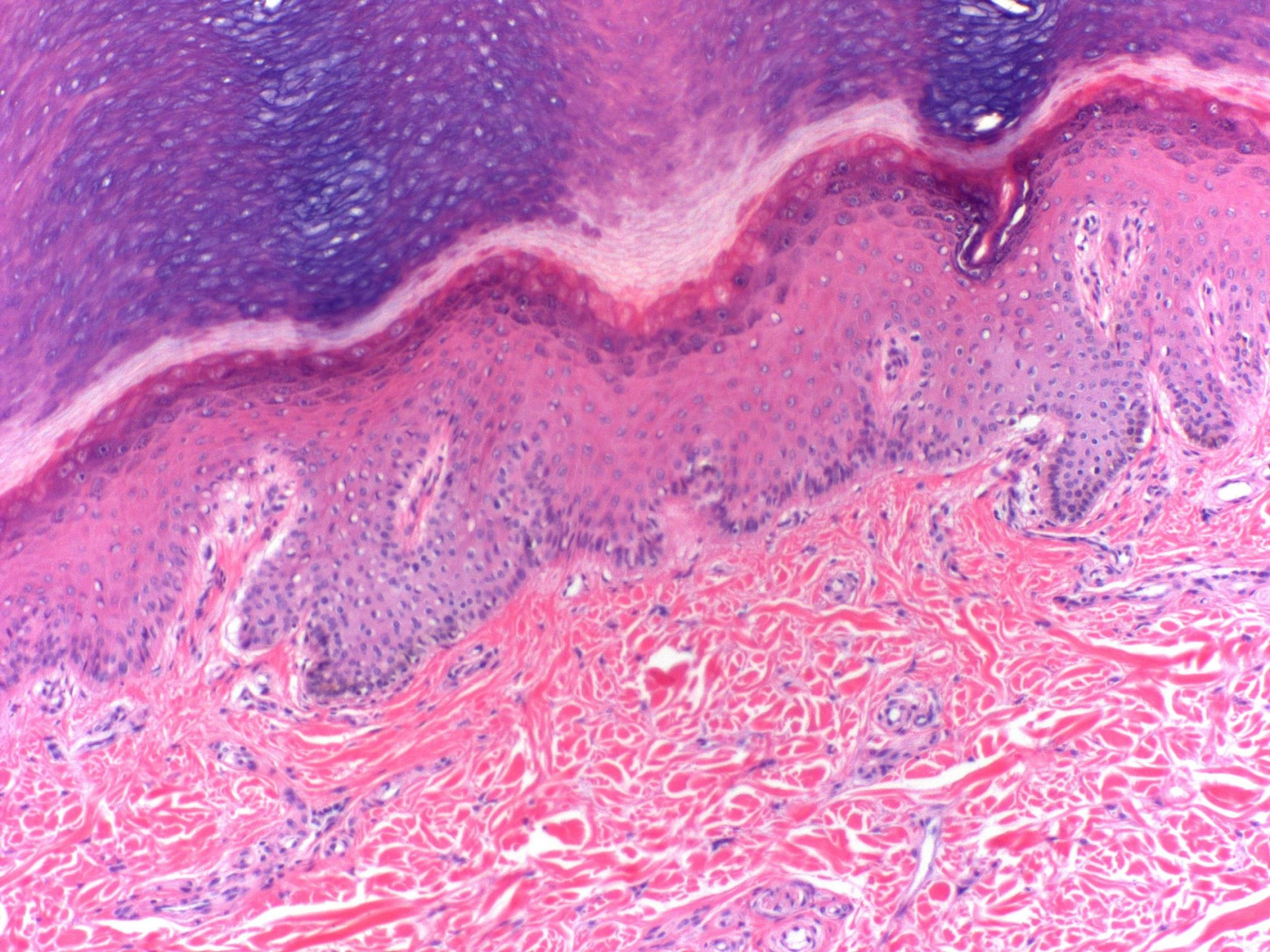


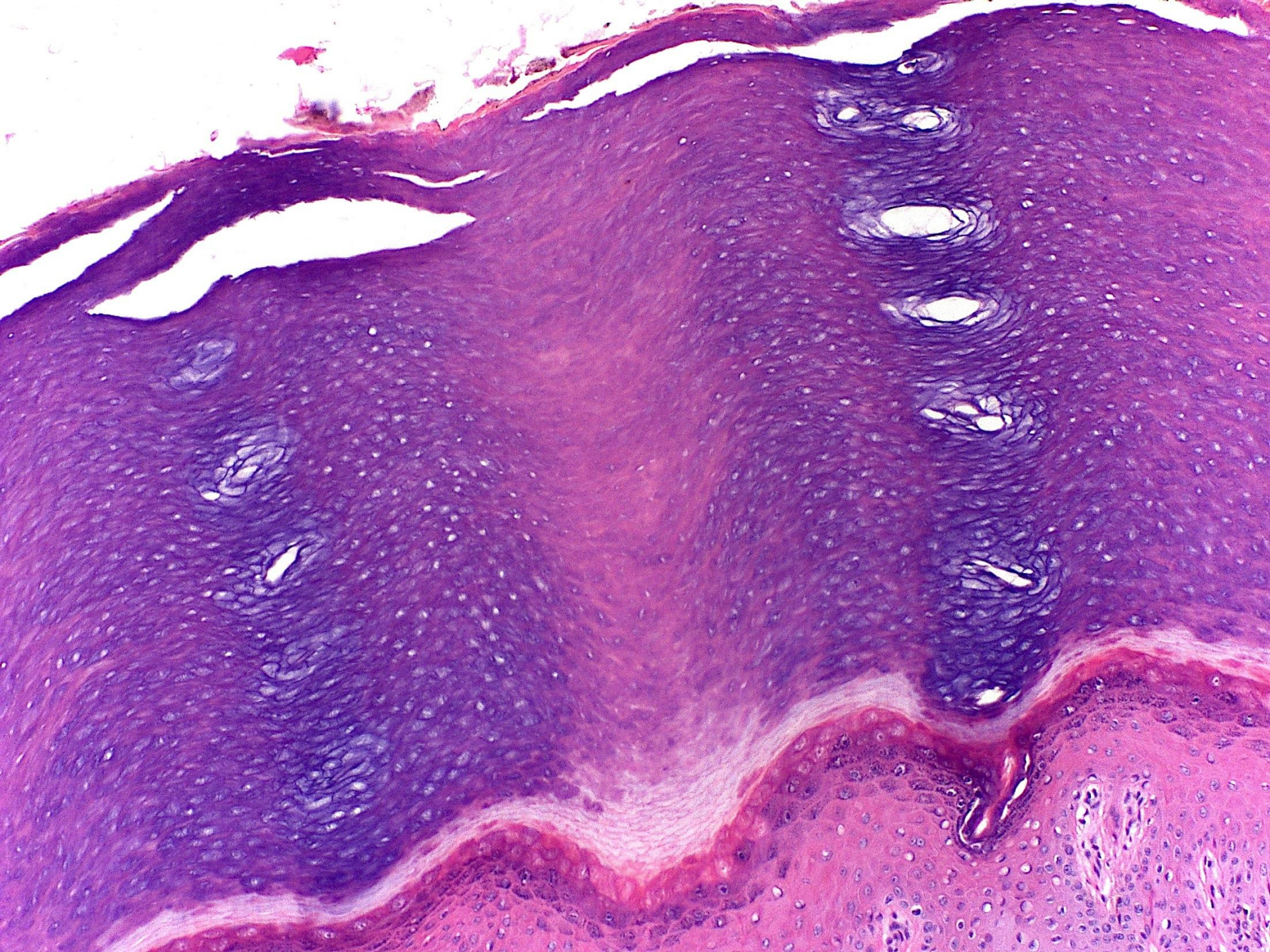


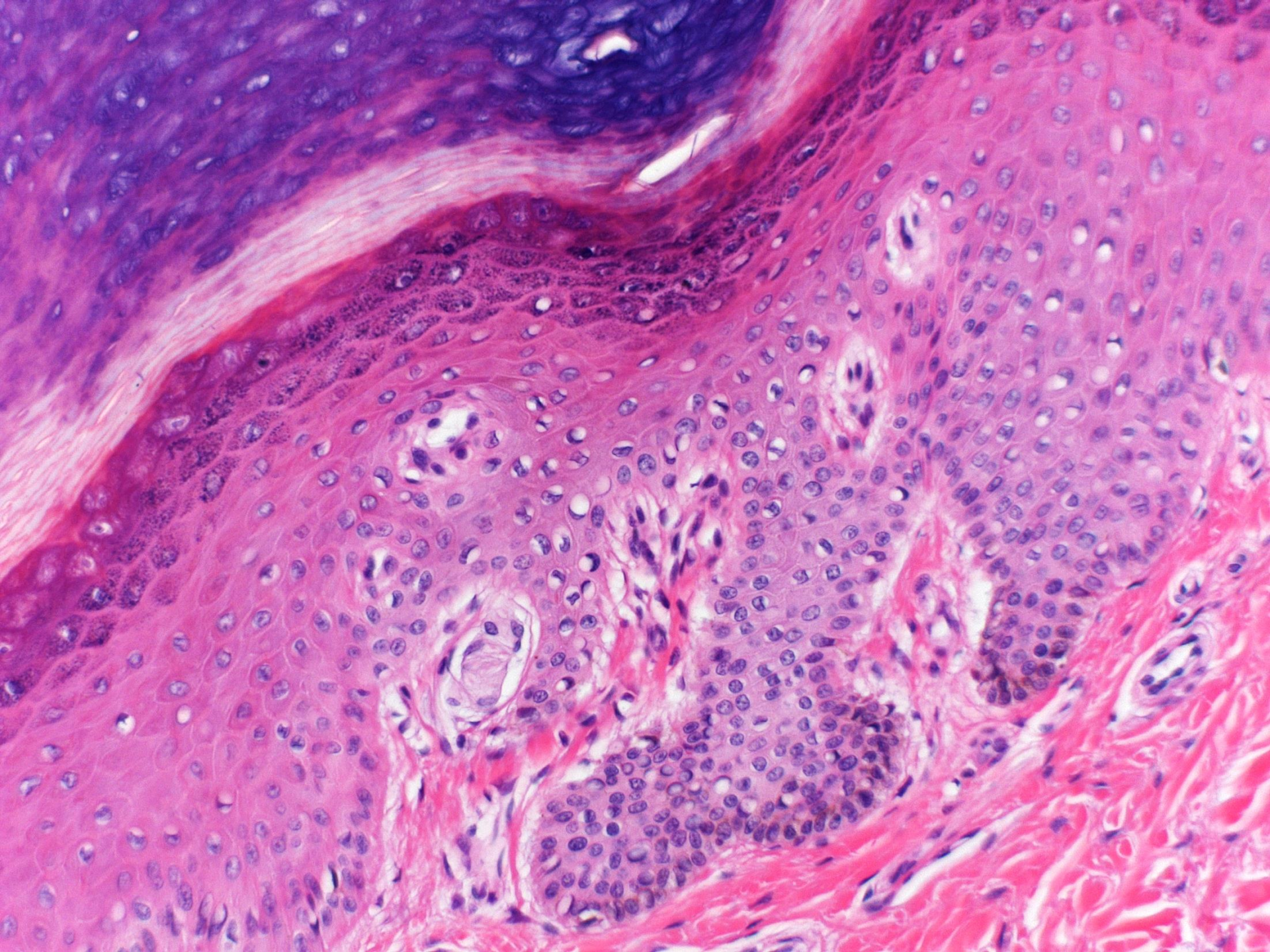




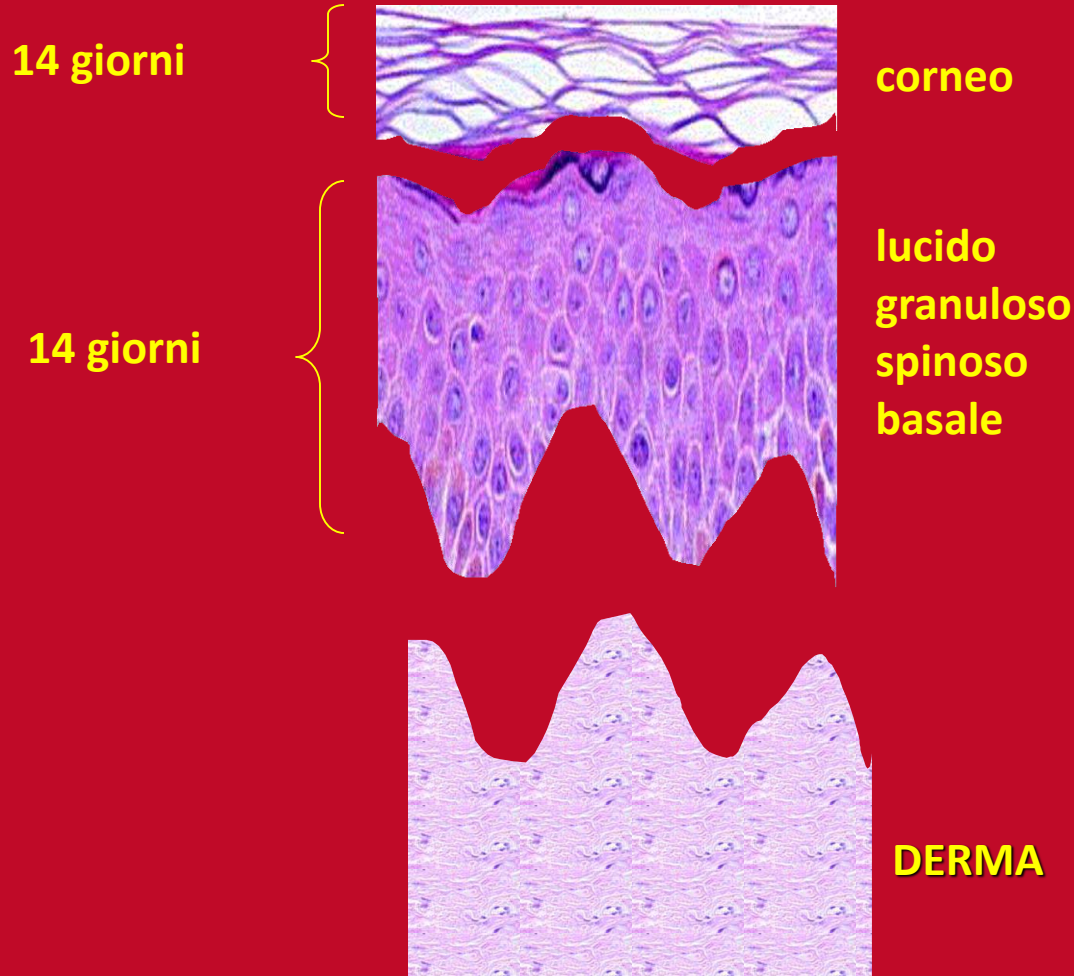








Turnover Epidermico (28 giorni)



- L'epidermide è un epitelio dinamico che si rinnova continuamente per tutta la vita

- Questo rapido turnover nell'adulto richiede la presenza di stem cell capaci di dare origine a cellule mature

Funzioni della cute

secretorie

- cheratinopoesi
- pigmentogenesi
- secr. sebacea
- secr. sudorifera

protettive

(flora residente)

- immunologiche
- meccaniche
- caloriche
- chimiche
- microrganismi

sensoriali

- meccanica
- dolore, prurito
- termorecezione

Localizzazione delle stem cells nella pelle

Epidermide: strato basale (organizzazione in clusters)
strato spinoso e granulare
follicoli piliferi (porzione esterna)

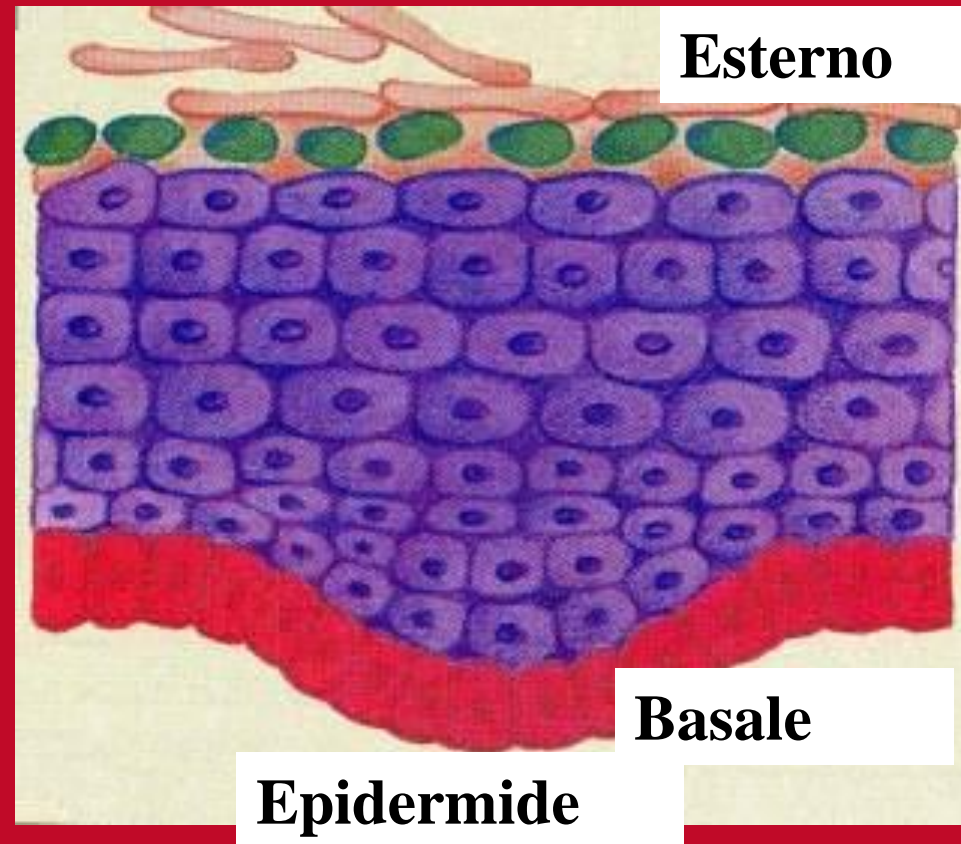
Derma: follicoli piliferi
cellule non follicolari

Le stem cells del derma possono dare origine:

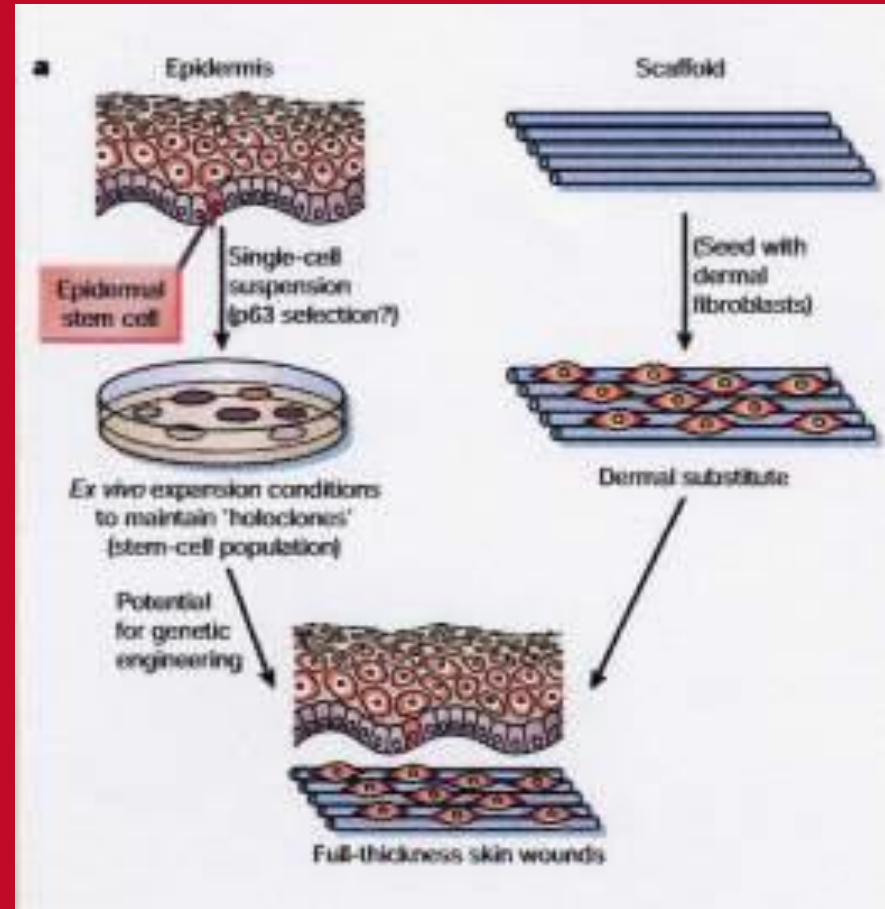
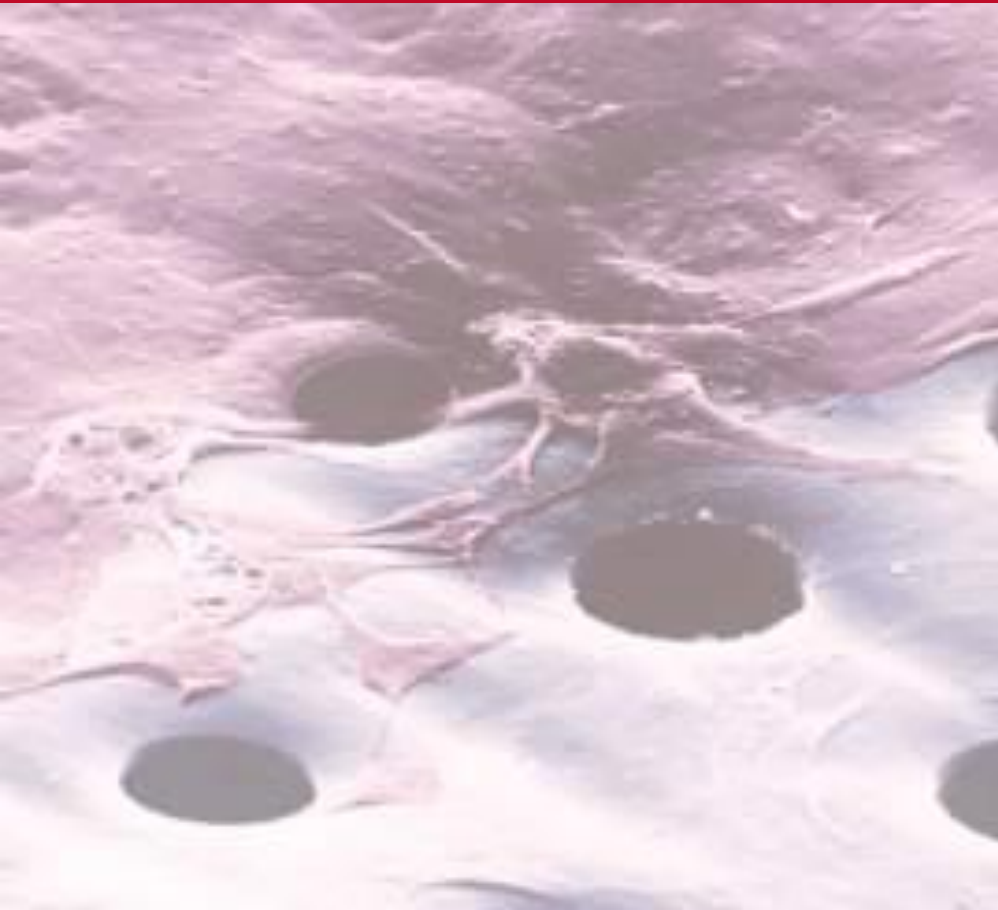
- cellule neuronali (neuroni e glia)
- derivati del mesoderma (osteociti, condrociti, cellule muscolari e adipociti)
- cellule ematopoietiche

Cellule Staminali dell'epidermide

Lo strato basale, contiene le cellule proliferanti che migrano verso gli strati superiori dove vanno incontro a differenziamento che porta alla formazione di cellule terminali prive di nucleo.



Cellule staminali ed ingegneria dei tessuti



- fibrina

+ fibrina

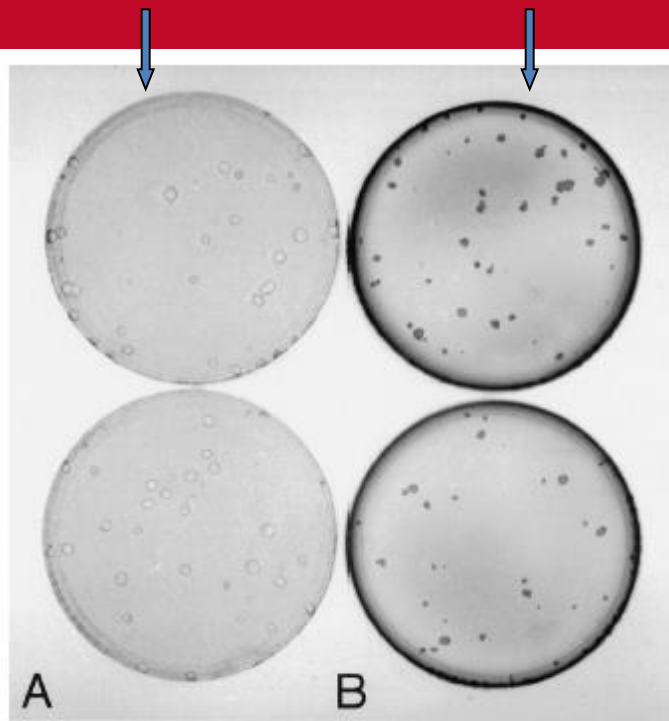


FIGURE 2. Colony-forming efficiency of human keratinocytes grown on a fibrin matrix. One hundred human keratinocytes (strain YF29, culture V) were plated onto 100-mm Petri dishes containing lethally irradiated 3T3 cells, in the absence (A) or presence (B) of a fibrin matrix. Cells were cultured for 12 days and were then fixed and stained with Rhodamine B (A) or Nile Blue (B). The presence of a fibrin matrix did not affect the clonogenicity of the keratinocytes. Similar numbers of colonies (54) were obtained with or without the fibrin matrix. A similar result was obtained with other strains of human keratinocyte. Colonies cultured on a fibrin matrix were less regular in shape and smaller than the control colonies.



FIGURE 1. Appearance of a cultured epithelium grown on a fibrin matrix. A cultured epithelium grown on a fibrin matrix was detached from the culture dish using two thin forceps. Note its transparency, its molding properties and the ease of handling.

LAMINE DI CHERATINOCITI

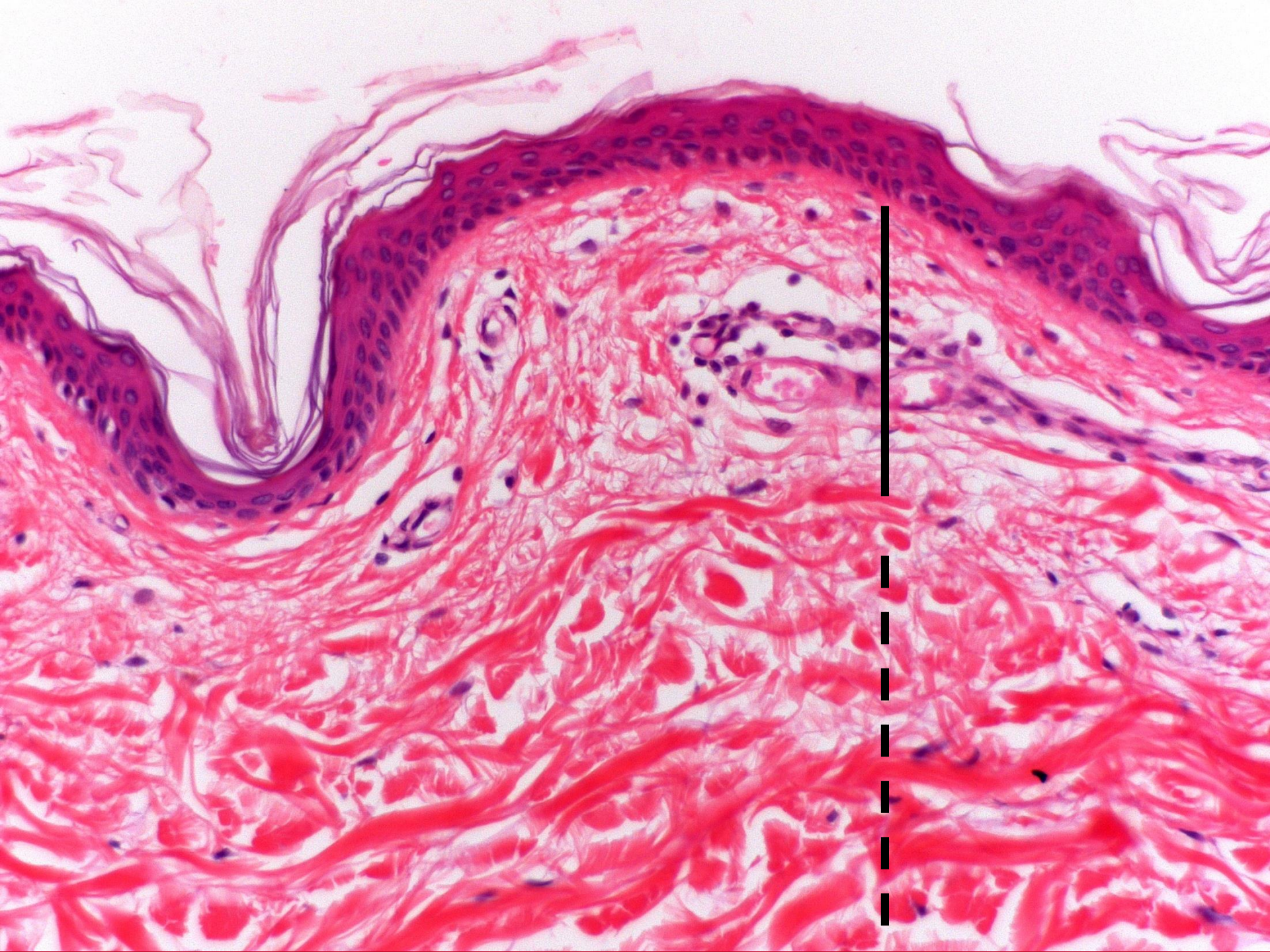
- **La sopravvivenza cellulare è direttamente correlata con le condizioni del letto ricevente.**

- **In particolare, la presenza di uno strato dermico ne incrementa la sopravvivenza**

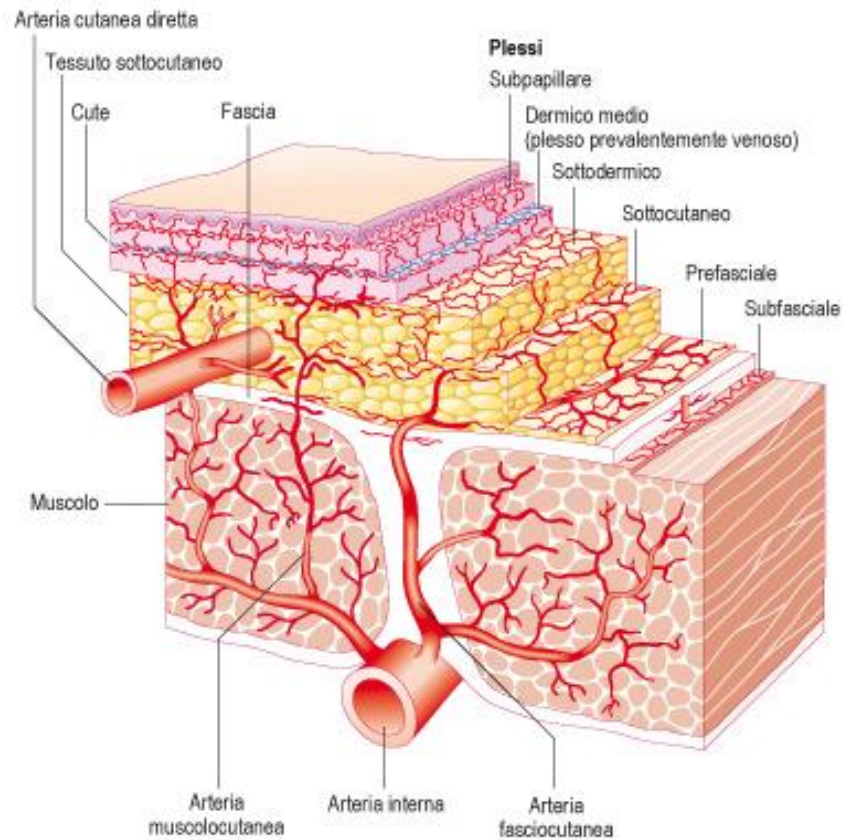
Fibroblasti

ECM

Micro-vascolarizzazione



A



B

