



Peels

**CLASSIFICATION OF A.TENENBAUM
AIME-AESTHETIC MEDICINE
02-2026**



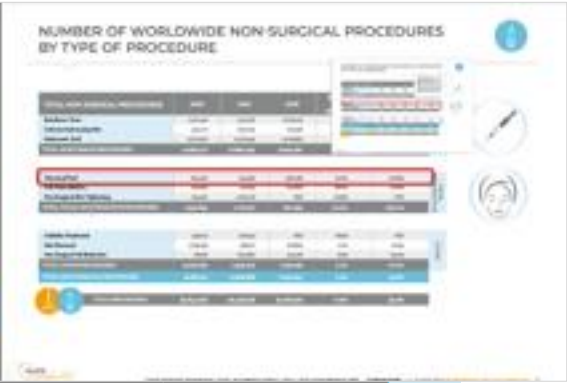
A.TENENBAUM, M.D., Ph.D., D.Sc
M.TIZIANI, RCSA



NUMBER OF WORLDWIDE NON-SURGICAL PROCEDURES BY TYPE OF PROCEDURE



TOTAL NON-SURGICAL PROCEDURES	2022	2021	2018	Pct. Change 2022 vs 2018
Botulinum Toxin	9,221,419	7,312,616	6,097,516	
Calcium Hydroxylapatite	350,716	290,095	129,038	
Hyaluronic Acid	4,312,037	5,279,344	3,729,833	
TOTAL INJECTABLES PROCEDURES	13,884,172	12,882,055	9,956,387	



Chemical Peel	844,616	534,831	408,485	57.9%	106.8%
Full Field Ablative	367,983	231,955	192,880	58.6%	90.8%
Non-Surgical Skin Tightening	734,257	1,003,731	N/A	-26.8%	N/A
TOTAL FACIAL REJUVENATION PROCEDURES	1,946,855	1,770,517	601,365	10.0%	223.7%



Cellulite Treatment	449,314	379,224	N/A	18.5%	N/A
Hair Removal	1,798,253	1,836,111	916,869	-2.1%	96.1%
Non-Surgical Fat Reduction	778,716	730,980	473,316	6.5%	64.5%
TOTAL OTHER PROCEDURES	3,026,284	2,946,316	1,390,185	2.7%	117.7%
TOTAL NON-SURGICAL PROCEDURES	18,857,311	17,598,888	11,947,937	7.2%	57.8%



TOTAL PROCEDURES	33,844,293	30,439,576	23,266,375	11.2%	45.5%
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What Is a Chemical Peel?

A clinical technique — not merely a skin flaking process

A chemical peel is a controlled clinical technique designed to

Improve and smooth facial and /or body skin structure through the application of a chemical solution.

Desquamation is not mandatory.

Clinical efficacy may occur with or without visible peeling.

- Progressive elimination of altered skin layers
- Regenerated skin appears smoother, more uniform, and less wrinkled

[Explore the Science of Chemical Peels >](#)



Le **pKa** peut être comparé
à un agresseur qui porte
un coup de poing.

La biologie représente la victime,
et l'outcome clinique
correspond aux conséquences finales —
bénignes, sévères ou incapacitantes.

Un coup de poing porté avec la même force n'aura
pas les mêmes conséquences biologiques ni cliniques
selon qu'il touche les fesses, le nez ou le sternum.

Ce n'est pas le coup en lui-même qui détermine
la gravité, mais le tissu touché, sa **structure**, sa
capacité d'adaptation et sa réponse biologique.

La biologie représente la victime, et l'outcome
clinique correspond aux conséquences finales —
bénignes, sévères ou incapacitantes.

pKa can be compared to
an aggressor delivering a punch.

Biology represents the victim, and
the clinical outcome reflects the
final consequences — mild,
severe, or incapacitating.

A punch delivered with the same force will **not**
produce the same biological effects or clinical
outcomes if it strikes the buttocks, the
nose, or the sternum.

Severity is **not** determined by the punch itself,
but by the tissue involved, its
structure, vulnerability, and biological response.

Pthe same way, **pKa** describes the chemical aggression,
but clinical efficacy and risk are determined
by tissue biology, not by chemical strength alone.

SCIENTIFIC PUBLICATIONS OF INTEREST



**THE ISSUES
CAUSED BY**

**ENDOWED
PERIODICALS**



BIG PHARMA



BUREAUCRATS



Endowed Periodicals

The issues caused by Endowed Periodicals

LIST OF ENDOWED PERIODICALS	
JOURNAL TITLE	ENDOWED BY
Developmental Medicine and Child Neurology	Mrs. Alex. Comfort, in memory of Dame Eileen Younghusband D.B.E., J.P.
European Journal of Clinical Pharmacology	Lilly Industries
Gut	Dr. A.H. James
Human Nutrition - Applied Nutrition	Mars Health Education Fund
Human Nutrition - Clinical Nutrition	Mars Health Education Fund
Immunology Today	Mr. N. Asherson
International Archives of Occupational and Environmental Health	Dr. E. T. Ruston
Journal of Investigative Dermatology	Stiefel Laboratories (UK) Limited
Journal of Laryngology and Otolaryngology	Mr. N. Asherson
Journal of Lipid Research	Bristol Myers Co. Limited

Archives of Dermatology	Stiefel Laboratories (UK) Limited
Archives of Environmental Health	Dr. E. T. Ruston
British Heart Journal	Boehringer Ingelheim Limited
British Journal of Clinical Pharmacology	Anonymous
British Journal of Dermatology	E.R. Squibb & Sons Limited
British Journal of Industrial Medicine	Dr. E. T. Ruston
British Journal of Ophthalmology	Beresford & Betty Hall-Parker
British Journal of Pharmacology	Leo Laboratories Limited
British Journal of Psychiatry	E.R. Squibb & Sons Limited
British Journal of Rheumatology	Air Commodore D. Stevenson
British Journal of Surgery	Henry Blacow Yates
Bulletin of the History of Medicine	Dr. A.S. Thorley
Cardiology	Florence Jackson Legacy
Cardiovascular Research	Florence Jackson Legacy
Caries Research	Mars Health Education Fund
Clinical Materials	Porter Nash Limited
Clinics in Developmental Medicine	Mrs. Alex. Comfort, in memory of Dame Eileen Younghusband D.B.E., J.P.

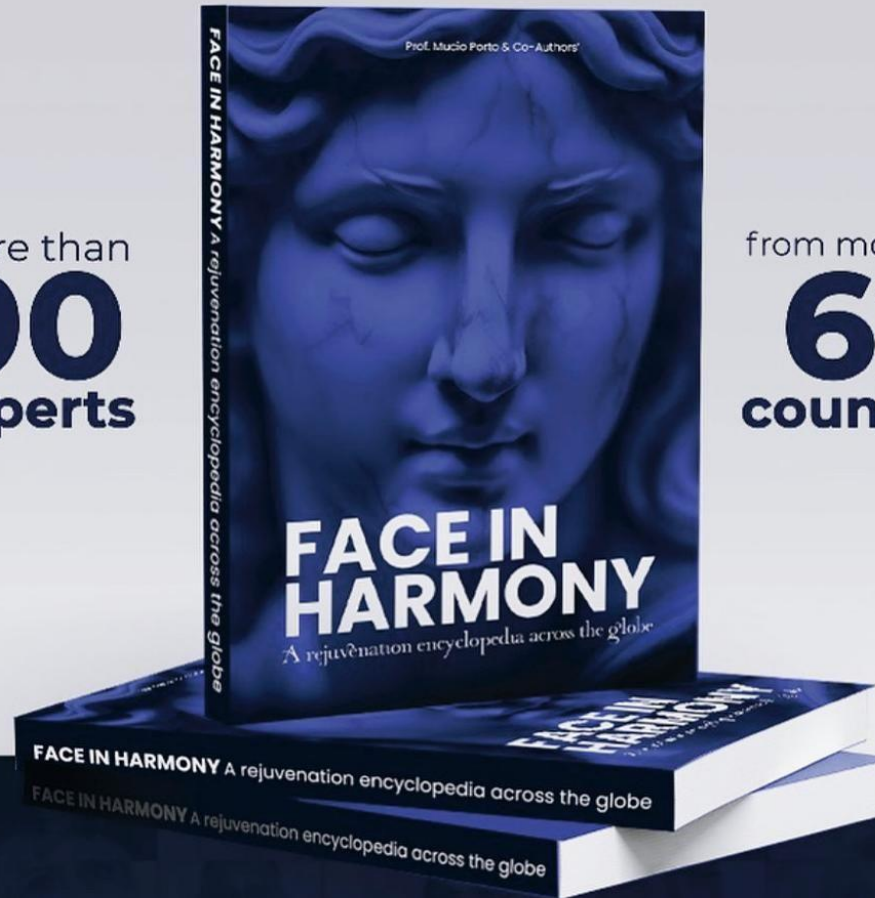
Endowed scientific periodicals, while often seen as a mark of quality, still require critical evaluation for objectivity. Their funding structure, while providing financial stability, can potentially influence editorial decisions and the types of research published. Objectivity in scientific publications means presenting facts and evidence without bias, which can be a challenge for any publication, regardless of funding



FACE IN HARMONY

A rejuvenation encyclopedia across the globe

more than
90
experts



from more than
62
countries

More than 1000 pages of studies, knowledge and
Techniques on Facial Rejuvenation across the globe



PROCEDURES IN COSMETIC DERMATOLOGY
Series editor: Jeffrey S. Dover
Associate editor: Murad Alam

Chemical Peels

Edited by Rebecca C Tung
Mark G Rubin

SAUNDERS
ELSEVIER

2ND EDITION



DERMATOLOGÍA ESTÉTICA
Editor de la serie: Jeffrey S. Dover
Editor asociado: Murad Alam

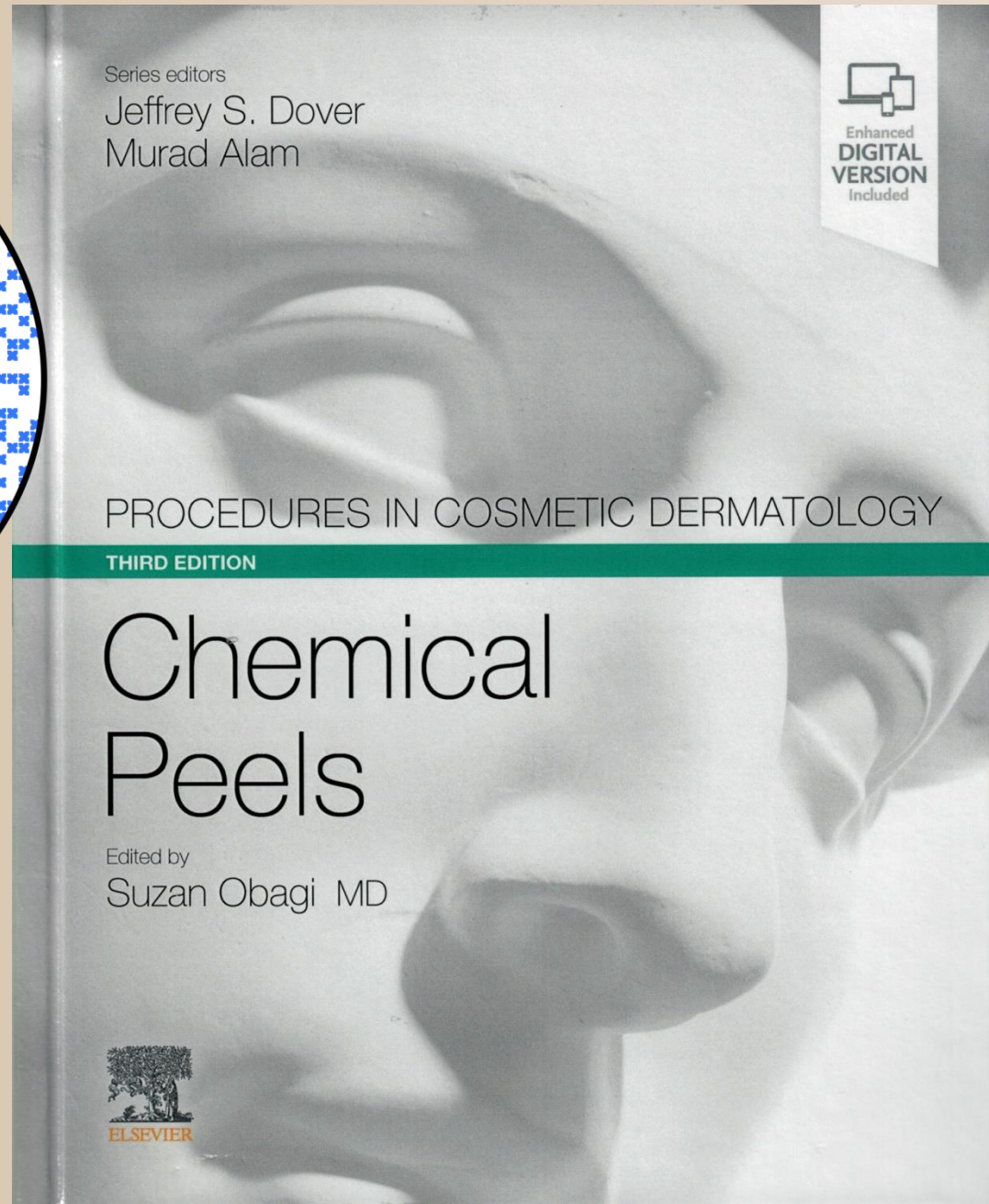
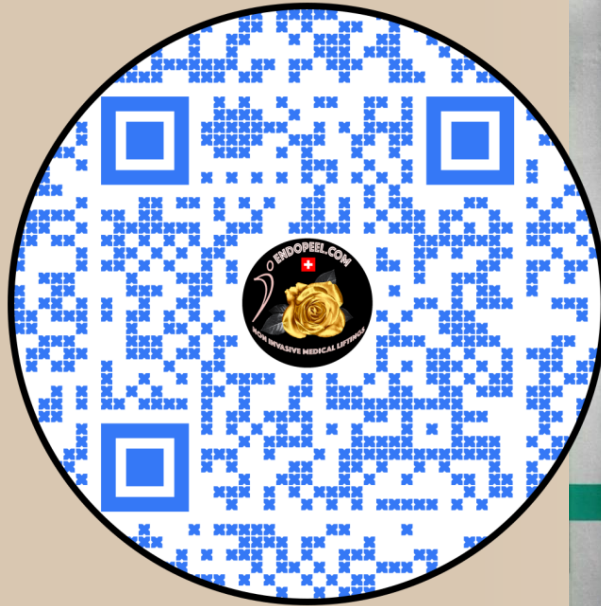
Exfoliación química

Editado por Rebecca C. Tung
Mark G. Rubin



2.^a
EDICIÓN






Why Perform Chemical Peels?

Peelings IMPROVE results of surgery, as non invasive procedures



Peelings make you different from other colleagues thinking just to do invasive, mini invasive, or non invasive treatments



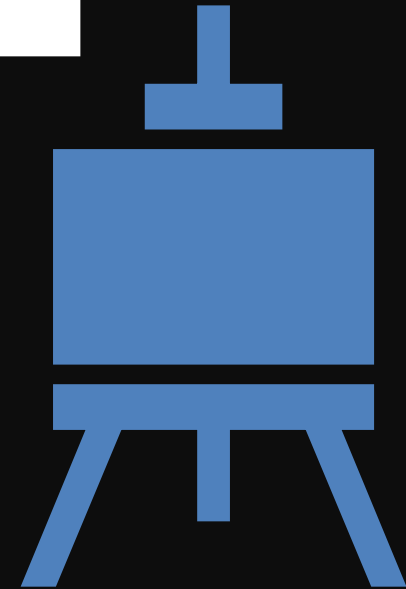
Peelings will bring you more patients



Patients judge us on their outlook.

TO WAKE UP
YOUR INTEREST

Peels Gallery



Take a Look

My personal case

I am not better , may be not so good as other surgeons in surgical procedures

But I can compete with other colleagues for my post surgical results thanks to Peelings

Surgical Rhinoplasty + Metabolic Peels

BEFORE

AFTER

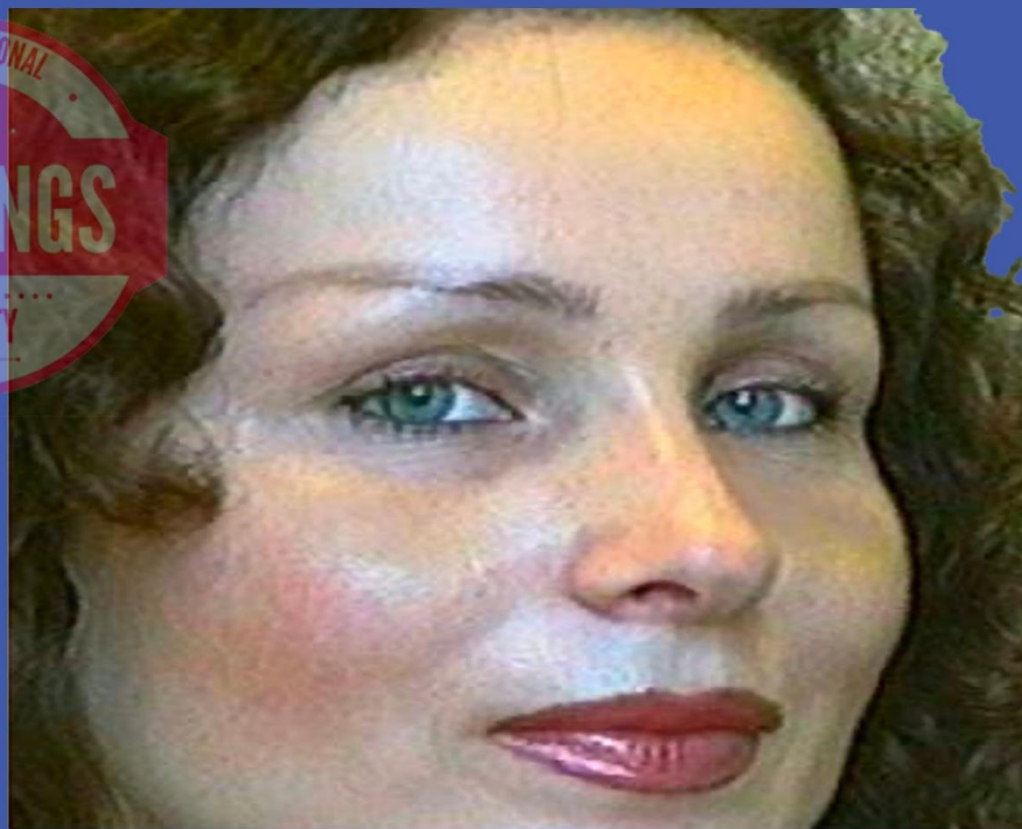


Courtesy of Dr. Alain Tenenbaum

Surgical Rhinoplasty + Metabolic Peels

BEFORE

AFTER



Courtesy of Dr. Alain Tenenbaum

Lipoplasty + Facelift + Blepharoplasty + Endo Peel + Metabolic Peels

BEFORE

AFTER



Courtesy of Dr. Alain Tenenbaum

Lipoplasty + Facelift + Blepharoplasty + Endo Peel + Metabolic Peels

BEFORE

AFTER



Courtesy of Dr. Alain Tenenbaum

Facelift + Blepharoplasty + Endo Peel + Metabolic Peels

BEFORE

AFTER



Courtesy of Dr. Alain Tenenbaum

Why Chemical Peels Must Be *Reconsidered Today*

Historical Paradigm

Chemical peels historically defined by:
-acid type,
-concentration, and
-depth

Current Limitation

Visible reaction
(erythema,
desquamation)
often mistaken for
therapeutic efficacy

Emerging Question

Does visible tissue injury truly
reflect
biological skin improvement?

For decades, we have evaluated peels by what we can see on the surface.
Today, the question is whether visible damage is still a valid marker of efficacy.

IMPORTANT TO KNOW

BOTH ARE PEELS

COMBINATION IS POSSIBLE

peel-off-flake-off
NO DESQUAMATION

30 Min Peel Off
No desquamation



Courtesy of Dr. Alain Tenenbaum



DO'S

Emerging paradigm
NO COMPLICATION

SLOUGH OFF
WITH DESQUAMATION

Desquamation 7 days after TCA on Asian Skin



Courtesy of Dr. Alain Tenenbaum & Mauro Tiziani



DON'TS

Traditional paradigm
HUGE RISK OF COMPLICATIONS

From Chemical Injury to Biological Regulation

Traditional Paradigm

- Chemical aggression
- Tissue injury
- Repair driven by inflammation
- Efficacy inferred from visible damage

The skin does not improve
because it is injured,
but because it adapts biologically.

Emerging Paradigm

- Biological signaling
- Metabolic adaptation
- Regulated inflammatory response
- Efficacy driven by cellular pathways

Historically, we relied on injury
to trigger repair.
Today, we understand that skin
quality improves when biological
regulation is restored, not
when damage is maximized.

DESQUAMATION DAY = DANGEROUS DAY STOP SOCIAL EVICTION (DOWNTIME) WITH METABOLIC PEELS



No cohesion of corneocytes

Easy Penetration of chemicals
Radiations etc

Damages following peelings
procedures are mostly dued
when patient is **at home** at
the moment of desquamation

Desquamation 7 days after TCA on Asian Skin



Courtesy of Dr.Alain Tenenbaum & Mauro Tiziani

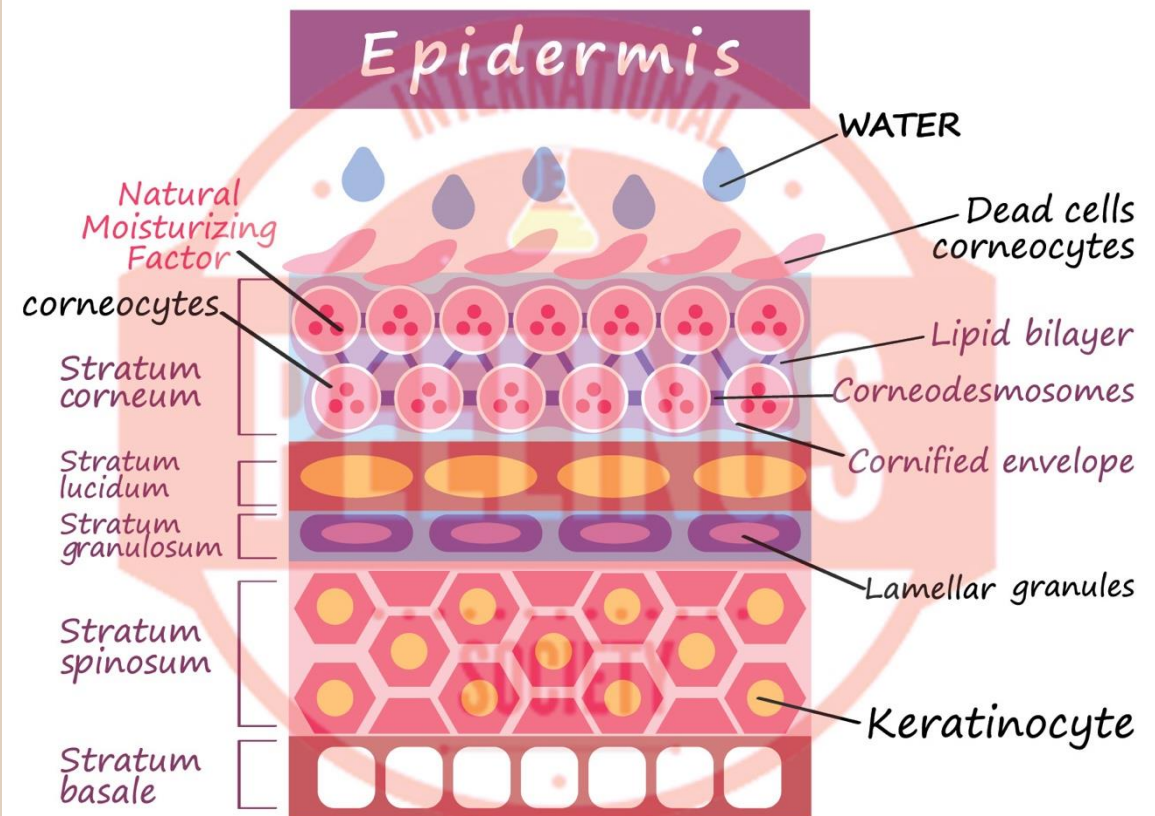


DESQUAMATION DAY = DANGEROUS DAY

ROLE OF CORNEOCYTES



Desquamation 7 days after TCA on Asian Skin



Courtesy of Dr. Alain Tenenbaum & Mauro Tiziani

DESQUAMATION DAY RECOMMENDATIONS

- No tap water, no mineral water for the skin
- Do not use alcohol or alcohol-based creams (like many sunscreens)
- Do not use hydroalcoholic solutions
- Do not use creams that contain metal ions (tattoo vs hyperchromia).





DESQUAMATION DAY
USE ONLY
DEMINERALIZED WATER



Two Distinct Concepts: Do Not Confuse Them

Metabolic Pathways

biological

- Endogenous biological processes
- Cellular energy regulation
- Signaling cascades
- Inflammatory modulation
- Tissue adaptation

Acidic Agents

chemical

- Exogenous chemical compounds
- Defined by pKa and formulation
- Encapsulated AHA
- True retinoic acid
- Delivery systems

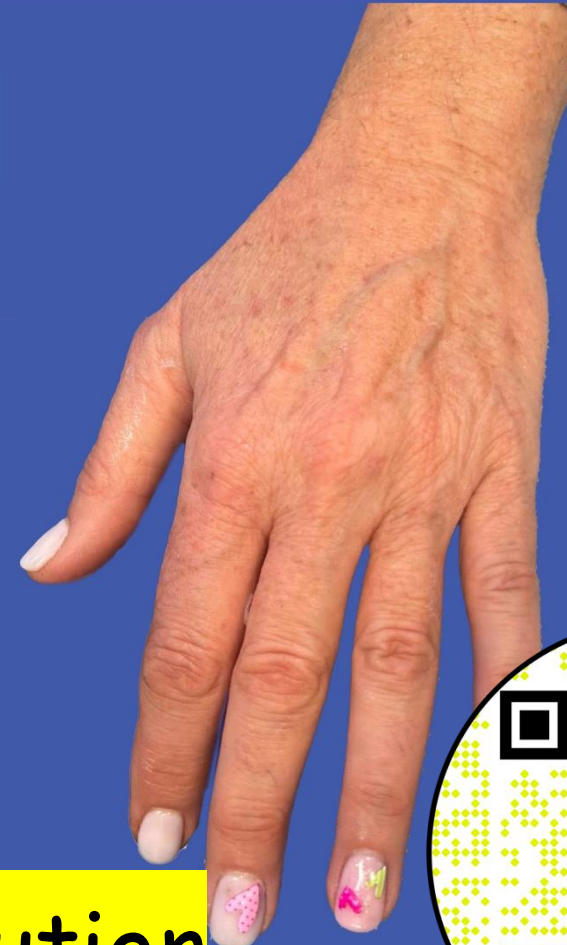
Metabolism is a biological response — not a property of an acid.

Hands Immediate Rejuvenation Using 30 Min Peel Off

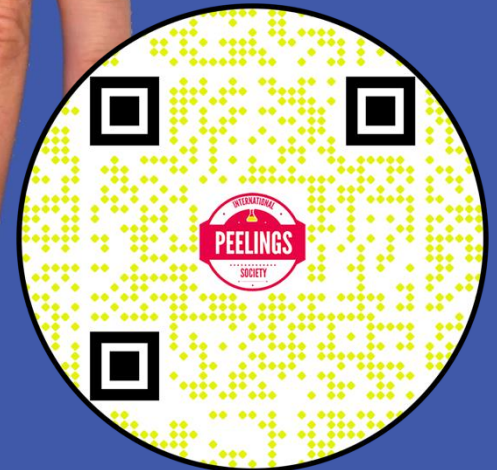
BEFORE



AFTER 45 MIN

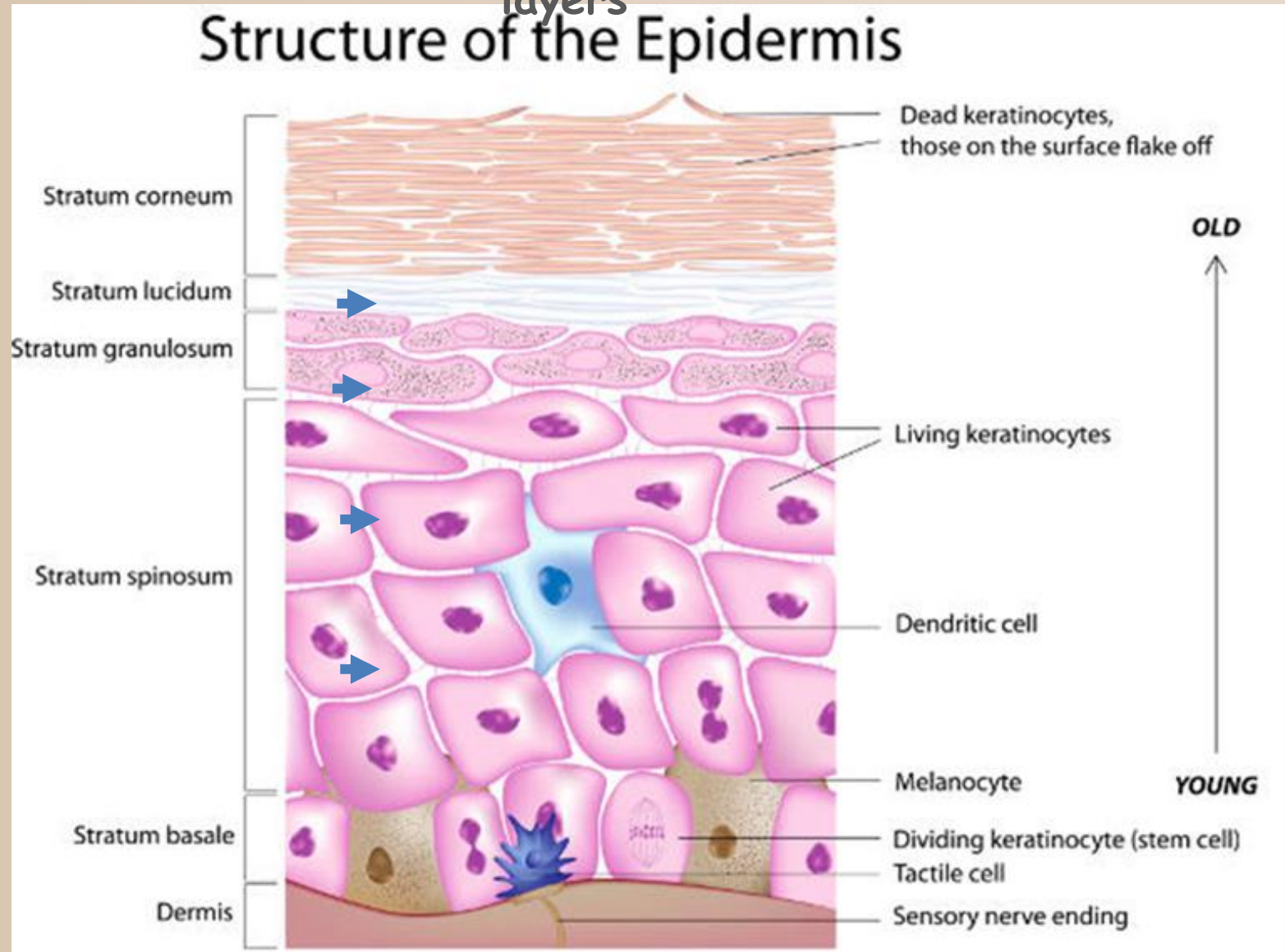


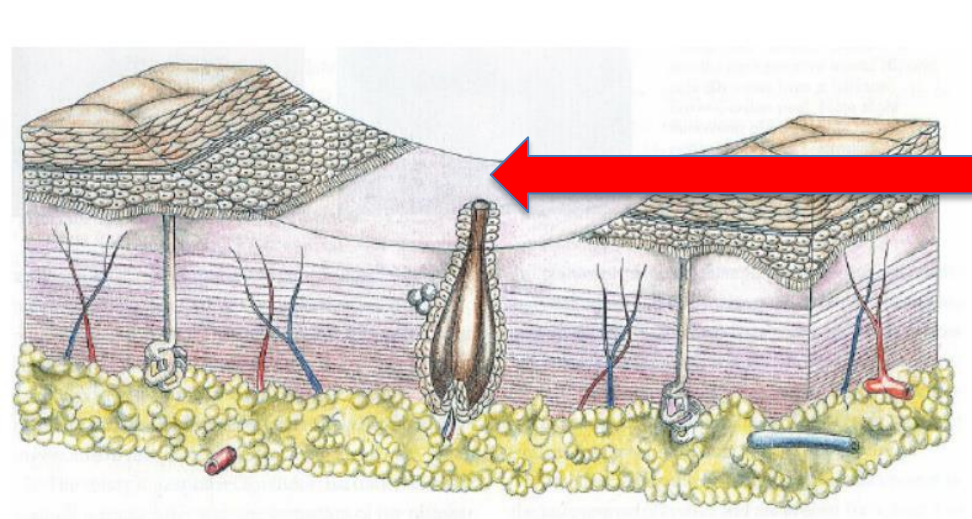
Metabolic Solution
courtesy of Mauro Tiziani



EPIDERMIS HISTOLOGY

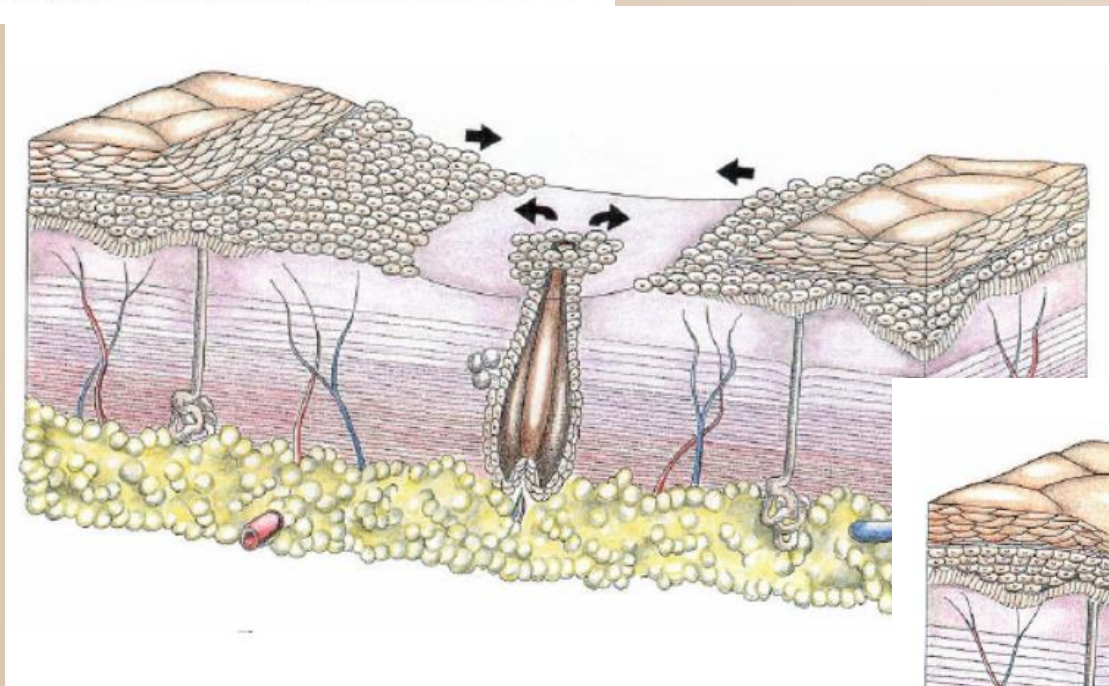
Most of the cells in the epidermis are keratinocytes, which are organized into 4 layers



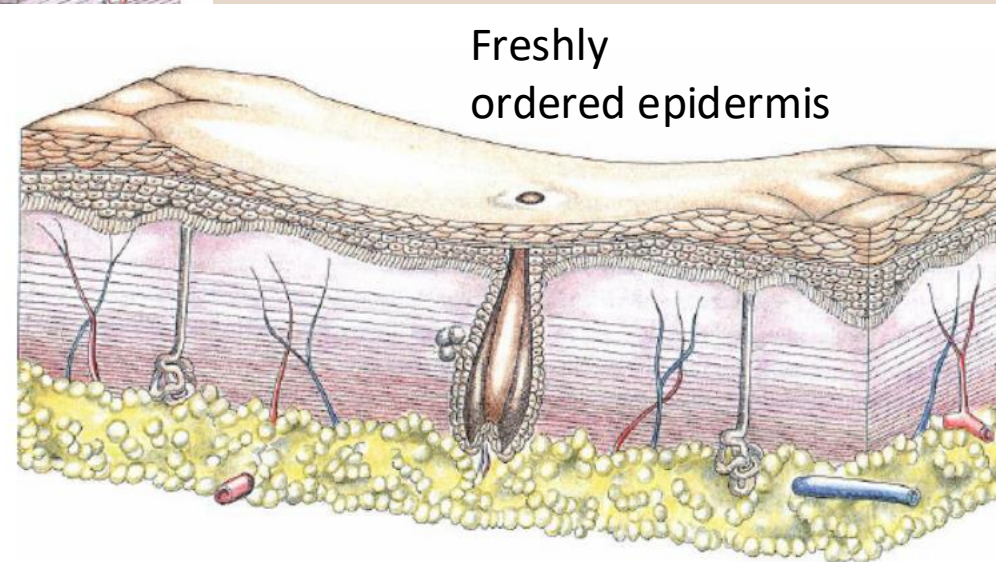


Acid-Induced Skin
Reaction

**Burn of the epidermis
and superficial dermis**



**Skin reparation after 2 to 5
days after the chemical burn
provoked by the peel**

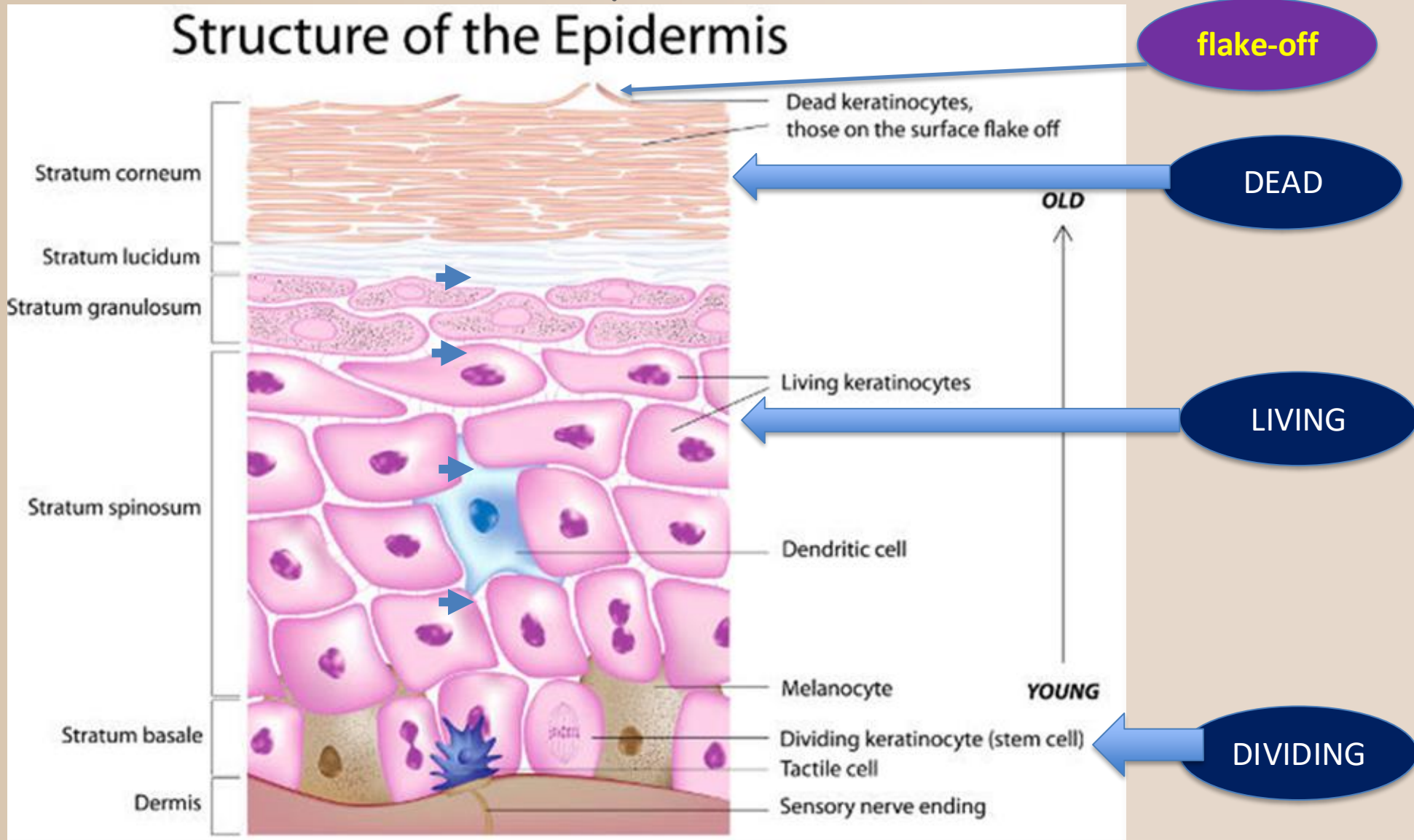


Freshly
ordered epidermis

HISTORICAL PARADIGM

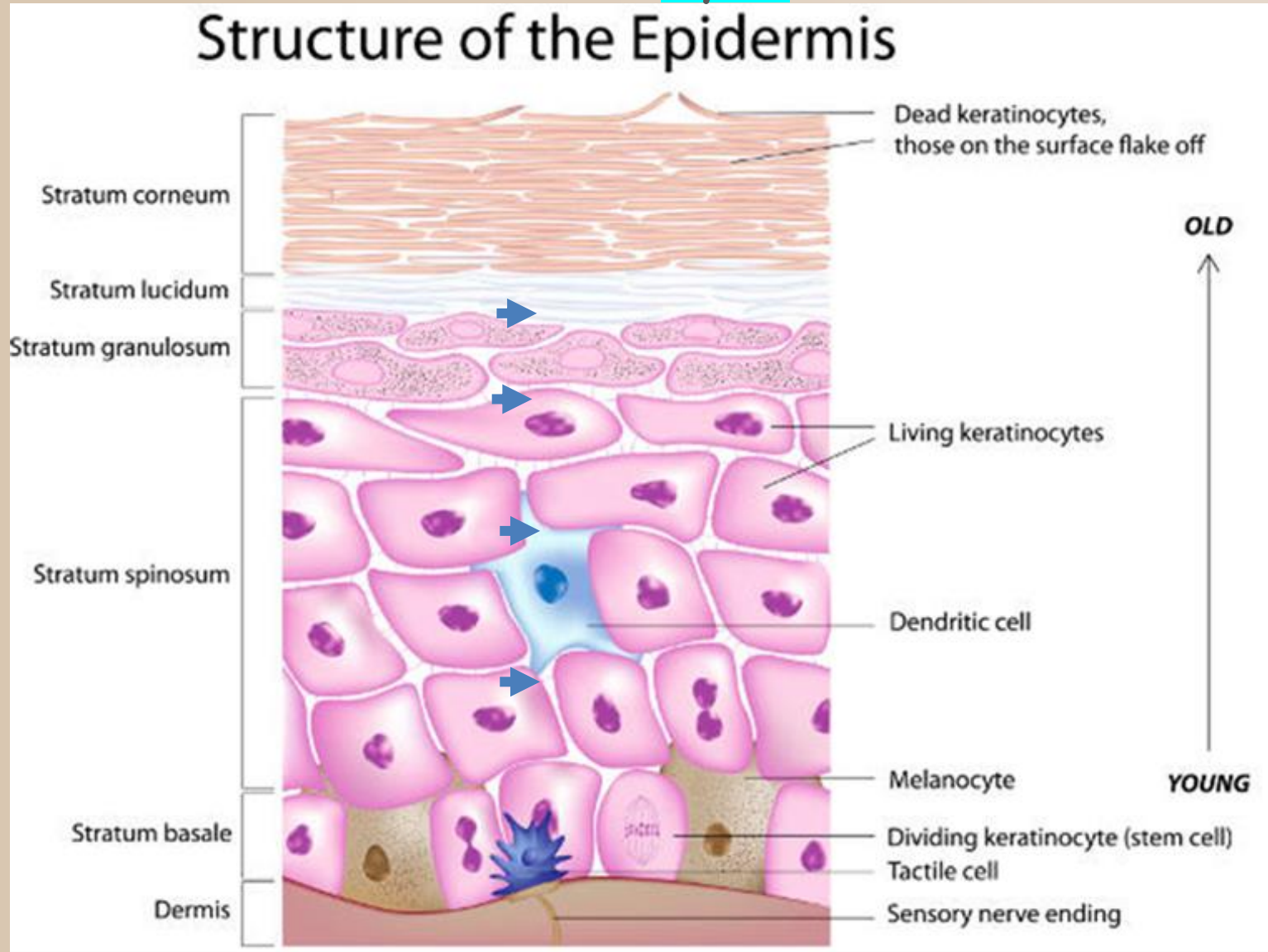
EPIDERMIS & KERATINOCYTES

Most of the cells in the epidermis are **keratinocytes**, which are organized into **4 layers**



EPIDERMIS

Most of the cells in the epidermis are **keratinocytes**, which are organized into **4 layers**



30 MIN peel-off

OTHER Peels

Acids Are Tools — Metabolism Is the Target

Acids

- External tools
- Defined by chemistry
- Act through delivery and formulation

Chemical agents trigger. →
Biological pathways respond.

Metabolism

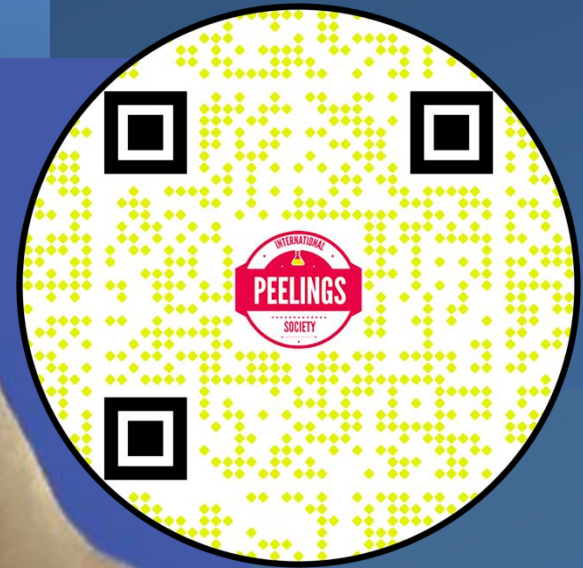
- Endogenous biological system
- Determines cellular response
- Defines clinical efficacy



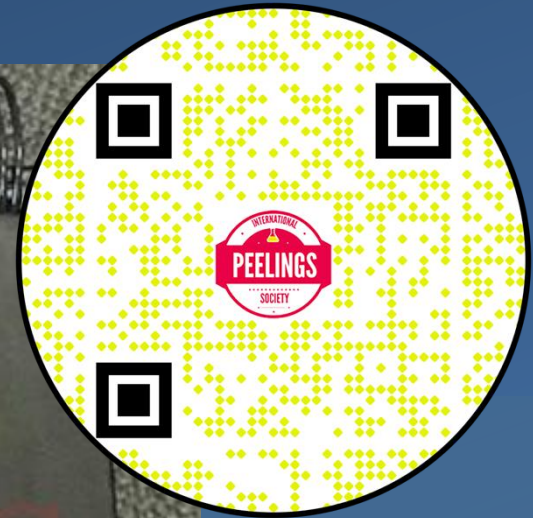
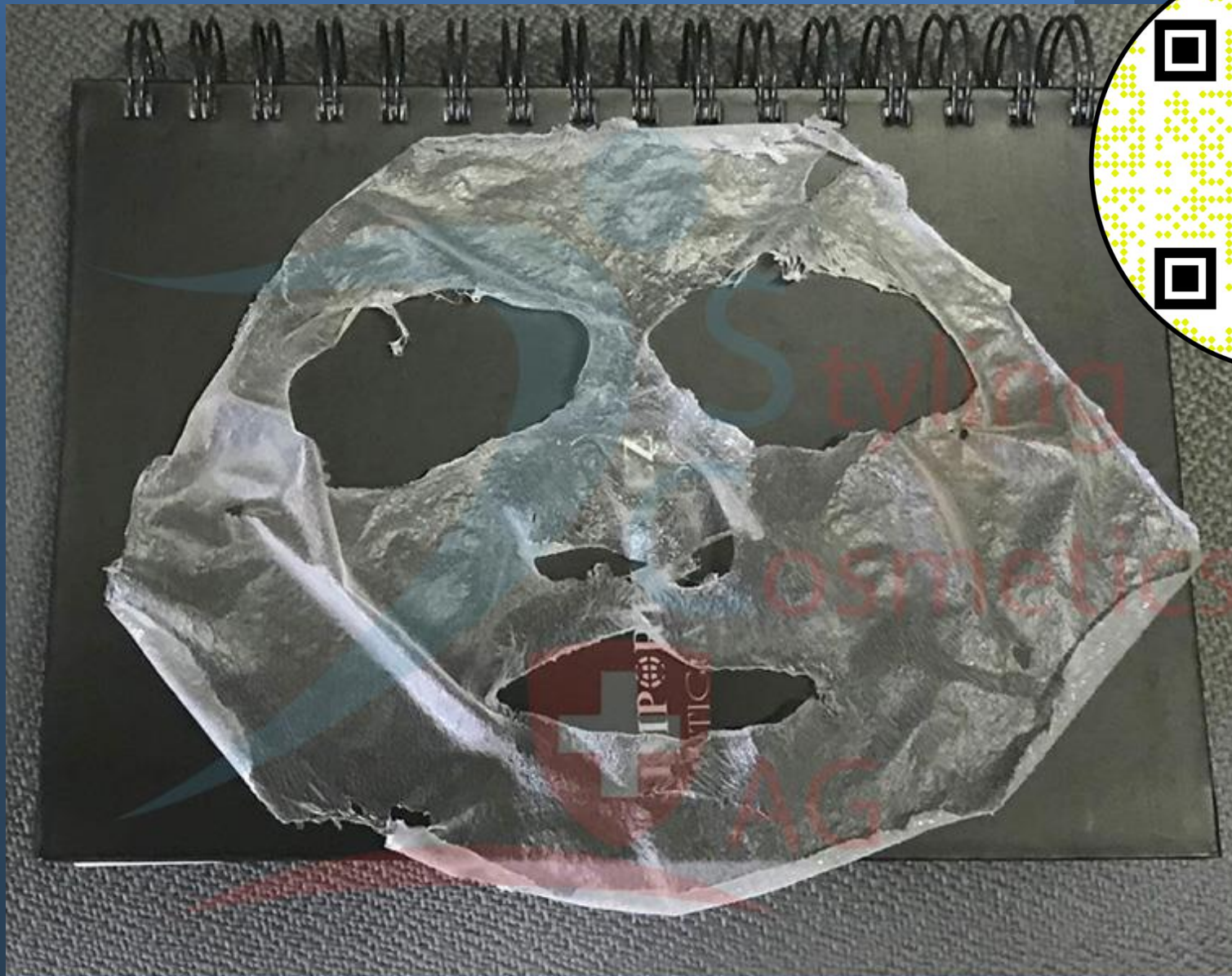
PEEL OFF-FLAKE OFF

VALID FOR ALL SKIN TYPES AT ANY TIME OF THE YEAR

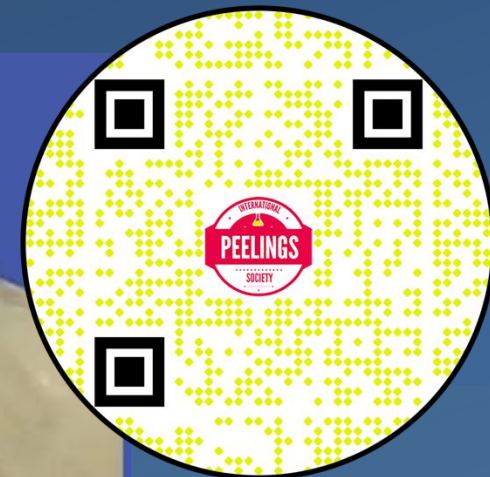




30 MIN PEEL OFF : REMOVE IT FROM TOP TO DOWN



30 MIN PEEL OFF : FULL MASK REMOVED



30 MIN PEEL OFF : BLACK POINT REMOVED

Clinical Protocol Logic

Correction • Stabilization • Maintenance



CORRECTION

Clinical objective:
Treat active **pathological** or
functional skin disorders

- Structural damage
- Dyschromias
- Acne, scars, inflammation

Role:

Active corrective
agents only

STABILIZATION

Clinical objective:
Consolidate results and
restore skin balance

- Barrier recovery
- Inflammation control
- Pigment regulation

Role:

Functional regulators
and buffers

MAINTENANCE

Clinical objective:
Maintain results and
prevent recurrence

- Long-term skin balance
- Anti-aging preservation
- Home-care continuity

Role:

Supportive and preventive
agents

Products are not interchangeable — they are selected according to their role within the protocol.

Protocol-Oriented Product Mapping

Each clinical protocol requires a sequence of dedicated products



CORRECTION PHASE

Targeted corrective products acting directly on the pathology

- Structural damage
- Dyschromias
- Acne, scars, inflammation

Clinical goal:

Active correction

STABILIZATION PHASE

Regulatory and buffering products to consolidate results

- Barrier recovery
- Inflammation control
- Pigment regulation

Clinical goal:

Result stabilization

MAINTENANCE PHASE

Supportive and preventive products for long-term control

- Skin balance preservation
- Anti-aging maintenance
- Home-care continuity

Clinical goal:

Prevention of recurrence

The protocol — not the product — is the clinical decision unit.

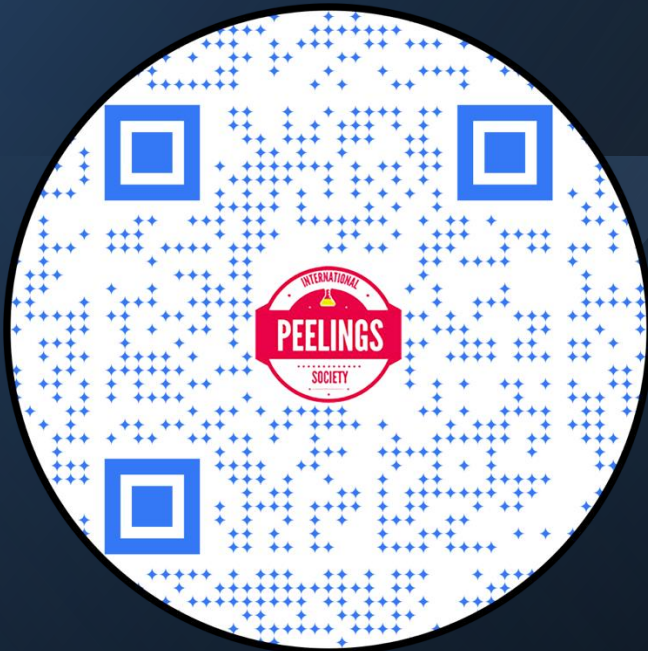


1.CORRECTION-
2.MODULATION/STABILIZATION
3.MAINTENANCE

Take a look at

<https://chemicalpeeling.com/main-protocols>

MAIN INDICATIONS FOR PEELS FACE, BODY, HANDS, FEET & INTIME AREA



- Anti Aging
- Acne
- Depigmentation
- Remove dead cells
- Skin regeneration
- Bleaching-whitening
- To improve the texture and tone of the skin
- Restore brightness and radiance to smokers' skin
- Scar Improvement
- Improve results of surgery
- Treatment of Complications dued to other Peels treatment



See More Indications on

<https://chemicalpeeling.com/clinical-indications>

COMPLICATIONS

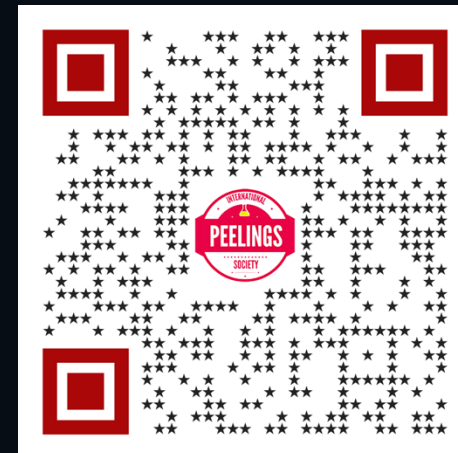


BRAND NAME

LACTIC COMPLICATION

- The issues caused by Big Pharma

Complications
of Brand Names Peels
with Big Marketing &
Poor Chemistry Knowledge
(Spain, Uruguay, Ecuador, Argentina
Korea, China, Poland, Austria...)





VICTIMS OF COVID RESTRICTIONS

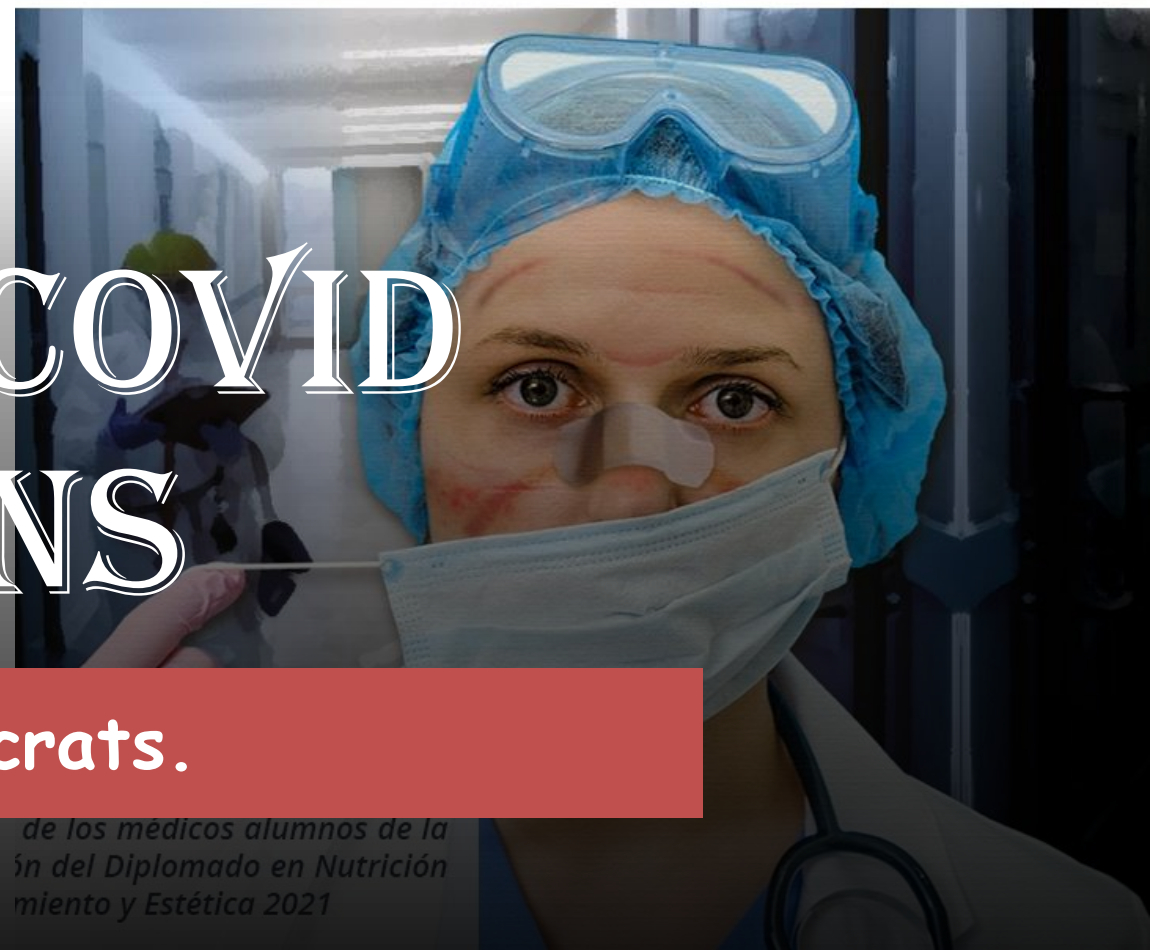
The issues caused by Bureaucrats.

NUTRICIÓN,
ANTI-ENVEJECIMIENTO
Y ESTÉTICA
Nº REGISTRO 17DAP04804



Secretaría
de Educación

Encuesta sobre Dermatitis facial
causada por el uso de cubrebocas en el personal
durante la pandemia de COVID-19 en México.



de los médicos alumnos de la
5ª del Diplomado en Nutrición
Anti-envejecimiento y Estética 2021

CONTRA INDICATIONS

<https://chemicalpeeling.com/contraindications>



Inflammation ≠ Efficacy

INFLAMMATION-DRIVEN THINKING

- **Underlying Assumption:**
Greater inflammation leads to greater efficacy
- **Biological Mechanism:**
 - Chemical or physical injury
 - Acute inflammatory cascade activation
 - Cytokine and prostaglandin release
- **Clinical Consequences:**
 - Erythema, edema, pain
 - Barrier disruption
 - Increased risk of PIH and complications
 - Unpredictable clinical outcomes
- **Clinical Limitation:**
 - Efficacy depends on tissue damage intensity

FUNCTION-DRIVEN THINKING

- **Underlying Principle:**
Clinical efficacy does not require inflammation
- **Biological Mechanism:**
 - Modulation of cellular metabolism
 - Regulation of signaling pathways
 - Controlled stimulation of renewal mechanisms
- **Clinical Consequences:**
 - Minimal or absent inflammation
 - Preserved barrier integrity
 - Higher safety and reproducibility
 - Improved patient compliance
- **Clinical Advantage:**
 - Efficacy achieved through functional regulation, not through inflammatory damage

Inflammation is a biological reaction to injury,
not a measure of therapeutic success.



LEARNING OBJECTIVES

- Which acids are dangerous and non-dangerous?
- Which acids are aggressive and not-aggressive?
- How to determine the penetration of an acid?
- Which parameters are variable and which are constant?
- What are the elements that distinguish one TCA from others?
- Do exist really deep, medium and superficial peels?
- How to treat complications of Peels? Why do they occur?
- What to avoid in a chemical peel treatment?
- Why do patients change their peeler?
- How to avoid a medico-legal case after exfoliation?
- Which classifications for chemical peels?
- What is the difference between physical, chemical, mechanical and thermodynamic peels?

From Chemistry to Biology:

What Really Determines Clinical Outcomes

Chemistry

- Describes molecular behavior
- Defines ionization and penetration
- Controls exposure and delivery

Biology

- Determines cellular response
Modulates inflammation
- Drives tissue adaptation and renewal

Clinical outcome

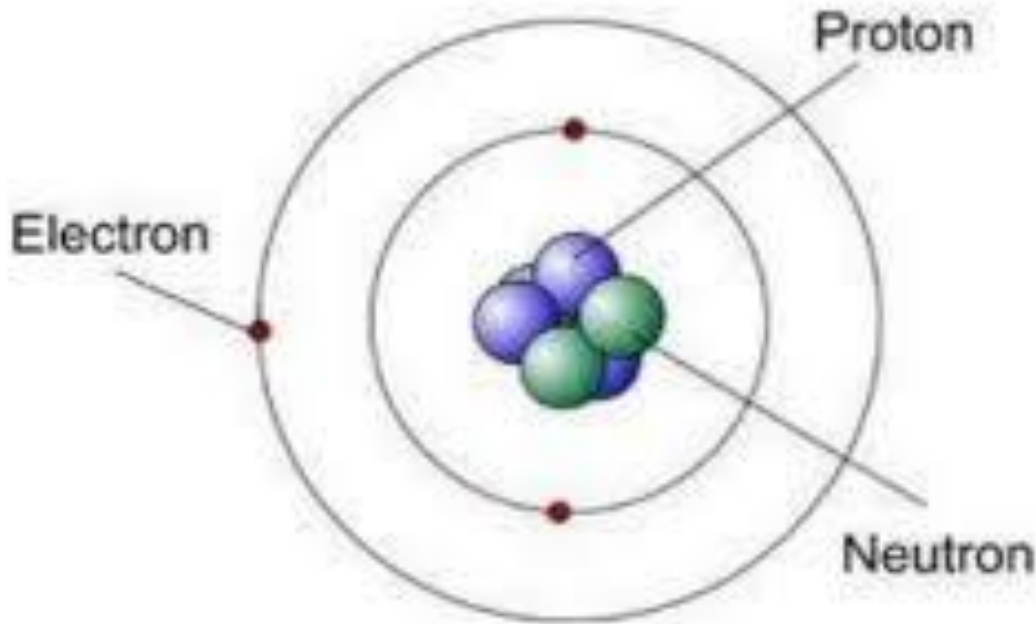
Emerges from biological regulation
Not from chemical strength alone
Not from visible injury

Chemistry initiates the stimulus.
Biology determines the result.

STRUCTURE OF ATOMS, MOLECULES

IONS : ANIONS + CATIONS

PROTONS AND NEUTRONS

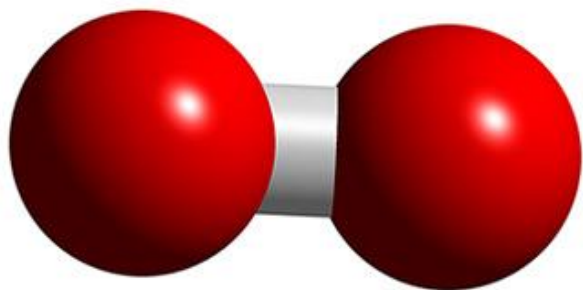


- Atoms are building blocks of matter.
- They consist of the atomic nucleus and the atomic shell.
- The atomic nucleus is composed of positively charged particles, the protons and the uncharged neutrons, which in turn consist of other elementary particles.

Atomic nucleus = protons + neutrons
Atomic shell = electrons

STRUCTURE OF MOLECULES

oxygen or O₂



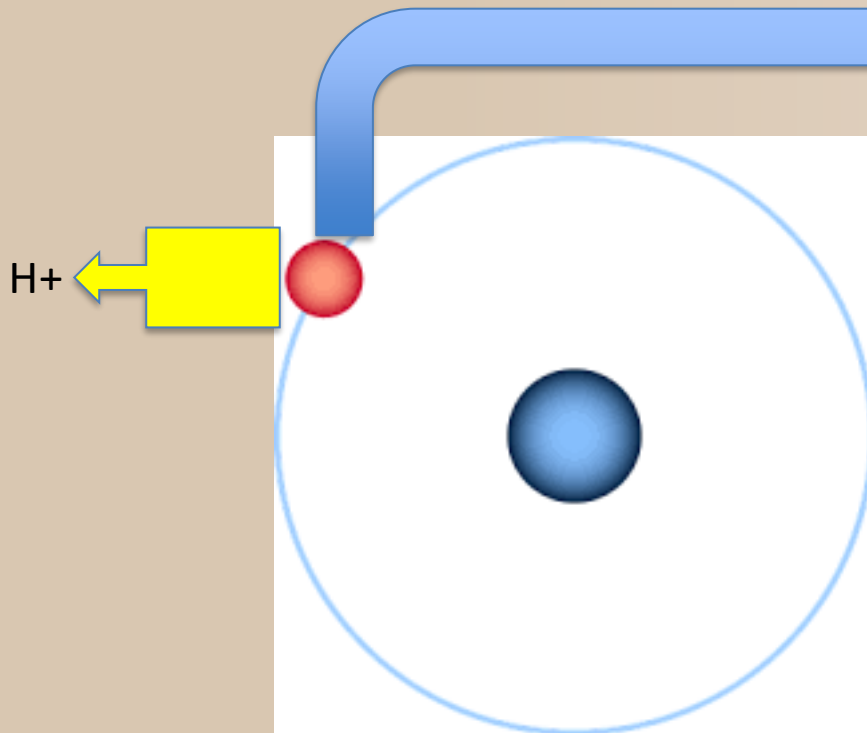
- Molecular structure or molecular geometry is the geometric, spatial relative arrangement of atoms in a molecule

Molecules \geq 2 atoms together (A.Tenenbaum)

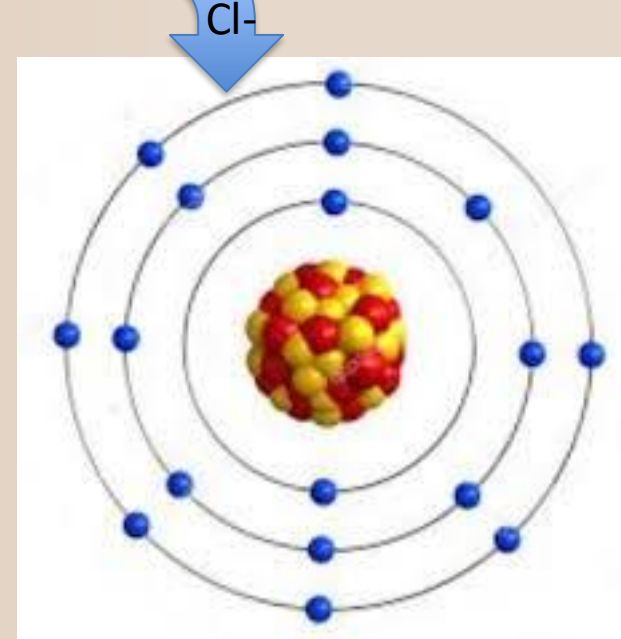
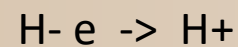
STRUCTURE OF IONS AND MOLECULES

ELECTRONS LOOSER = POSITIVE IONS = CATIONS

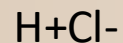
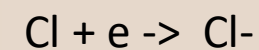
ELECTRONS WINNER = NEGATIVE IONS = ANIONS



HYDROGEN



CHLORINE



MENDELEEV'S PERIODIC TABLE

1

1,0079

H

Wasserstoff

2,20,09

3

6,9675

Li

Lithium

0,980,53

4

9,0122

Be

Beryllium

1,571,85

11

22,990

Na

Natrium

0,930,97

12

24,305

Mg

Magnesium

1,311,74

19

39,098

K

Kalium

0,820,86

20

40,078

Ca

Calcium

1,01,55

21

44,956

Sc

Scandium

1,362,98

37

85,468

Rb

Rubidium

0,821,53

38

87,620

Sr

Strontium

0,952,63

39

88,906

Y

Yttrium

1,224,47

55

132,906

Cs

Caesium

0,791,90

56

137,328

Ba

Barium

0,893,59

57

138,906

La

Lanthan

1,16,15

58-71

siehe unten

87

(223)

Fr

Francium

0,7?

88

(226)

Ra

Radium

0,895,5

89

(227)

Ac

Actinium

1,110,1

90-103

siehe unten

Legende

Symbol

schwarz = Feststoff
blau = Flüssigkeit
rot = Gas

Ordnungszahl

Atommasse

Symbol

Name

Elektronegativität

Serie (Flächenfarbe)

Alkalimetalle
Erdalkalimetalle
Übergangsmetalle
Lanthanoide
Actinoide

Metalle
Halbmetalle
Nichtmetalle
Halogene
Edelgase
unbekannt

Dichte

rot = kg / m³
schwarz = kg / dm³
grau = unbestimmt

Schraffur
durchgehend = natürliches Element
schraffiert = künstliches Element

Gruppe

13

14

15

16

17

18

2

4,0026

He

Helium

0,18

10

20,180

Ne

Neon

0,90

18

39,948

Ar

Argon

0,821,78

36

83,798

Kr

Krypton

3,75

54

131,294

Xe

Xenon

2,65,90

86

(222)

Rn

Radon

9,73

5

10,813

B

Bor

2,042,46

13

26,981

Al

Aluminium

1,612,70

21

44,956

Sc

Scandium

1,362,98

39

88,906

Y

Yttrium

1,224,47

57

138,906

La

Lanthan

1,16,15

72

178,49

Hf

Hafnium

1,313,3

90-103

siehe unten

6

12,011

C

Kohlenstoff

2,552,26

14

28,085

Si

Silicium

1,92,34

22

47,867

Ti

Titan

1,544,50

40

91,224

Zr

Zirkonium

1,336,50

58-71

siehe unten

7

14,007

N

Stickstoff

3,041,25

15

30,974

P

Phosphor

2,192,69

23

50,941

V

Vanadium

1,636,11

41

92,906

Nb

Niob

1,68,57

59

140,908

Pr

Praseodym

1,136,48

73

180,948

Ta

Tantal

1,516,65

91

231,036

Pa

Protactinium

1,313,54

8

15,999

O

Sauerstoff

3,441,43

16

32,067

S

Schwefel

2,582,07

24

51,996

Cr

Chrom

1,667,14

42

95,95

Mo

Molybdän

2,1610,28

60

144,242

Nd

Neodym

1,147,01

74

183,84

W

Wolfram

2,3619,25

92

238,029

U

Uran

1,3618,95

9

18,998

F

Fluor

3,981,70

17

35,451

Cl

Chlor

3,163,21

25

54,938

Mn

Mangan

1,557,43

43

(97)

Tc

Technetium

1,911,50

61

(145)

Pm

Promethium

1,137,22

75

186,207

Re

Rhenium

1,921,0

93

(244)

Pu

Plutonium

1,319,82

10

58,693

Ni

Nickel

1,888,91

28

58,693

Ni

Nickel

1,888,91

36

83,798

Kr

Krypton

3,75

54

131,294

Xe

Xenon

2,65,90

82

207,2

Pb

Blei

2,3311,35

110

(281)

Ds

Darmstadtium

??

11

63,546

Cu

Kupfer

1,98,92

29

63,546

Cu

Kupfer

1,98,92

47

107,868

Ag

Silber

1,9310,49

65

65,380

Zn

Zink

1,657,14

83

208,980

Bi

Bismut

2,029,75

111

(281)

Rg

Roentgenium

??

12

65,380

Zn

Zink

1,657,14

30

65,380

Zn

Zink

1,657,14

48

112,414

Cd

Cadmium

1,698,64

66

162,50

Dy

Dysprosium

1,228,55

84

209,98

Lv

Livermorium

??

112

(285)

Cn

Copernicium

??

13

26,981

Al

Aluminium

1,612,70

31

69,723

Ga

Gallium

1,815,90

49

114,818

In

Indium

1,787,31

67

164,930

Ho

Holmium

1,249,05

85

(210)

At

Astat

2,2?

113

(286)

Nh

Nihonium

??

14

28,085

Si

Silicium

1,92,34

32

72,631

Ge

Germanium

2,015,32

50

118,711

Sn

Zinn

1,967,26

68

167,259

Er

Erbium

1,249,05

86

(222)

Rn

Radon

9,73

114

(289)

Fl

Flerovium

??

15

30,974

P

Phosphor

2,192,69

33

74,922

As

Arsen

2,185,73

51

121,760

Sb

Antimon

2,056,70

69

168,934

Tm

Thulium

1,259,32

87

(223)

Fr

Francium

0,7?

115

(288)

Mc

Moscovium

??

16

32,067

S

Schwefel

2,582,07

34

78,972

Se

Selen

2,554,82

52

127,60

Te

Tellur

2,666,25

70

173,045

Yb

Ytterbium

1,016,97

88

(226)

Ra

Radium

0,895,5

116

(293)

Lv

Livermorium

??

17

35,451

Cl

Chlor

3,163,21

35

79,904

Br

Brom

2,963,12

53

126,904

I

Iod

2,14,94

71

174,967

Lu

Lutetium

1,016,97

89

(227)

Ac

Actinium

1,110,1

117

(294)

Ts

Tenness

??

18

39,948

Ar

Argon

0,821,78

37

85,468

Rb

Rubidium

0,821,53

55

132,906

Cs

Caesium

0,791,90

73

183,84

W

Wolfram

2,3619,25

91

231,036

Pa

Protactinium

1,313,54

118

(294)

Og

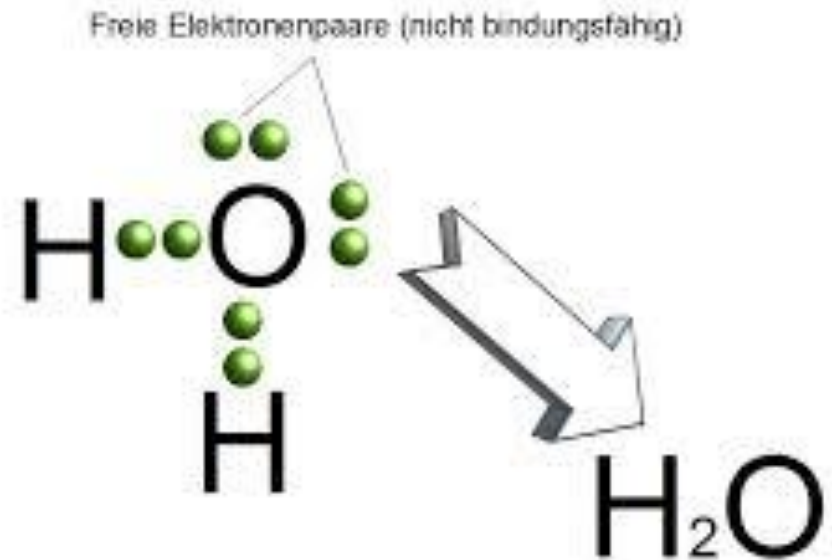
Oganesson

??

BINDING AND 2(NXN) COVALENCE

- Atomic number Z
- H Z=1 1
- O Z=8 (2+6)

Lewis-Struktur von Wasser H₂O





water

THE
WATER
H₂O

Anzahl der Valenzelektronen

2



He

5



N

3



Al

6



S

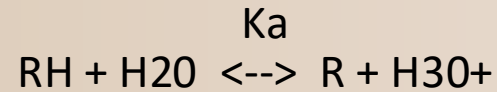
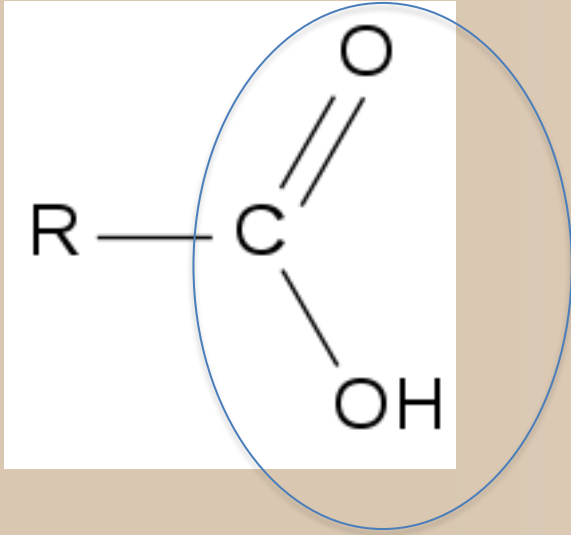
2



Ca

Was uns interessiert	VALENZ	
C	4	vierwertig
H	1	einwertig
O	2	zweiwertig
Cl	1	einwertig

THE ACID FUNCTION



R = A⁻ = Anion

H₃O⁺ = Cation

K_a = constant of dissociation of an acid in solution

$$\text{pK}_a = -\log(\text{K}_a)$$



H₃O⁺ = HYDRONIUM ION or acid ion



R-COO⁻ = Anion

$$\text{pH} = -\log(\text{H}_3\text{O}^+)$$

PRACTICAL EXERCISE

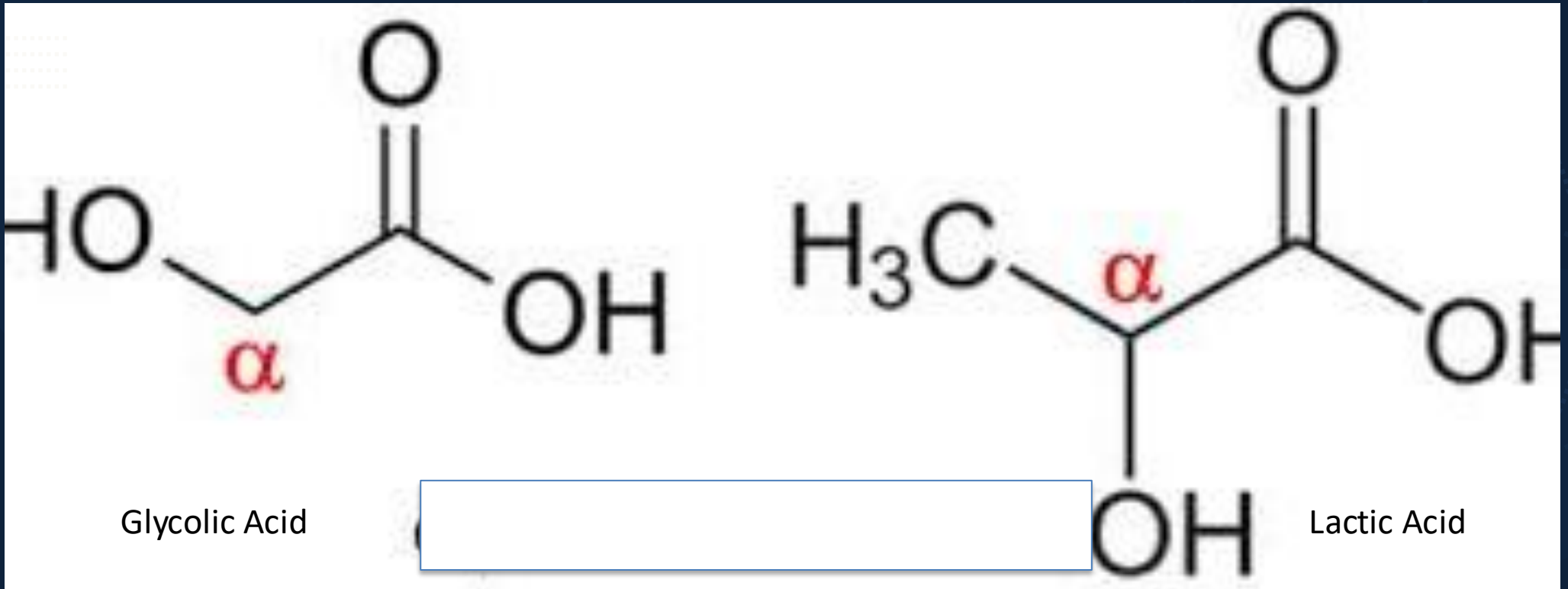
the acid
function -
COOH

- H₂O Water
- Glycolic acid C₂H₄O₃
- Lactic acid C₃H₆O₃
- TCA Trichloroacetic acid
C₂Cl₃HO₂
- Acetic acid C₂H₄O₂

THE CATEGORY OF AHA OR ALPHA HYDROXY ACIDS

Glycolic acid and lactic acid
Fruit acids and alpha
hydroxy acids (AHA)

- the hydroxy radical -OH
- Alpha carbon
- the acid function -COOH



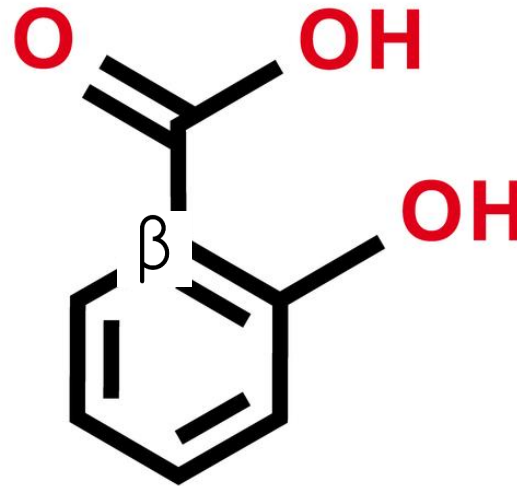
THE CATEGORY OF BHA OR BETA HYDROXY ACIDS



the hydroxy radical -OH

β Beta carbon

the acid function -COOH

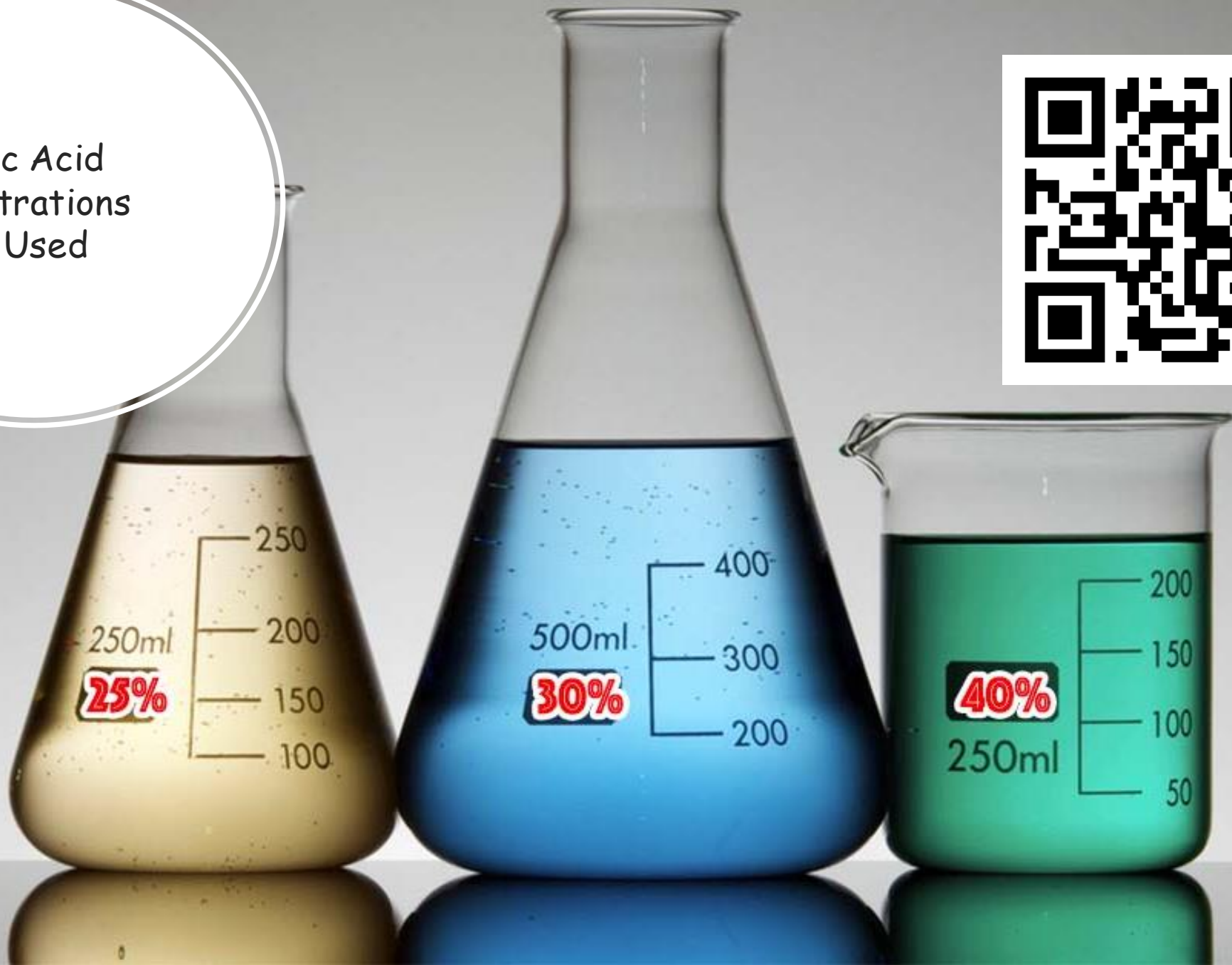


SALICYLIC ACID



SALICYLIC ACID CONCENTRATIONS

Salicylic Acid
Concentrations
Mostly Used



AHA vs. BHA: Solubility and Clinical Application



AHA

Alpha Hydroxy Acid

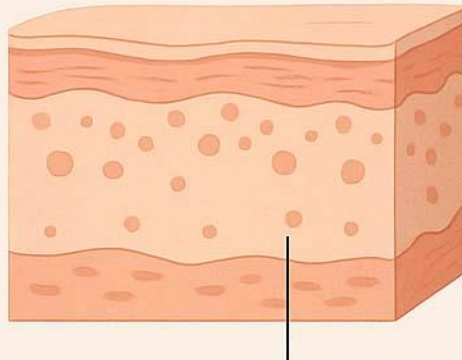
WATER-SOLUBLE



BHA

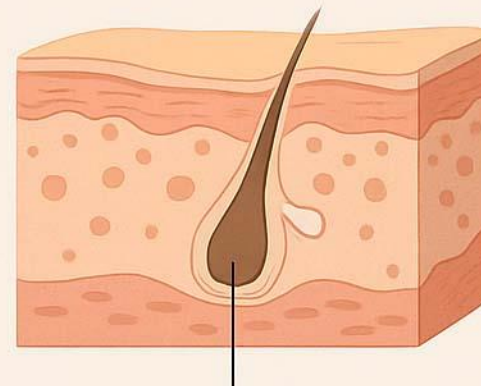
Beta Hydroxy Acid

OIL-SOLUBLE



Surface exfoliation

Dry, sun-damaged skin



Deeper into pores

Acne-prone, oily skin



Comparison of AHA vs BHA

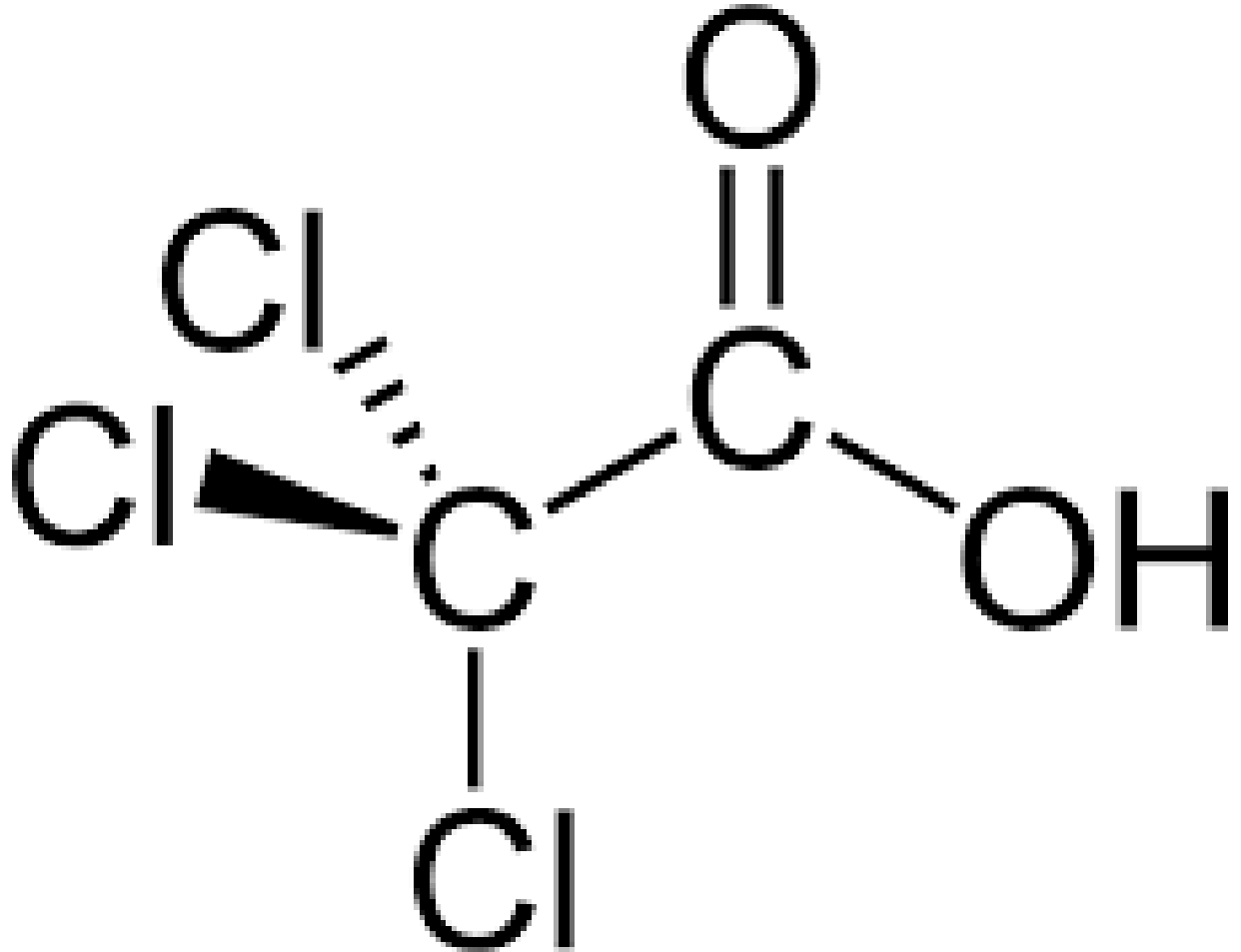
	AHA	BHA
1 Acid	<ul style="list-style-type: none">• Citric Acid• Lactic Acid	Salicylic Acid
2 Solubility	Water Soluble	Oil Soluble
3 Target	Surface Exfoliation	Penetrates deeper into the Pores
4 Actions	<ul style="list-style-type: none">• On Skin surface• Remove Skin Dead cells• Improve Texture• Hydrate the Skin (citric acid)	<ul style="list-style-type: none">• Dissolve Sebum• Clear Clogged Pores• Reduce Inflammation• Comedolytic Properties
5 Effectiveness	<ul style="list-style-type: none">• On Dry Skin• On Sun Damaged Skin• On Mature Skin	<ul style="list-style-type: none">• On Acne Prone• On Oily Skin• On Black Heads
6 For Dark Skin Types		Better Tolerated

TOPOGRAPHICAL MAPPING OF DRY AND OILY FACIAL SKIN

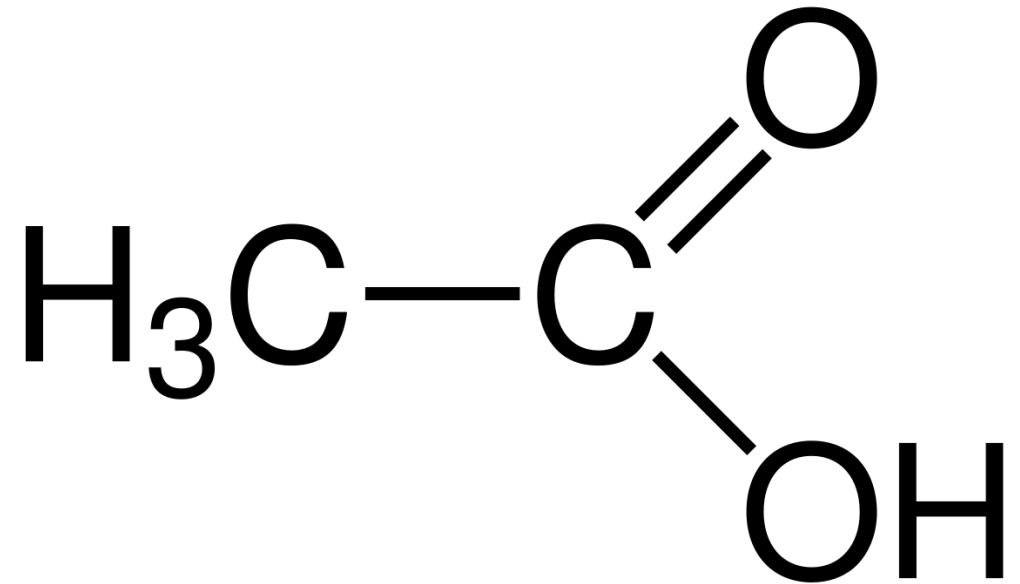


Courtesy of Dr. Alain TENENBAUM & Mauro TIZIANI

NOTE: TCA
IS NEITHER
AN AHA,
NOR A BHA

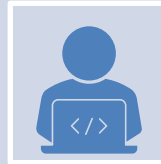


ACETIC ACID IS
NEITHER
AN AHA NOR A BHA



DECIMAL LOGARITHMS PH & PKA

- $\text{Log } 1 = 0$
- $\text{Log } 10 = 1$
- $\log 100 = 2$
- $\log 1000 = 3$
- $\log 100000000 = 7$



$$\text{pH} = -\log (H^+) \\ = -\log (H_3O^+)$$

$$\text{pK}_a = -\log K_a$$

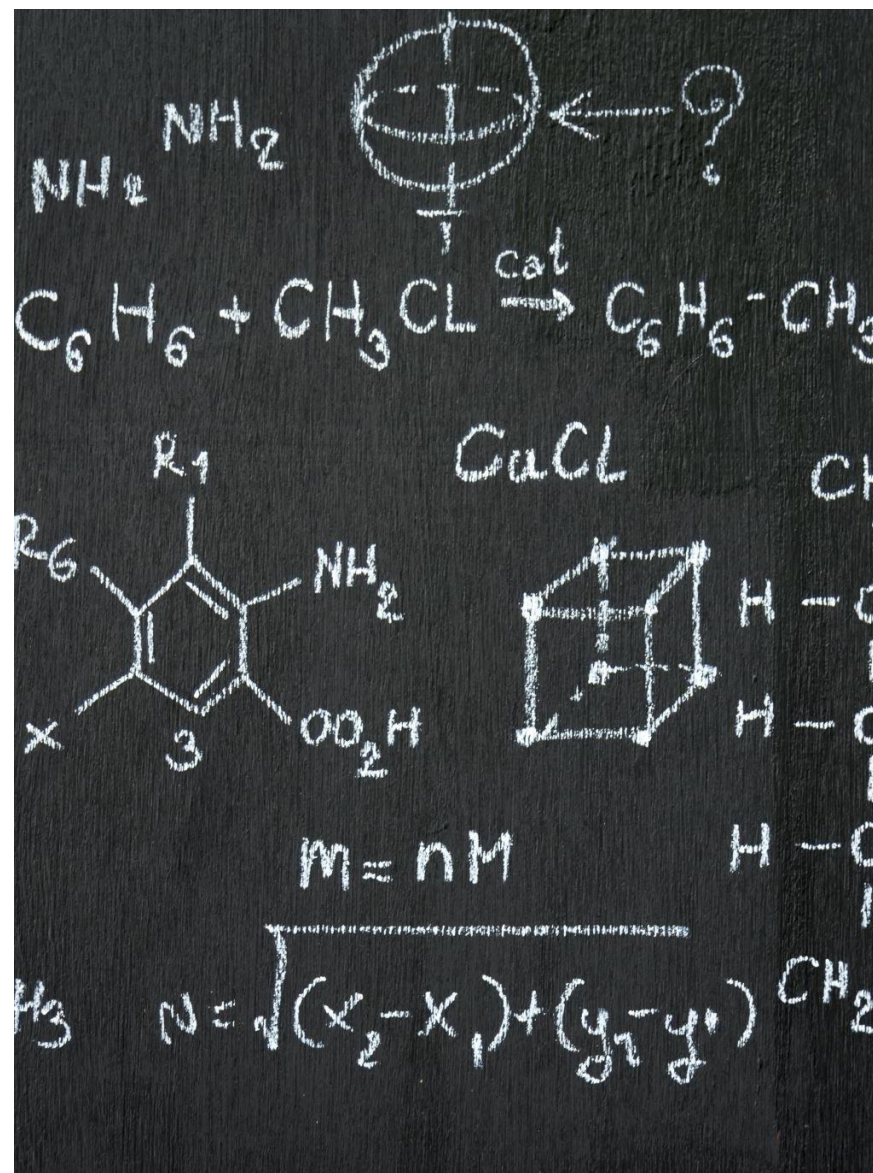


$$\text{pH} = -\log (H_3O^+)$$

UNIT OF CONCENTRATION OF AN ACID

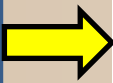
MOL/L VS G/L

- EXAMPLE : HCl Chloric Acid
- Sum of atomic masses (Mendeleiev)
- $H = 1$ $Cl = 35$ ie $HCl = 36$
- 1 mol HCl weighs 36g i.e.
- 0.1 mol weighs 3.6g
- HCl 0.1 mol/l = 3.6g crystal HCl for 1l H_2O
- pH of 0.1 mol/l HCl = $-\log(1/10)=1$ extremely acidic
- $pH = -\log(H^+) = -\log(H^+)$

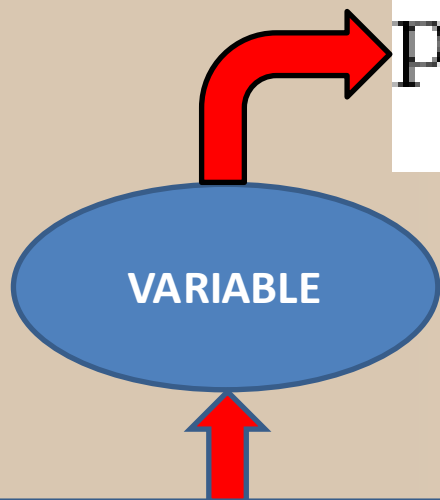




CONSTANT



$$K_a = \frac{[H^{+}][A^{-}]}{[HA]}$$



CONSTANT

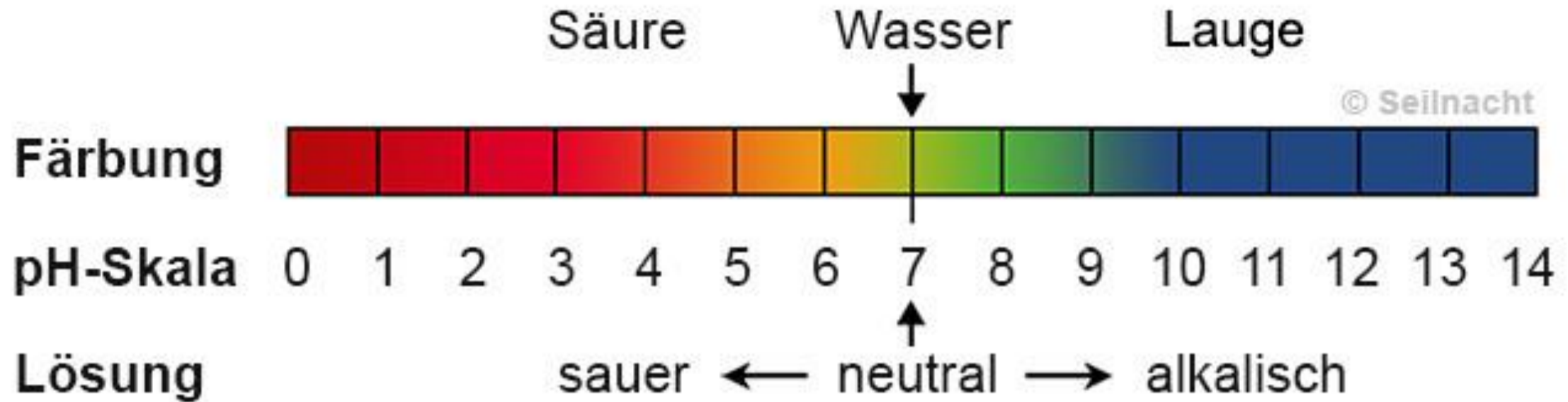


$$pH = pK_a + \log_{10} \left(\frac{[A^{-}]}{[HA]} \right)$$

Equation of
Henderson
Hasselbach

Choose the acid f(pKa)

THE PH OF AN ACID IS VARIABLE FOR A SAME CONCENTRATION



Your task will be to play with the pH of the chosen acid
once after selecting the acid (A.TENENBAUM)

pH is function of temperature, altitude, hygrometry



PH AND PKA

Choose your acidity

pKa

Modulate your acidity pH

- ☐ pKa = aggressiveness

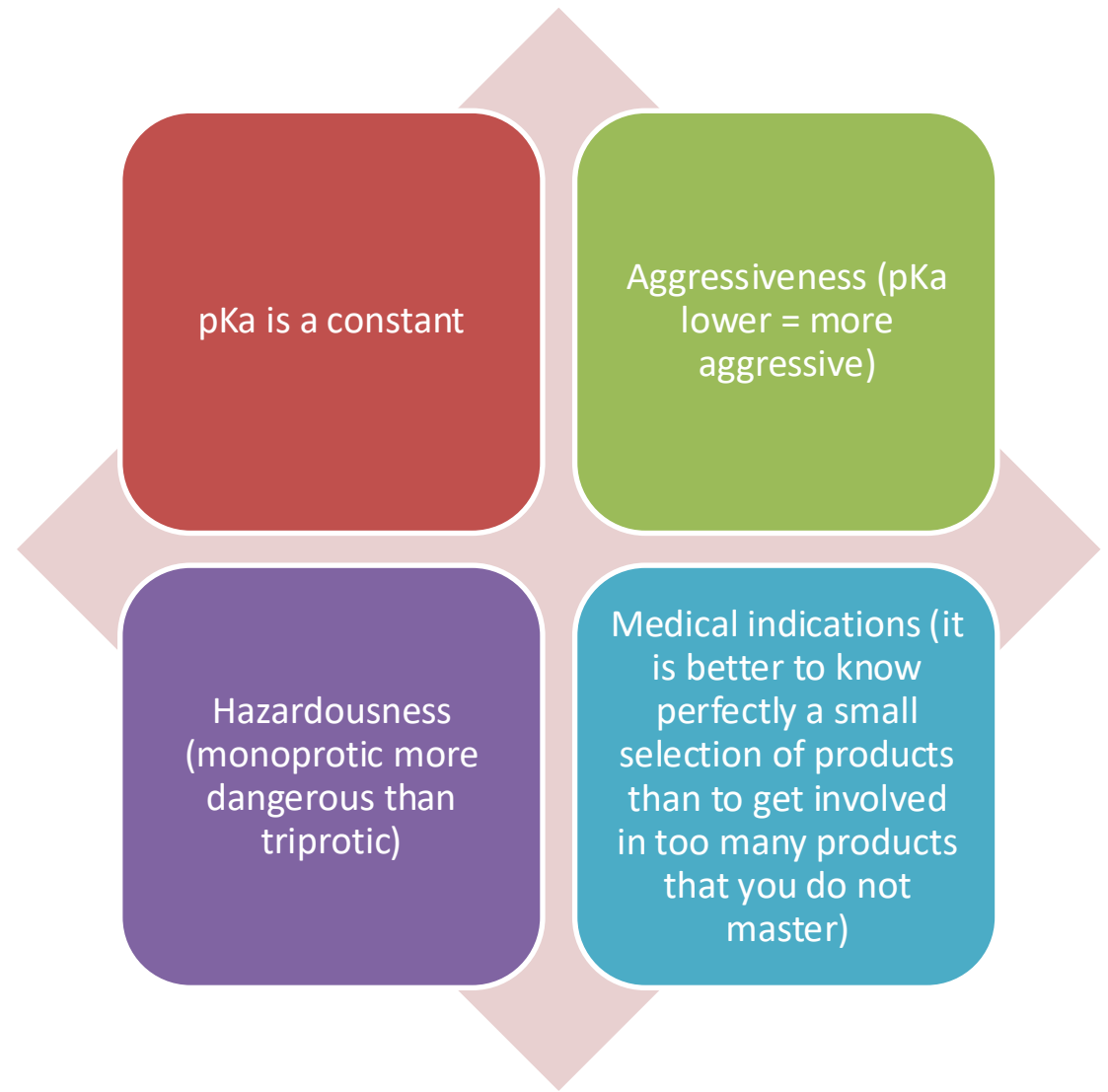
- ☐ pKa = Constant

- ☐ pH = penetration

- ☐ pH = variable

How to
choose an
acid and
modulate it

playing with
pH and pKa



How to **MODULATE** an acid Playing with **PH** and pKa

- **pH** is a variable

The pH is lowered

1. for a higher **acid concentration** in mol / l, g / l, w / v
2. the **number of applications or coats** with the same acid on the skin in 1 session
3. **Repeating sessions**, especially on flaky skin or skin in desquamation.

PLAYING WITH THE PH USING CONCENTRATIONS

Example : TCA (2C + 2O + 3 Cl) Mol mass : 127g

In Europe weight/weight

- TCA 10% w/w = 10g TCA in 90g H₂O

In USA weight/volume

- TCA 10% w/v = 10g TCA in 100 g(ml) H₂O

How to transform your TCA 10% into TCA 5%

We want to get 5g TCA for 95ml
H₂O (5%)

This is same as 10 g for 190 ml H₂O
Then just add 100 ml

5g TCA in 100 ml is like 10g TCA in 200 ml
Then add 100 ml

10g in 190 ml (EU) is more concentrated than 10g in 200 ml (US)

Conclusion : pH Acid x with y % w/w < pH same Acid x with y% w/v



A. Tenenbaum working in a self-constructed temporary laboratory

Safety and protective measures are paramount.

Instruments, devices, and laboratory glassware required for the emergency preparation of TCA outside a conventional laboratory environment.

Raw materials and water



HOW TO CHOOSE A TCA



1. The density of the steam.
2. The degree of purity.
3. The quality (analytical indication of the pH value)
4. Refractive index
5. The boiling point per liter
6. The density in g /ml bei 25 ° C.
7. Residual traces of anions and/or cations, if they are still present, can cause tattoos. (differential diagnosis with dyschromia) in case of deep penetration associated with pH. For this reason, it is not recommended to use TCA, which is regulated or neutralized with ordinary water, as it contains metal ions.
8. Other chemical residues: whether they should be considered ignored or not, such as SO₄.
9. The flash point (A high flash point provides more safety).
10. Any impurities, e.g. insoluble materials, etc.
11. Solubility in water in "moles" at 20 C° with the clarity or lack of color of the solution obtained
12. Turbidity
13. Vapor Pressure (For sealing and lubricating at low vapor pressure in high vacuum applications.) Vapor expression, Pa at 51 ° C: 133.
14. Stability when offered in gel



TCA AND PACKAGING

Liquid

TCA GEL
STABLE till 18%w/w

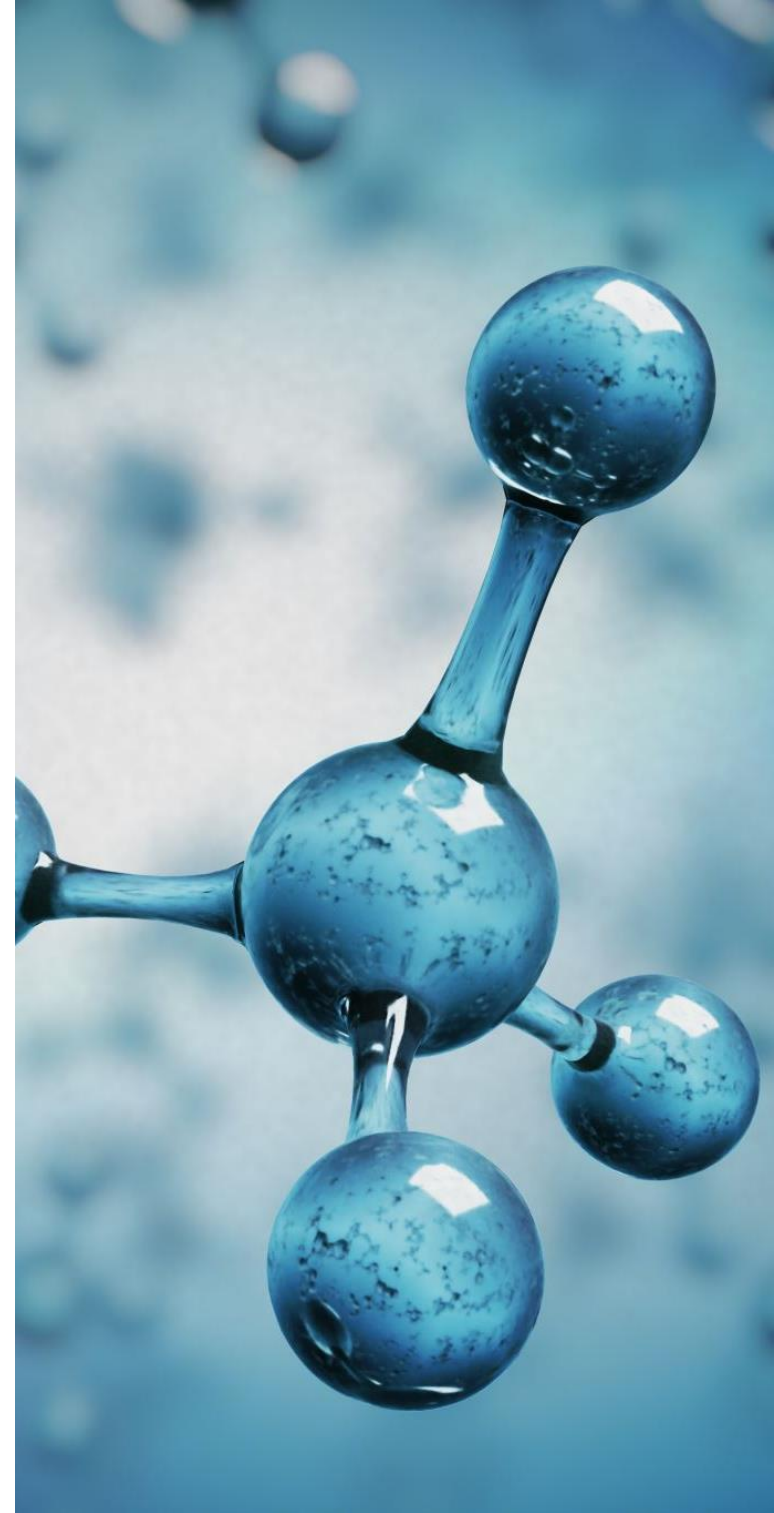


Airless

**RECOMMENDED
METHODS FOR
HELPING
INGREDIENTS TO
PENETRATE INTO
THE SKIN OR
DECREASING THE
PH (MODULATE
THE PH)**



- Q or exothermic reaction
(Cream 1 + Cream 2)
- Increasing the concentration of the acid
- More coats applied on same area with same acid
- Lipoic acid(A.TENENBAUM + M.TIZIANI)
- The fewer ingredients a cream contains, the easier it is to penetrate due to its low molecular weight (M.TIZIANI)
- For severely dehydrated skin, use a low molecular weight moisturizer + Lipoic acid in a sequential way



Classification of Chemical Peels

— A. Tenenbaum —



pKa Describes Chemical Behavior — Not Biological Efficacy

What pKa tells us

- Degree of ionization
- Chemical reactivity
- Predictable penetration behavior
- Acid–base equilibrium

What pKa does NOT tell us

- Cellular signaling pathways
- Metabolic adaptation
- Inflammatory modulation
- Long-term tissue response

Where efficacy really emerges


Interaction between
formulation and tissue
Cellular and metabolic
pathways
Biological regulation, not
chemical strength

pKa informs chemistry. **Biology defines efficacy.**


After this chemical section, it is important to **be very clear**:


pKa helps us understand acids, but it does not predict **biological efficacy**.


Clinical results emerge from how skin biology responds to a stimulus, not from pKa values alone.

Acid category	Acid subcategory	pKa>3 rising	pKa=3	pKa<3	pKa1	pKa2	pKa3	L. Dewandre	Number of reactions
Alpha Hydroxy	Aliphatic		Wine A (tartaric)		3.04	4.37		Metabolism	2
		Citric A			3.15	4.77	6.40	Metabolism	3 Triprotic
		Apples A (malic)			3.40	5.13		Metabolism	2 Diprotic
		Glycol A			3.83			Metabolism	1 monoprotic
CLASSIFICATION DR.ALAIN TENENBAUM		Milk.A (lactic)			3.86			Metabolism	1
	Aromatic Benzene ring		Almond.A mandelic		3.37			Metabolism	1
Alpha Keto				Grapes Acid (pyruvic)	2.49			Not available	1
Bicarboxylic acid		Azelain a			4.55	5.59		Metabolism	2
Beta-Hydroxid			Salicylic a		2.97			poisonous	1
TCA				TCA	0.54			caustic	1
Phenol	Aromatisch	Phenol			9.95			poisonous	Alcohol-Base

Acid category	Acid subcategory	pKa>3 rising	pKa=3	pKa<3	pKa1	pKa2	pKa3	L. Dewandre	Number of reactions
Alpha Hydroxy	Aliphatic		Wine A (tartaric)		3.04	4.37		Metabolism	2
		Citric A			3.15	4.77	6.40	Metabolism	3 Triprotic
		Apples A (malic)			3.40	5.13		Metabolism	2 Diprotic
		Glycol A			3.83			Metabolism	1 monoprotic
<div>CHOOSE AN ACID f(pKa)</div>		Milk.A (lactic)			3.86			Metabolism	1
	Aromatic Benzene ring		Almond.A mandelic		3.37			Metabolism	1
Alpha Keto				Grapes Acid (pyruvic)	2.49			Not available	1
Bicarboxylic acid		Azelain a			4.55	5.59		Metabolism	2
Beta-Hydroxid			Salicylic a		2.97			poisonous	1
TCA				TCA	0.54			caustic	1
Phenol	Aromatisch	Phenol			9.95			poisonous	Alcohol-Base

Acid category	Acid subcategory	pKa>3 rising	pKa=3	pKa<3	pKa1	pKa2	pKa3	L. Dewandre	Number of reactions
Alpha Hydroxy	Aliphatic		Wine A (tartaric)		3.04	4.37		Metabolism	2
		Citric A			3.15	4.77	6.40	Metabolism	3 Triprotic
		Apples A (malic)			3.40	5.13		Metabolism	2 Diprotic
		Glycol A			3.83			Metabolism	1 monoprotic
<div> <div>EU LAW</div> <div>Cosmeticians cant use any acid with pKa <3</div> </div>		Milk.A (lactic)			3.86			Metabolism	1
	Aromatic Benzene ring		Almond.A mandelic		3.37			Metabolism	1
Alpha Keto				Grapes Acid (pyruvic)	2.49			Not available	1
Bicarboxylic acid		Azelain a			4.55	5.59		Metabolism	2
Beta-Hydroxid			Salicylic a		2.97			poisonous	1
TCA				TCA	0.54			caustic	1
Phenol	Aromatisch	Phenol			9.95			poisonous	Alcohol-Base

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		Citric A			3.15	4.77	6.40	Metabolism	3 Triprotic
		Apples A (malic)			3.40	5.13		Metabolism	2 Diprotic
	<div>Aggressiveness</div> <div>e.g. citric A>glycolic A</div> <div>pKa 3.15<3.83</div>		Glycol A		3.83			Metabolism	1 monoprotic
			Milk.A (lactic)		3.86			Metabolism	1
	Aromatic Benzene ring		Almond.A mandelic		3.37			Metabolism	1
Alpha Keto				Grapes Acid (pyruvic)	2.49			Not available	1
Bicarboxylic acid		Azelain a			4.55	5.59		Metabolism	2
Beta-Hydroxid			Salicylic a		2.97			poisonous	1
TCA				TCA	0.54			caustic	1
Phenol	Aromatisch	Phenol			9.95			poisonous	Alcohol-Base

Acid category	Acid subcategory	pKa>3 rising	pKa=3	pKa<3	pKa1	pKa2	pKa3	L. Dewandre	Number of reactions
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		Citric A			3.15	4.77	6.40	Metabolism	3 Triprotic
		Apples A (malic)			3.40	5.13		Metabolism	2 Diprotic
	<div> DANGER Low Nr of reactions e.g. glycolic A vs citric A Mono vs Triprotic </div>	Glycol A			3.83			Metabolism	1 monoprotic
		Milk.A (lactic)			3.86			Metabolism	1
	Aromatic Benzene ring		Almond.A mandelic		3.37			Metabolism	1
Alpha Keto				Grapes Acid (pyruvic)	2.49			Not available	1
Bicarboxylic acid		Azelain a			4.55	5.59		Metabolism	2
Beta-Hydroxid			Salicylic a		2.97			poisonous	1
TCA				TCA	0.54			caustic	1
Phenol	Aromatisch	Phenol			9.95			poisonous	Alcohol-Base

Anti Aging Peelings on Phototype 3
Treatment : Citric Acid each day x 15 days
+ 2 times metabolic peels 1 x week

BEFORE



AFTER 15 DAYS



Protocol : Go on Peeling (citric acid) on the skin even during desquamation
Patient can go to the beach (Cancun) each day

Courtesy of Dr. Alain Tenenbaum

Anti Aging Peelings on Phototype 3
Treatment : Citric Acid each day x 15 days
+ 2 times metabolic peels 1 x week

BEFORE



AFTER 15 DAYS



MORE SESSIONS > MORE AGGRESSION

CUMULATIVE IMPROVEMENT

WITHOUT HIGH-RISK AGGRESSION



LOW-INTENSITY, REPEATABLE PEELS OUTPERFORM "ONE-SHOT" AGGRESSIVE APPROACHES



TO REMEMBER

Compared with glycolic acid, citric acid is generally regarded as less hazardous, yet it can induce more aggressive tissue reactions and is commonly more cost-effective.

HOW TO AVOID SOCIAL DOWNTIME


- It is preferable to repeat multiple treatment sessions using a **safe and sufficiently active acid** rather than performing a single session with a **highly aggressive and potentially hazardous acid**.


(e.g., TCA vs citric acid)

- It is preferable to repeat sessions with a **lower acid concentration** rather than using a **hyperconcentrated acid** in one or two sessions
(not applicable to hyperchromias).

- Example with TCA: **4-3-2-2-...**
instead of **1 × 4**

Acid category	Acid subcategory	pKa>3 rising	pKa=3	pKa<3	pKa1	pKa2	pKa3	L. Dewandre	Number of reactions
Alpha Hydroxy	Aliphatic		Wine A (tartaric)		3.04	4.37		Metabolism	2
		Citric A			3.15	4.77	6.40	Metabolism	3 Triprotic
		Apples A (malic)			3.40	5.13		Metabolism	2 Diprotic
KERATOREGULATORS		Glycol A			3.83			Metabolism	1 monoprotic
		Milk.A (lactic)			3.86			Metabolism	1
	Aromatic Benzene ring		Almond.A mandelic		3.37			Metabolism	1
Alpha Keto				Grapes Acid (pyruvic)	2.49			Not available	1
Bicarboxylic acid		Azelain a			4.55	5.59		Metabolism	2
Beta-Hydroxid			Salicylic a		2.97			poisonous	1
TCA				TCA	0.54			caustic	1
Phenol	Aromatisch	Phenol			9.95			poisonous	Alcohol-Base

Acid category	Acid subcategory	pKa>3 rising	pKa=3	pKa<3	pKa1	pKa2	pKa3	L. Dewandre	Number of reactions
Alpha Hydroxy	Aliphatic		Wine A (tartaric)		3.04	4.37		Metabolism	2
		Citric A			3.15	4.77	6.40	Metabolism	3 Triprotic
		Apples A (malic)			3.40	5.13		Metabolism	2 Diprotic
	MOISTURIZERS	Glycol A			3.83			Metabolism	1 monoprotic
		Milk.A (lactic)			3.86			Metabolism	1
	Aromatic Benzene ring		Almond.A mandelic		3.37			Metabolism	1
Alpha Keto				Grapes Acid (pyruvic)	2.49			Not available	1
Bicarboxylic acid		Azelain a			4.55	5.59		Metabolism	2
Beta-Hydroxid			Salicylic a		2.97			poisonous	1
TCA				TCA	0.54			caustic	1
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	KERATOREGULATORS+ MOISTURIZERS		Glycol A		3.83			Metabolism	1 monoprotic
		Milk.A (lactic)			3.86			Metabolism	1
	Aromatic Benzene ring		Almond.A mandelic		3.37			Metabolism	1
Alpha Keto				Grapes Acid (pyruvic)	2.49			Not available	1
Bicarboxylic acid		Azelain a			4.55	5.59		Metabolism	2
Beta-Hydroxid			Salicylic a		2.97			poisonous	1
TCA				TCA	0.54			caustic	1
Phenol	Aromatisch	Phenol			9.95			poisonous	Alcohol-Base

From Chemistry to Biology:

What Really Determines Clinical Outcomes

Chemistry

- Describes molecular behavior
- Defines ionization and penetration
- Controls exposure and delivery

Biology

- Determines cellular response
Modulates inflammation
- Drives tissue adaptation and renewal

Clinical outcome

Emerges from biological regulation
Not from chemical strength alone
Not from visible injury

Chemistry initiates the stimulus.
Biology determines the result.

Le **pKa** peut être comparé
à un agresseur qui porte
un coup de poing.

La biologie représente la victime,
et l'outcome clinique
correspond aux conséquences finales —
bénignes, sévères ou incapacitantes.

Un coup de poing porté avec la même force n'aura
pas les mêmes conséquences biologiques ni cliniques
selon qu'il touche les fesses, le nez ou le sternum.

Ce n'est pas le coup en lui-même qui détermine
la gravité, mais le tissu touché, sa **structure**, sa
capacité d'adaptation et sa réponse biologique.

La biologie représente la victime, et l'outcome
clinique correspond aux conséquences finales —
bénignes, sévères ou incapacitantes.

pKa can be compared to
an aggressor delivering a punch.

Biology represents the victim, and
the clinical outcome reflects the
final consequences — mild,
severe, or incapacitating.

A punch delivered with the same force will **not**
produce the same biological effects or clinical
outcomes if it strikes the buttocks, the
nose, or the sternum.

Severity is **not** determined by the punch itself,
but by the tissue involved, its
structure, vulnerability, and biological response.

Pthe same way, **pKa** describes the chemical aggression,
but clinical efficacy and risk are determined
by tissue biology, not by chemical strength alone.

METABOLIC PEELS

CLASSIFICATION OF L.DEWANDRE

REVISED & UPDATED BY A.TENENBAUM



Metabolic Peels are next generation skin peels that incorporate Chirally Correct AHA and encapsulated Retinoic Acid to stimulate the skins natural renewal processes.



Peels without downtime or irritation,



Metabolic Peels target the living layer of the skin(Stratum Germinativum or Basal) and increase cell renewal.

From Chemistry to Biology:

What Really Determines Clinical Outcomes

Chemistry

- Describes molecular behavior
- Defines ionization and penetration
- Controls exposure and delivery

Biology

- Determines cellular response
Modulates inflammation
- Drives tissue adaptation and renewal

Clinical outcome

Emerges from biological regulation
Not from chemical strength alone
Not from visible injury

Chemistry initiates the stimulus.
Biology determines the result.

Metabolic Peels vs Traditional Chemical Peels

METABOLIC PEELS

- Therapeutic philosophy:
Support and modulate skin metabolism.
- Primary target:
Living epidermal layers (stratum basale / germinativum)
- Mechanism of action:
 - Metabolic activation
 - Controlled stimulation of crenewal
 - Respect of physiological skin proceses
- Tissue response:
 - No forced tissue destruction
 - Minimal or no inflammation
 - No obligatory desquamation
- Clinical approach:
 - Non-aggressive

Primary goal:

Restore skin function and long-term balance

TRADITIONAL CHEMICAL PEELS

- Therapeutic philosophy:
Force tissue repair through chemical injury.
- Primary target:
Stratum corneum and superficial dermis
- Mechanism of action:
 - Protein coagulation
 - Chemical burn-induced injury
 - Inflammatory-oxidative activation
- Clinical approach:
 - Ablative or semi-ablative
 - Aggression-dependent efficacy
 - Depth-oriented classification
- Downtime:
 - Variable, often significant

Primary goal:

Induce repair through controlled injury.

Metabolic Peels work with skin physiology, while traditional chemical peels work through controlled injury.

Biological Targets of Metabolic Peels

Target: Epidermal Cell Renewal

- Primary biological target:
- Basal keratinocytes (stratum basale / germinativum)
- Mechanism:
 - Activation of cell proliferation
 - Regulation of differentiation pathways
 - Acceleration of physiological epidermal turnover
- Clinical relevance:
 - Improved skin texture and tone
 - Progressive normalization without forced desquamation

Target: Cellular Metabolism and Signaling

- Primary biological target:
- Intracellular metabolic pathways
- Mechanism:
 - Modulation of enzymatic activity
 - Regulation of oxidative balance
 - Optimization of cellular energy metabolism
- Clinical relevance:
 - Improved skin vitality
 - Reduced dysfunctional signaling
 - Enhanced response to subsequent treatments

OXIDATION AND EFFECT



SUFFIXE - AL for ALDEHYDE

Retinoic acids work best when they are stable
Peeling de Luxe Plus contains stable retinoic acids

Retinoic acids are more expensive than retinol



Chemical Reactions: Lab vs. Skin Biology

Laboratory Chemistry

Carboxylic Acid \rightarrow Alcohol \rightarrow R-CH₂-OH
(Strong Reduction)

Alcohol \rightarrow Aldehyde \rightarrow Carboxylic Acid



(Oxidation)



Controlled Lab Reactions

In Vivo
Enzymatic Pathways

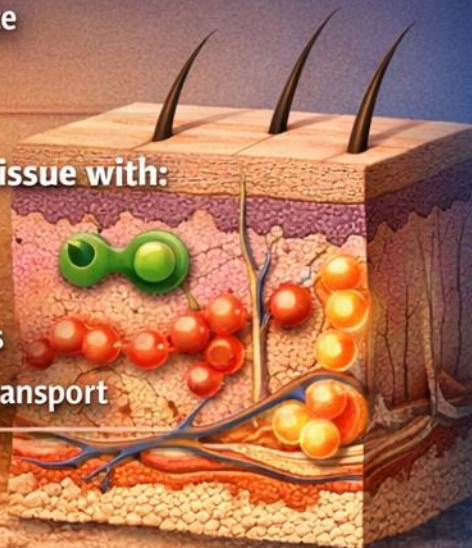
Biological Metabolism

Molecules Applied to Skin:

- Retinol
- Retinyl Palmitate
- Retinal

Enter a Living Tissue with:

- Enzymes
- Cofactors
- Redox Systems
- Intracellular Transport



The Skin is Not a Beaker

RETINOIC PATHWAY

Acid category	Acid subcategory	pKa>3 rising	pKa=3	pKa<3	pKa1	pKa2	pKa3	L. Dewandre	Number of reactions
Alpha Hydroxy	Aliphatic		Wine A (tartaric)		3.04	4.37		Metabolism	2
		Citric A			3.15	4.77	6.40	Metabolism	3 Triprotic
		Apples A (malic)			3.40	5.13		Metabolism	2 Diprotic
		Glycol A			3.83			Metabolism	1 monoprotic
CLASSIFICATION DR.L.DEWANDRE		Milk.A (lactic)			3.86			Metabolism	1
	Aromatic Benzene ring		Almond.A mandelic		3.37			Metabolism	1
Alpha Keto				Grapes Acid (pyruvic)	2.49			Not available	1
Bicarboxylic acid		Azelain a			4.55	5.59		Metabolism	2
Beta-Hydroxid			Salicylic a		2.97			poisonous	1
TCA				TCA	0.54			caustic	1
Phenol	Aromatisch	Phenol			9.95			poisonous	Alcohol-Base

Positioning of Metabolic Peels Within the Protocol

Metabolic Peels are not defined by intensity, but by their role within the protocol sequence

BLOCK 1 — CORRECTION PHASE

Primary role:

Active metabolic correction

Clinical indications:

- Dyschromias
- Acne and post-acne sequelae
- Photoaging with functional impairment
- Altered epidermal renewal

Mechanism:

- Stimulation of cell renewal from the basal layer
- Metabolic activation without forced tissue injury



Clinical objective:

Correct pathological signaling while minimizing trauma

STABILIZATION PHASE

Primary role:

Functional normalization and consolidation

Clinical indications:

- Post-correction phase
 - Barrier recovery
- Reduction of inflammatory rebound
- Prevention of pigment relapse

Mechanism:

- Support of physiological regeneration
- Regulation of epidermal metabolism
 - Buffering of previous corrective interventions



Clinical objective:

Stabilize results and restore skin homeostasis

MAINTENANCE PHASE

Primary role:

Long-term metabolic support

Clinical indications:

- Maintenance of clinical results
 - Anti-aging prevention
- Long-term pigment control
 - Home-care integration

Mechanism:

- Controlled stimulation without downtime
- Support of natural skin renewal cycles



Clinical objective:

Preserve results and prevent recurrence over time

Chiral Correction

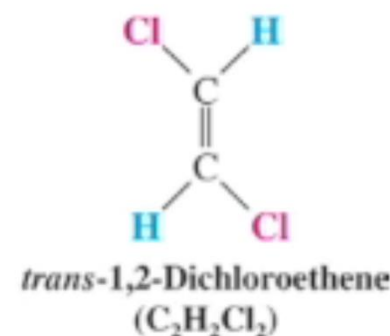
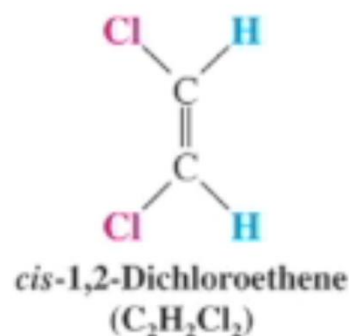
What is Chiral Correction?

In a Nut Shell - Chiral-correction is the purification on a molecular level to ensure optimal interaction with your body.

The Science - Some molecules are like hands. Left and right hands are mirror images, but are not superimposable.

A molecule that is not superimposable is said to be **chiral**.

Chemically the same ingredient, but structurally different. An L-or D-prefix denote the chiral version.



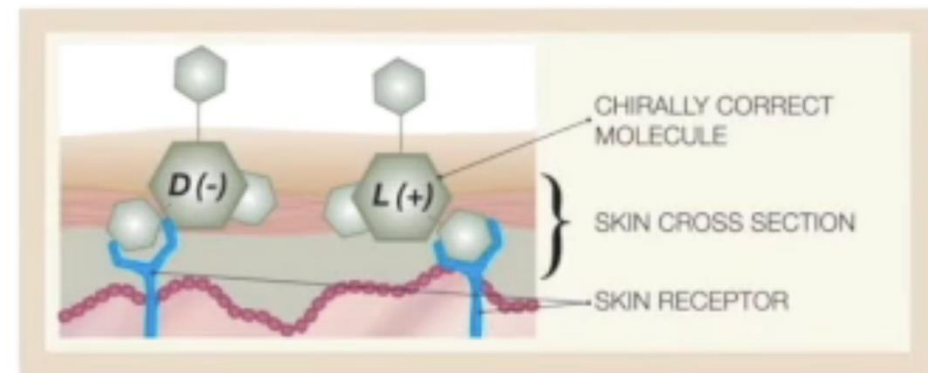
Chiral Correction 2

Implications for the Skin

Although chemically two forms of an ingredient may be identical, because their shape is different they will interact with the skin differently.

Like a **key in a lock**, only the chirally-correct form of an ingredient can interact with the appropriate cell receptor and trigger the right cellular response. The wrong form of the ingredient can collect on the skin's surface and can cause irritation.


How a Chiral Molecule Works



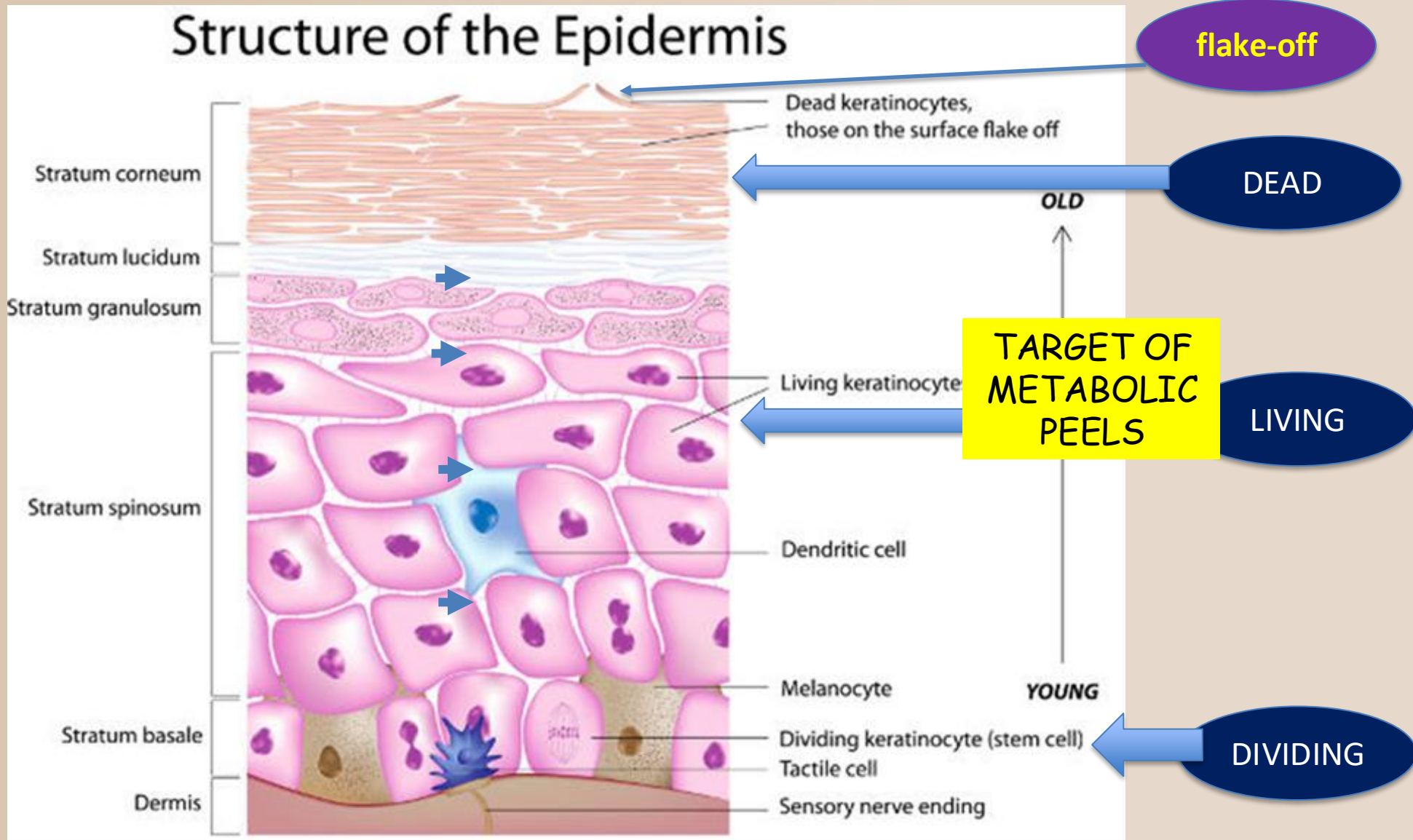
CosMedix ingredients contain only the correct-fitting molecules for the skin's receptors

- L-Lactic Acid - Boosts ceramide production by 50%
- D-Lactic Acid - Has no effect on ceramide production and causes irritation.

Ceramides are fats or lipids that are found in skin cells. They make up 30% to 40% of your outer skin layer, or epidermis. Ceramides are important for retaining your skin's moisture and preventing the entry of germs into your body.

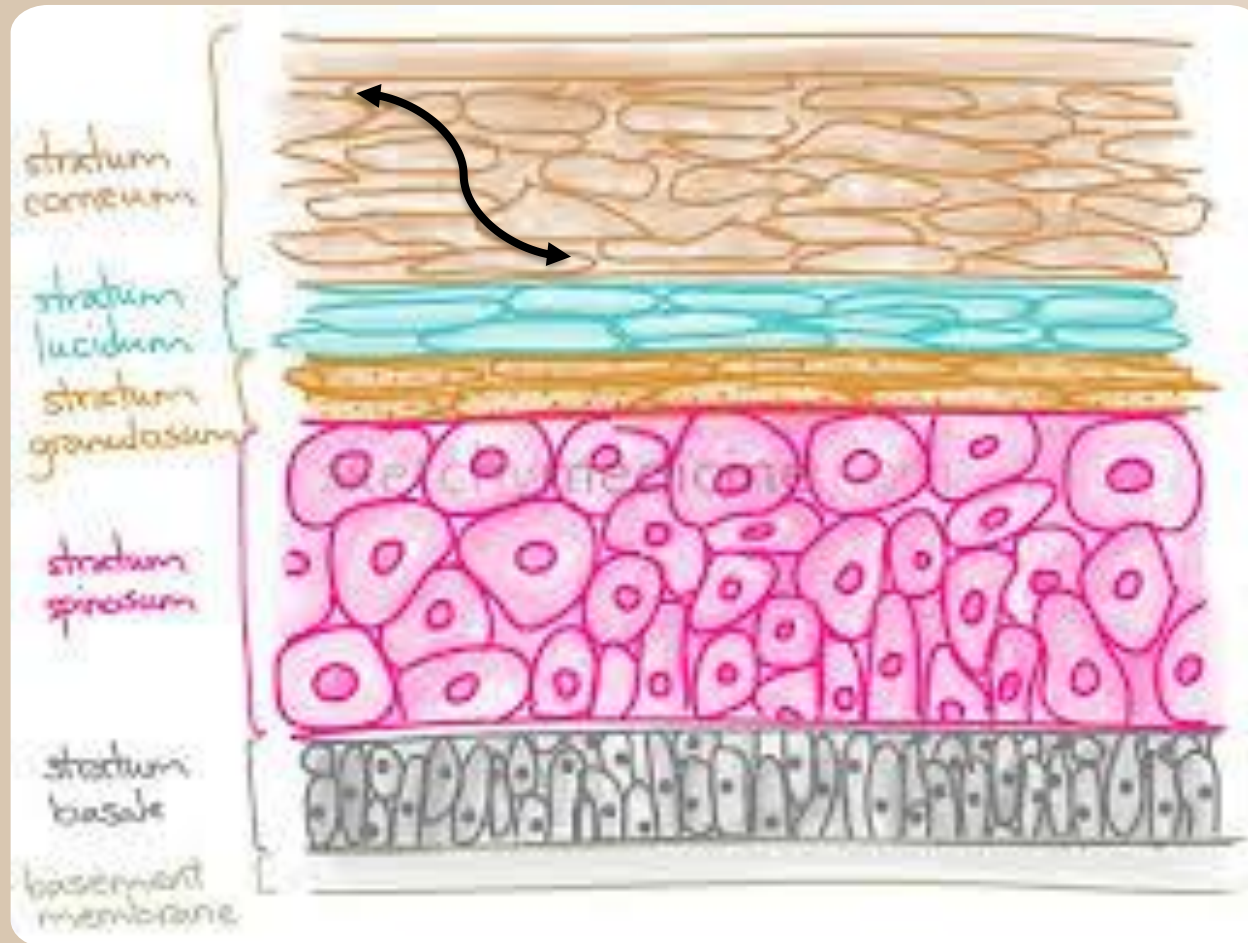
Acid category	Acid subcategory	pKa>3 rising	pKa=3	pKa<3	pKa1	pKa2	pKa3	Tenenbaum ENANTIOMERS	Number of reactions
Alpha Hydroxy	Aliphatic		Wine A (tartaric)		3.04	4.37		Chiral	2
		Citric A			3.15	4.77	6.40	Chiral	3 Triprotic
		Apples A (malic)			3.40	5.13		Chiral	2 Diprotic
CLASSIFICATION Chiral Acids A.TENENBAUM		Glycolic A	Skin does not have a receptor site for glycolic acid.		3.83			Not available	1 monoprotic
		Milk.A (lactic)			3.86			Chiral	1
	Aromatic Benzene ring		Almond.A mandelic		3.37			Chiral	1
Alpha Keto				Grapes Acid (pyruvic)	2.49			Not available	1
Bicarboxylic acid		Azelain a			4.55	5.59		Not available	2
Beta-Hydroxid			Salicylic a		2.97			Chiral Aspirin not chiral	1
TCA				TCA	0.54			caustic	1

MAIN TARGET OF METABOLIC PEELS



TRADITIONAL CHEMICAL PEELS

-> FORCING REPAIR



- Exfoliate the skin from the stratum corneum downwards



PH AND PKA

BUFFER IN DERMATOLOGY- SKIN HYDRATION

PH SKIN = 5.5

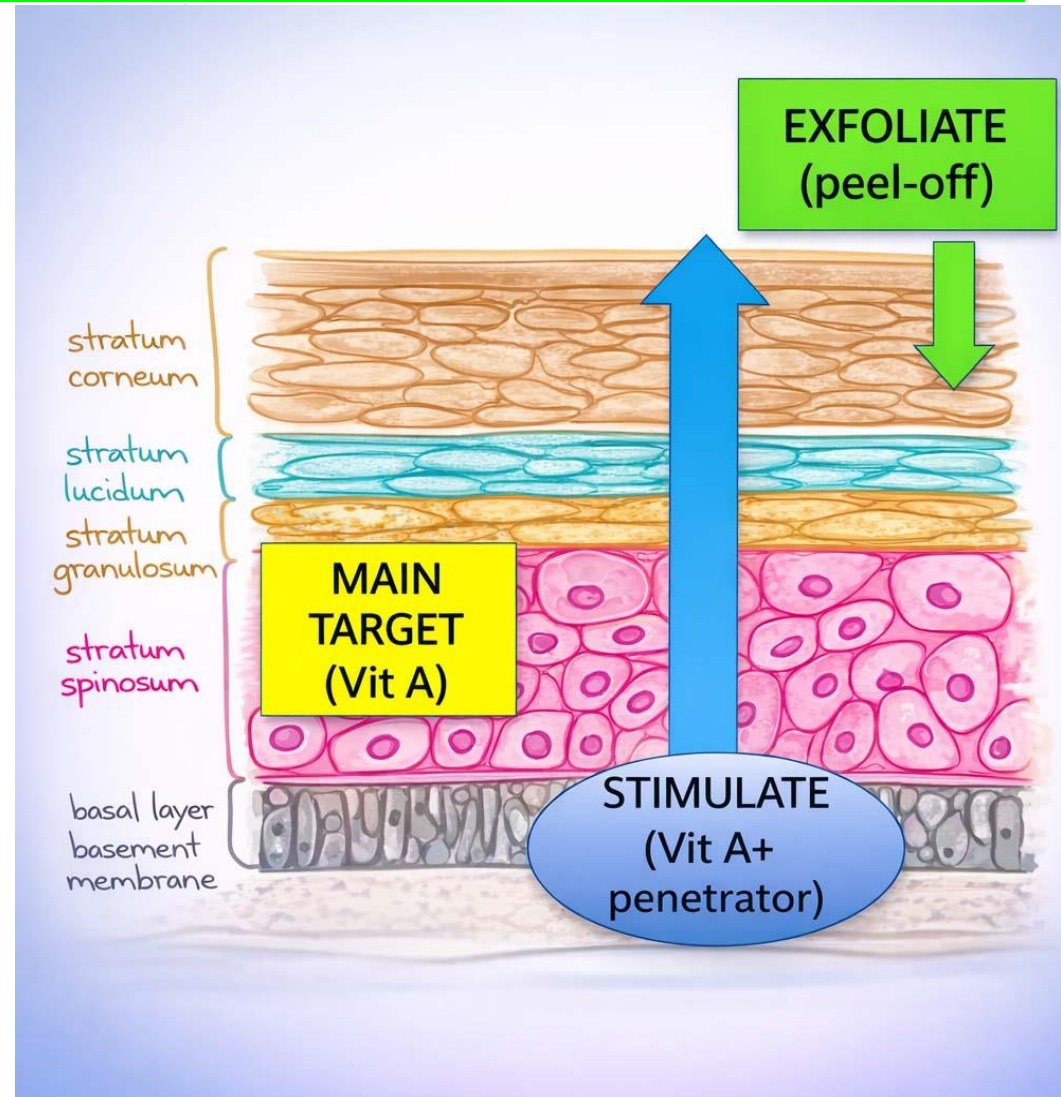
- $\text{pH skin (5.5)} < \text{pKa} < 7$ Moisturizing and less corrosive. (hydratant-MOISTURIZER)
- $0 < \text{pKa} < \text{pH skin (5.5)}$ keratoregulators and promote desquamation (desquamation + cell regeneration)
- $\text{pKa} = \text{pH (5.5) skin}$ Ideal (citric acid - triprotic) moisturizing + desquamation

Die Haut muss nach einem Peeling mit einer keratoregulatorischen Säure ($\text{pKa} < 5.5$) systematisch mit Feuchtigkeit versorgt werden

Peeling de Luxe Plus is the ideal buffer for any acid that needs to be buffered.
Peeling de Luxe Plus is not a moisturizer.

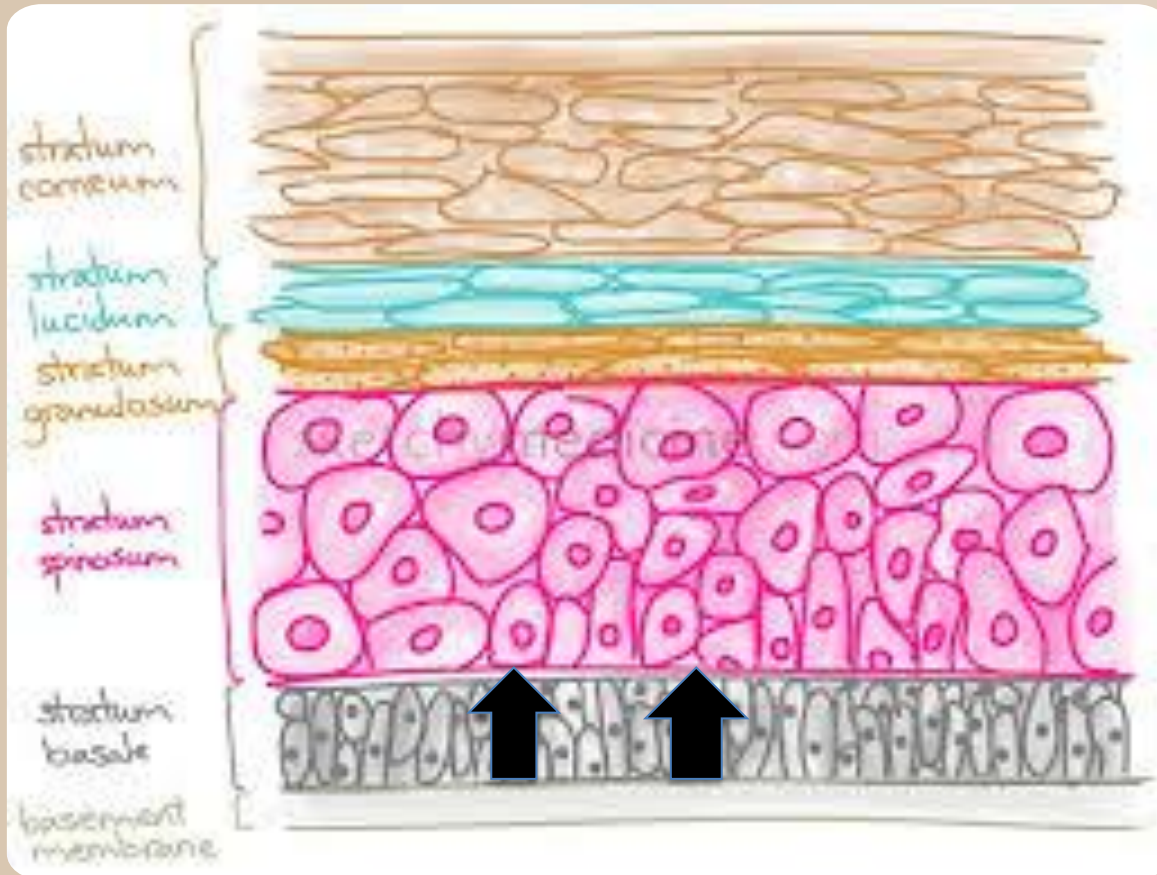
METABOLIC PEELS VS CHEMICAL PEELS

- Metabolic Peels contain Vit A that targets the living layer of the skin and increases cell renewal. (Ex *Peeling de Luxe Plus*)
- Traditional Chemical Peels use Acids to exfoliate the skin from the stratum corneum downwards (Ex 30 min peel-off)
- Metabolic Peels use encapsulated Retinoic Acid to stimulate renewal from the stratum germinativum upwards, working with the skins natural renewal instead of forcing repair.
- This is a non ablative, non traumatic approach to skin peeling that supports skin function



Stratum basal = Stratum Germinativum

METABOLIC PEELS -> SKIN NATURAL RENEWAL SUPPORTING SKIN FUNCTION



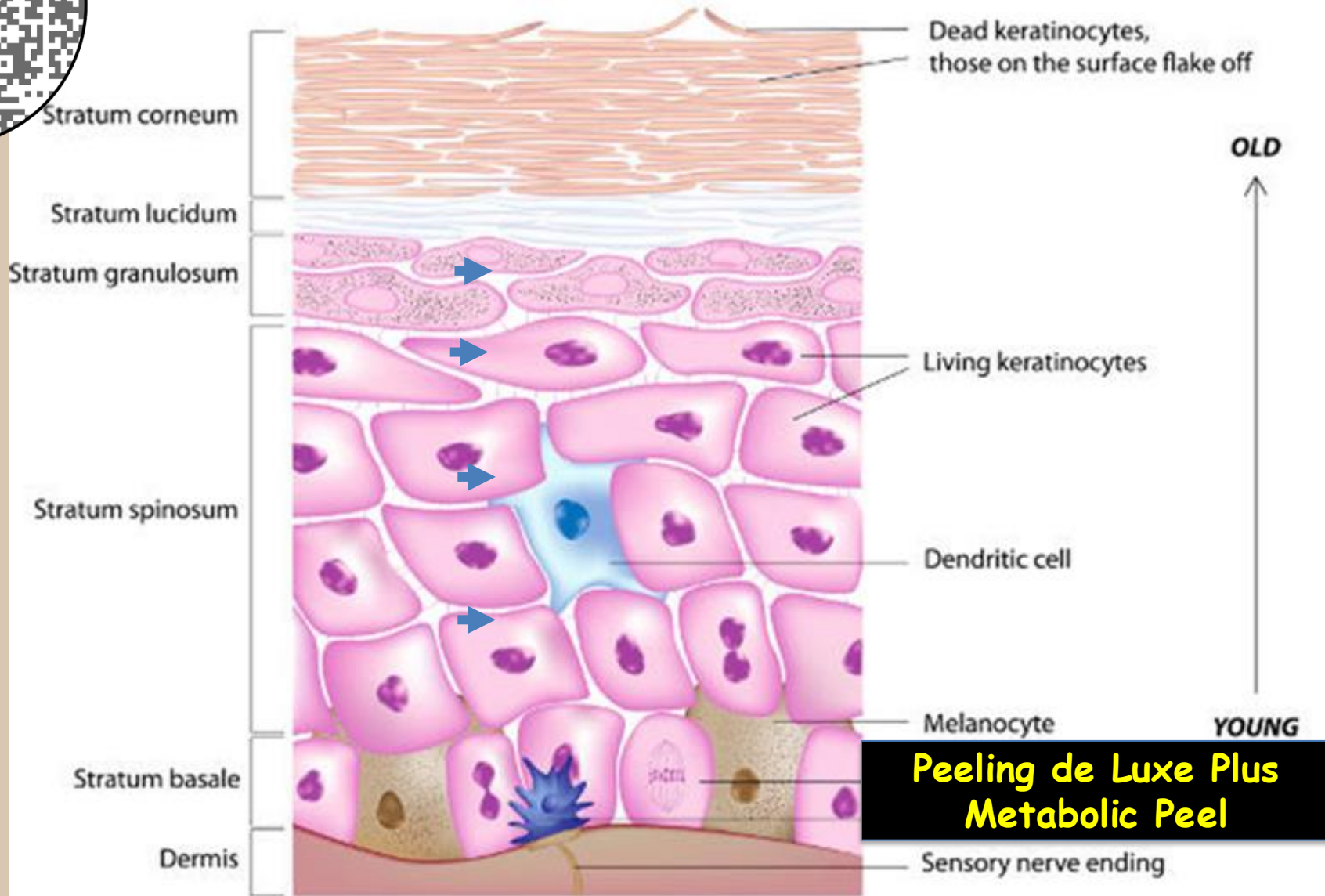
- Stimulate renewal from the stratum basal or germinativum upwards

DIFFERENT TARGETS OF EPIDERMIS

METABOLIC PEEL TARGETS STRATUM BASAL AND INCREASES CELL RENEWAL

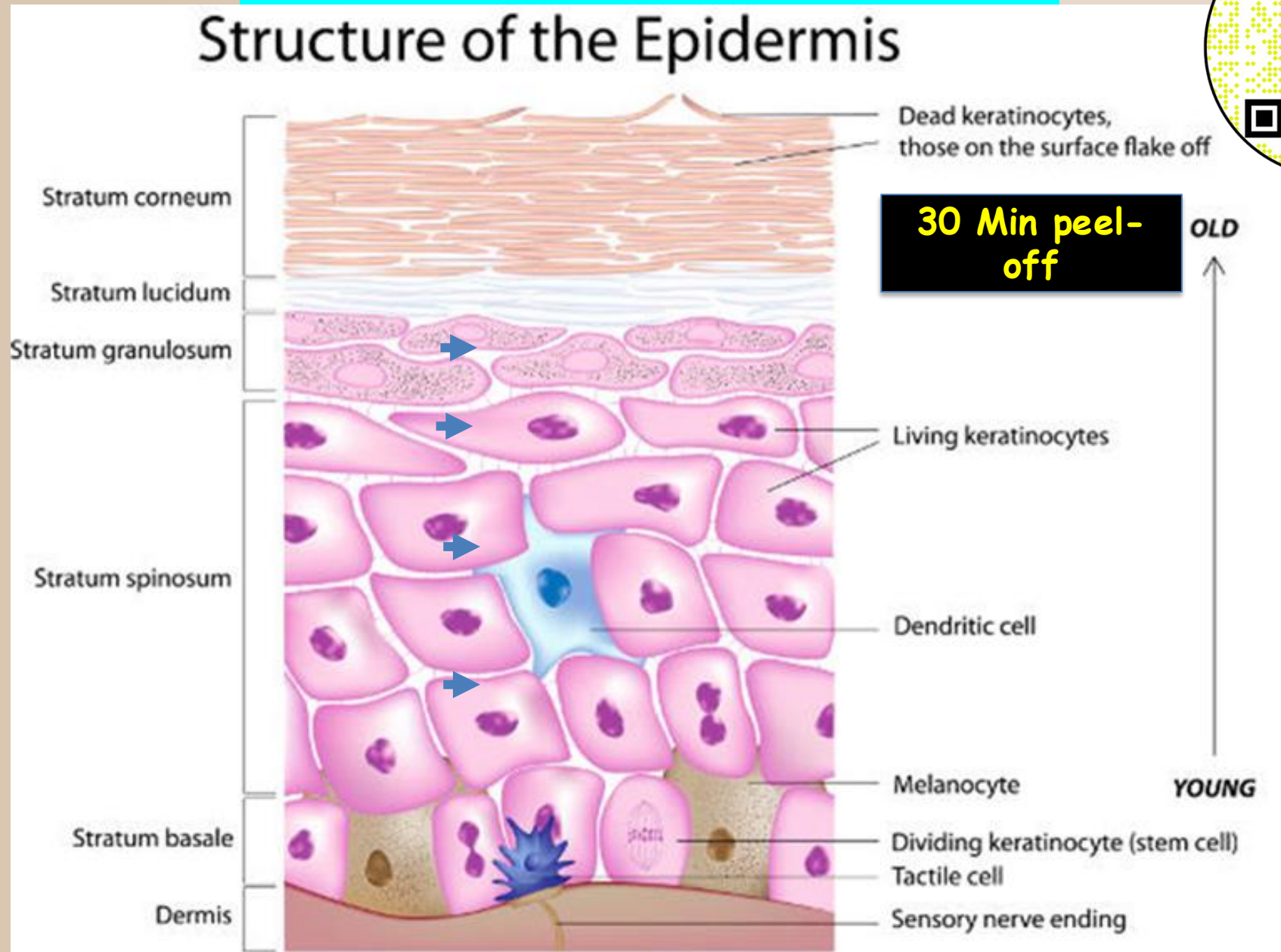


Structure of the Epidermis

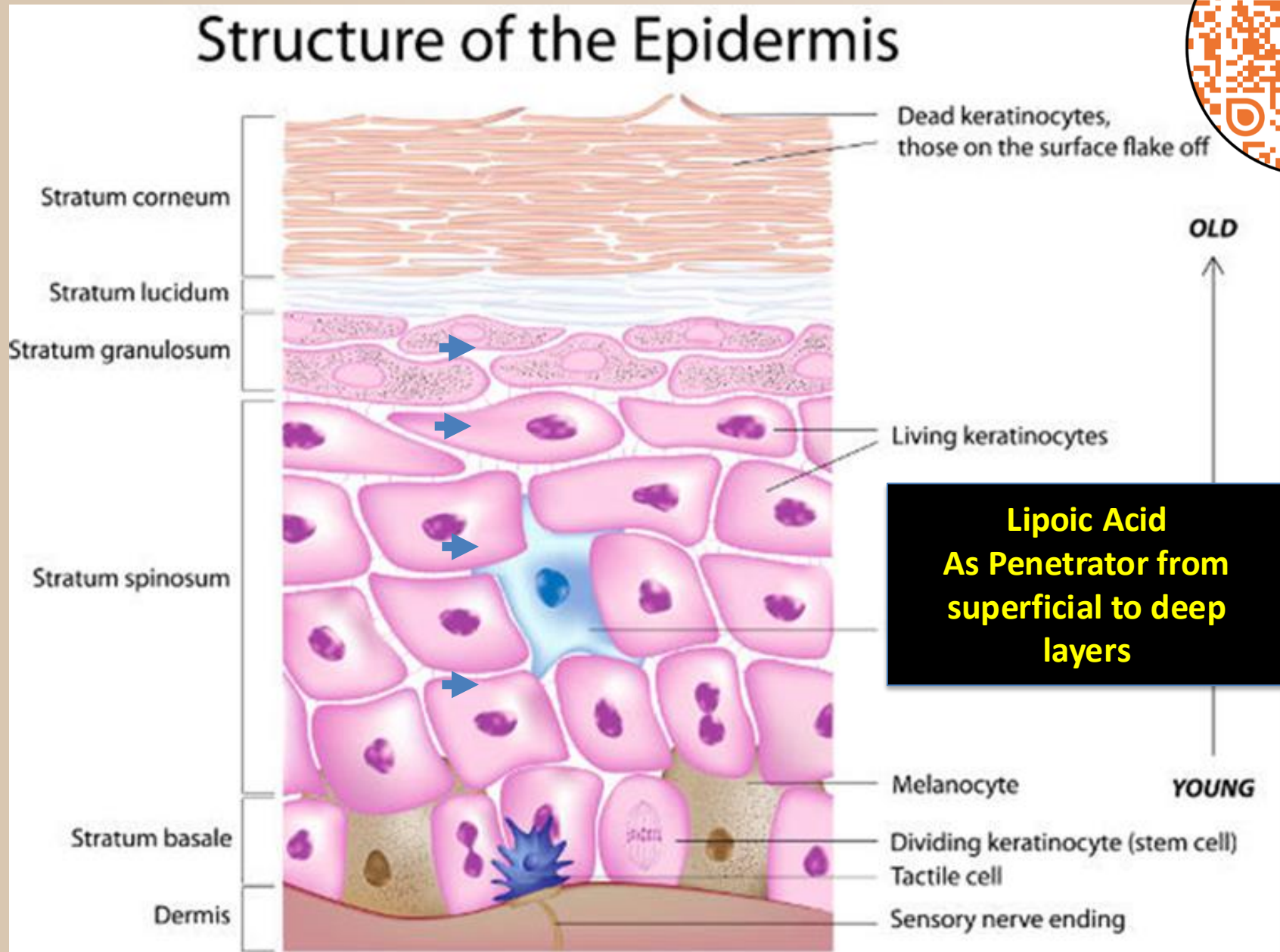


DIFFERENT TARGETS OF EPIDERMIS

SUPERFICIAL TARGET



Different Targets of Epidermis



Why Desquamation Is Not Mandatory

DESQUAMATION-BASED APPROACH

- Underlying assumption:
Visible peeling equals clinical efficacy
- Biological mechanism:
 - Chemical injury to the stratum corneum
 - Protein coagulation and necrosis
 - Inflammatory response
- Clinical consequences:
 - Epidermal disruption
 - Barrier impairment
 - Downtime and social eviction
- Clinical consequences:
 - Epidermal disruption
 - Barrier impairment
 - Increased risk of PIH and complications
- Clinical limitation:
 - Efficacy depends on the degree of tissue damage

FUNCTION-BASED APPROACH






- Underlying principle:
Clinical efficacy does not require visible peeling
- Biological mechanism:
 - Activation of epidermal metabolism
 - Stimulation of cell renewal from living layers
 - Regulation of skin signaling pathways
- Clinical consequences:
 - Preservation of the skin barrier
 - Minimal inflammation
 - No mandatory downtime
 - Improved safety profile
- Clinical advantage:
 - Efficacy achieved through functional modulation, not through tissue destruction
- Clinical advantage:
 - Efficacy achieved through functional modulation, not through tissue destruction

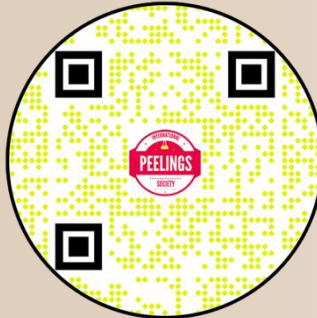
TREATMENT PROTOCOL OF MAURO TIZIANI

3 TARGETS = 3 PRODUCTS

STIMULATE, EXFOLIATE, PENETRATE

THE METABOLIC PEELS SET

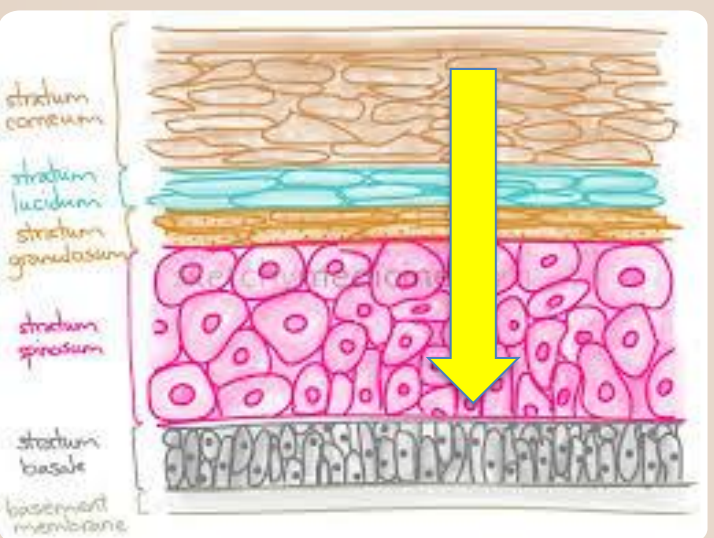
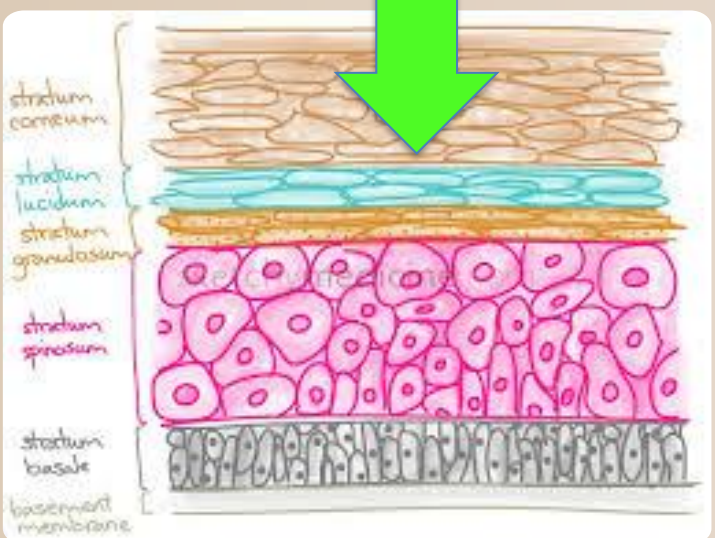
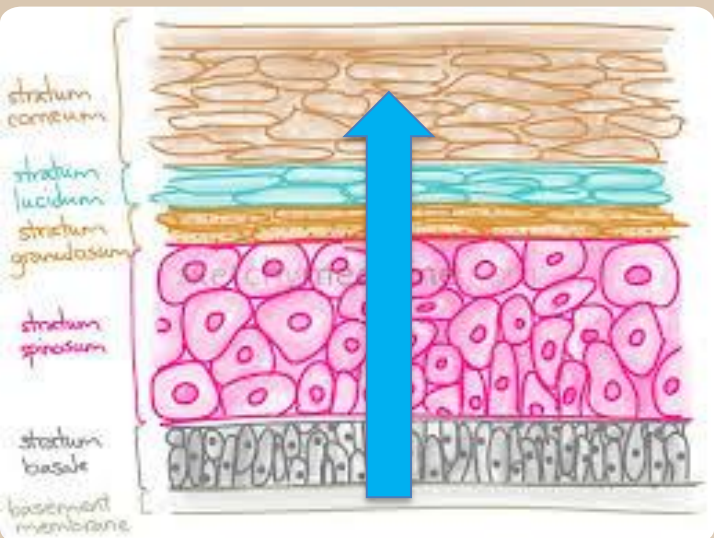
STIMULATE	EXFOLIATE	PENETRATE
		
PEELING DE LUXE PLUS PURE METABOLIC PEEL	30 MIN PEEL OFF	LIPOIC ACID
 <p>PEELING DE LUXE PLUS</p> <p>CLINIC SIZE 1 KG</p>	 <p>30 MIN PEEL OFF</p> <p>CLINIC SIZE 1 KG</p>	 <p>LIPOIC ACID CREAM</p> <p>CLINIC SIZE 1 KG</p>



STIMULATE

EXFOLIATE

PENETRATE



PEELING DE LUXE PLUS

30 MIN PEEL OFF

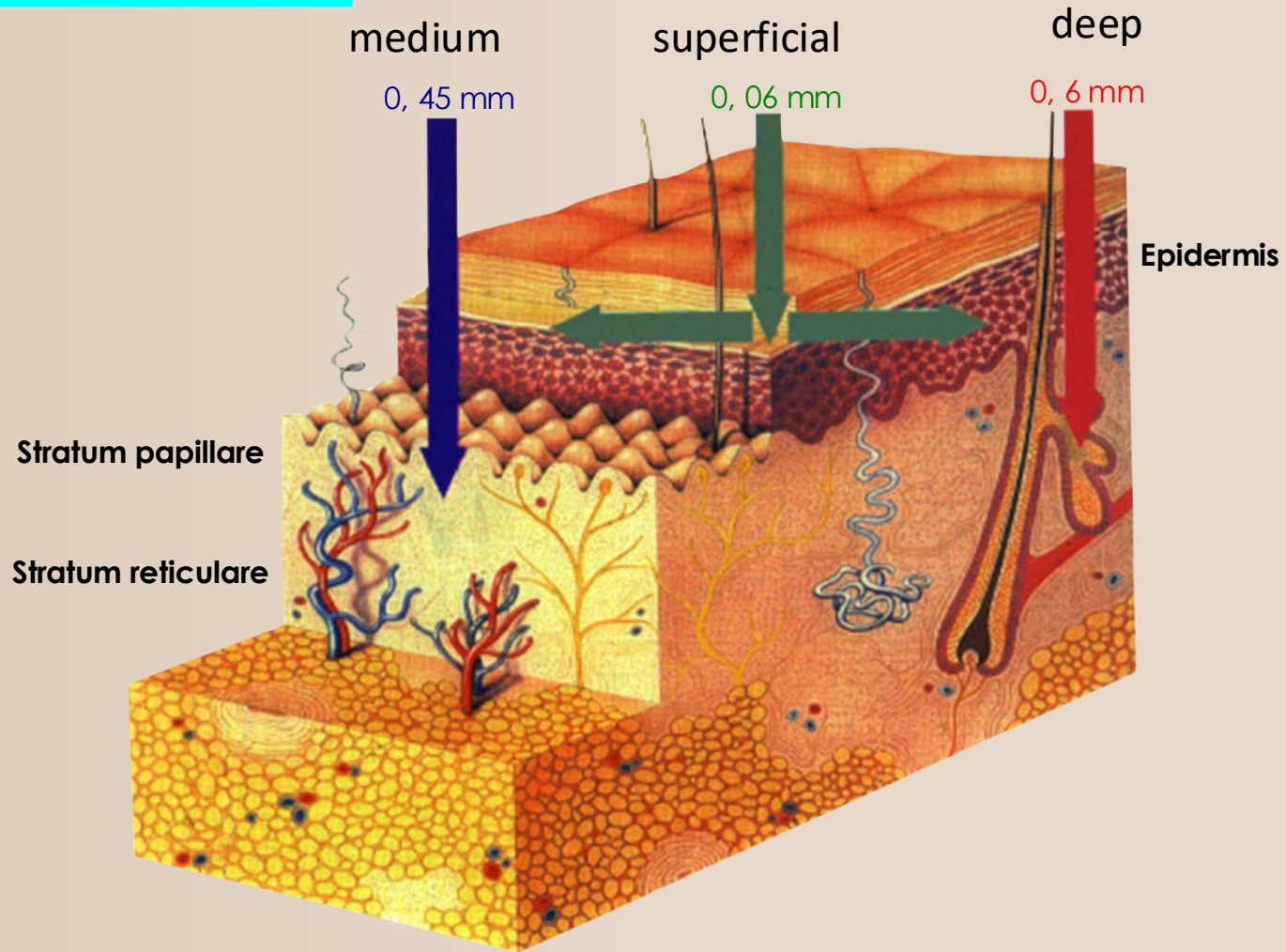
LIPOIC ACID



SUPERFICIAL, MEDIUM & DEEP PEELS

PENETRATION DEPENDS OF PH MODULATION

NEW CONCEPT OF MAURO TIZIANI



Get deeper with Lipoic Acid without modulating the
pH
(A.TENENBAUM)



CONCEPT OF MAURO TIZIANI

- Chin
- Perioral area
- Forehead
- Nasal Tip
- Cheeks / nose
- Frown
- Periocular area
- Neck

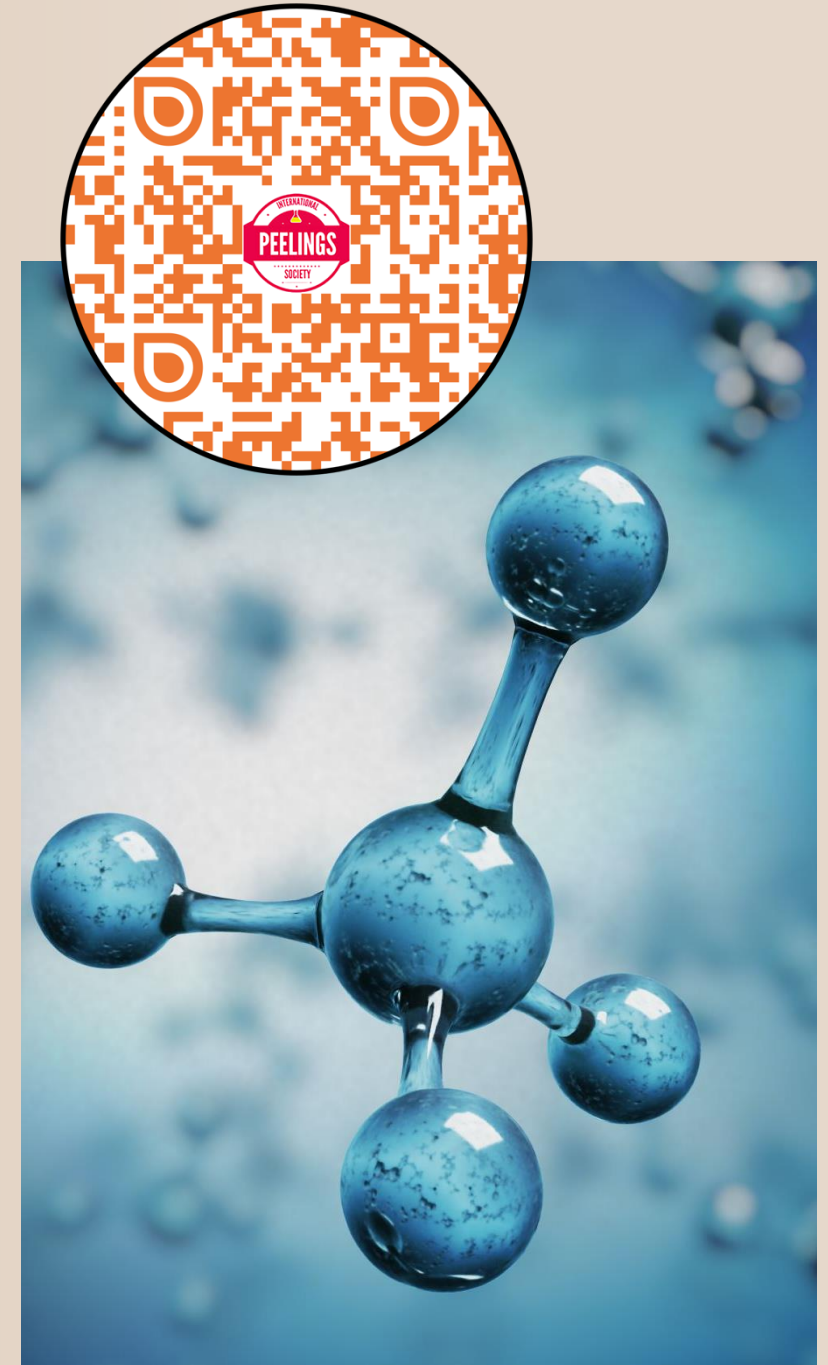


APPLICATION
PRESSURE



RECOMMENDED METHODS FOR HELPING INGREDIENTS TO PENETRATE INTO THE SKIN OR DECREASING THE PH (MODULATE THE PH)

- Q or exothermic reaction
(Cream 1 + Cream 2)
- Increasing the concentration of the acid
- More coats applied on same area with same acid
- Lipoic acid(A.TENENBAUM + M.TIZIANI)
- The fewer ingredients a cream contains, the easier it is to penetrate due to its low molecular weight (M.TIZIANI)
- For severely dehydrated skin,
use a low molecular weight moisturizer + Lipoic acid in a sequential way



HOW TO ALLOW AN ACID TO PENETRATE DEEPLY INTO THE SKIN LAYERS

CHOOSE AN AGGRESSIVE ACID (PKA MIN) AT
LOWER CONCENTRATION IS THE BEST TOOL

1. High acid concentration
2. Number of coats on the skin
3. Time on the skin before neutralization or defrosting
4. From Rosé Frosting to White Frosting
5. Repeat sessions (especially during desquamation)
6. Best Option: Add Lipoic Acid



LIPOLIC ACID



PRACTICAL TRENDS OF A. TENENBAUM

Do peels without
downtime or social
eviction, for all skin types
in any seasons

*Better a long ,,invisible,,
desquamation than a
short visible one (
downtime) A.Tenenbaum*

Choose best tools to
convert a superficial peel
into a medium or deep
peel (Q, lipoic acid ..)
reducing the side effects



FORBIDDEN COMBINATION OF CHEMICAL PEELS

- With not chiral molecules
- With alcohols I, II
- With bases
- With L+D mixed molecules
- With D molecules

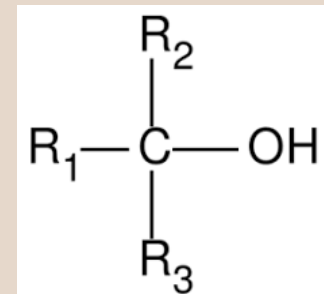
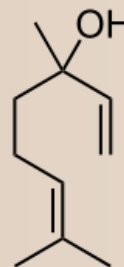


THE 3 TYPES OF PRIMARY, SECONDARY AND TERTIARY ALCOHOL

- **primary alcohol** $R-CH_2OH$ ex Ethanol CH_3-CH_2OH
- **Secondary** Alcohol $R_1-CHOH-R_2$
zB Isopropanol $CH_3-CHOH-CH_3$

Ethanol and isopropanol are disinfectants

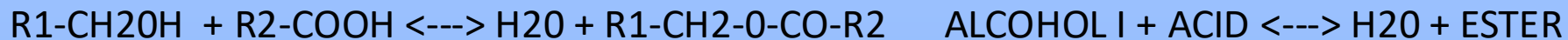
- **Tertiary** Alcohol ex Linalol oder 3,7-Dimethyl-1,6-octadien-3-ol is a tertiary alcohol with a floral and fresh smell.



ESTERIFICATION REACTIONS

THE ESTERS

- Alcohol I or II + acid \rightleftharpoons ester + H₂O
- Alcohol III : No esterification reaction with acids



For us, this means that the skin should never be disinfected with alcohol I or II before applying an acid to the patient's skin. Use better a wipe with cetrimonim (Aseptiskin)

Similarly, after exfoliation, the patient should not wear perfumed clothing or use perfume.

ESTERIFICATION REACTIONS

THE ESTERS

Ethanol as Desinfection
TCA after



Esterification Reaction on the Jowl
Alcohol as Skin Desinfectant + Acid



Courtesy of Dr. Alain Tenenbaum

PRIMARY & SECONDARY ALCOHOLS IN MOST SKINCARE SUNSCREENS SPF30!



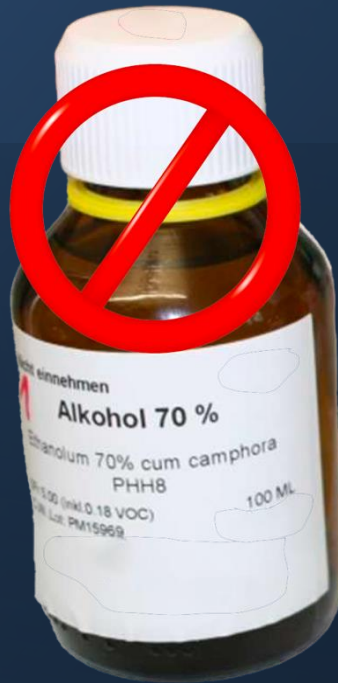
- **Cetyl Alcohol used as emollient**

- primary alcohol
- High molecular weight
- $\text{CH}_3-(\text{CH}_2)_{14}-\text{CH}_2\text{OH}$

- **Butylene Glycol used as Solvent**

- primary alcohol & secondary alcohol
- $\text{CH}_2\text{OH}-\text{CHOH}-\text{CH}_3$

ETHANOL IS UNFORTUNATELY MOSTLY USED AS
SKIN DESINFECTANT –SKIN TONICS- SKIN CLEANSERS
AND TOO AS „ DENATURED ALCOHOL !,,



A denatured alcohol is made unfit for human consumption by adding additives

Esterification Reaction on the Neck Perfume (Alcohol) + Acid

**ALCOHOL & PERFUME
ON TREATED SKIN**

**COMMON CAUSE OF
POST-PEEL IRRITATION**



Courtesy of Dr. Alain Tenenbaum

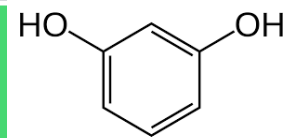
ONE MIXTURE
COMBINATION TO KNOW
JESSNER PEEL

OLD FASHION PEEL

100 mg of 95% ethanol !!



14 g of resorcinol (chiral).



It is neither a primary nor a secondary alcohol

14 g of salicylic acid (chiral)
BHA

and 14 ml of lactic acid (chiral)
AHA

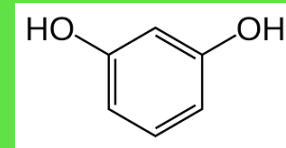
NOT SEQUENTIAL COMBINATION

THIS MODIFIED JESSNER IS FOR ME LOGIC

ONE MIXTURE
COMBINATION TO
KNOW
JESSNER PEEL
MODIFIED

5% W/V citric acid (chiral)


20% W/V of resorcinol
(chiral)



10% W/V lactic acid
(chiral)

NOT SEQUENTIAL COMBINATION

Modern PEEL

Acid category	Acid subcategory	pKa>3 rising	pKa=3	pKa<3	pKa1	pKa2	pKa3	L. Dewandre Modified by AT ENANTIOMERS	Number of reactions
Alpha Hydroxy	Aliphatic		Wine A (tartaric)		3.04	4.37		Chiral	2
 <div> <div>CLASSIFICATION</div> <div>DR.L.DEWANDRE</div> <div>modified by</div> <div>A.TENENBAUM</div> </div>		Citric A			3.15	4.77	6.40	Chiral	3 Triprotic
		Apples A (malic)			3.40	5.13		Chiral	2 Diprotic
		Glycolic A	Skin does not have a receptor site for glycolic acid.		3.83			Not available	1 monoprotic
		Milk.A (lactic)			3.86			Chiral	1
	Aromatic Benzene ring		Almond.A mandelic		3.37			Chiral	1
Alpha Keto				Grapes Acid (pyruvic)	2.49			Not available	1
Bicarboxylic acid		Azelain a			4.55	5.59		Not available	2
Beta-Hydroxid			Salicylic a		2.97			Chiral Aspirin not chiral	1
TCA				TCA	0.54			caustic	1

COMBINATION CHEMICAL PEELS VS SINGLE CHEMICAL PEELS IN MILD MODERATE ACNE

Single Chemical Peels

Combination Chemical Peels

TCA 30% W/V	Jessner followed by TCA 20% W/V- SEQUENTIAL
Salicylic 30% W/V	Salicylic 20% W/V + Mandelic 10% W/V MIXTURE

Side Effects !!!

Treatment Protocol : 6 SESSIONS WITH 2 WEEKS INTERVAL
FOLLOW UP 3 MONTHS AFTER LAST SESSION

OLD FASHION TREATMENT OF ACNE

SEQUENTIAL COMBINATION > MIXTURE COMBINATION

Mild Acne
treatment
with no side
effects since
26 years !



1-Preparation: Aseptiskin-PrePeel-
2-In Office: TCA 10-Peeling de Luxe Plus-30 min Peel Off-Lipoic Acid
3-Homecare/Suncare

**NEUTRALIZATION OF AN ACID
PH REGULATION
AND EXOTHERMIC REACTION (RELEASE OF THERMAL ENERGY)**



Hydrochloric acid + caustic soda \rightarrow table salt + water + Q



Products which are pH Regulators of Peels acids

NaHCO_3 (sodium bicarbonate salt) instead of NaOH (base) (IMPORTANT) is not a frosting stopper but increases the pH

PEELING DE LUXE PLUS
(frosting stopper) for your emergency equipment stops the duration of the low pH effect of the peelings acid

NEVER ADD WATER
because $\text{Acid} + \text{H}_2\text{O} = \text{Q}$
(combustion)

PH REGULATION AND NEUTRALIZER



NaHCO_3

Not frosting stopper

Use for acid with $\text{pKa} > 3$



PEELING DE LUXE



PLUS

FROSTING STOPPER

for your emergency equipment

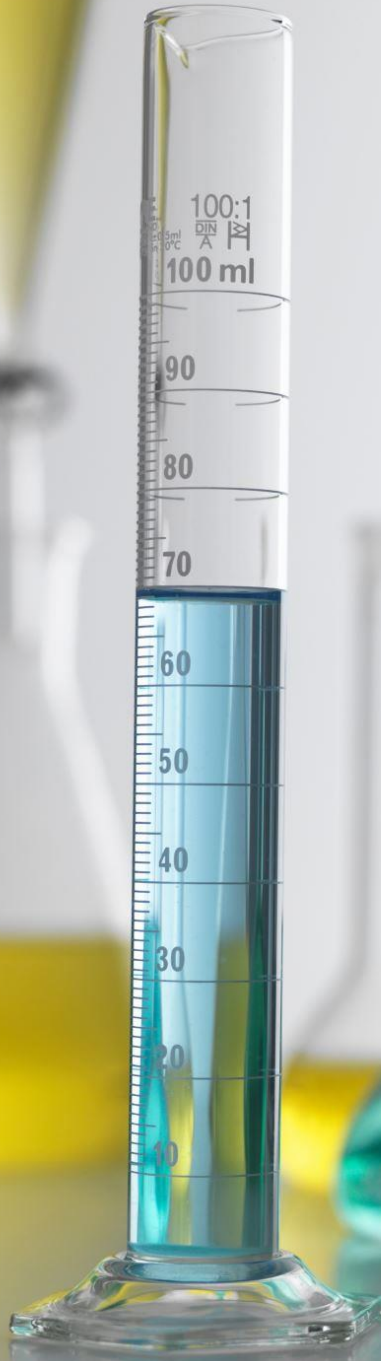
Use for acid with $\text{pKa} < 3$



BUFFER OR TAMPON

$\text{PKA} = \text{PH}$

- A buffer solution is used to limit fluctuations in pH.
- If you want to buffer a „weak,, acid , you need to use a strong base salt
- Best : NaHCO_3 and as well as peeling de luxe plus
- It's up to you to buffer, but do not buy products containing acid + buffer in the same bottle
- Some companies sell, "buffered TCA »which is responsible of pigmentary rebounds !



Wavy lines



**COSMETIC
CREAMS
SHOULD
NOT
CONTAIN**



1. Primary or secondary alcohols (esterification)
 2. Acids with $pK_a < 3$
 3. Phenol
 4. Aldehydes without antioxidants
 5. Comedogenic agents
 6. Allergic INGREDIENTS
 7. Acne pathogens
 8. Not INCI Ingredients
 9. Animal collagen (sheep, beef, chicken, pig)
 10. Huge amounts of preservatives (parabens..)
- Grid of dots



COMEDOGENIC INGREDIENTS

in post-peel cosmetic formulations

HIGHLY COMEDOGENIC INGREDIENTS

- Lanolin / Lanolin Alcohol
- Isopropyl Palmitate
- Isopropyl Myristate
- Butyl Stearate
- Octyl Stearate
- Laureth-4 / Laureth-23
- Ethylhexyl Palmitate

Clinical note: post-peel barrier recovery increases susceptibility to comedogenic flares.

COMEDOGENIC REACTION

Post-peel cosmetic formulation



Potentially comedogenic depending on concentration

- Petrolatum
- Paraffinum liquidum
- Mineral oil
- Dimethicone (certain grades)

Always check the INCI list before using a cosmetic product after a peel.

CHOOSE PRODUCTS WITH LOWEST MOLECULAR MASS



Home care
must include

1. Anti-UVA, Anti-UVB without Alcohols (no sunscreens) -> melasma story
2. Antioxidants
3. Anti free radicals
4. Hydratant factors
5. Vitamin factors
6. Depigmentation agents (choose them)
7. Triprotic or Diprotic Acids with high pKa, which continue the peeling effect
8. Sea vegetable collagen
9. Avoid parabens

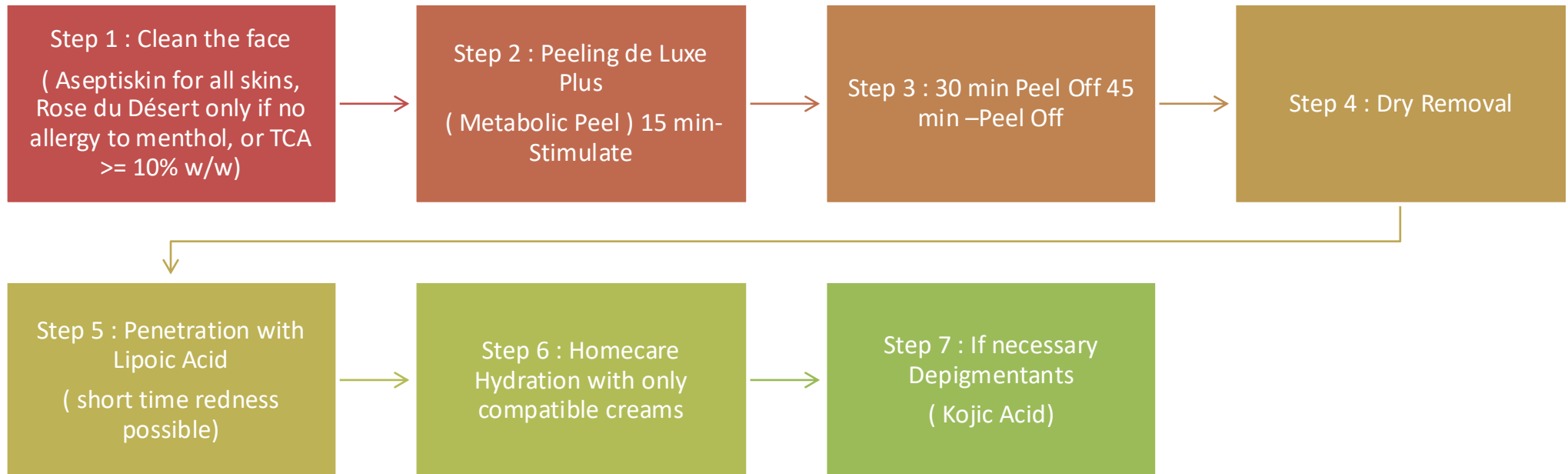




MEDICATIONS THAT CAUSE ACNE

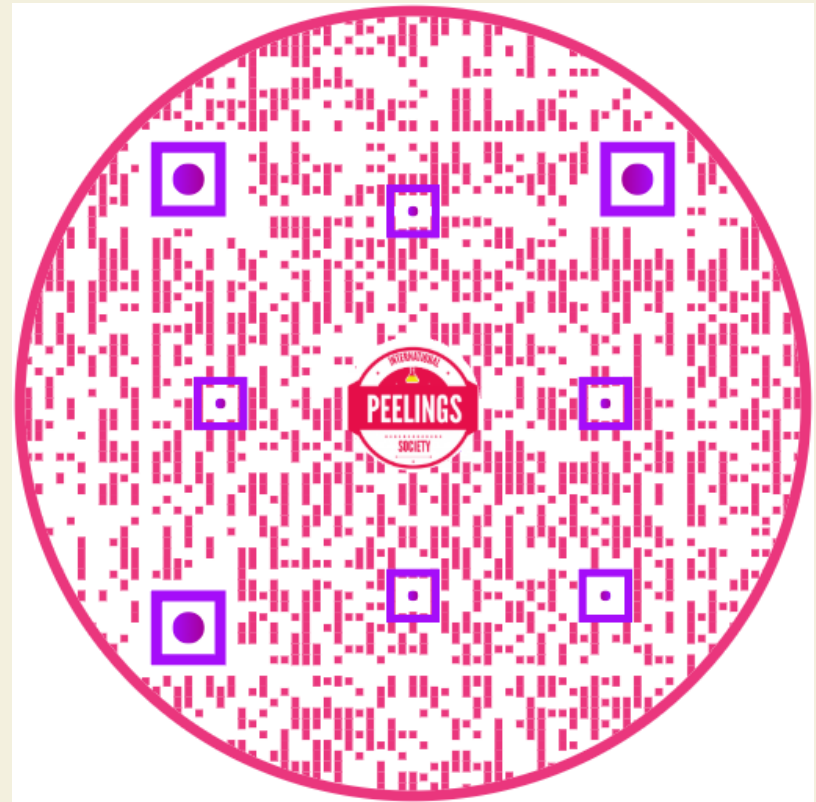
- ☐ Steroids
- ☐ ACTH hormone
- ☐ Antiepileptics:
phenytoin, phenobarbital
- ☐ Vitamins: B1, B6, B12
- ☐ Halogens: fluorine,
bromine, iodine
- ☐ Androgens: anabolic,
danazol
- ☐ Tetracyclines
- ☐ Isoniazid
- ☐ Lithium
- ☐ Quinidine
- ☐ Amineptine
- ☐ Thyroid hormone
- ☐ Ammonium salts
- ☐ Chemotherapy:
actinomycin D.
- ☐ Thiourea

TREATMENT PROTOCOL OF MAURO TIZIANI , MOLECULAR BIOLOGIST FOR MOST INDICATIONS, VALID FOR ALL SKIN TYPES IN ANY SEASONS



MAIN PROTOCOL METABOLIC PEELS

[Protocol Metabolic Peels](#)



Peeling on Black Skin
Metabolic Peels-Protocol of Mauro Tiziani

BEFORE



AFTER



Courtesy of Dr. Alain Tenenbaum

Peeling on Black Skin
Metabolic Peels-Protocol of Mauro Tiziani

BEFORE

AFTER



Courtesy of Dr. Alain Tenenbaum

Peeling on Latino (Argentina) Skin
Metabolic Peels-Protocol of Mauro Tiziani

BEFORE



AFTER



Courtesy of Dr. Alain Tenenbaum

Peeling on Latino (Argentina) Skin
Metabolic Peels-Protocol of Mauro Tiziani

BEFORE



AFTER



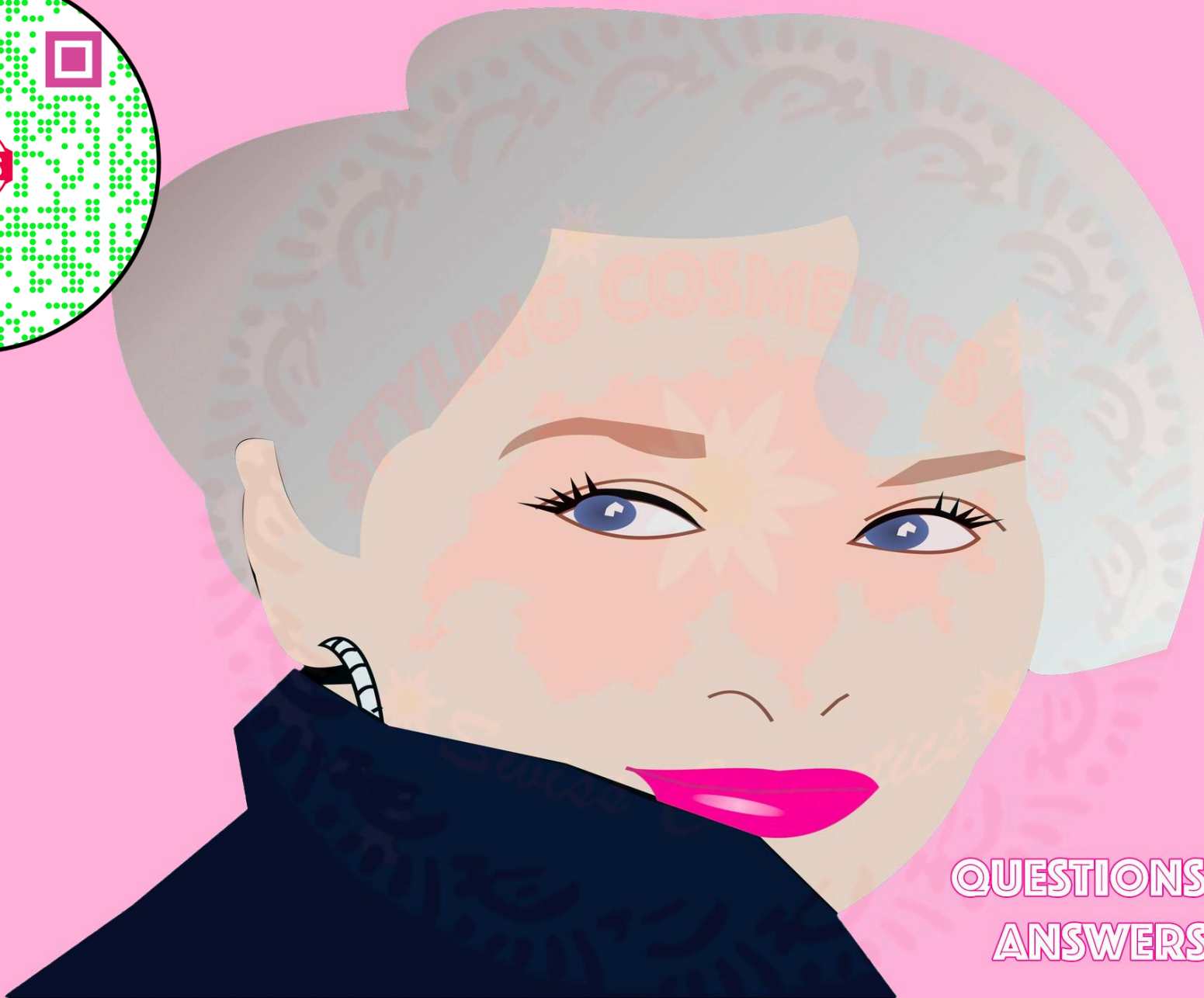
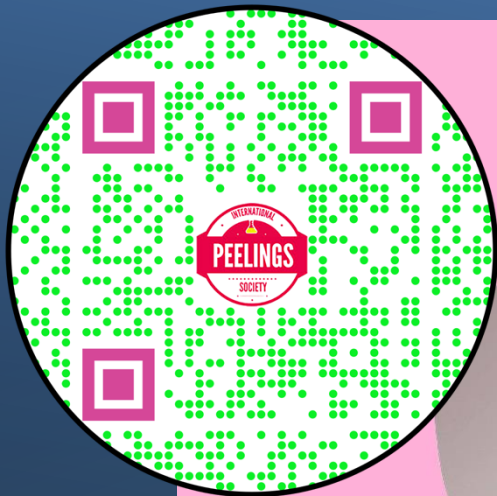
Courtesy of Dr. Alain Tenenbaum

AESTHETIC TREATMENTS

MAINLY USED BY COSMETOLOGISTS

Milchsäure (Lactic Acid)	Azelainsäure Azelaic Acid	Mandelsäure Mandelic Acid	Kojisäure (in Cremes) Kojic Acid	Phytinsäure Phytic acid	Glykolsäure Glycolic acid
<ul style="list-style-type: none"> - Keratoregulator ++ — Skin lightener - Bacteriostatics - Stimulates cell differentiation — Moisturizing 	<ul style="list-style-type: none"> - Skin lightener - Bacteriostatics - Anti-oxidant - Keratoregulator + - Moisturizer 	<ul style="list-style-type: none"> - Keratoregulator +++ - Bacteriostatics - Moisturizing 	<ul style="list-style-type: none"> -Tyrosinase-Inhibitor -Depigmentation 	<ul style="list-style-type: none"> — Melanin-Formation-Blocker - Anti-oxidant ?? 	<ul style="list-style-type: none"> - Good results (epidermal enzymes) — Dispersion of melanin of the basal layer - Keratoregulator +++ — Moisturizing

Medical supervision is helpful in reducing the risk of post-inflammatory hyperpigmentation PIH that has been observed with the use of these products.



QUESTIONS &
ANSWERS

CHOOSING THE RIGHT ACID :

LACTIC ACID VS. TCA VS. SALICYLIC ACID

Property	Lactic Acid	TCA (Trichloroacetic Acid)	Salicylic Acid
Chemical Type	AHA	Medium-depth chemical acid	BHA
Main Actions	Gentle exfoliation	Resurfacing, pigmentation, texture	Keratolytic, sebo regulating, anti-acne prone skin
Penetration Depth	Superficial	Superficial to medium	Superficial
Skin Type Suitability	Dry, sensitive, dehydrated skin	Photodamaged, aged, thick skin	Oily, acne-prone skin
Risk of PIH	Moderate to high	Moderate to high (especially in dark skin)	Moderate
Downtime	Moderate to significant	Moderate to significant	Mild to moderate
Best for	Brightening	Deep pigmentation, lines, texture	Acne Prone Skin, comedones, enlarged pores
Hydrating	No	No	No
Common Side Effects	Temporary redness, stinging	erythema	Dryness, stinging

- Lactic Acid is ideal for gentle, no-downtime treatments in sensitive or dry skin. But for my own opinion never use it.
 - TCA offers deeper results but must be used with caution, especially on darker skin types.
 - Salicylic Acid is the go-to for „ oily,, acne prone skin and oily skin management.
- Tip: Always consider skin type, medical history, and desired outcomes when choosing a peeling agent.

BRAND NAME

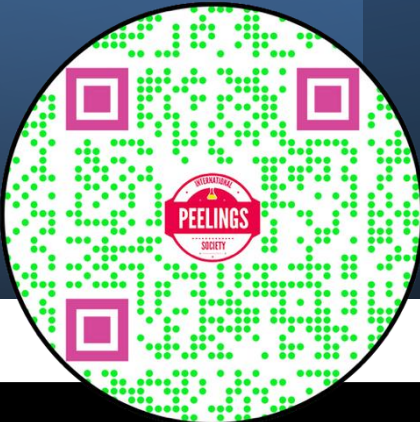
LACTIC COMPLICATION

- The issues caused by Big Pharma

Complications
of Brand Names Peels
with Big Marketing &
Poor Chemistry Knowledge
(Spain, Uruguay, Ecuador, Argentina
Korea, China, Poland, Austria...)



PEELS VS LASERS

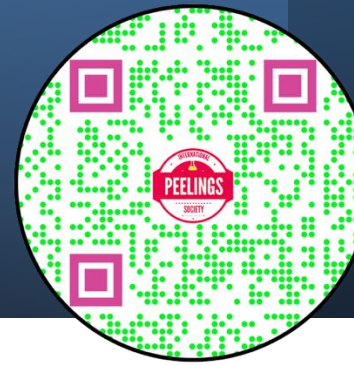


	Peelings	Lasers
Goals	Peels involve applying a chemical solution to the skin, which exfoliates the <u>outer layers</u> , promoting cell turnover and revealing fresher, smoother skin.	Laser treatments use focused light to target <u>specific layers</u> of the skin, stimulating collagen production and improving skin texture.
Advantages	<ul style="list-style-type: none">• Effective for texture issues like fine lines, acne scars, and pigmentation.• No need for expensive equipment.• Minimal risk when used correctly.• Generally less expensive than lasers.	<ul style="list-style-type: none">• Targeted treatment for specific concerns like pigmentation, vascular issues, and deeper wrinkles.• Stimulates collagen production for long-term skin improvements.• Can offer more precise results with less risk of post-inflammatory pigmentation (depending on the type of laser).
Disadvantages	<ul style="list-style-type: none">• Potential for side effects like irritation, redness, or pigmentation changes (especially with deep peels).• Downtime can vary depending on the depth of the peel.	<ul style="list-style-type: none">• More expensive than peels.• Can involve more downtime, especially with more aggressive lasers.• Potential for side effects like hyperpigmentation or scarring (especially if not done correctly).



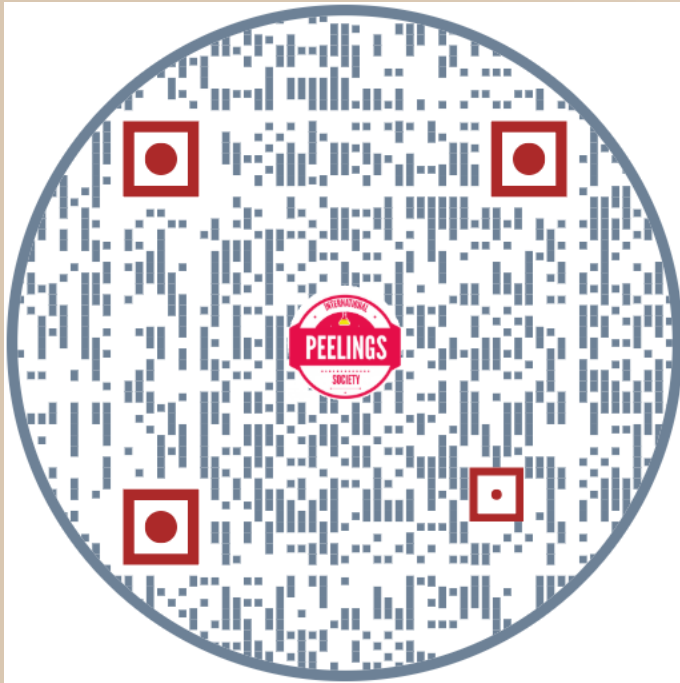
	Peelings. (M.D)	Hydrafacial (Cosmetician)
How it works	Chemical peels involve applying a chemical solution to the skin, which causes the top layers to exfoliate and peel off .	Hydrafacial is a non-invasive treatment that combines cleansing , exfoliation, extraction, hydration, and antioxidant protection. It uses a specialized device that vacuums out impurities and infuses the skin with hydrating serums.
Benefits	<ul style="list-style-type: none">• Improves skin texture and tone.• Reduces fine lines and wrinkles.• Treats acne by unclogging pores and reducing oil production.• Brightens pigmentation (e.g., melasma, age spots).• Results are typically more dramatic	<ul style="list-style-type: none">• Instant hydration and glow.• Gentle exfoliation• Improves skin texture and appearance.• No downtime• Targets a wide range of concerns, such as acne, fine lines, hyperpigmentation, and dullness.• No risk of peeling as it's more a "deep cleansing" than a full exfoliation.
Intensity	<ul style="list-style-type: none">• more intense• longer-lasting results	<ul style="list-style-type: none">• Gentler
Downtime	<ul style="list-style-type: none">• Peeling• redness post-treatment.	<ul style="list-style-type: none">• No downtime
Customization	<ul style="list-style-type: none">• higher degree of customization based on skin type and concerns	<ul style="list-style-type: none">• Customization , especially in terms of serums used, but are more focused on hydration and exfoliation (gommage)
Maintenance		<ul style="list-style-type: none">• hydration, quick glow, maintain a perfect hydrafacial

Peels vs Microneedling



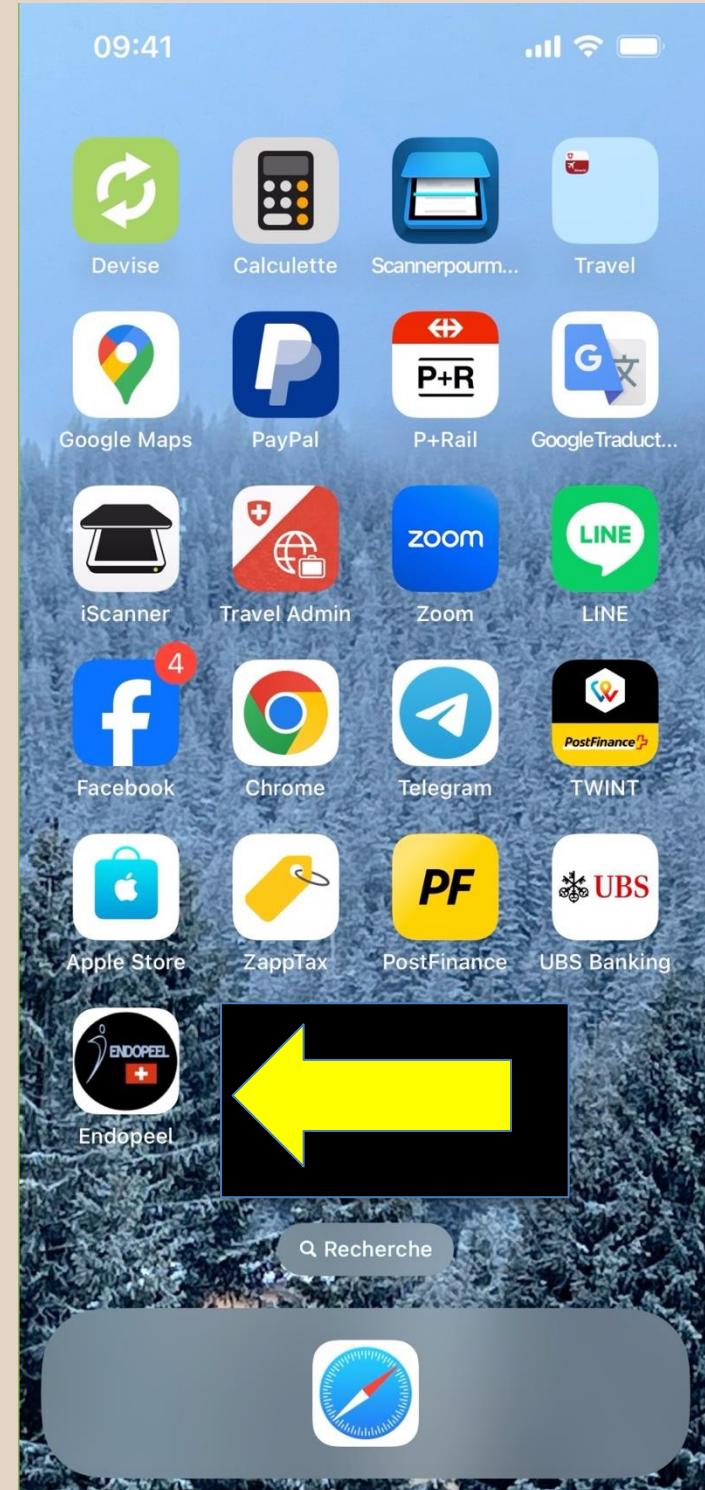
	Peelings	Microneedling
Mechanisms of Action	<ul style="list-style-type: none">• <u>Chemical</u> Exfoliation• Skin Regeneration	<ul style="list-style-type: none">• <u>Physical</u> Stimulation to get Collagen Production through• Micro Injuries
Treatment Depth	<ul style="list-style-type: none">• All layers but superficial layers are mostly diffused	<ul style="list-style-type: none">• Deeper Layers
Downtime	<ul style="list-style-type: none">• Desquamation• Redness for deep ones	<ul style="list-style-type: none">• <u>No desquamation</u>• Redness• <u>Swelling</u>
Indications	<ul style="list-style-type: none">• Pigmentation Issues• Fine Lines• Texture• <u>Immediate</u> Results	<ul style="list-style-type: none">• Collagen Boosting• Tackling Scars• Advanced Aging Concerns
Recommandations	<ul style="list-style-type: none">• Use Lipoic Acid to allow a better penetration	<ul style="list-style-type: none">• Can be done after a light peeling to enhance results• To get deeper use bestens Lipoic Acid

emergency-skin-rash-kit

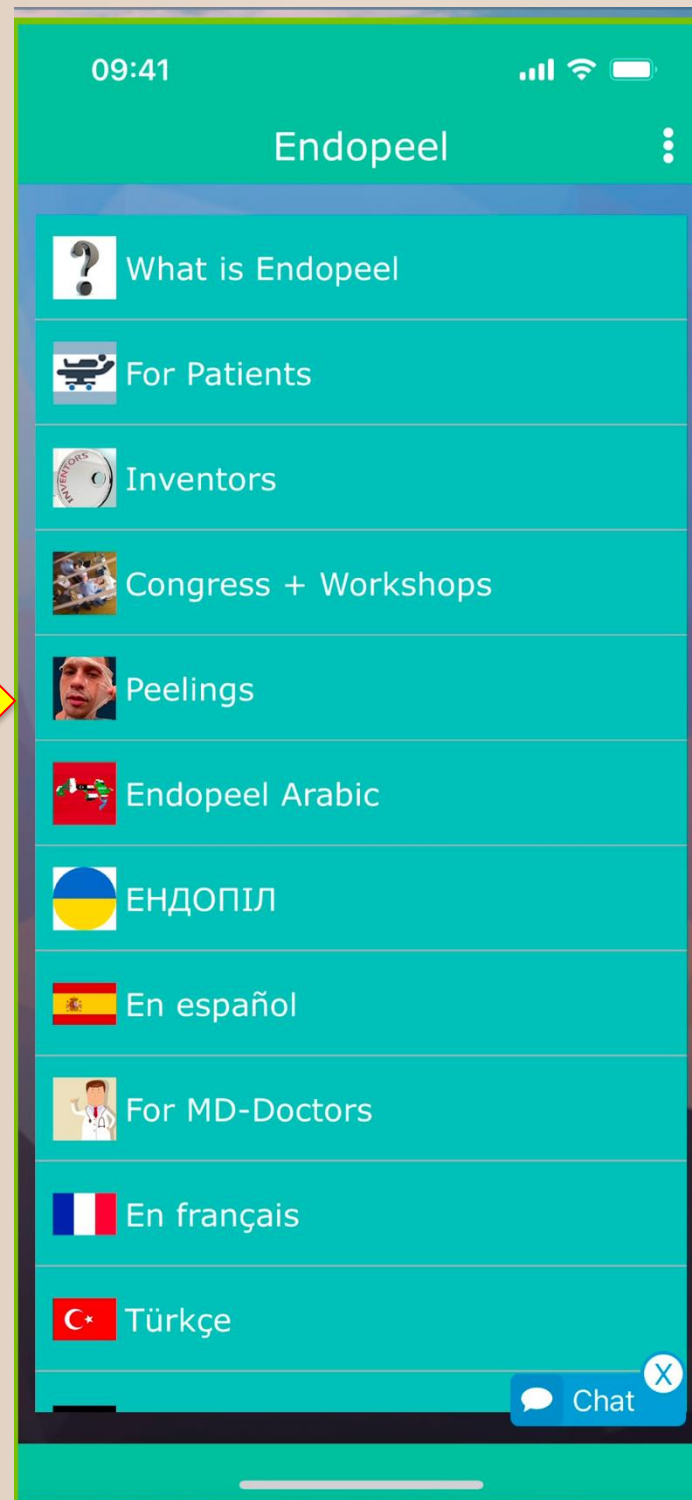


APP „ ENDOPEEL,,
VALID TILL SUMMER 2026

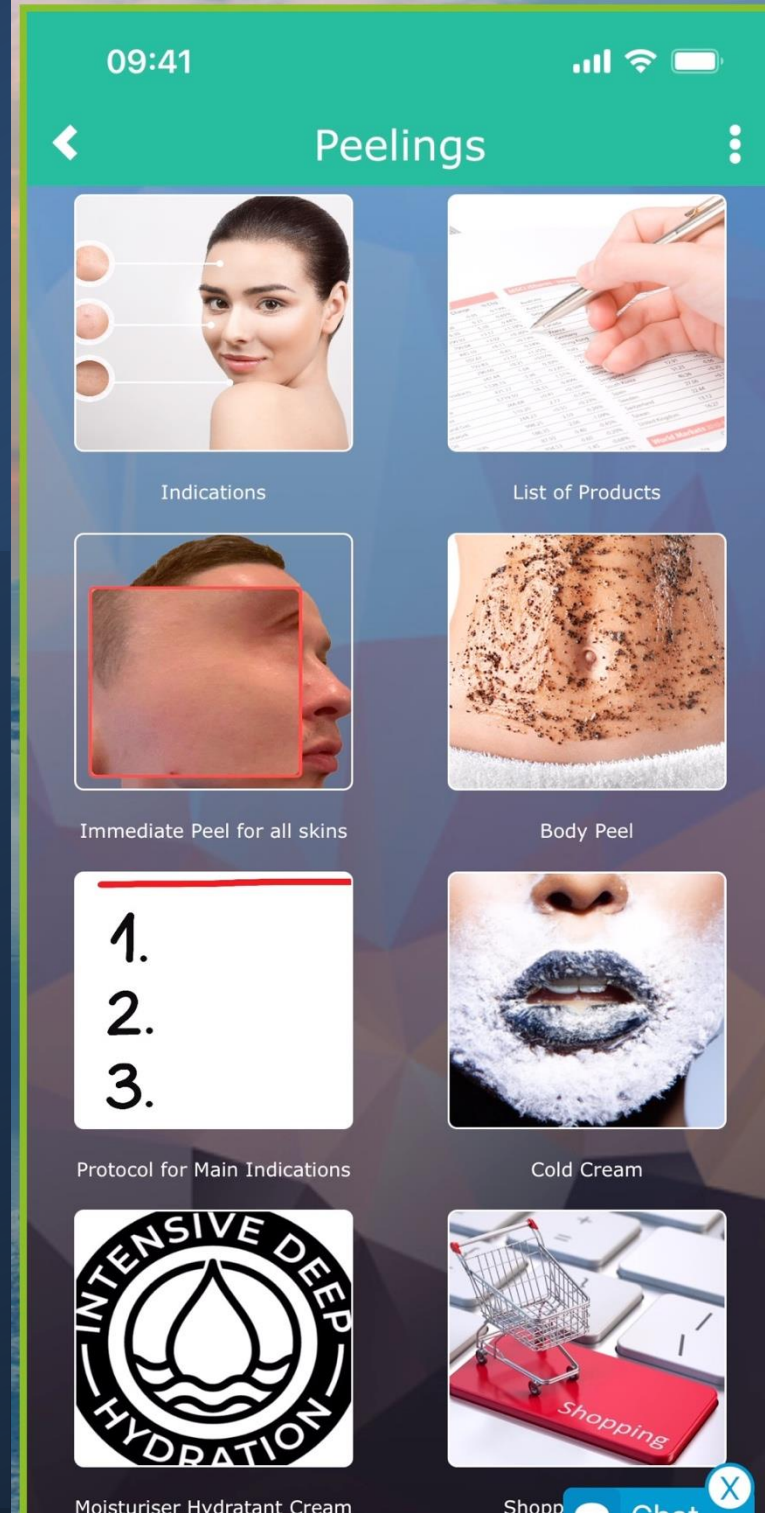
ON APPLESTORE
& GOOGLEPLAY



APP „ENDOPEEL„
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App,,Endopeel,,
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AesthetiMedica

SOON BEFORE
SUMMER 2026
NEW APP :

AESTHETIMEDICA



Home



For MD-Doctors



For Patients



shop



Education



Ecosystem

BRING HOME THIS MESSAGE

Peelings should be done the day of sutures removals,
even on the scars

If peelings are done without previous surgery, **OUR**
peelings can be done by your MPA

Do not use „ cocktail peelings,, made in countries
without knowledge of chemistry

Do not use phenol peelings which can be lethal, with
stricted limited indications, which need OP room with
anaesthesist and which cant be controlled as surgery



GET QUICKLY TANNED WITHOUT BURNS

How to get
quickly and
deep tanned
without burns



Next Workshops in Zürich

WORKSHOP
FULL DAY - FOUNDATIONS + TARGETED AREAS
ENDOPEEL FACE & BODY – COMPREHENSIVE PROGRAM
ZÜRICH-SWITZERLAND : MARCH 21 ST, 9 AM-5 PM

WITH
MAURO TIZIANI
DR. ALAIN TENENBAUM

SCAN ME

<https://aesthetic.events/workshops/next-workshops>

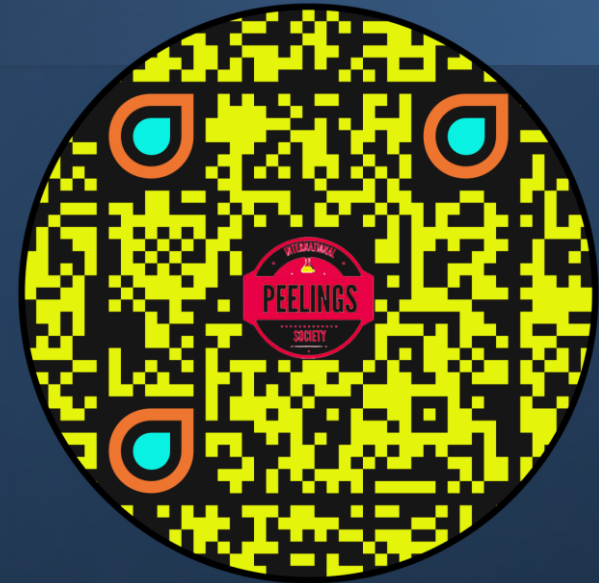
PRACTICAL METABOLIC PEELS WORKSHOP
HANDS ON ONLY

ZÜRICH-SWITZERLAND :
MARCH 20 TH, 2026 : 1.30 PM-5 PM

WITH
MAURO TIZIANI
DR. ALAIN TENENBAUM

SCAN ME

<https://aesthetic.events/workshops/next-workshops>



DURING
COFFEE BREAK
I CAN HELP YOU
FOR



Become a
Member

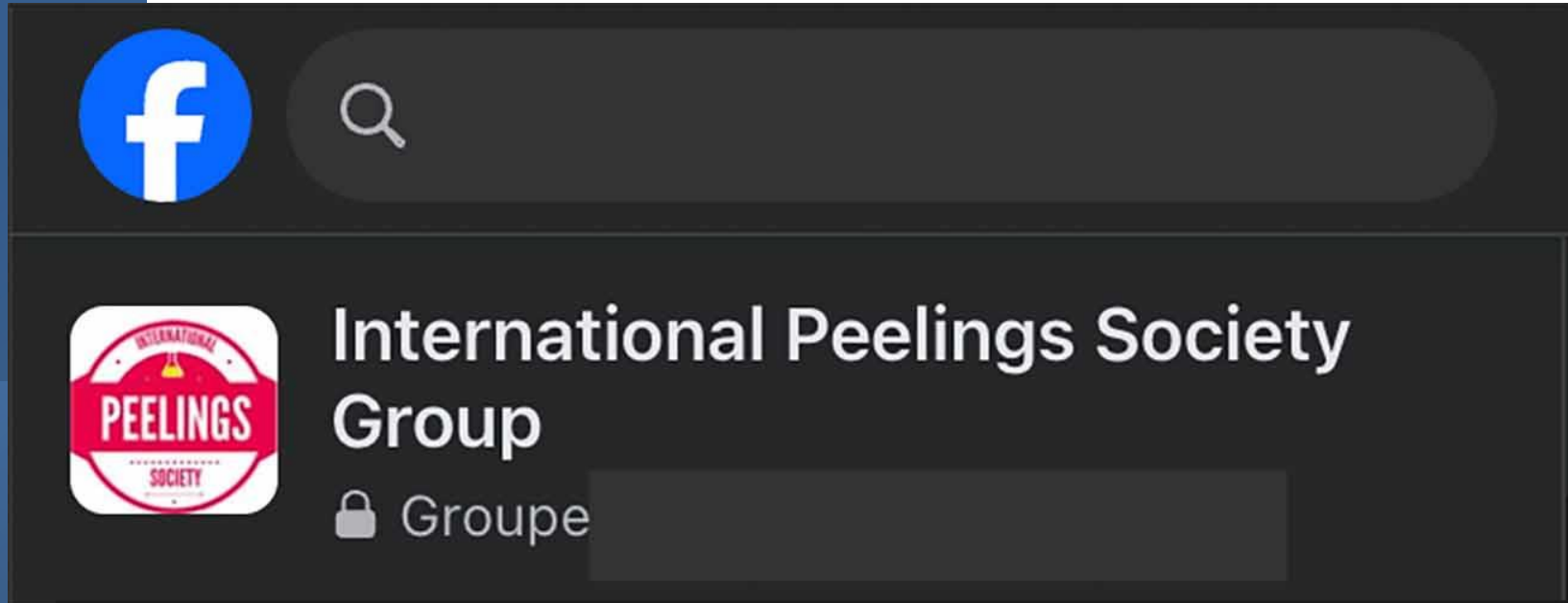
MEMBERSHIP



DURING THE COFFEE BREAK



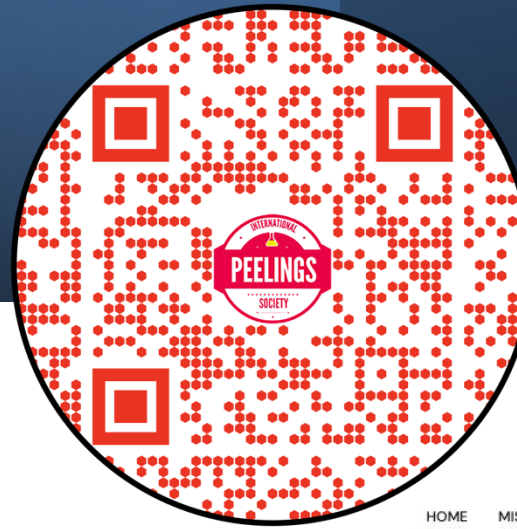
FACEBOOK GROUP



[ask to be inside this group](#)

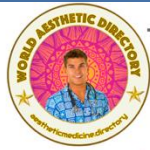


[link to website](#)



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DR ALAIN TENENBAUM

41813

endopeel is the best procedure in aesthetic medicine For plastic surgeons, gynecologists, cosmetic dermatologists and aesthetic medicine practitioners.



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Quote

Request



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CONTACT PERSON

RESEARCH &
DEVELOPMENT

ALAIN
TENENBAUM

DETAILS

CUSTOM GALLERY

MEMBERSHIPS

OVERVIEW

Endopeel Board Certified Medical Doctors

SACDAM Members

IPSC Members (International Peelings Society)

Aesthetic Medicine Specialists

endopeel

retensadcutaneo

pectoroplasty

slipacks

medical facelift

medical rhinoplasty

abdominopexy

trapezoplasty

biceps projection

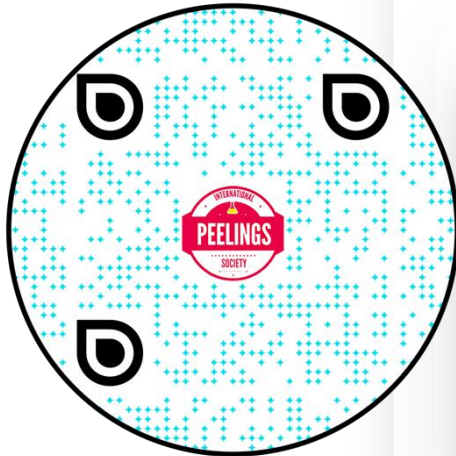
arm lift

<https://aestheticmedicine.directory/your-name>



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sixpacks

medical facelift

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trapezoplasty

biceps projection

CONCLUSION



*CHEMISTRY PROVIDES
TOOLS.*



*BIOLOGY DEFINES
OUTCOMES.*



*CLINICAL RESPONSIBILITY IS
KNOWING THE DIFFERENCE.*



NEXT LECTURE



HYPERPIGMENTATION

Facial Hyperchromy



Introduction to Hyperpigmentation

Hyperpigmentation refers to the darkening of the skin due to an increase in melanin production or deposition. It can occur in localized areas or over larger areas of the body. This condition is commonly seen in clinical practice and can result from a variety of intrinsic and extrinsic factors.

Definition

- **Hyperpigmentation** is the condition in which certain areas of the skin become darker than the surrounding skin due to an excess production of melanin. It can be either focal (localized) or diffuse (spread over a larger area) and is often a cosmetic concern for patients.
Hyperpigmentation is not a single disorder but rather a clinical manifestation of various underlying pathophysiological processes.

Specific types of hyperpigmentation

- *melasma*
- *post-inflammatory hyperpigmentation*
- *Lentigines*

CONCEPTS OF A.TENENBAUM & M.TIZIANI

Targeting the pigment directly
(ColorKiller – non-cytotoxic metabolic approach)



- Acts on **abnormal melanin accumulation**, not on normal melanocytes
- Targets **pathologic pigment retention** within the eidermis
- Does **not inhibit physiological melanogenesis**



By inducing controlled protein coagulation affecting abnormal pigment retention

Complete frosting is required on the hyperpigmented lesion itself;
peri-lesional areas must be managed separately to prevent demarcation lines.

Addressing the 3 key compartments of melanin metabolism

- **Production**: *normalization , not suppression* (Modifying the melanogenesis process to reduce melanin synthesis.)
- **Transport**: *limiting abnormal melanosome transfer* (Interfering with the movement of melanin within the skin to prevent uneven distribution.)
- **Destruction**: *Control removal of excess pigment* (Enhancing the breakdown and removal of excess melanin from the skin.)

Colorkiller is a metabolic regulator, not a chemical depigmenting agent

FROSTINGS

*Targeting the
pigment
(ColorKiller)*



ROSE FROSTING



Superficial protein
coagulation
(early epidermal response)

GREY FROSTING



Intermediate coagulation
with pigment interaction
(active ColorKiller phase)

WHITE FROSTING



Dense protein coagulation
*(maximum pigment coverage
on lesion)*



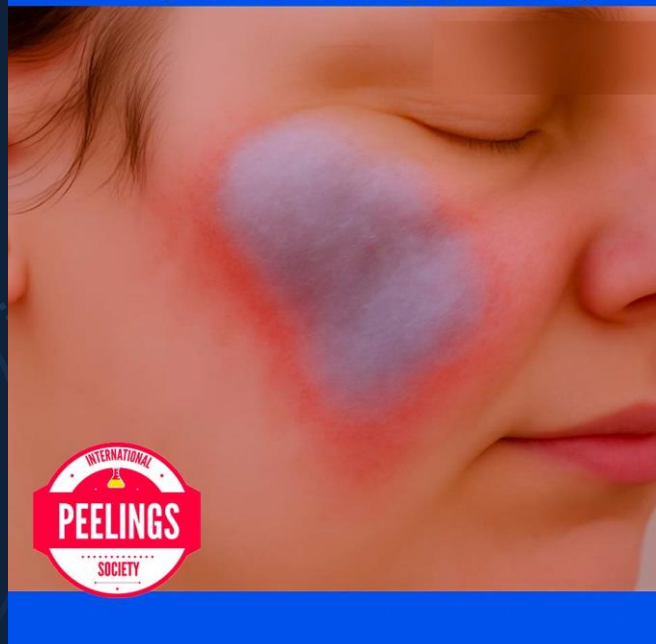
FROSTINGS & TCA

*Targeting the
pigment
(ColorKiller)*

ROSE FROSTING



GREY FROSTING



WHITE FROSTING



Frosting intensity reflects protein coagulation patterns,
not the depth of tissue injury nor the TCA concentration alone.



Targeting the pigment (ColorKiller)

The frosting has to completely cover the hyperpigmented areas



Incomplete frosting may lead to insufficient correction or post-inflammatory hyperpigmentation

Residual untreated pigment areas can act as triggers for rebound pigmentation.

FROSTING DOESN'T COMPLETELY COVER THE HYPERPIGMENTED AREAS ON THIS ASIAN MALE PATIENT



Courtesy of Dr. Alain Tenenbaum



White Frosting
appearing
over existing
Crusts
at 2nd or 3rd
Session

The goal is pigment control, not
mechanical crust removal.

FROSTING APPEARING OVER EXISTING CRUSTS



Courtesy of Dr. Alain Tenenbaum

POST CHEMO-DERMABRASION DEMARCATIION LINE

Complication
Observed After
Chemodermabrasion
Performed by a
Dermatologist

Demarcation lines result from
abrupt transitions between
treated and untreated skin.

They reflect insufficient peri-
lesional management rather than
excessive treatment.”



Courtesy of Dr. Alain Tenenbaum

Targeting the pigment (ColorKiller)

Degressive Concentrations
(from higher to lower)
around the lesion are the
Best Option to avoid
Demarcation Lines.

Don't forget to treat
WHOLE FACE even for a
small loco-regional lesion !
(A.TENENBAUM)

The TCA Mosaic Approach allows
selective pigment correction while
preserving progressive transitions
between treated and untreated
skin.

THE TCA MOSAIC APPROACH FOR TREATING A DEMARCATION LINE



Courtesy of Dr. Alain Tenenbaum

CORRECTION-STABILIZATION-MAINTENANCE

THE TCA MOSAIC APPROACH FOR TREATING A DEMARCATION LINE

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Courtesy of Dr. Alain Tenenbaum

Targeting the pigment (ColorKiller)

Peri-lesional frosting must be recognized and controlled, not ignored.

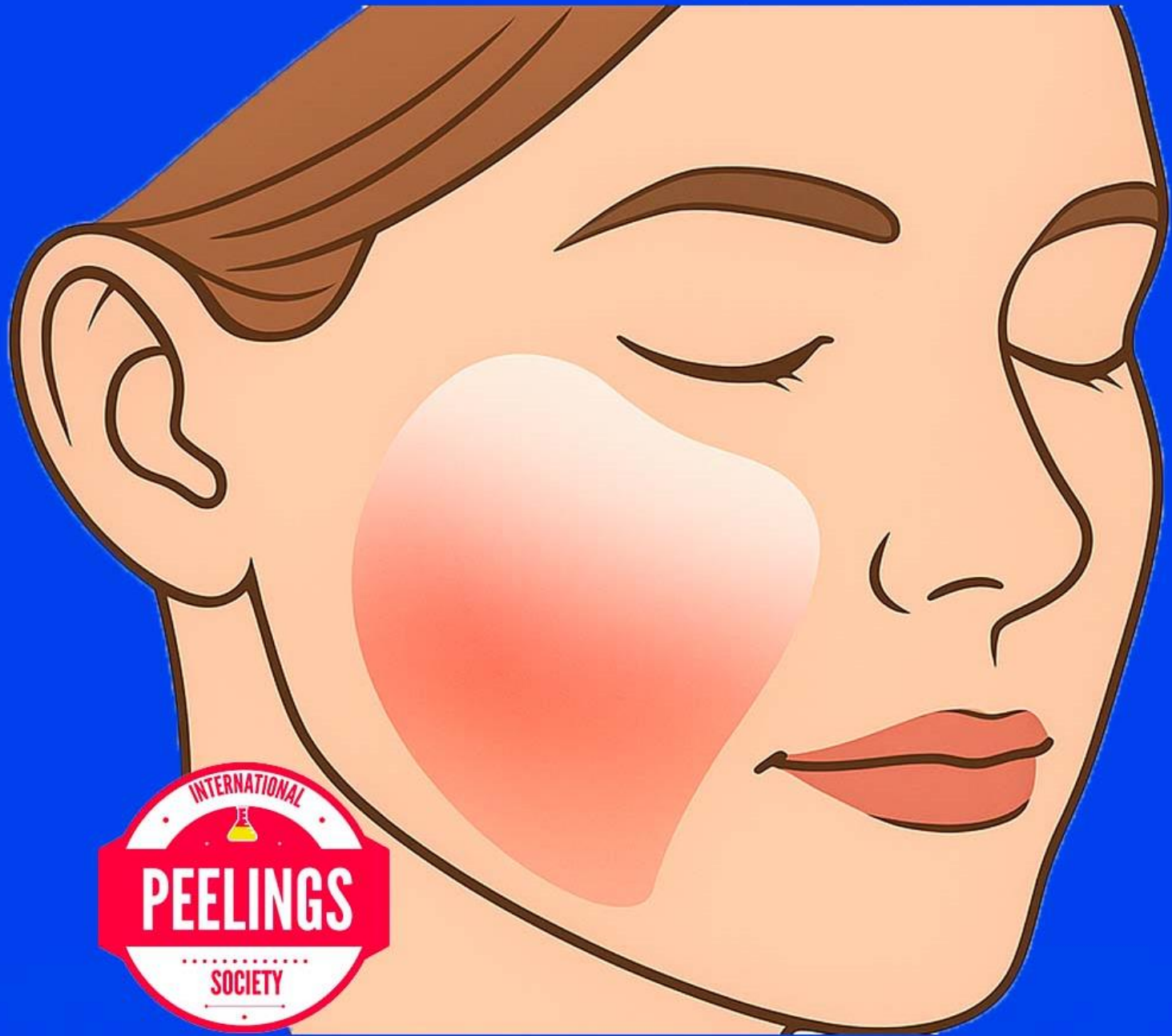
**THE WHITE FROSTING EXTENDS BEYOND THE LESION
INTO THE PERILESIONAL AREA**



Courtesy of Dr. Alain Tenenbaum

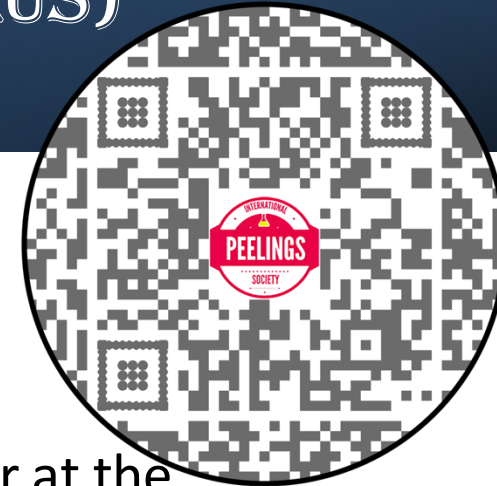
DEFROSTING FOLLOWING WHITE FROSTING

*Targeting
the pigment
(ColorKiller)*



Targeting the pigment (ColorKiller)

WHEN TO APPLY THE FROSTING STOPPER (PEELING DE LUXE PLUS)



Condition

White frosting appears on the lesion

Frosting turns from grey to white on the perilesion (external limits of the lesion)

Frosting turns from red to grey on the perilesion (near the lesion but not in direct contact)

Action

Apply the frosting stopper at the moment of defrosting following the white frosting on the lesion.

Apply the frosting stopper just juxtaposing the external limits of the lesion.

Apply the frosting stopper not directly juxtaposing the lesion, but at the moment of defrosting.

*Targeting
the pigment
(ColorKiller)*

When to
apply the
frosting
stopper
(Peeling de
Luxe Plus)

Timing determines outcome more
than concentration

WHEN TO USE THE FROSTING STOPPER (PEELING DE LUXE PLUS)



WAIT THE DEFROSTING AFTER
COMPLETE WHITE FROSTING



Courtesy of Dr. Alain Tenenbaum

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By inducing controlled protein coagulation affecting abnormal pigment retention

Complete frosting is required on the hyperpigmented lesion itself;
peri-lesional areas must be managed separately to prevent demarcation lines.




Addressing the 3 key compartments of melanin metabolism

- **Production**: *normalization , not suppression* (Modifying the melanogenesis process to reduce melanin synthesis.)
- **Transport**: *limiting abnormal melanosome transfer* (Interfering with the movement of melanin within the skin to prevent uneven distribution.)
- **Destruction**: *Control removal of excess pigment* (Enhancing the breakdown and removal of excess melanin from the skin.)

Colorkiller is a metabolic regulator, not a chemical depigmenting agent

HYPERPIGMENTATION REQUIRES COMPARTMENT-SPECIFIC TARGETING, NOT UNIFORM TREATMENT.

COMPARTMENT		GOAL/AIM	CATEGORY OF TREATMENT	INGREDIENTS	PRODUCT
1	Production	Reduce the production of melanin	Prevention + Maintenance Treatments	<ul style="list-style-type: none">TretinoinTranexamic Acid	Peeling de Luxe Plus
2	Circulation	Lower the melanin transfer through melanosomes	Maintenance Treatment	<ul style="list-style-type: none">Topical Melatonin Regulates Melanin (see below)	Clarté de Lune
3	Destruction	Treat the hyperchromy or hyperchromies	Attack Treatment	<ul style="list-style-type: none">Azelaic AcidArbutinKojic AcidVit C	StretchPeel



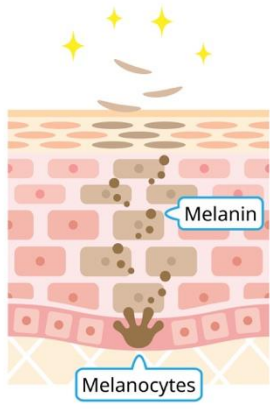
Addressing the 3 Key Compartments of Melanin Metabolism



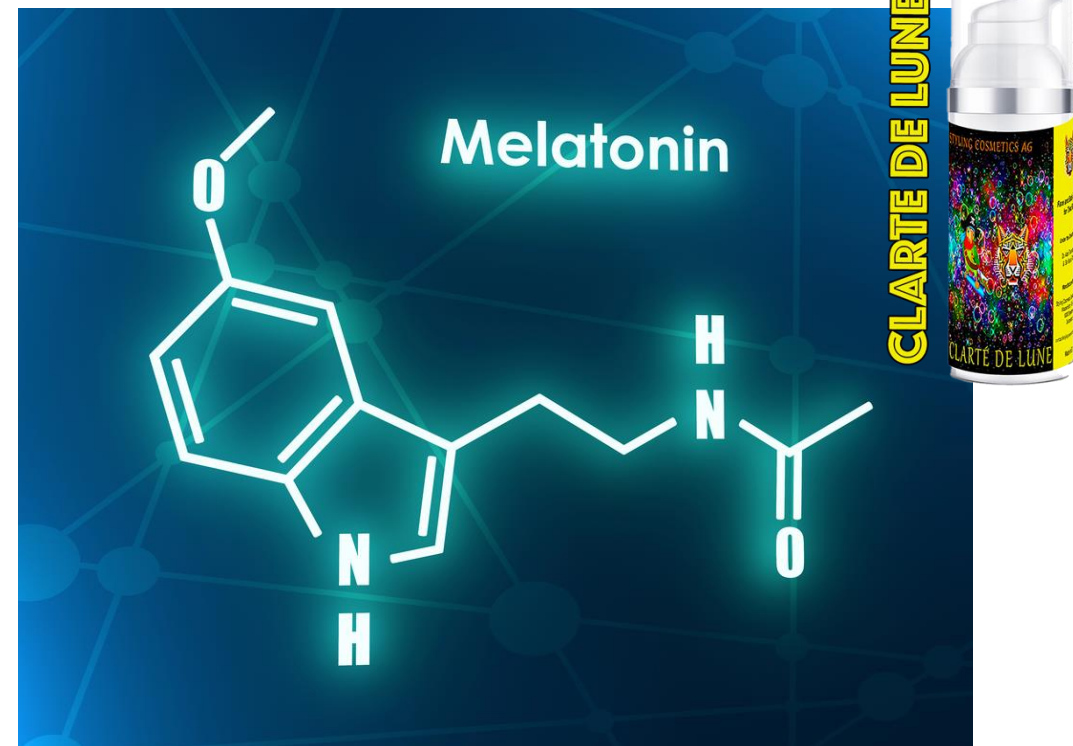
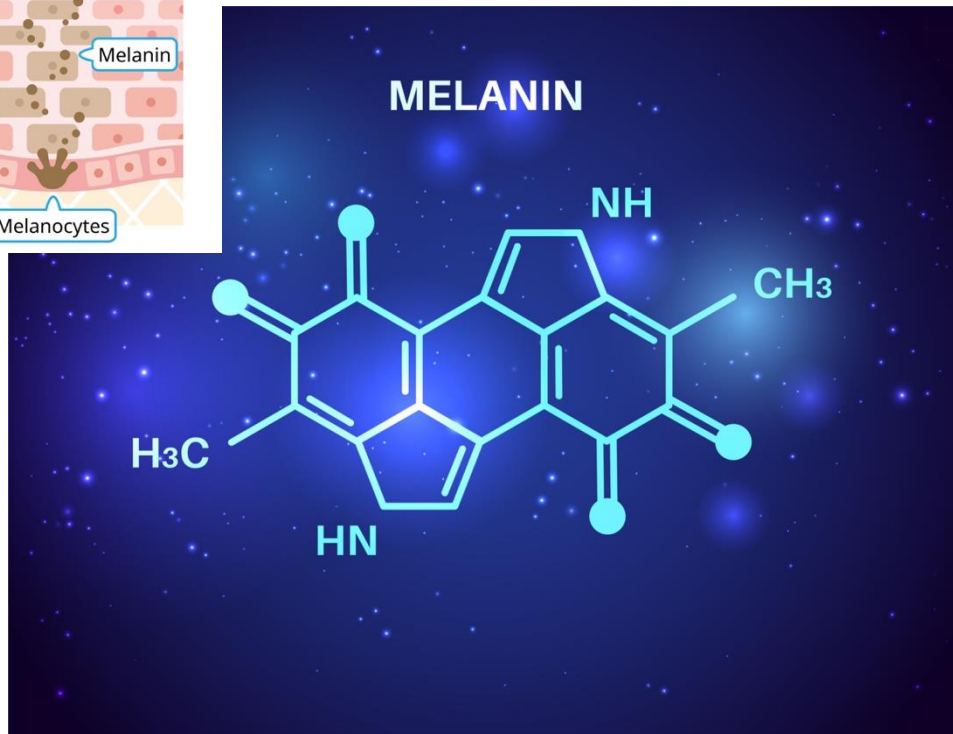
PROTOCOL HYPERPIGMENTATION

Protocol
Hyperpigmentation

No Confusion



Addressing the 3 key compartments of
melanin metabolism



Treatment of Cervical Hyperchromy
Post Chemodermabrasion
TCA + Metabolic Peels + Depigmentants
Protocol of A.Tenenbaum

BEFORE



AFTER 4 SESSIONS



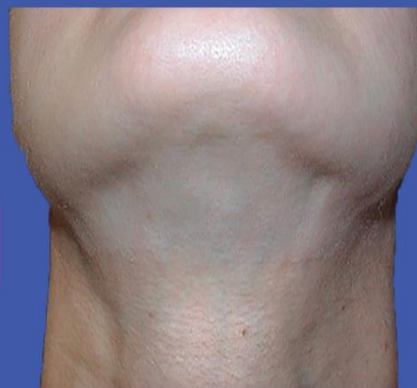
Courtesy of Dr. Alain Tenenbaum

Treatment of Cervical Hyperchromy
Post Chemodermoabrasion
TCA + Metabolic Peels + Depigmentants
Protocol of A.Tenenbaum : 4 sessions-1 session/week

D0



D365 stable



D60



Courtesy of Dr.Alain Tenenbaum

Treatment of Cervical Hyperchromy
Post Chemodermoabrasion
TCA + Metabolic Peels + Depigmentants
Protocol of A.Tenenbaum : 4 sessions-1 session/week

D0



D365 stable



D60



Courtesy of Dr.Alain Tenenbaum

Treatment of Facial Hyperchromy
dued to a peelings complication (TCA + Glycolic Acid)
Fixed with TCA + Metabolic Peels + Depigmentants

BEFORE

3 weeks after TCA + Glycolic Acid



AFTER 4 MONTHS



Protocol Hyperchromy of Dr. Alain Tenenbaum

Treatment of Facial Hyperchromy
due to a peelings complication (TCA + Glycolic Acid)
Fixed with TCA + Metabolic Peels + Depigmentants

LIFE
GOES
ON

Never give up

before

1 month

2 months

3 months

4 months



Protocol Hyperchromy of Dr. Alain Tenenbaum



2.ADDRESSING THE 3 KEY COMPARTMENTS OF MELANIN METABOLISM

MECHANISM OF ACTION OF TOPICAL MELATONIN IN SKIN PIGMENTATION REGULATION

Key Ingredients

- Melatonin: neurohormone with cutaneous activity
- Tocopherol (Vitamin E): antioxidant synergy
- Glutamic Acid & Citric Acid: support skin metabolism and pH balance

Mechanisms of Action

Mechanism	Description
Antioxidant Activity	Neutralizes free radicals and reduces oxidative stress that stimulates melangensis
Tyrosinase Downregulation	Inhibits tyrosinase enzyme, reducing melanin synthesis
Melatonin Receptor Antivation	Binds MT1/MT2 receptors o melanocytes, modulating melanogenic signaling pathways
Inhibition of Melanin Transfer	Modulates keratinocyte-melanocyte communication, decreasing melanosome transfer
Gene Expression Regulation	Influences MITF and related genes controlling melanin production
Anti-inflammatory Effects	Reduces post-inflammatory hyperpigmentation through skin calming and repair

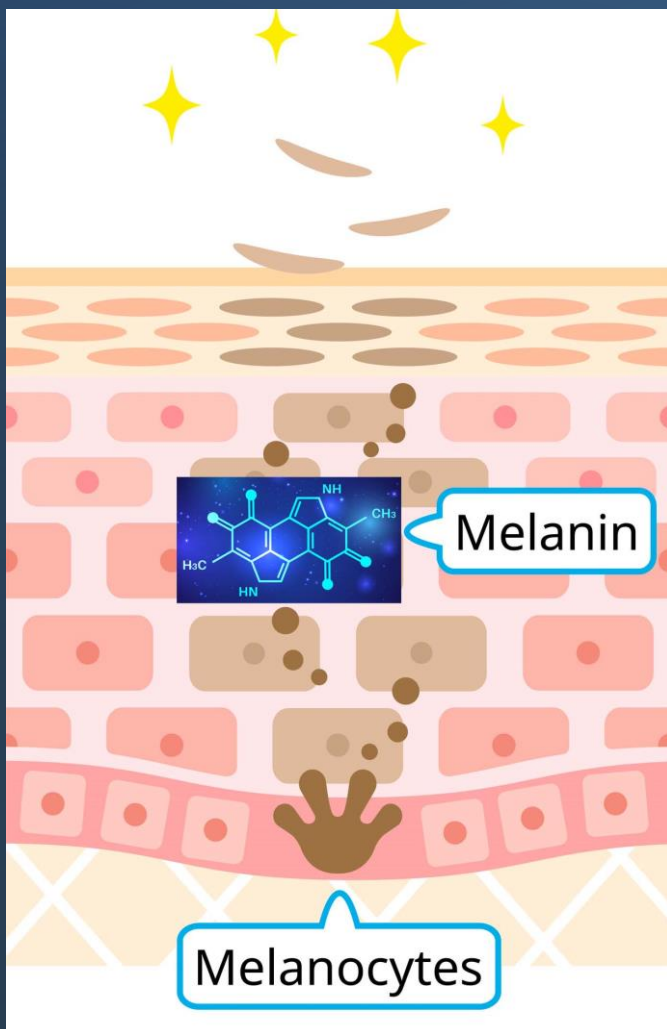
Clinical Effects

- Reduction of hyperchromia and dark spots
- Improved skin tone uniformity
- Enhanced skin protection against UV-induced pigmentation

Application Tips

- Can be combined with metabolic peels for synergistic depigmentation

CLARTE DE LUNE



DEFINITION OF MELASMA

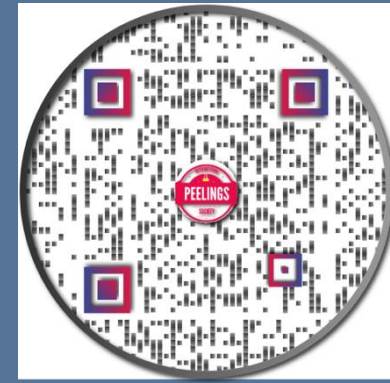
Melasma on Asian Female



- Melasma is a common skin condition characterized by the development of brown or grayish-brown patches, typically on the face.
- It is often found on areas that are exposed to the sun, such as the
 - -cheeks
 - -forehead
 - -nose and
 - -upper lip.
- The condition is more common in **women**, especially during **pregnancy** or when using **birth control pills**, due to hormonal changes.
- It can also be triggered or worsened by **sun exposure**, certain **medications**, or skin irritation.
- The patches are usually symmetrical, and melasma is typically more noticeable in individuals with darker skin tones.

MELASMA

Histo-Pathophysiology of Melasma



Melanocyte hyperactivity:

Melasma is not due to an increased number of melanocytes, but rather their increased **functional activity**, leading to excess melanin production.

Epidermal melasma: Melanin is predominantly found in the basal and suprabasal layers.

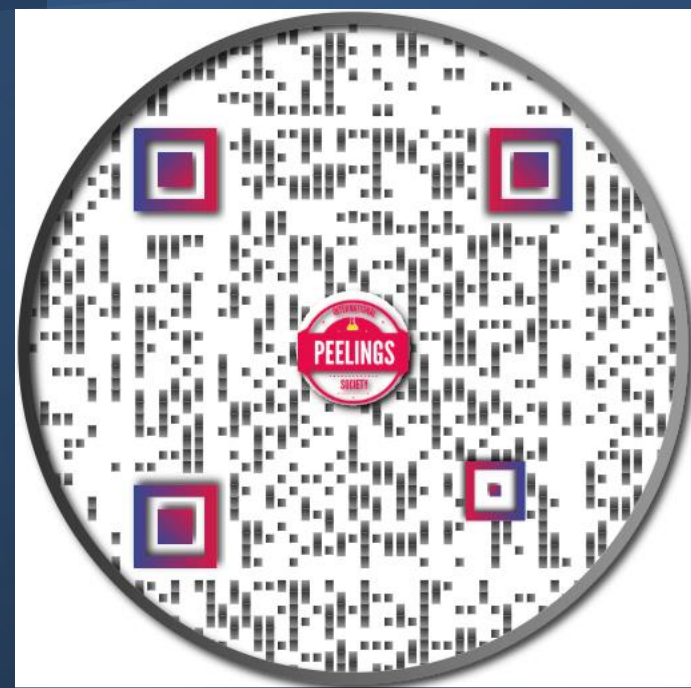
Dermal melasma: Melanin is present in melanophages in the dermis, often due to melanin leakage (pigmentary incontinence).

Normal skin

Epidermal melasma

Dermal melasma

MELASMA HAS BECOME
MORE PREVALENT
IN RECENT YEARS

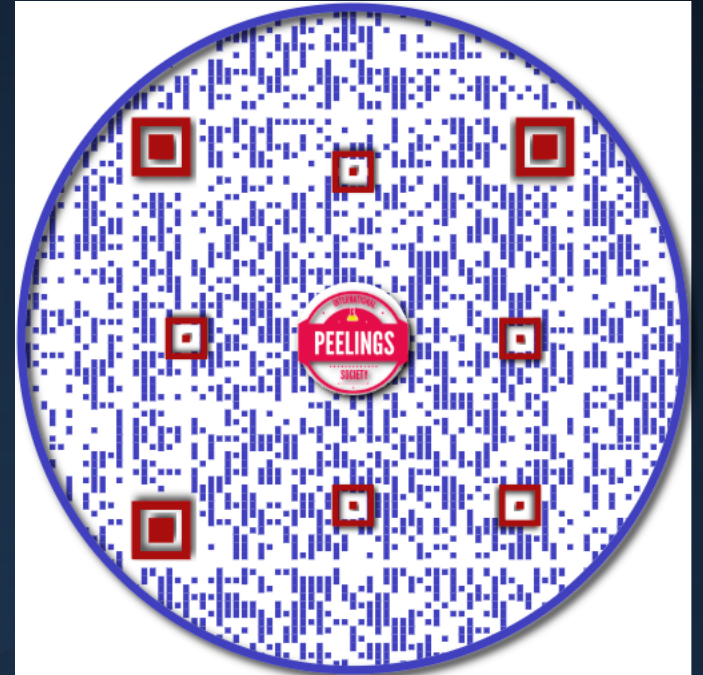


Melasma

MELASMA

Treatment must
be aggressive
Any "soft"
treatment will
only worsen the
results

PROTOCOL MELASMA



Protocol & Treatment Melasma

CLINICAL PITFALLS

CLINICAL PITFALL IN MELASMA INCORRECT VS CORRECT ENDPOINT

7 DAYS AFTER TCA
INCORRECT ENDPOINT



Insufficient frosting
Color-killer not achieved

7 DAYS AFTER TCA
CORRECT ENDPOINT



Complete color-killer coverage
Melasma adequately treated

Correction of Insufficient Treatment of infraocular Melasma with TCA 30% w/w + Metabolic Peels + Depigmentants

7 DAYS
AFTER TCA 18% w/w
UNCORRECT



Korean Female 29Y



7 DAYS
AFTER TCA 30% w/w
CORRECT



Courtesy of Dr. Alain Tenenbaum

D0



D21+3



D8+3



D15+3



**EXPECTED DOWNTIME
DURING TCA +
METABOLIC PROTOCOL**

**APPEARANCE MAY WORSEN
AFTER EARLY SESSIONS —
CONTINUATION IS ESSENTIAL**

[Patient Leaflet to avoid Panic](#)

Patient Leaflet to Avoid Panic



A close-up photograph of a person's hands. One hand is holding a clear, cylindrical tube of sunscreen, and the other hand is open, palm up, with a small amount of white sunscreen being applied to the center. The background is blurred, showing a person wearing a blue shirt. The overall lighting is soft and natural.

Which Sunscreen without Alcohol ?

Melasma Maintenance

STRETCHPEEL AS SUNPROTECTOR

2.Addressing the 3 Key Compartments of Melanin Metabolism

Stretchpeel is helping to make heller hyperchromies and is acting as well as sunprotector without alcohol.

Stretchpeel is *completely free of chemical filters like oxybenzone,*

avobenzone, and octinoxate.

These chemical filters can sometimes lead to hormonal disruptions or photosensitivity in some people, possibly contributing to melasma

Stretchpeel cream is a powerful Sunprotector Depigmenting agent and multivitamin cream.

STRETCHPEEL



Melasma Maintenance

COMPLEX ACNE SCARS PROTOCOLS





COMPLEX ACNE SCARS PROTOCOLS



TCA 30% w/w on Asian Skin for Deep Acne Scars
1 Session Only
Protocol : TCA + Metabolic Peel + Depigmentant + Moisturizer

D0

D7



Courtesy of Dr. Alain Tenenbaum

TCA 30% w/w on Asian Skin for Deep Acne Scars
1 Session Only
Protocol : TCA + Metabolic Peel + Depigmentant + Moisturizer



D0

D7



D1

D3

D4



Courtesy of Dr. Alain Tenenbaum

COMBINATION EBD+
PEELS
„PEELING SUSHI,, OF
A. TENENBAUM

Combination EBD (Erb Yag) & TCA 30% w/w
No waterphotodetersion ! Only with TCA

D0



1 ST SESSION

Korean Male



D7



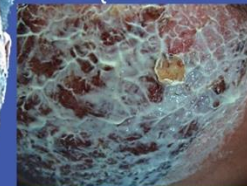
D1



D4



Peeling Sushi or
Laser Spot
Desquamation



Courtesy of Dr. Alain Tenenbaum

Combination EBD (Erb Yag) & TCA 30% w/w
No waterphotodetersion ! Only with TCA

2nd SESSION

No need to go
on with a 2nd
session.

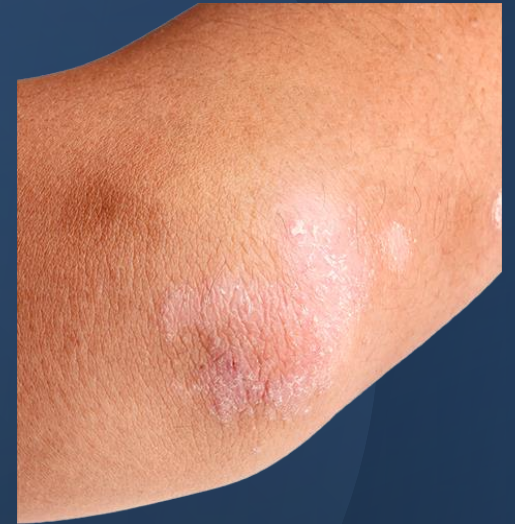
It s better to
go on with
metabolic peels
without TCA



More sessions ≠ better results
if the strategy is wrong.

Courtesy of Dr.Alain Tenenbaum

INDICATIONS FOR *KERATOTIC ZONES* USE OF SALICYLIC ACID IN ETHANOL



<https://chemicalpeeling.com/products-list/superficial-peels/salicylic-acid>



IS SALICYLIC ACID A COLOR KILLER?

2.Targeting the pigment (ColorKiller)

- Frosting with salicylic acid is purely physical, not biochemical
- Therefore, the **colorkiller** effect is more about **optical masking** or synergistic combination with agents like TCA or lactic acid for pigmentation correction.
- Unlike TCA, salicylic acid does not produce true frosting through protein coagulation.
- Instead, it creates a **pseudofrosting** due to **crystallization** of the acid on the skin as the solvent evaporates—often mistaken for "white frost."
- That said, we can still classify **clinical effects and visual appearances** by concentration

COMPARISON TABLE : TCA VS. SALICYLIC ACID FOR
HYPERPIGMENTATION USED AS ,, COLOR KILLERS,,

S (Salicylic) for S (Superficial)

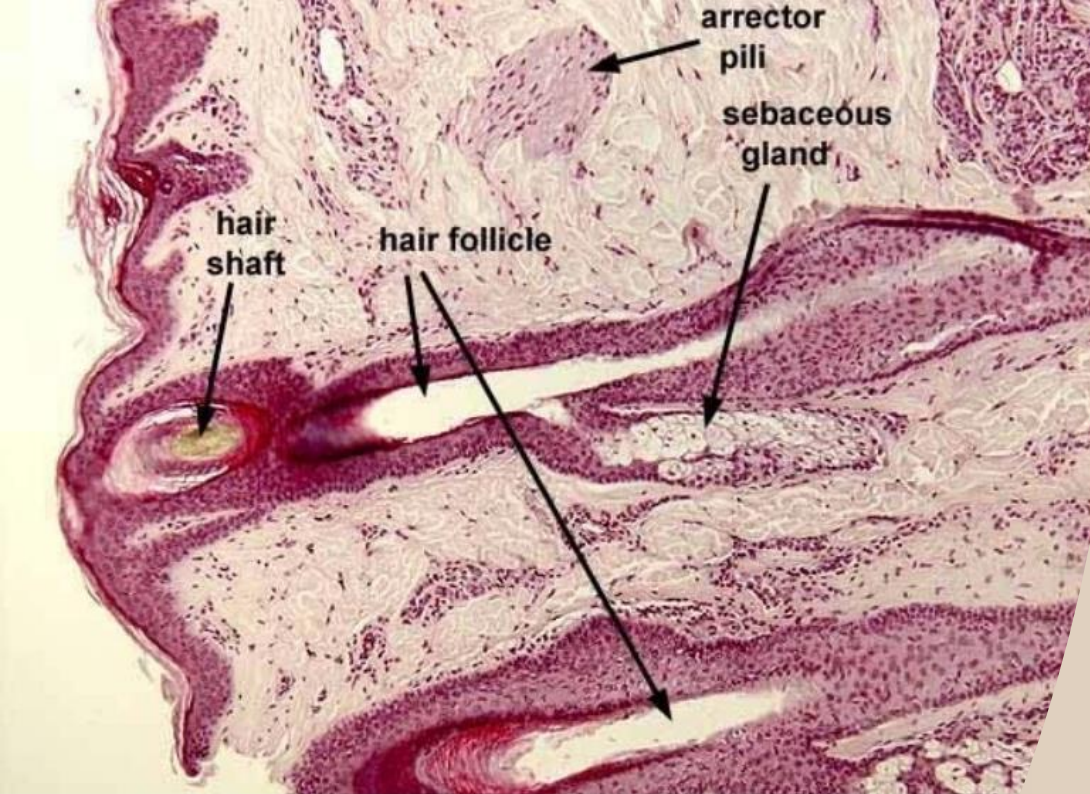
2.Targeting the
pigment
(ColorKiller)

Feature	TCA (Trichloroacetic Acid)	Salicylic Acid
Type of Acid	TCA	Beta-hydroxy acid (BHA)
Mechanism of Action	Medium-depth peel that exfoliates and stimulates collagen production	Superficial exfoliation, oil-soluble, penetrates pores
Target Area	Deeper pigmentation (e.g., melasma, sun spots)	Superficial pigmentation (e.g., PIH)
Penetration Depth	Medium depth	Superficial to medium depth
Suitable for Skin Types	Thicker skin types or severe pigmentation	Best for oily, acne-prone skin with superficial ,,light,,pigmentation
Frequency of Treatment	4 treatments spaced 1 /week or 2 weeks	Can be used more frequently (bi-weekly, monthly)
Main Benefits	Reduces deeper pigmentation, stimulates collagen production, improves skin texture	Prevents clogged pores, brightens skin
Side Effects	Redness, swelling, scabbing (can be intense)	Mild redness less intense than TCA

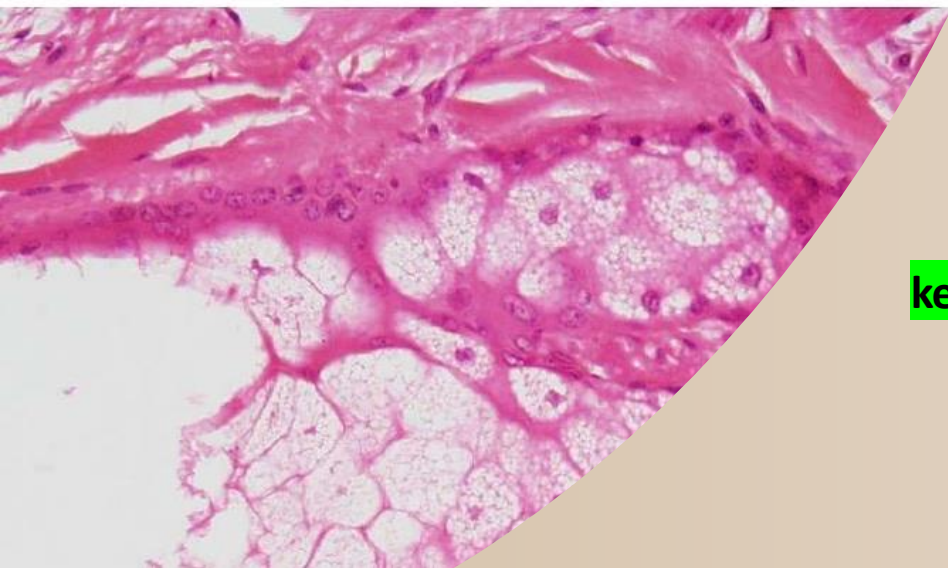
Clinical Effects & Visual Appearance

2.Targeting the pigment (ColorKiller)

SALICYLIC CONCENTRATIONS & VISUAL EFFECTS		ROSE (BLUSH)	GREY (PATCHY)	WHITE DENSE
1	Physical Effect	Appearance	Crystallization	Pseudo Frosting
2	Concentrations w/v	25% or less	25-30%	30-40%
3	Effects	<ul style="list-style-type: none">Mild erythemano visible frostingno visible crystallization	<ul style="list-style-type: none">Visible pseudofrosting in some zonespartial precipitation of crystals	<ul style="list-style-type: none">Thick white filmcrystallized salicylic acidnot true protein coagulation
4	Depth	<ul style="list-style-type: none">Very superficial	<ul style="list-style-type: none">Superficial peel	<ul style="list-style-type: none">Superficial epidermal peel
5	Indications	<div><ul style="list-style-type: none">Sensitive skin</div> <div><ul style="list-style-type: none">maintenance treatments</div>	<ul style="list-style-type: none">Acne-prone skincomedonesoily zones	<ul style="list-style-type: none">Acneseborrheic skinkeratosis pilaristhick skin areas (nose, chin, back)



TO PEEL OR NOT TO PEEL ARMPITS



keratinized stratified squamous epithelium. = Keratotic Zone

Hyperpigmentation of Axillary Hollow
Treatment : Salicylic Acid + Metabolic Peels

BEFORE

AFTER



Courtesy of Dr. Alain Tenenbaum

TCA + Metabolic Peels
alternate with Salicylic Acid + Metabolic Peels

BEFORE

AFTER 3 SESSIONS



WHAT DO YOU NEED?

- Disposable clothing for patients and peelers (gown-e.g. non-sterile)
- Monouse gloves (S-M-L) non-sterile
- Robust cotton swabs non-sterile
- non-sterile wooden tongue depressor
- Monouse hoods
- Multicompresses 10x10 cm made of cotton non-sterile (no swabs)
- Porcelain, glass container-(No metal containers)
- **Products**



PEELS IN INTIMATE AREAS

Anatomical, histological and clinical approach

**A.TENENBAUM,
M.D.,Ph.D., D.Sc**



M.TIZIANI, RCSA

drpeeling@bluewin.ch



INTIMATE AREAS WHERE SKIN CARE IS IMPORTANT

For Men

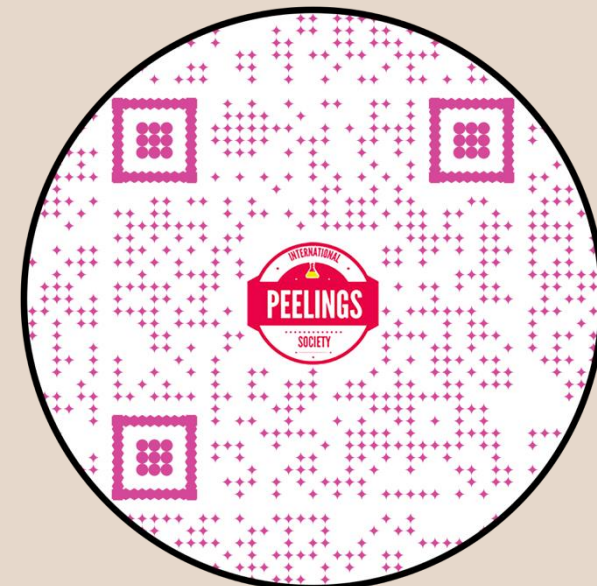
Perianal
Area

For Women

Perianal
Area

External
Genital Skin

Inner Thighs



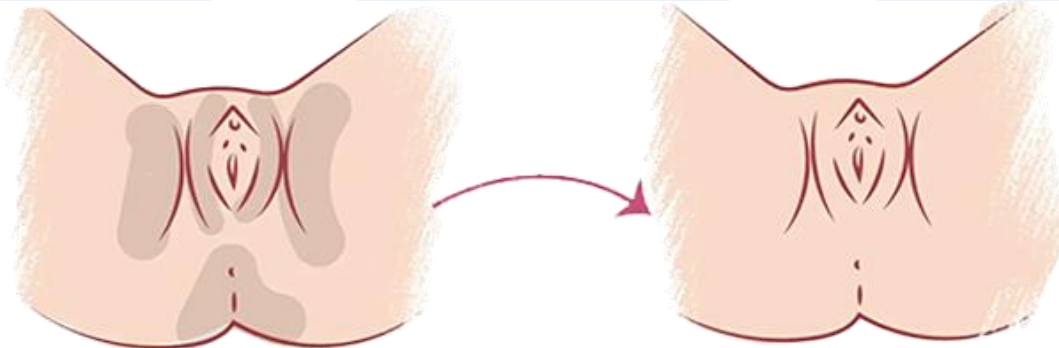
COMMON INDICATIONS

Common Aesthetic & Functional Indications

Pigmentation
Issues

Post Pregnancy
Skin Changes

Age-related
textural &
pigmentary
changes



Specific Medical Indication (Expert Use Only)

Lichen Sclerosus

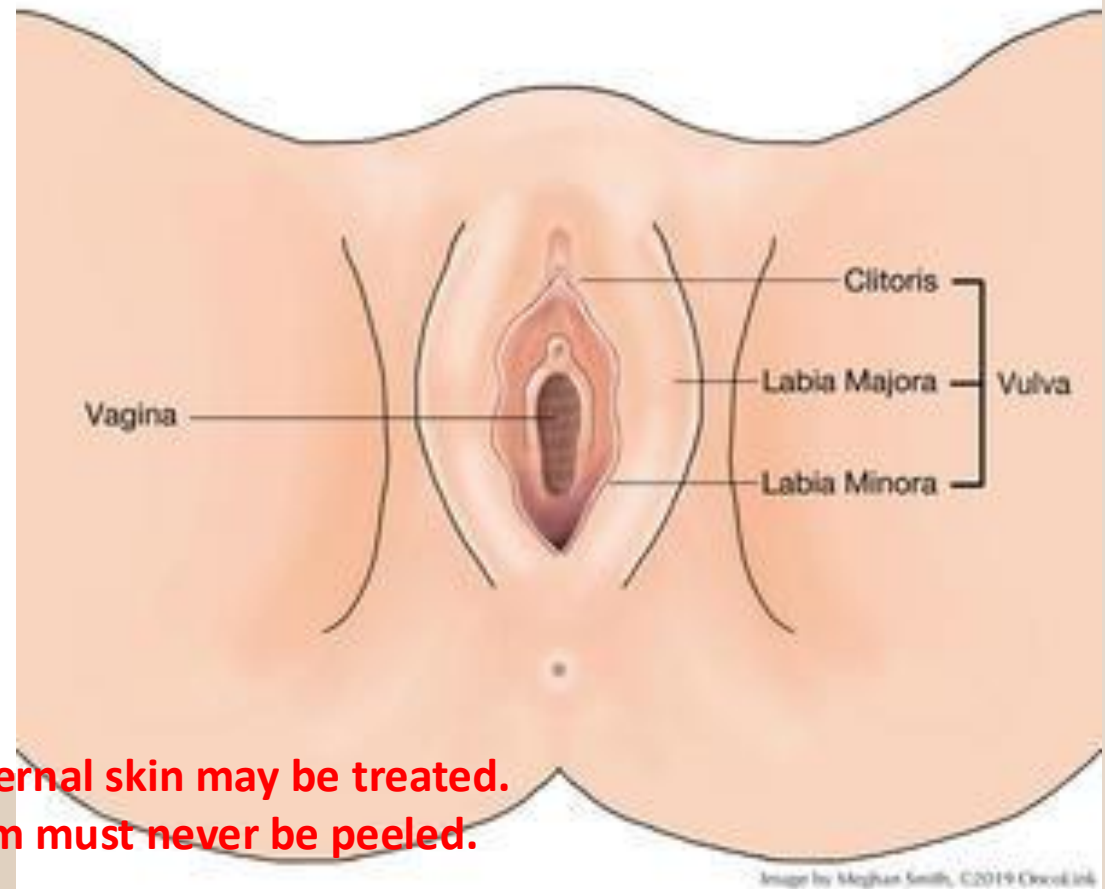
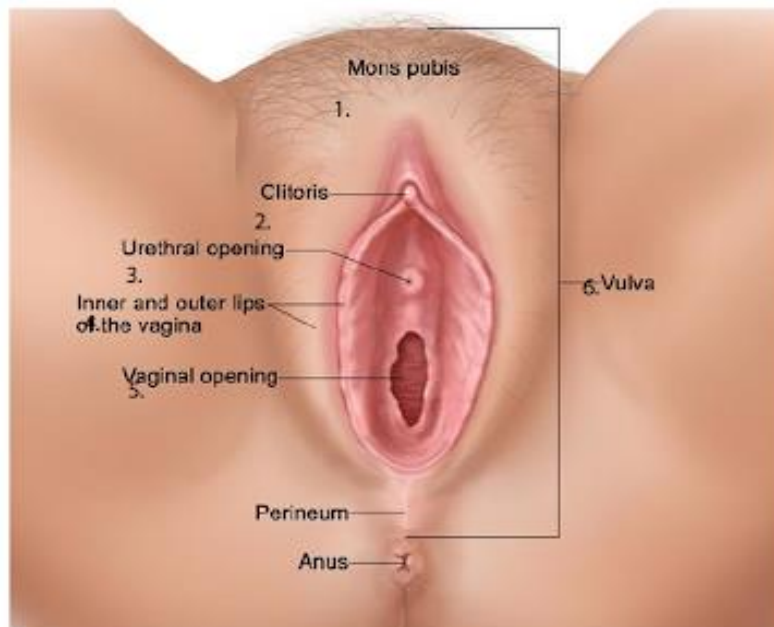


INTIMATE AREAS – EXTERNAL FEMALE ANATOMY

STRUCTURES OF THE VULVA

NEVER TREAT INTERNAL STRUCTURES
OF THE VULVA WITH PEELS

External Reproductive Anatomy



**Only keratinized external skin may be treated.
Mucosal epithelium must never be peeled.**

WHERE TO PEEL

IN BLUE : EXTERNAL FEMALE GENITALIA+ PERIANAL AREA

IN GREEN : INNER THIGHS



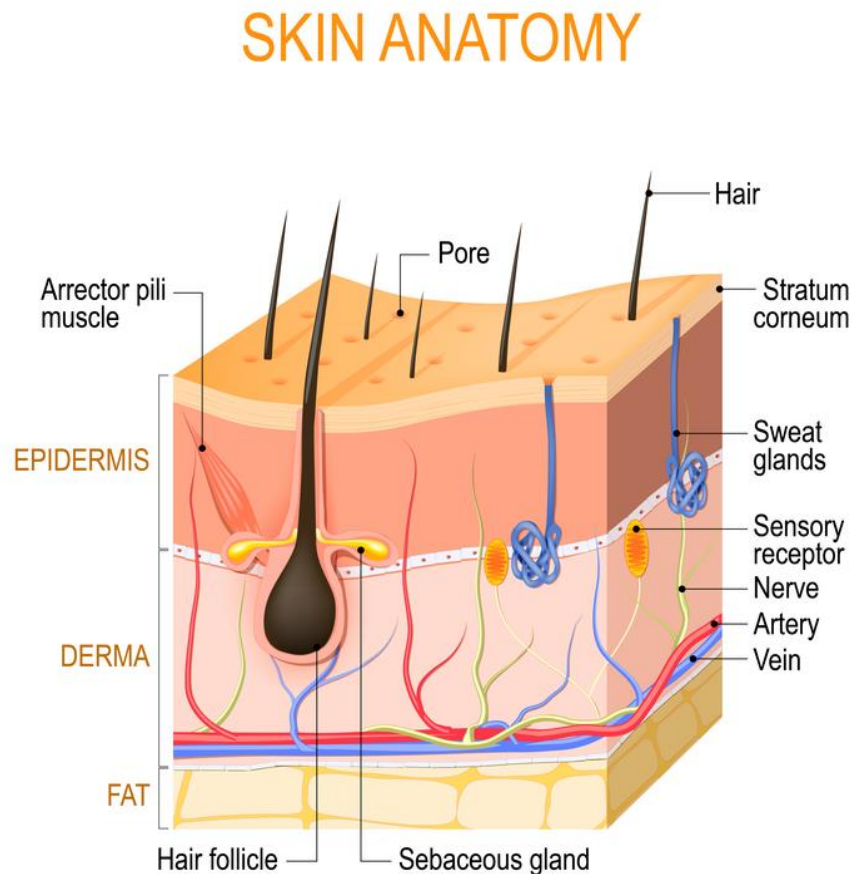
Color coding is based on epithelial type, not on anatomy alone.



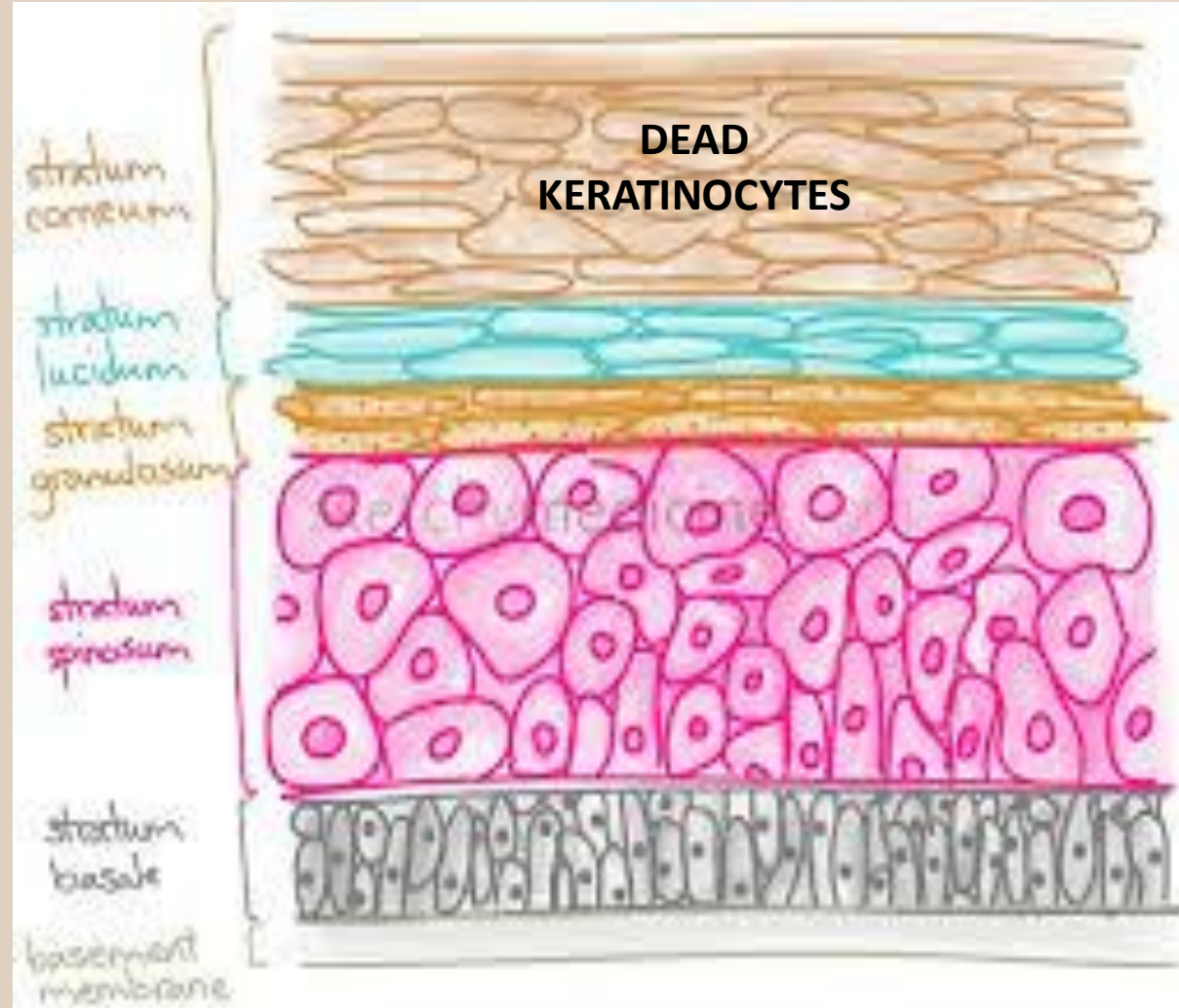
HISTOLOGY OF THE SKIN AND EPIDERMIS

Skin in Intimate Areas is thinner with higher sensitivity and potential for Hyperpigmentation

3 Layers

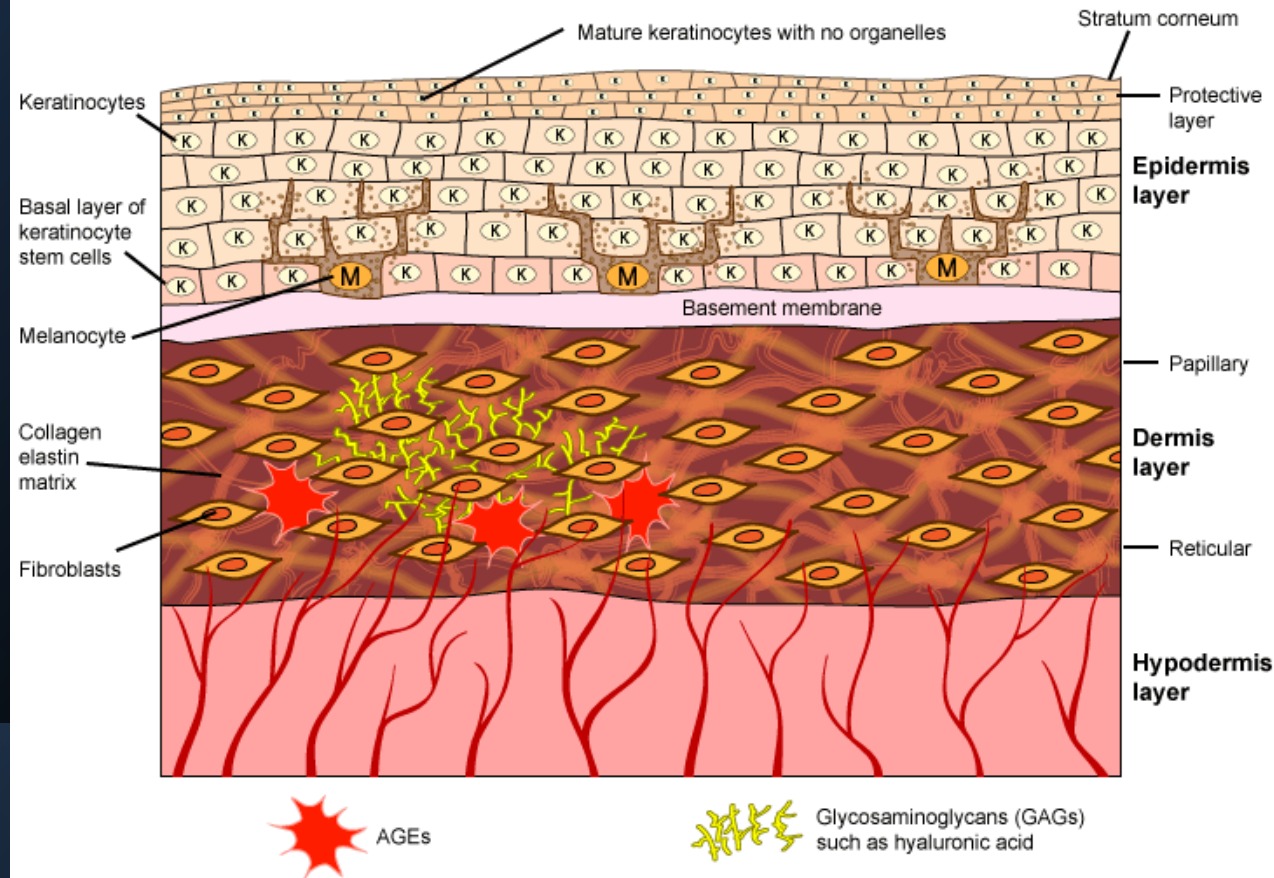


5 Epidermal layers



PIGMENT DISTRIBUTION INVOLVES EPIDERMIS, BASEMENT MEMBRANE AND DERMIS

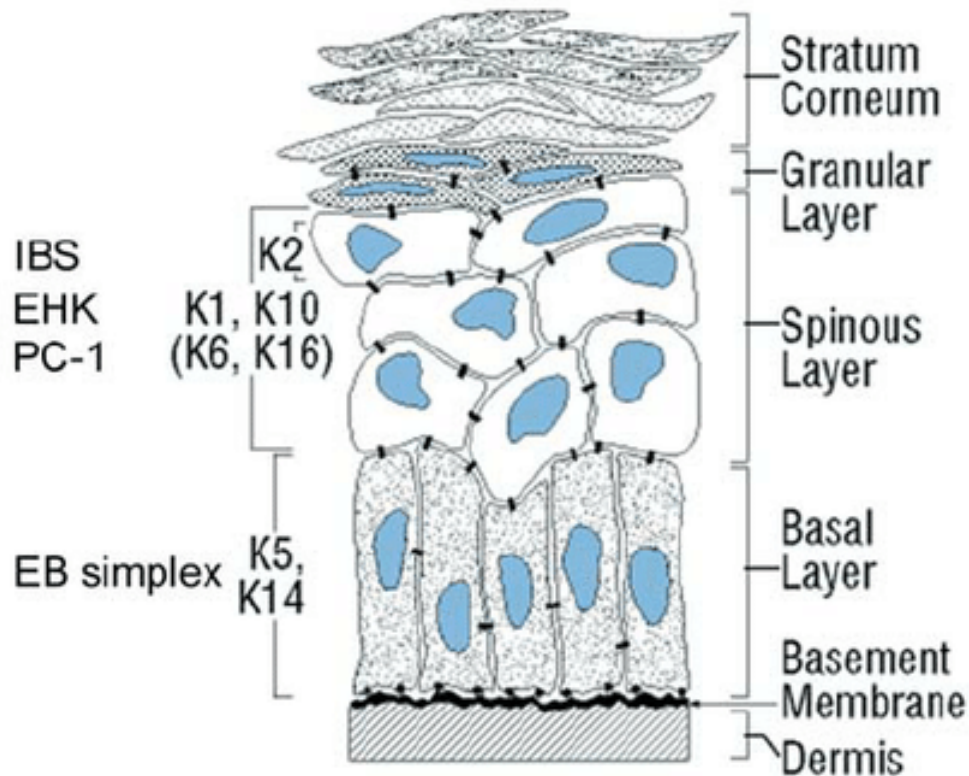
Superficial treatments
alone are insufficient
when pigment extends
beyond the epidermis.



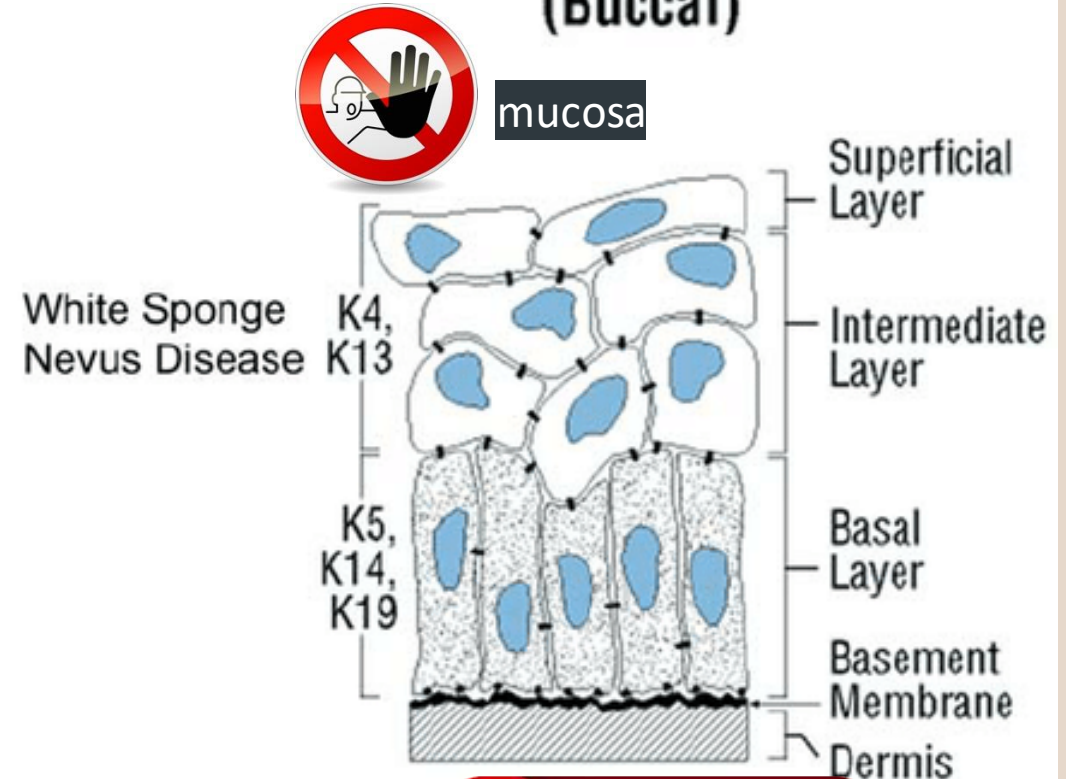
KERATINIZED AND NON KERATINIZED STRATIFIED SQUAMOUS EPITHELIUM

Only keratinized stratified squamous epithelium
can be safely peeled.

Epidermis, Oral Keratinized



**Oral Non-Keratinized
(Buccal)**



TO PEEL OR NOT TO PEEL THAT IS THE QUESTION

Anatomy is only a proxy - epithelial histology is the rule.



keratinized stratified squamous epithelium

MONS PUBIS

LABIA MAJORA

EXTERNAL ANAL MARGIN

PERIANAL AREA

INNER THIGHS



NOT keratinized stratified squamous epithelium
(Mucosa)

CLITORIS

LABIA MINORA

VESTIBULE

ANUS

VAGINA

TO PEEL OR NOT TO PEEL THAT IS THE QUESTION



CLITORIS

NOT keratinized stratified squamous epithelium
(Mucosa)

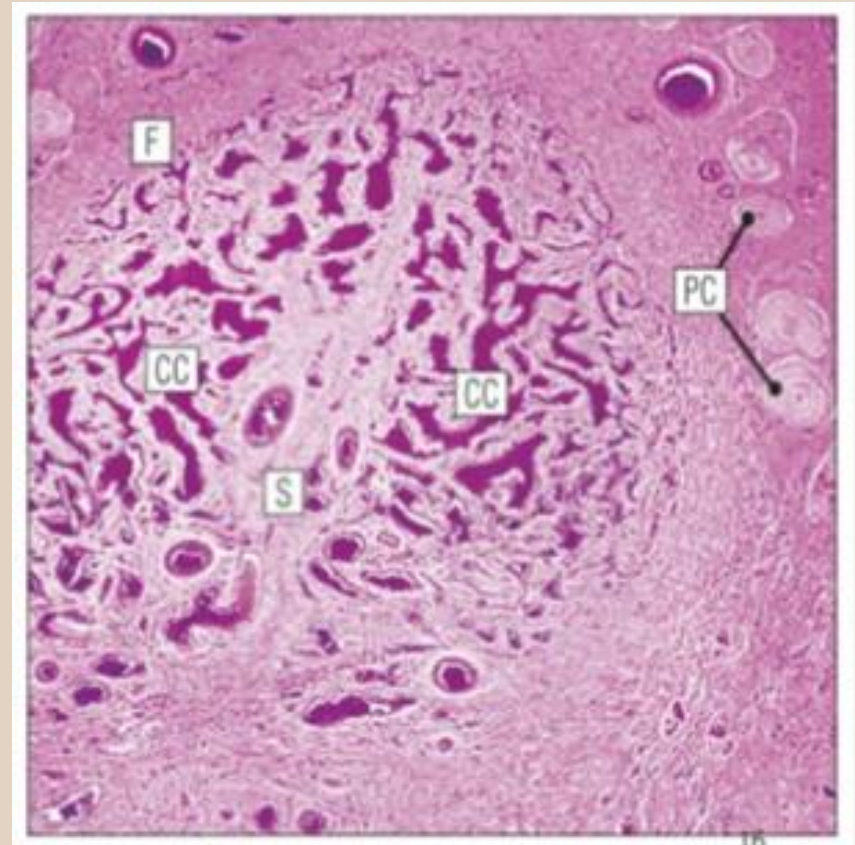
CLITORIS

LABIA MINORA

VESTIBULE

ANUS

VAGINA



No hair follicles are observed in this section

TO PEEL OR NOT TO PEEL THAT IS THE QUESTION



LABIA MINORA

NOT keratinized stratified squamous epithelium
(Mucosa)

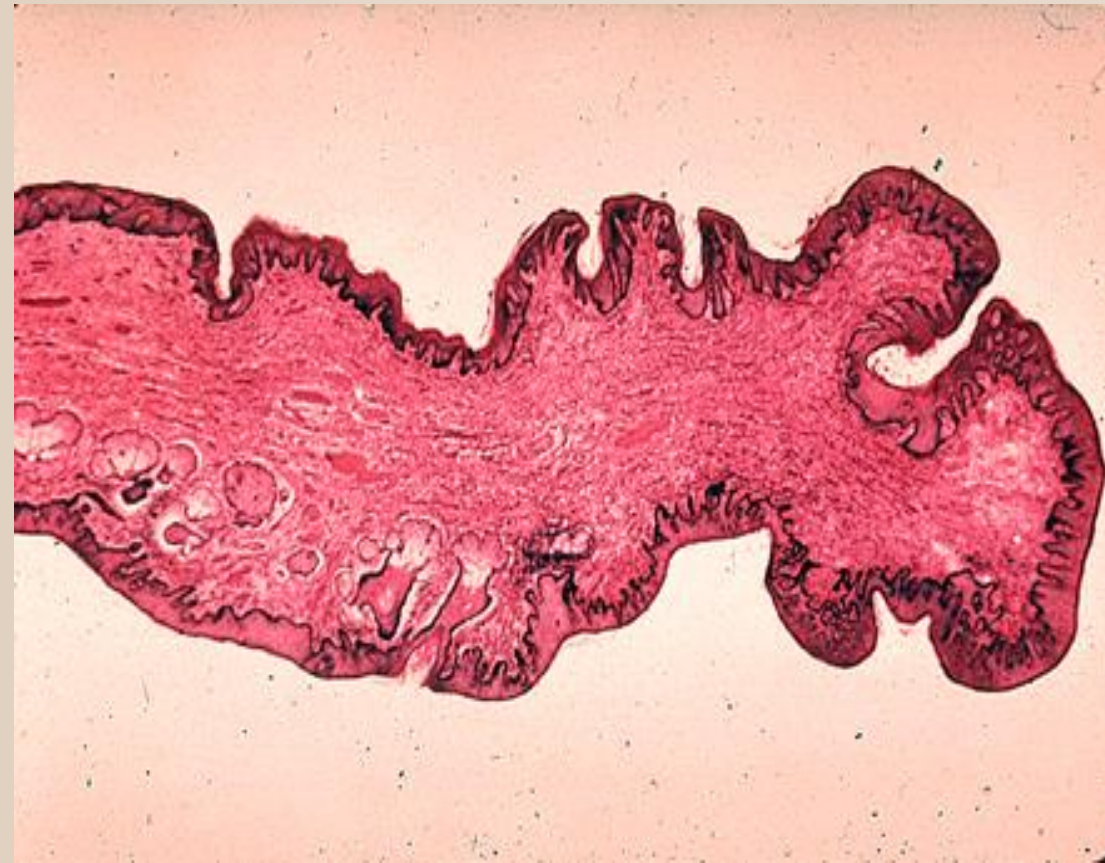
CLITORIS

LABIA MINORA

VESTIBULE

ANUS

VAGINA



No hair follicles are observed in this section

TO PEEEL OR NOT TO PEEEL THAT IS THE QUESTION



VESTIBULE

NOT keratinized stratified squamous epithelium
(Mucosa)

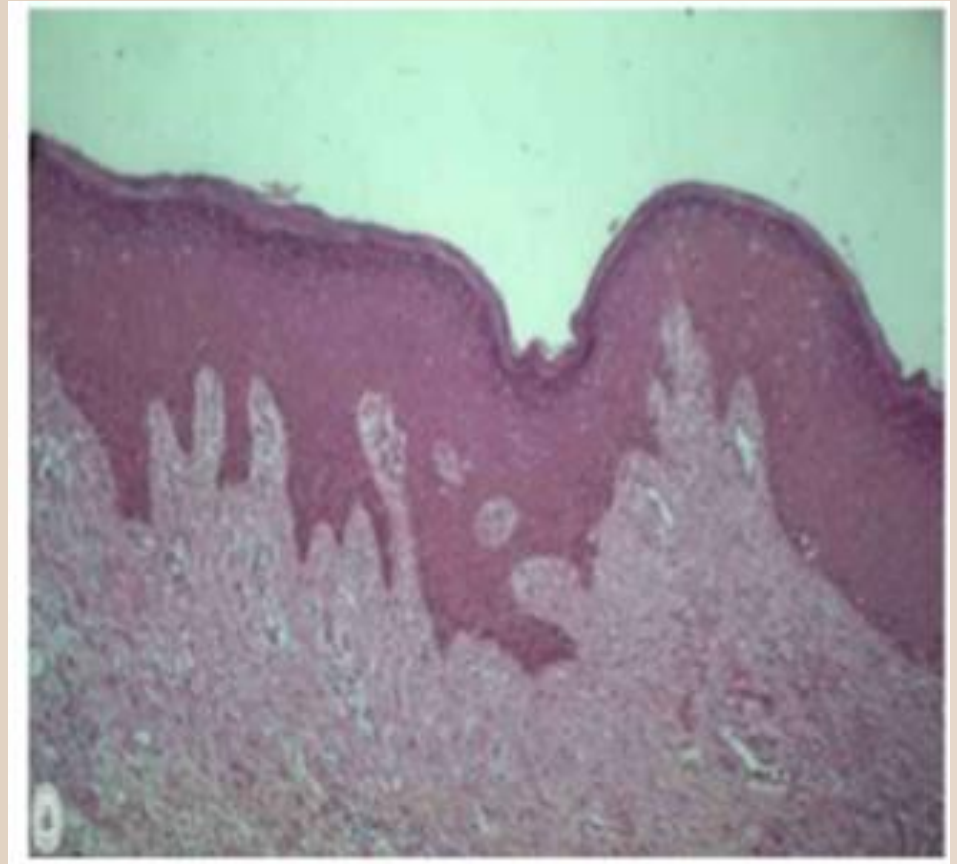
CLITORIS

LABIA MINORA

VESTIBULE

ANUS

VAGINA



No hair follicles are observed in this section

TO PEEL OR NOT TO PEEL THAT IS THE QUESTION



RECTUM

NOT keratinized stratified squamous epithelium
(Mucosa)

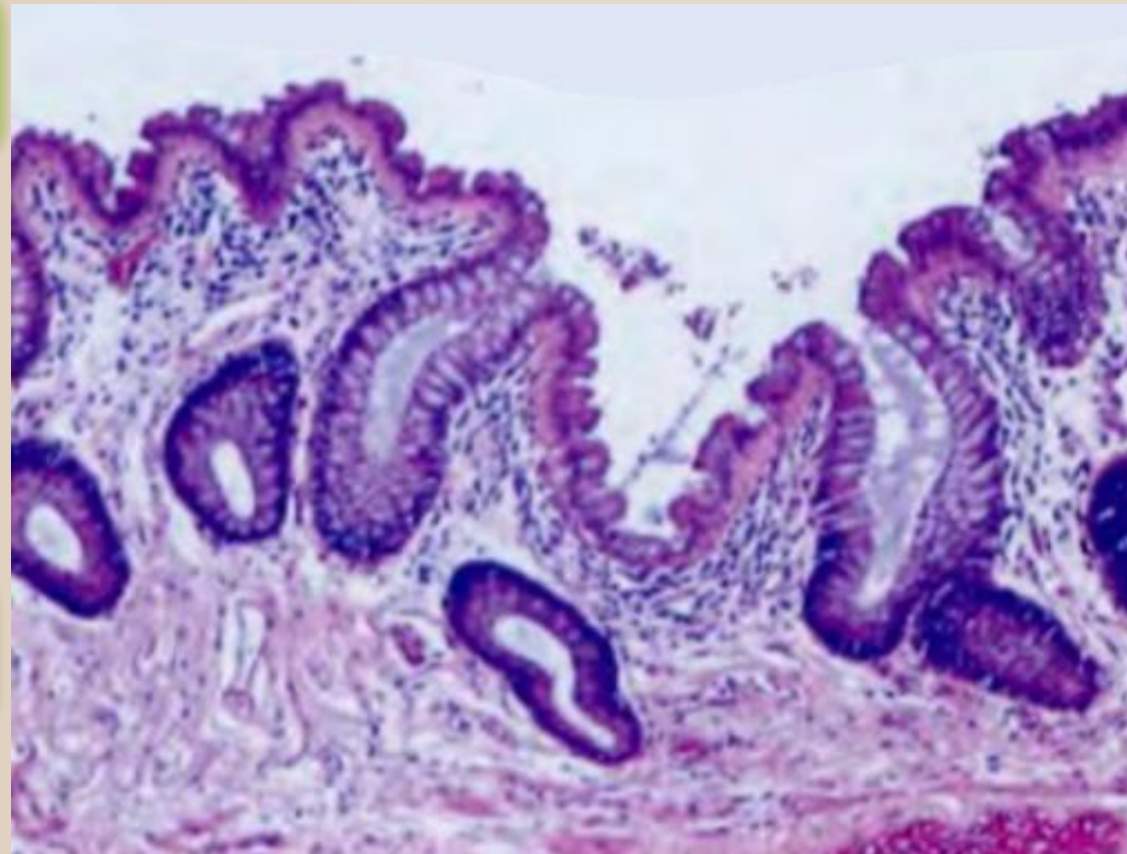
CLITORIS

LABIA MINORA

VESTIBULE

ANUS

VAGINA



No hair follicles are observed in this section

TO PEEL OR NOT TO PEEL THAT IS THE QUESTION



VAGINA

NOT keratinized stratified squamous epithelium
(Mucosa)

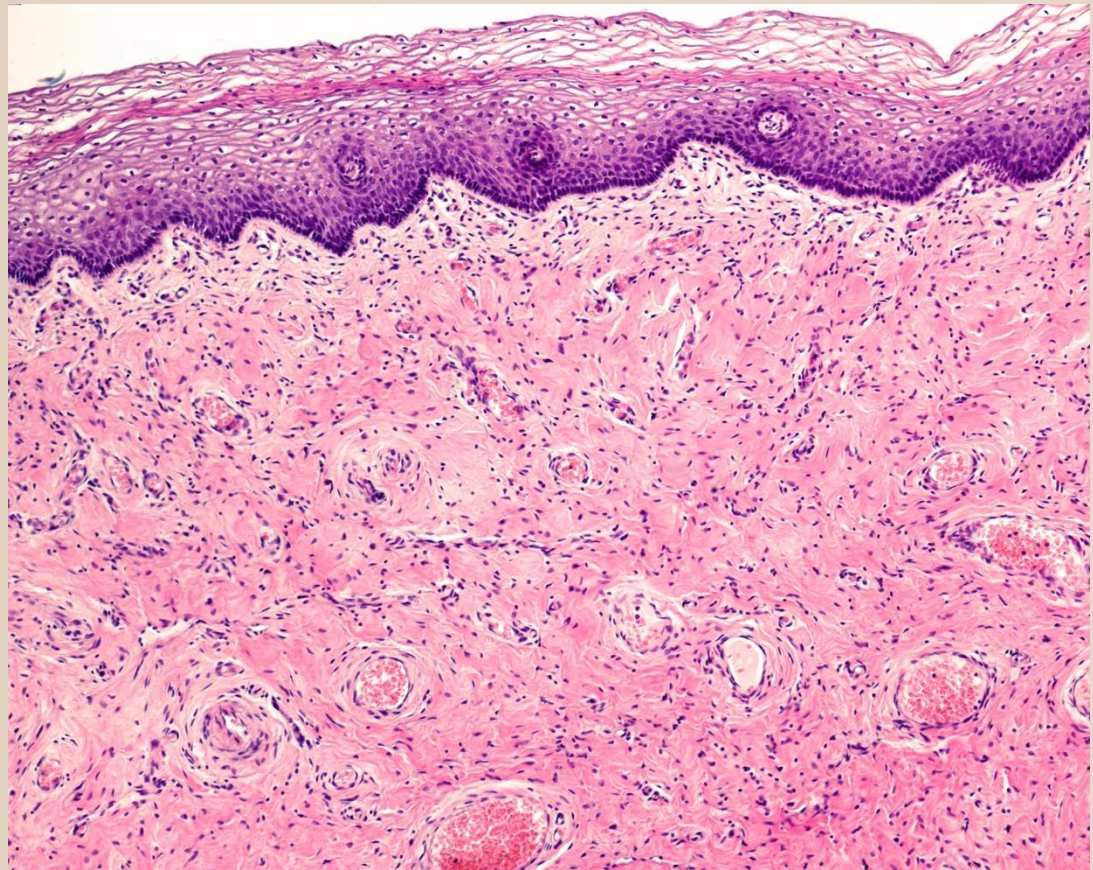
CLITORIS

LABIA MINORA

VESTIBULE

ANUS

VAGINA



No hair follicles are observed in this section

TO PEEL OR NOT TO PEEL THAT IS THE QUESTION



keratinized stratified squamous epithelium

MONS PUBIS

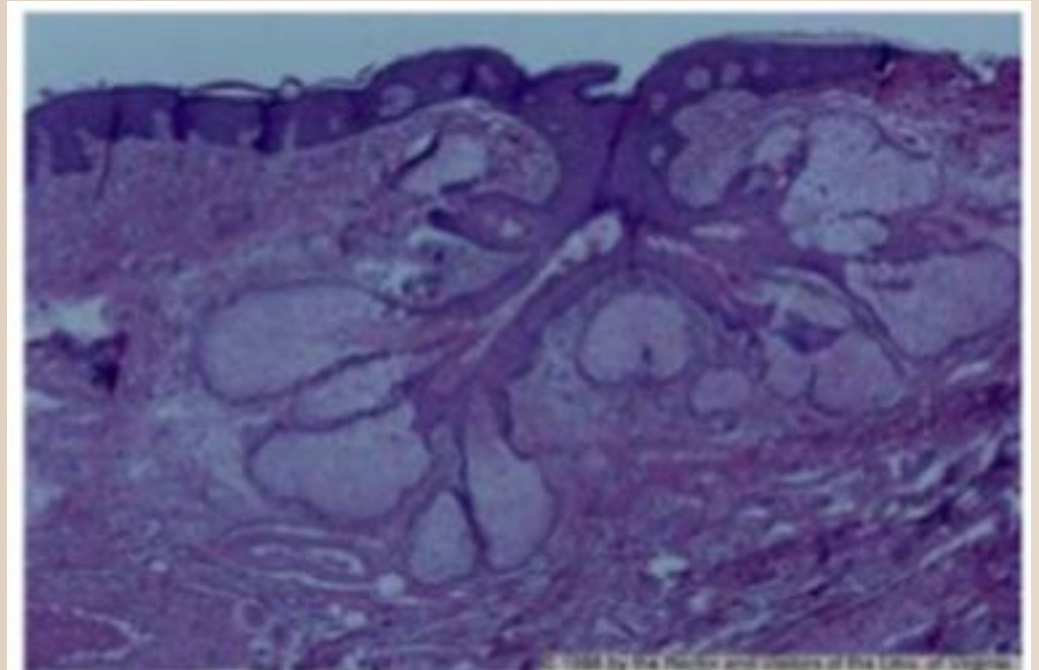
LABIA MAJORA

EXTERNAL ANAL MARGIN

PERIANAL AREA

INNER THIGHS

MONS PUBIS



Hair follicles are observed in this section

**Only keratinized external skin is suitable for peeling.
This does not include mucosal or transitional epithelium.**

TO PEEL OR NOT TO PEEL THAT IS THE QUESTION



keratinized stratified squamous epithelium

MONS PUBIS

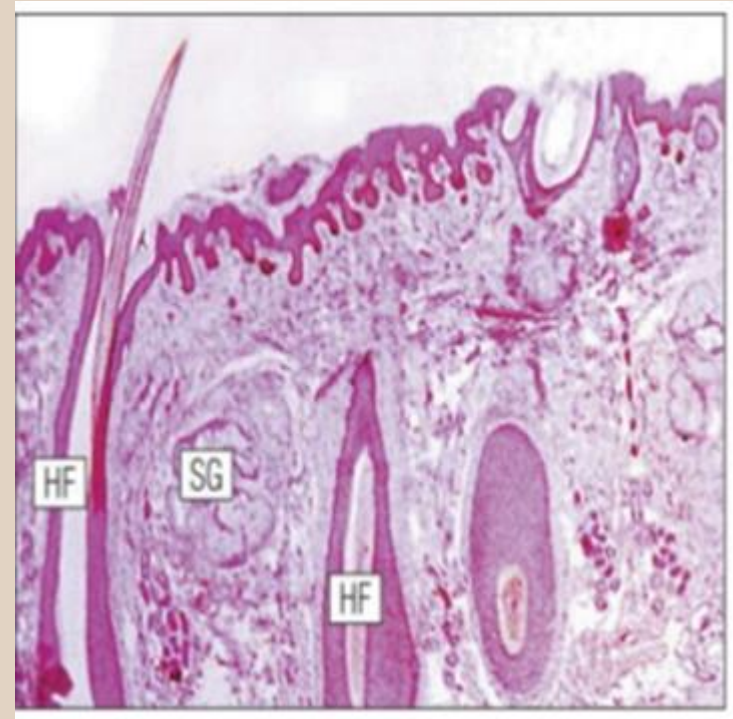
LABIA MAJORA

EXTERNAL ANAL MARGIN

PERIANAL AREA

INNER THIGHS

LABIA MAJORA



Hair follicles are observed in this section

TO PEEL OR NOT TO PEEL THAT IS THE QUESTION



keratinized stratified squamous epithelium

MONS PUBIS

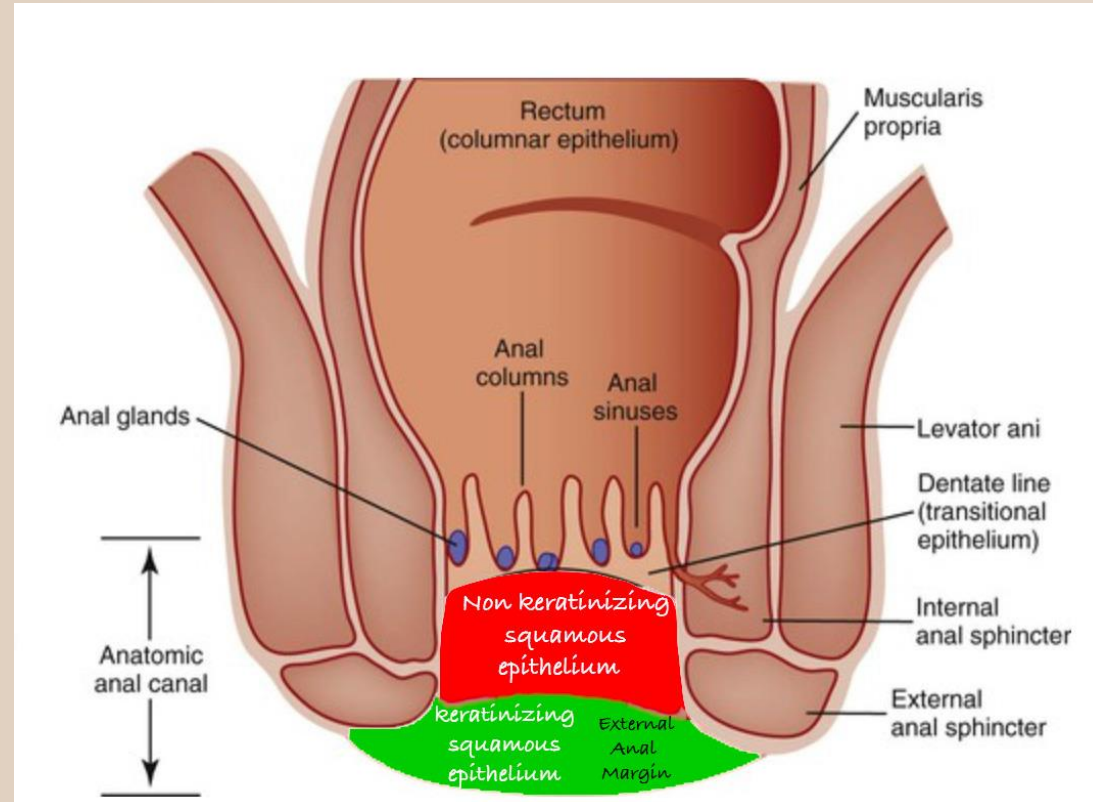
LABIA MAJORA

EXTERNAL ANAL MARGIN

PERIANAL AREA

INNER THIGHS

EXTERNAL ANAL
MARGIN
TRANSITION AREA



Only the external keratinized anal margin is suitable for peeling.
The anal canal and internal mucosa are strictly contraindicated.

TO PEEL OR NOT TO PEEL THAT IS THE QUESTION



keratinized stratified squamous epithelium

MONS PUBIS

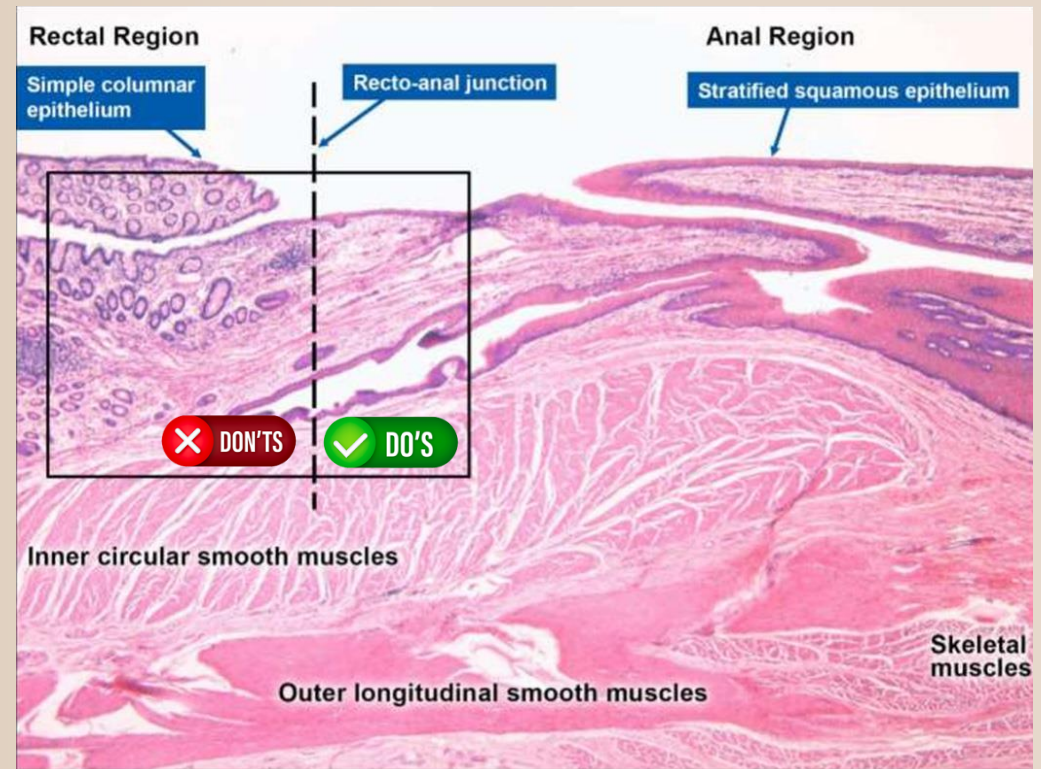
LABIA MAJORA

EXTERNAL ANAL MARGIN

PERIANAL AREA

INNER THIGHS

EXTERNAL ANAL MARGIN TRANSITION ZONE



No hair follicles are seen in this section,
although they are typically present in the
peri anal skin

TO PEEL OR NOT TO PEEL THAT IS THE QUESTION



keratinized stratified squamous epithelium

MONS PUBIS

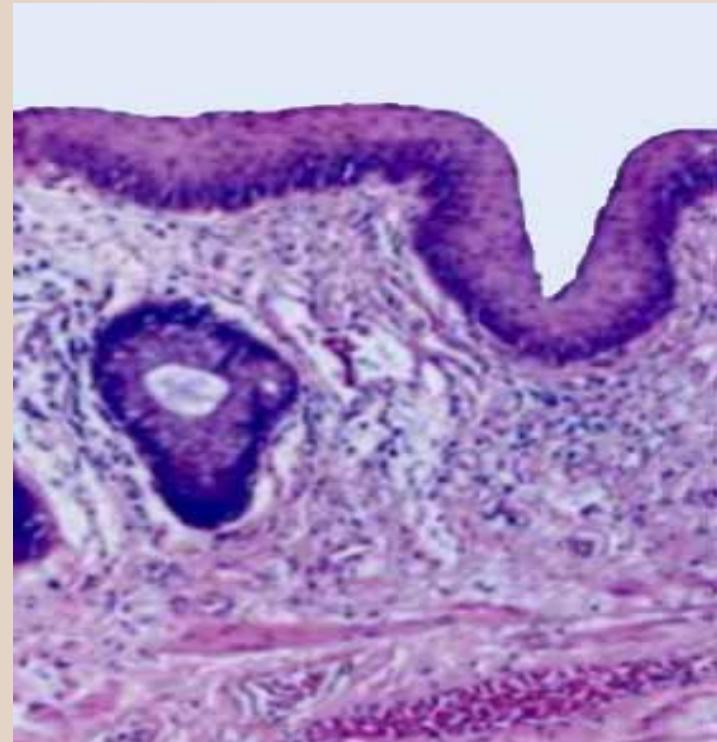
LABIA MAJORA

EXTERNAL ANAL MARGIN

PERIANAL AREA

INNER THIGHS

PERIANAL AREA



No hair follicles are seen in this section,
although they are typically present in this region

ETIOLOGY OF HYPERPIGMENTATION IN INTIMATE AREAS

- **DRUGS & CHEMICALS**

- Hydroquinon->hypopigmentation
- Antibiotics :Tetracyclins,Rifampicin,AZT
- Antimalaria
- Cytostatics
- Psychomedicines : phenothiazin
- Antiepileptics : Phenytoin
- Antiarythmics: Cordaron
- Psoralens
- Melanocyte Stimulating Hormon
- Arsenic
- Bergamote

- **DERMA-PATHOLOGIES**

- PIH
- Psoriasis
- Pityriasis Versicolor
- LED
- Lichens
- Ephelides
- Lentigines
- Neurofibromatosis
- Naevus (Becker)
- Peutz-Jeghers Syndrome (gastro intestinal disease)
- Hemochromatosis (Fe)

- **BAD HABITS**

- -UV or solarium
- -jeans too tight
- -underwear too tight
- -pantyhose
- -check laundry products
- -check buttons, metals ... of clothes & underwears
- -depilation, shaving, other products

- **HORMONAL DISORDERS**

- -Addison
- -ACTH
- -Testosteron in excess
- -Anabolisants
- -Pregnancy
- -Oral Contraceptives
- -Post Pubertal Melanosis

Chronic irritation and inflammation are primary drivers

CONCEPTS OF A.TENENBAUM & M.TIZIANI APPLIED TO INTIMATE HYPERPIGMENTATION

1.Targeting the pigment directly Or the „ColorKiller,, of A.Tenenbaum



by inducing a form of *„photo,,damage or „photo,,necrosis* to effectively destroy the colors responsible for the clinical expression of hyperpigmentation

WHITE FROSTING



By using the frosting effect ensuring that it *completely covers the hyperpigmented area and effectively obscures the discoloration.*

2.Addressing the 3 key compartments of melanin metabolism

- **Production**: Modifying the melanogenesis process to reduce melanin synthesis.
- **Transport**: Interfering with the movement of melanin within the skin to prevent uneven distribution.
- **Destruction**: Enhancing the breakdown and removal of excess melanin from the skin.

1 + 2 = Aggressive Treatment with Downtime

(Protocol by A. Tenenbaum)

2 = Gentle Treatment Without Downtime

(Protocol by M. Tiziani)

EPIDERMAL THICKNESS OF INTIMATE AREAS

Reduced epidermal thickness increases penetration depth and inflammatory response.

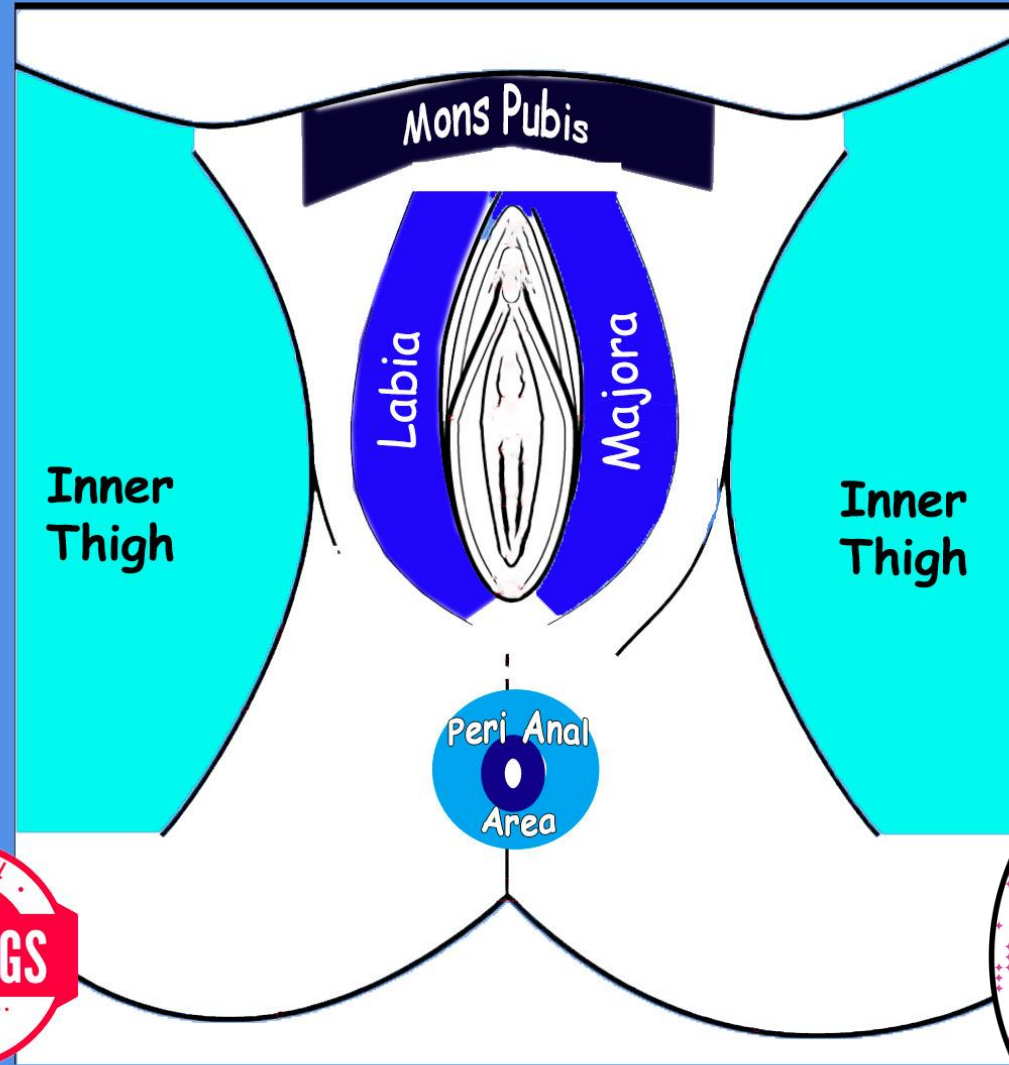
100-120
(μm)

90-100
(μm)

70-90
(μm)

60-75
(μm)

50-70
(μm)



These differences explain why standard peel concentrations cannot be transposed.



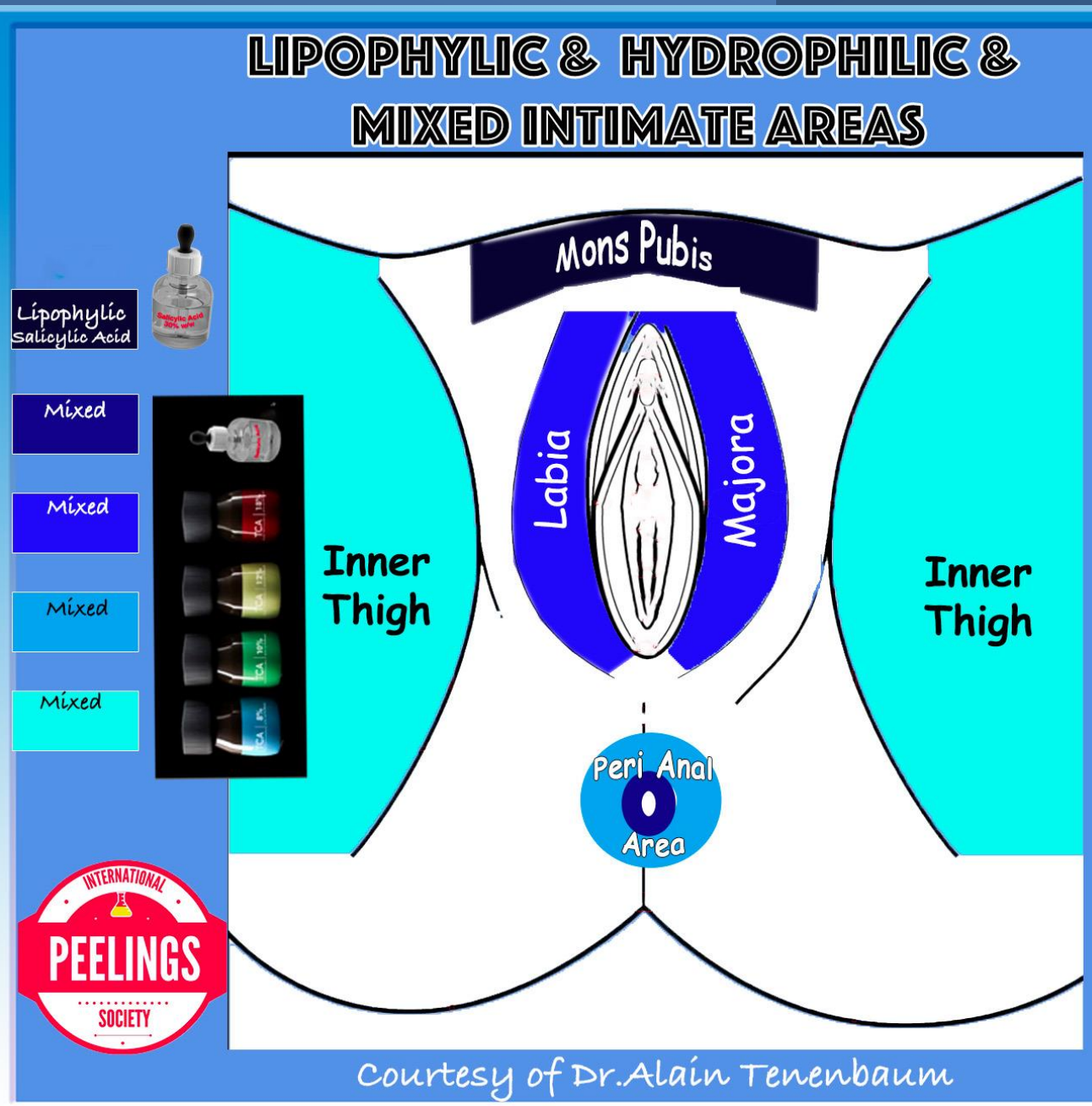
Courtesy of Dr. Alain Tenenbaum



**1.TARGETING THE PIGMENT
DIRECTLY
OR THE „COLORKILLER,, OF
A.TENENBAUM**

Why solvent choice matters in
intimate areas

Penetration is driven more by the
solvent than by the acid
concentration alone.



Targeting the pigment directly
Or the „ColorKiller,, of A.Tenenbaum

WHICH SALICYLIC ACID TO USE ?



Comparison of Salicylic Acids

Different salicylic acid formulations behave very differently on thin, reactive skin.

**TARGETING THE PIGMENT
DIRECTLY
OR THE „COLORKILLER,, OF
A.TENENBAUM**

Solvent and delivery
system are the primary
determinants of safety.

Lower concentration does
not automatically mean
safer treatment.

SALICYLIC ACIDS IN ETHANOL VS GLYCEROL FOR INTIMATE AREAS

Lipophylic Salicylic Acid

Mixed

Mixed

Mixed

Mixed

Salicylic Acid in Glycerol

Inner Thigh

Mons Pubis

Labia

Majora

Inner Thigh

Peri Anal Area

**INTERNATIONAL
PEELINGS
SOCIETY**

Courtesy of Dr. Alain Tenenbaum

TYPICALLY POST-
INFLAMMATORY
AND FRICTION-
RELATED
PIGMENTATION.

Clinical appearance
alone is not sufficient
to decide treatment.

HYPERPIGMENTATION OF FEMALE GENITAL SKIN



Courtesy of Dr. Alain Tenenbaum

**TARGETING THE PIGMENT
DIRECTLY**
OR THE „COLORKILLER,, OF
A.TENENBAUM

**INITIAL APPEARANCE OF
WHITE (PSEUDO) FROSTING
ON THE LABIA MAJORA**



Courtesy of Dr. Alain Tenenbaum

Pseudo-frosting ≠ protein coagulation.

Thin epidermis may transiently
appear white without tissue
damage.

PERIANAL AREA: THICK, DRY, KERATINIZED SKIN

The Fitzpatrick Scale					
					
TYPE I Light pale white Always burns never tans	TYPE II White, fair Usually burns, tans with difficulty	TYPE III Medium white to olive Sometimes mild burn, gradually tans to olive	TYPE IV Olive, moderate brown Rarely burns, tans easily to moderate brown	TYPE V Brown, dark brown Very rarely burns, tans very easily	TYPE VI Black, very dark brown to black Never burns, tans easily, always hyperpigmented

Male Perianal Area Phototype 3/4

The patient must be depilated 3 days before the procedure and cleaned with chlorhexidine or cetrimide without alcohol before the procedure

PERIANAL SKIN HYPERPIGMENTATION



COURTESY OF DR. ALAIN TENENBAUM





PERIANAL AREA IS A THICK DRY AREA

The perianal area is typically considered a dry area, as it has a low concentration of sebaceous (oil-producing) glands.

However, it can become moist due to sweat, mucus, or hygiene-related factors.

The skin in this area can be sensitive, so it's important to maintain proper hygiene without causing irritation.

Only the external anal margin is keratinized and treatable.
The anal canal is mucosal and must never be peeled.

EXTERNAL ANAL MARGIN IS THICKER AND DRIER VS PERIANAL AREA



The external anal margin is generally thicker and drier compared to the rest of the perianal skin.

This area has a tougher skin texture, which helps protect it from friction and abrasion.

It's also less oily because it has fewer sebaceous glands.

However, like the rest of the perianal region, it can still be affected by moisture, such as sweat, or irritation from hygiene practices or stool. The skin is also more prone to dryness, particularly if there's excessive wiping or use of harsh products.

Keratinization allows treatment, but does not eliminate PIH risk.



PROTOCOL EXTERNAL ANAL MARGIN & PERI ANAL AREA

[take a look](#)

CONCEPTS OF A.TENENBAUM & M.TIZIANI

EXTERNAL ANAL MARGIN & PERI ANAL AREA

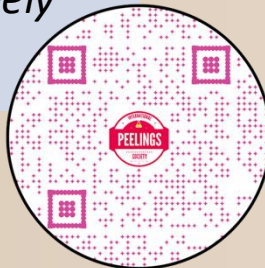
Targeting the pigment directly
Or the „ColorKiller,, of
A.Tenenbaum

COLORKILLER

by inducing a form of *photodamage* or '*photonecrosis*' to effectively destroy the colors responsible for the clinical expression of hyperpigmentation

WHITE FROSTING

By using the frosting effect ensuring that it *completely covers the hyperpigmented area and effectively obscures the discoloration.*



Results shown are obtained on selected keratinized skin under controlled protocols.

Early improvement does not replace long-term stabilization.

Attack Treatment Protocol for Peri Anal Area



- Alternate TCA , Salicylic acid in ethanol and Salicylic acid in glycerol to get a frosting 1 session each 2 weeks x 4-6 weeks
- Then as usual

Attack Treatment Protocol for External Anal Margin



- **Dont use TCA liq because the frosting could reach the rectum !**
- Then as usual

Concepts of A.Tenenbaum & M.Tiziani

External Anal Margin & Peri Anal Area

1.Targeting the pigment directly Or the „ColorKiller,, of A.Tenenbaum



COLORKILLER

by inducing a form of *photodamage* or '*photonecrosis*' to effectively destroy the colors responsible for the clinical expression of hyperpigmentation



WHITE FROSTING

By using the frosting effect ensuring that it *completely covers the hyperpigmented area and effectively obscures the discoloration.*

2.Addressing the 3 key compartments of melanin metabolism

- **Production**: Modifying the melanogenesis process to reduce melanin synthesis.
- **Transport**: Interfering with the movement of melanin within the skin to prevent uneven distribution.
- **Destruction**: Enhancing the breakdown and removal of excess melanin from the skin.



1 + 2 = Aggressive Treatment with Downtime

(Protocol by A. Tenenbaum)

2 = Gentle Treatment Without Downtime

(Protocol by M. Tiziani)

MAINTENANCE ACIDS AIM AT STABILIZATION, NOT ACTIVE DEPIGMENTATION.

Which acids can be used for maintenance

These acids are suitable for maintenance only, on selected keratinized skin and under strict conditions

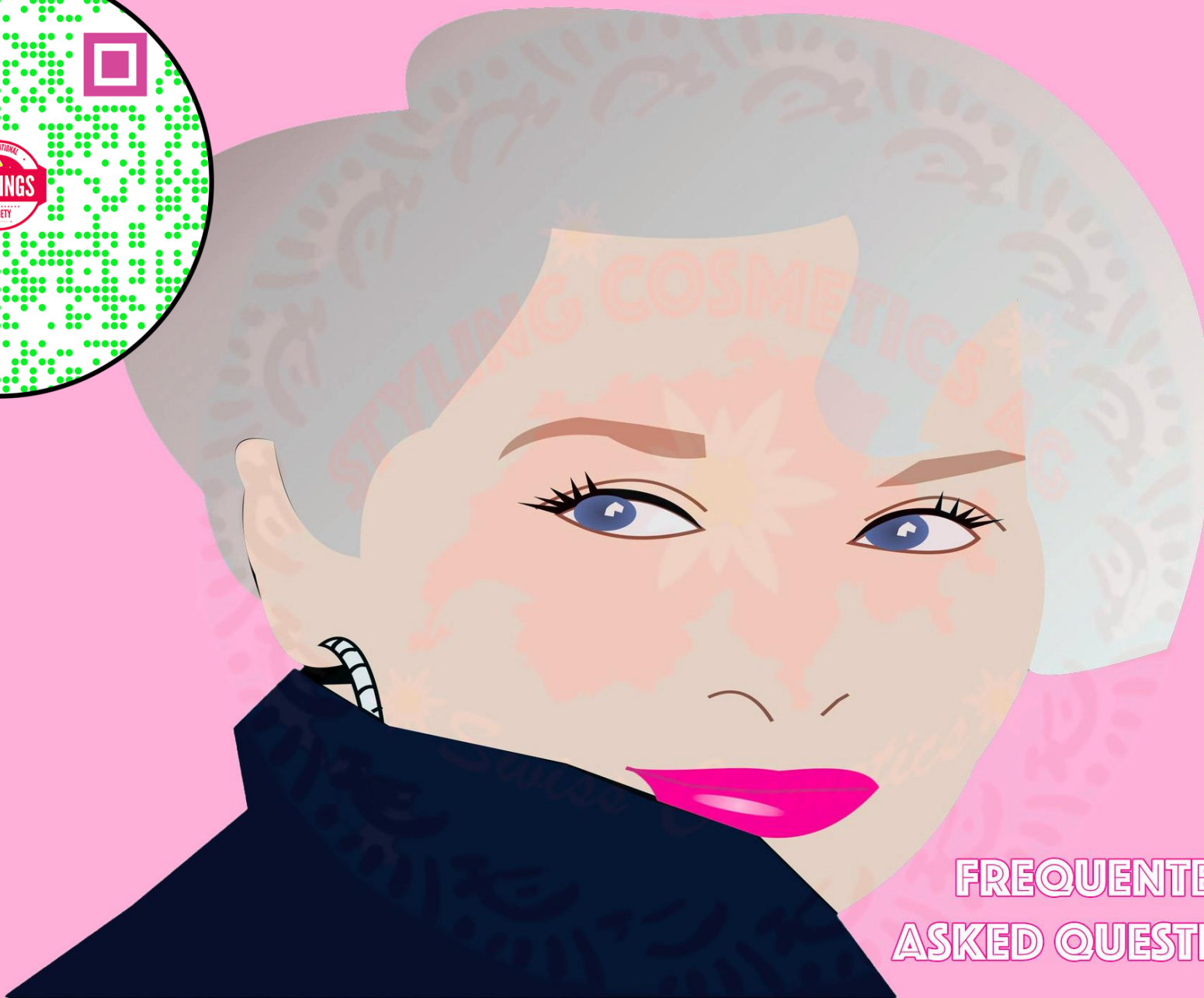
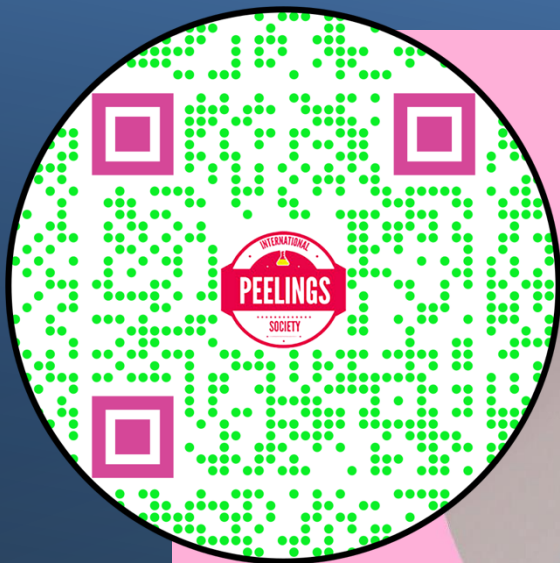
Acids	Mandelic	Glycolic	Lactic	Azelaic	Kojic	Phytic
Action	Promotes surface lifting of excess pigment	NO Dispersion of Melanin in basal membrane layer IRRELEVANT	Skin Lightening ??? Not bleaching agent IRRELEVANT	Skin Lightening	Tyrosinase Inhibitor	Melanin formation blocker though chelation ??? IRRELEVANT
pKa	3.37	3.83	3.86	4.55-5.59 DIPROTIC	9.40	HEXAPROTIC 1.1-3.2-5.2 8.0-9.2-12.0
Kerato regulator	YES	YES	YES	YES	NO	?
Moisturizer	NO	NO	NO	MINIMUM	NO	?
AHA						

SPECIAL RECOMMENDATIONS FOR WOMEN



- Beware of menstrual blood (contains iron and can therefore create a tattoo)
- That is why **Peeling de Luxe Plus** is used during the menstrual period.
- Contraindications: pregnancy, hormones (birth control pills) ,anal sex.
- Sex is possible when desquamation is over

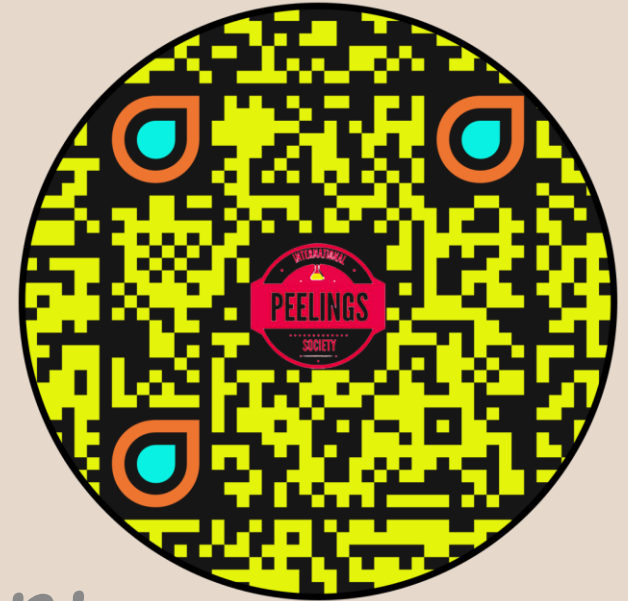




FREQUENTED
ASKED QUESTIONS

CLINICAL COMPETENCE REQUIRES MORE THAN KNOWLEDGE

- Academic training provides structure and understanding.
- Clinical competence emerges only through supervised application.
- For physicians wishing to go beyond theory, I offer **restricted-access hands-on clinical sessions**, independent from AIME.
- Participation is limited and subject to clinical prerequisites.



Further information:

<https://aesthetic.events/next-workshops>



NEXT WORKSHOPS IN ZÜRICH

MARCH 20-21, 2026 IN 1 MONTH !

PRACTICAL METABOLIC PEELS WORKSHOP
HANDS ON ONLY

ZÜRICH-SWITZERLAND :
MARCH 20 TH, 2026 : 1.30 PM-5 PM

WITH
MAURO TIZIANI
DR. ALAIN TENENBAUM

PEELINGS SOCIETY

SCAN ME

<https://aesthetic.events/workshops/next-workshops>

WORKSHOP

FULL DAY - FOUNDATIONS + TARGETED AREAS
ENDOPEEL FACE & BODY – COMPREHENSIVE PROGRAM

ZÜRICH-SWITZERLAND : MARCH 21 ST, 9 AM-5 PM

WITH
MAURO TIZIANI
DR. ALAIN TENENBAUM

SCAN ME

<https://aesthetic.events/workshops/next-workshops>





Peels COURSE

Its time to wake up



A.TENENBAUM, M.D.,Ph.D., D.Sc
M.TIZIANI, RCSA