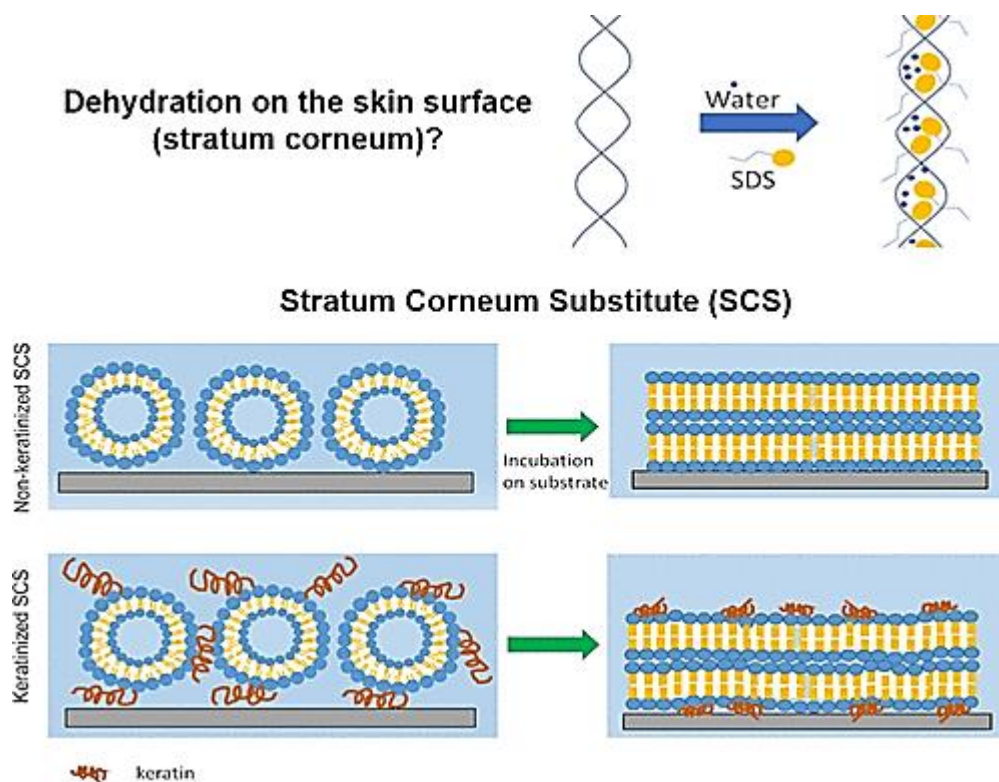


Effect of Surfactant–Keratin Hydrolysate Interactions on the Hydration Properties of a Stratum Corneum Substitute

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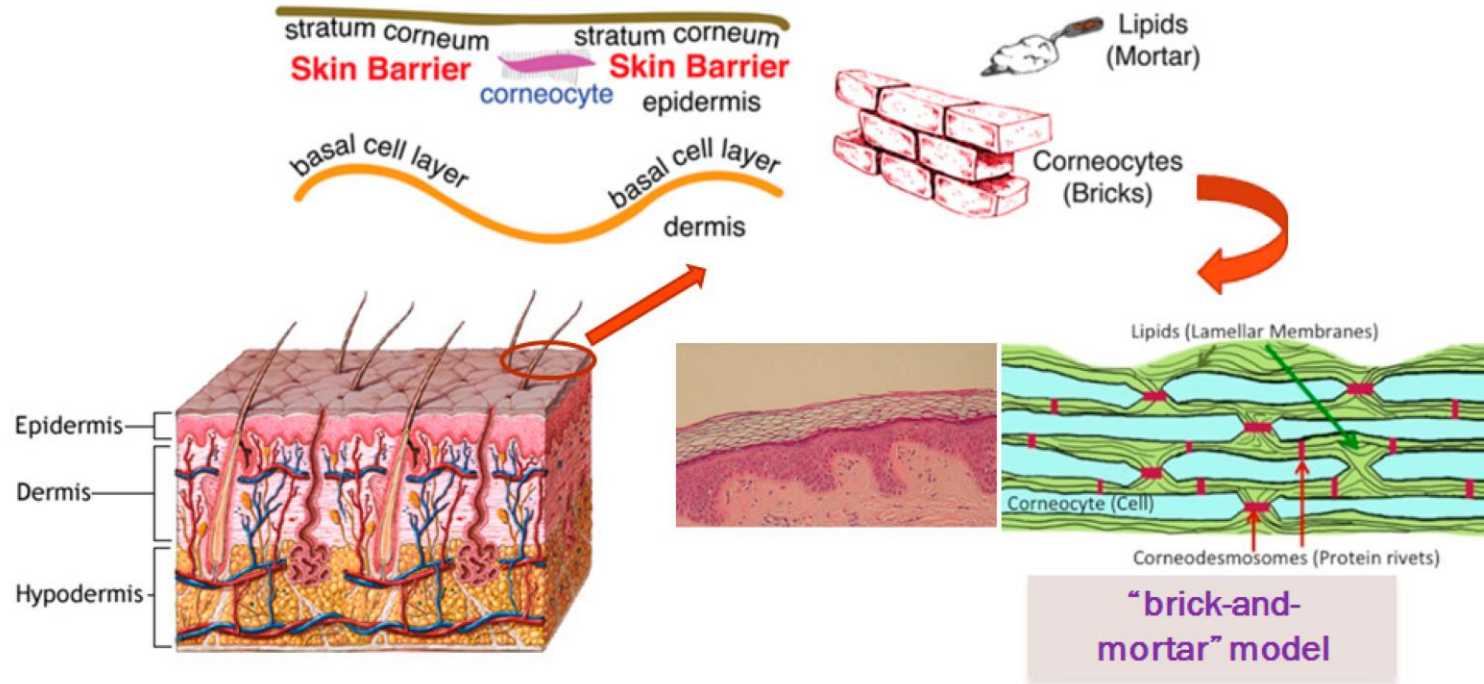


Tanvi Gupte

02.05.2020

Why this paper?

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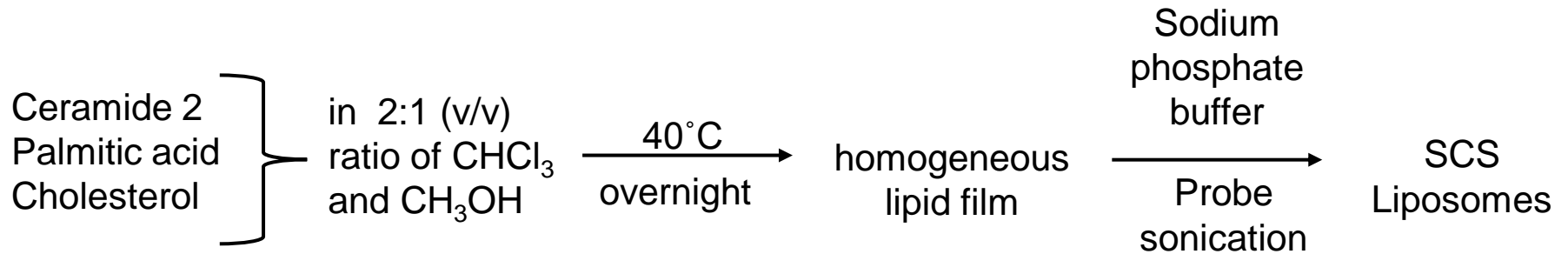
- Stratum corneum (SC): outermost skin layer, consisting of corneocytes and lipid bilayers.
- Corneocytes: keratinocytes cell organelles. Major constituents of corneocytes are keratin filaments.
- Stratum corneum substitute (SCS): skin model developed in this research.

Introduction

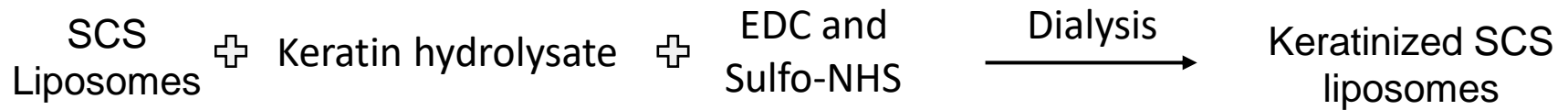
- A novel stratum corneum substitute (SCS) has been developed, to study the fundamental mechanism of the dehydration process.
- Changes in hydration of the SC are a suitable indicator to evaluate the irritation of surfactants on the skin.
- It is evident that both surfactant–protein and surfactant–lipid interactions can greatly influence the dehydration rate.
- **Significance of the paper:** an adequate skin model that helps in understanding the fundamentals of surfactant interactions with the SC at different time scales.

Synthesis of SCS

Liposomes



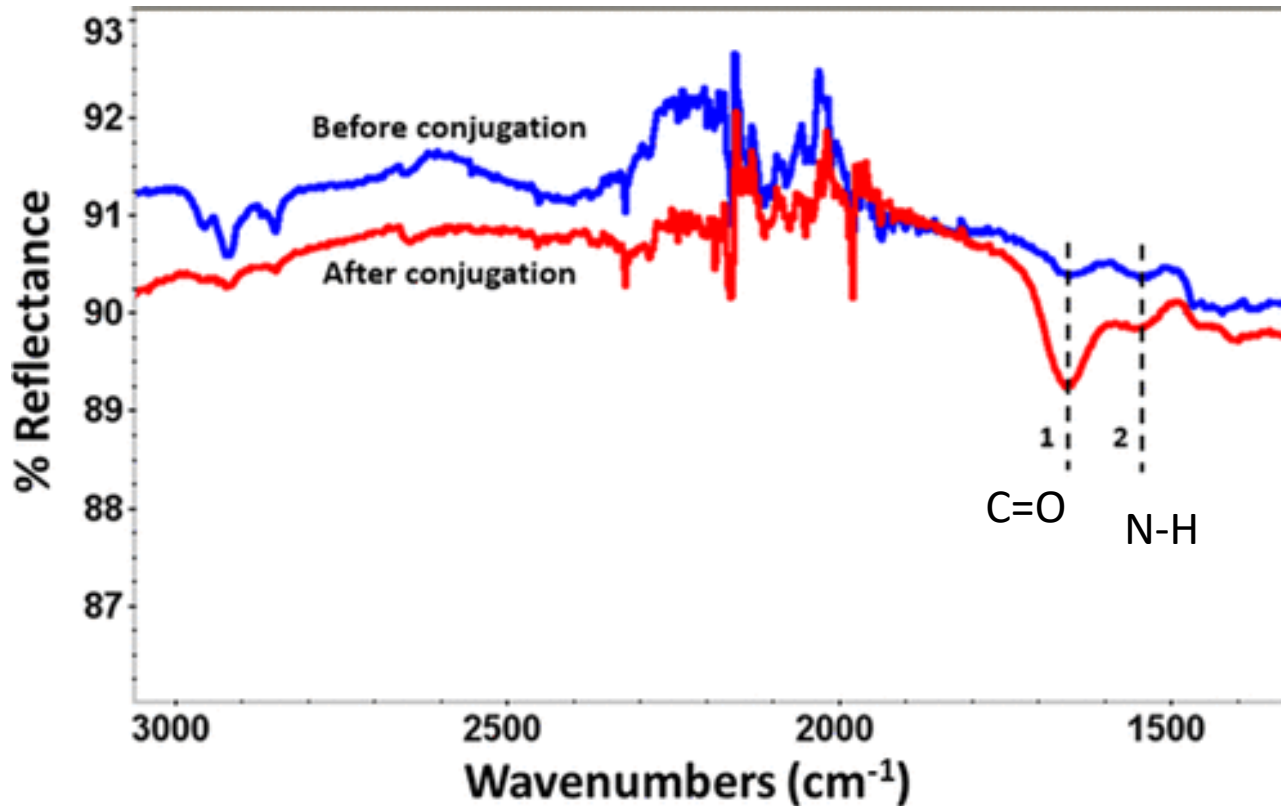
Keratinized SCS liposomes



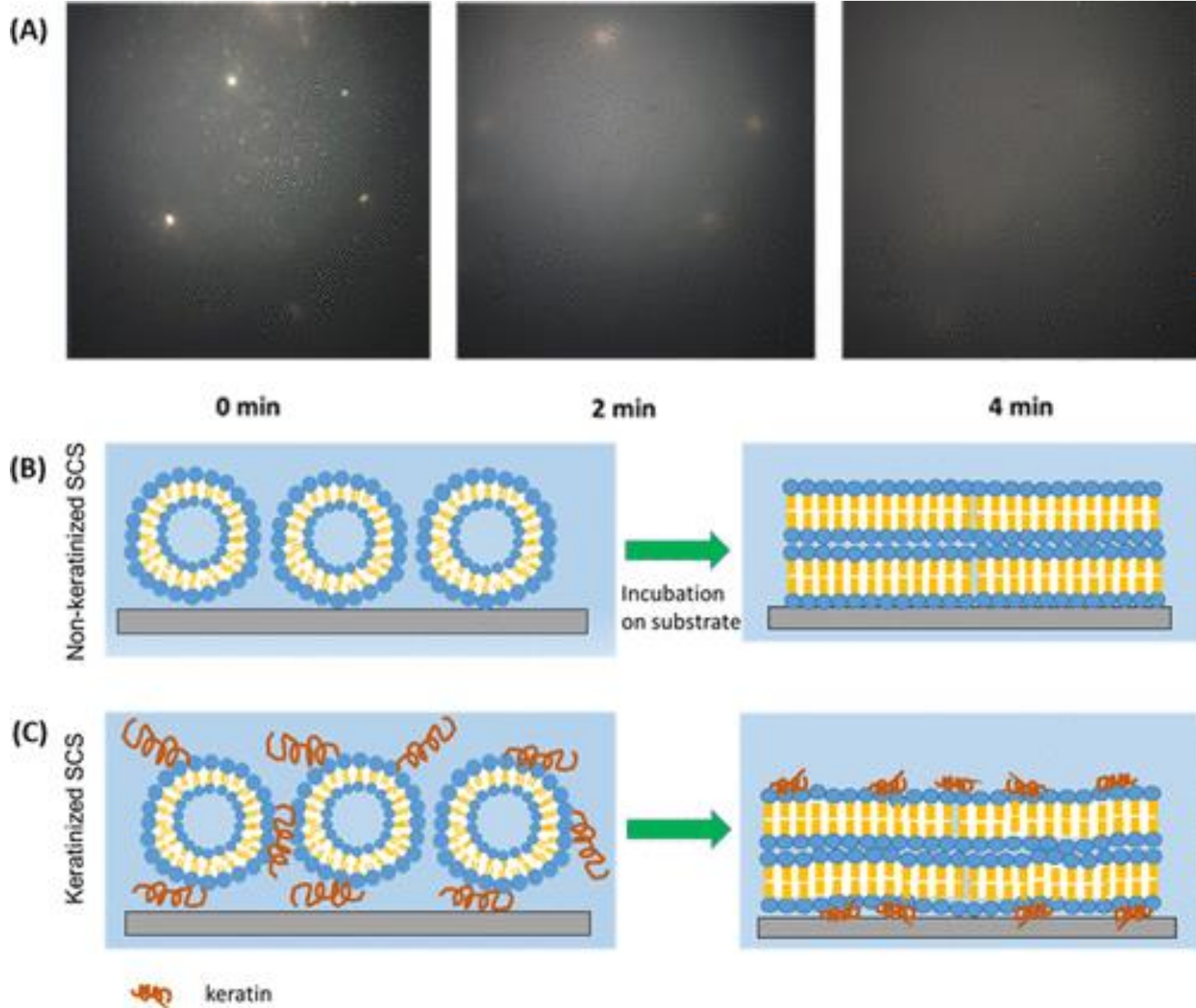
Non-keratinized/keratinized SCS lipid bilayer

Liposomes or Keratinized liposomes incubated on hydrophilic substrate

Characterization of Non-Keratinized and Keratinized SCS Liposomes.

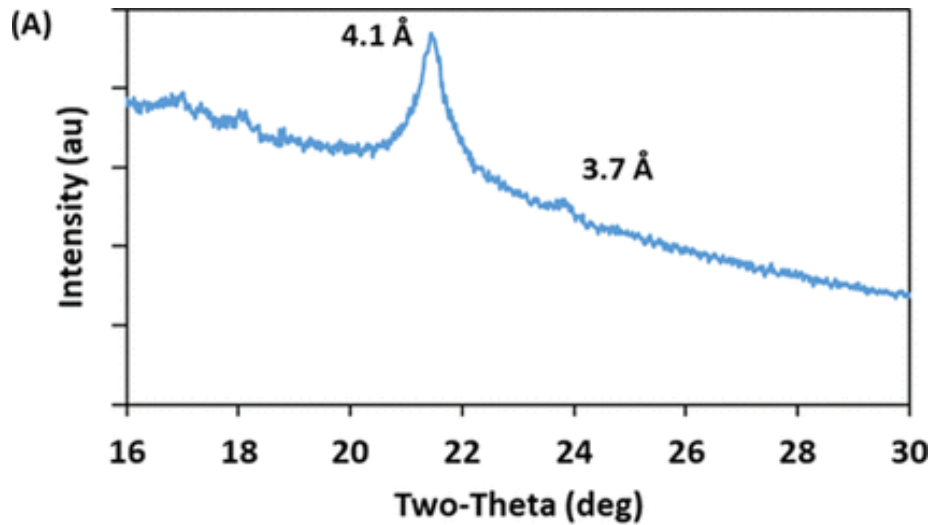


FT-IR data for non-keratinized SCS (blue) and keratinized SCS (red). The peak 1 near 1650 cm⁻¹ and peak 2 near 1547 cm⁻¹ correspond to amide bonds I and II, respectively.

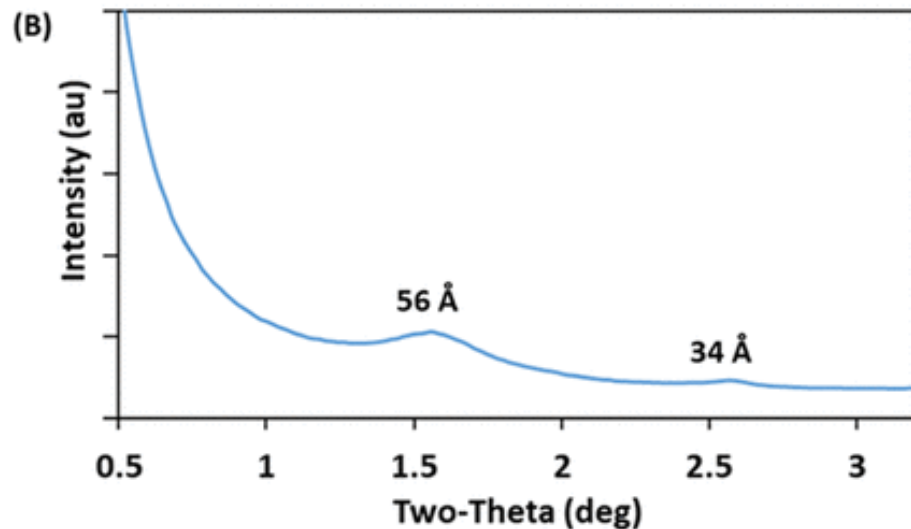


(A) Time-lapse dark-field images of SCS liposomes on a plasma-treated glass slide and schematics of the SCS layer formation for (B) non-keratinized SCS and (C) keratinized SCS on a hydrophilic substrate.

Characterization of Non-Keratinized and Keratinized SCS Lipid Bilayers



Lateral
packing

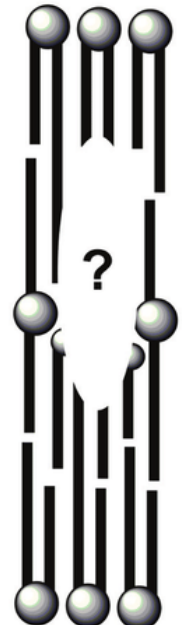


Lamellar
organization

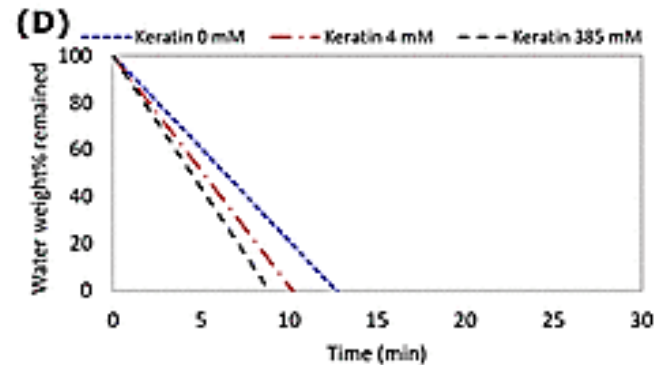
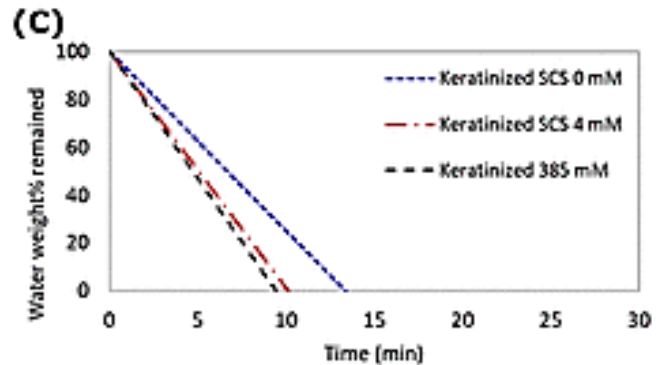
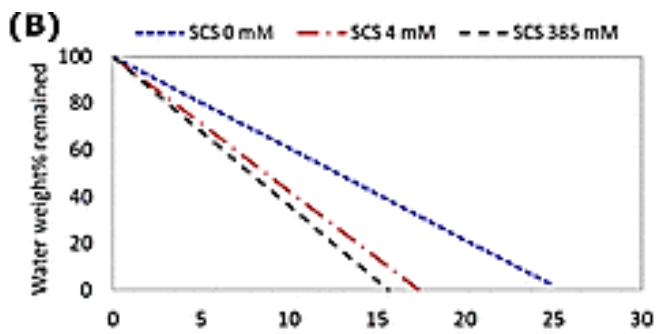
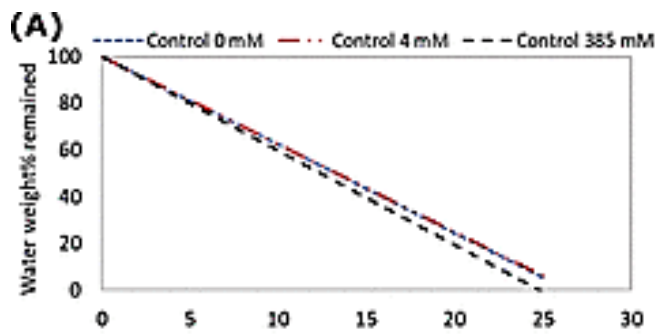
short periodicity
phase (5-6 nm)



long periodicity phase (12-13 nm)



(A) Lateral packing: WAXS of non-keratinized SCS. (B) Lamellar phase: SAXS of non-keratinized SCS.



Dehydration rates (% wwr vs time) for (A) control, (B) non-keratinized, (C) keratinized lipid bilayer, and (D) pure keratin hydrolysate samples and (E) summary of the slope

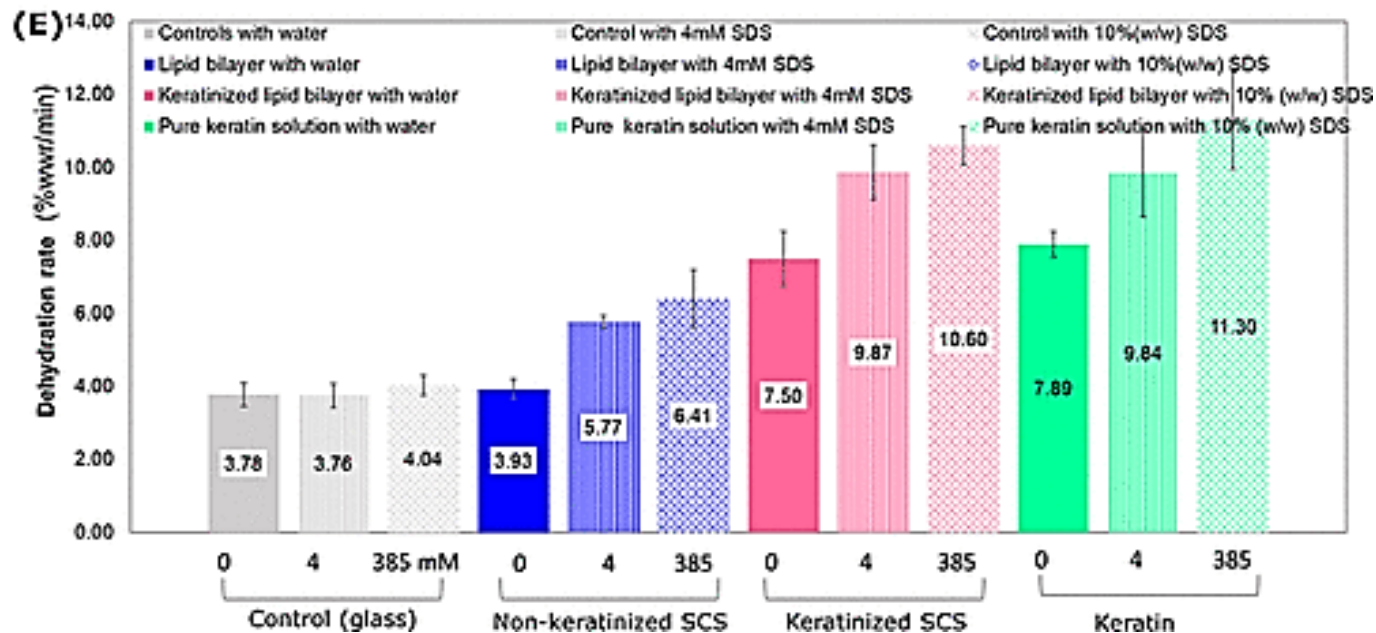
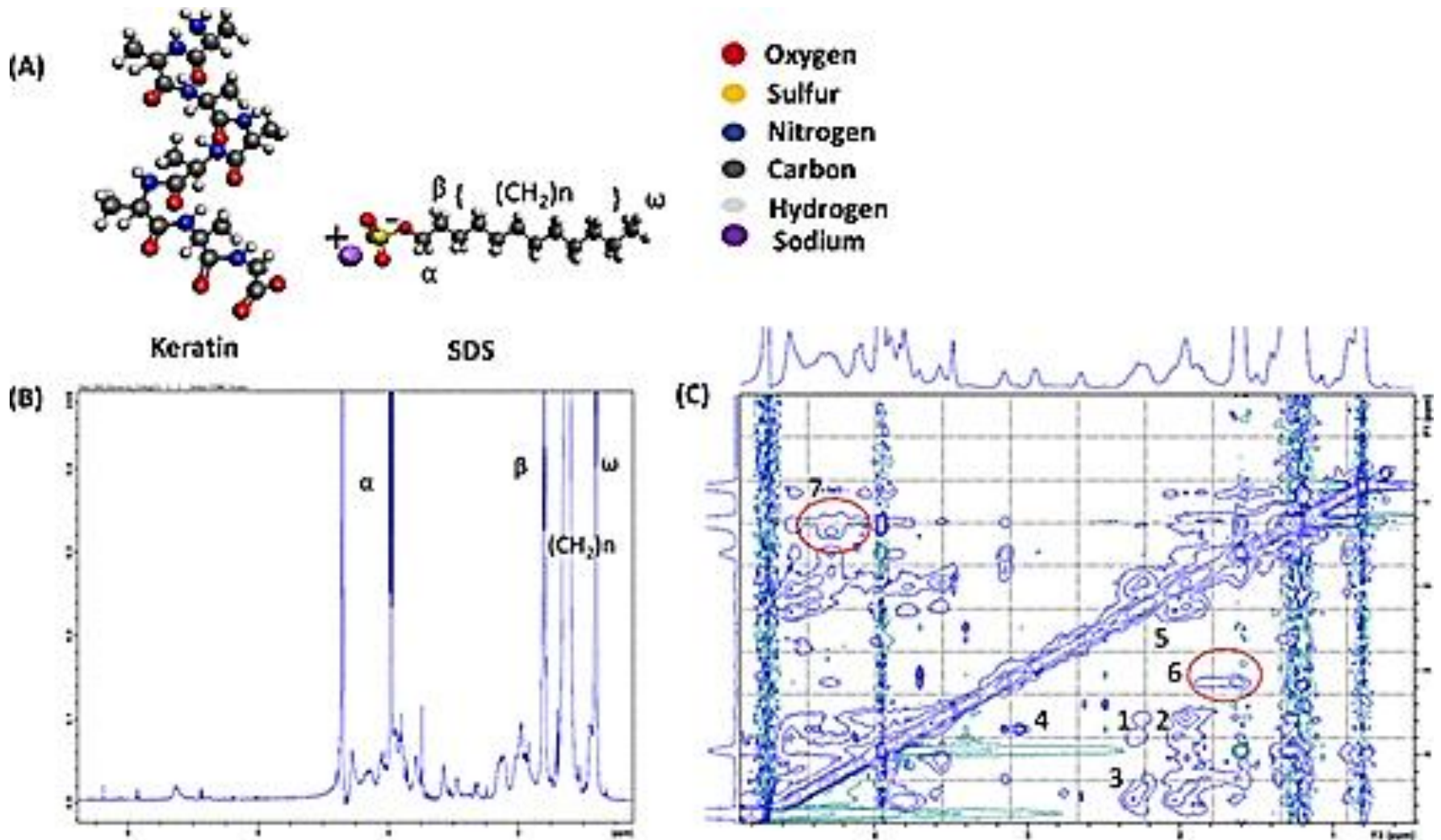


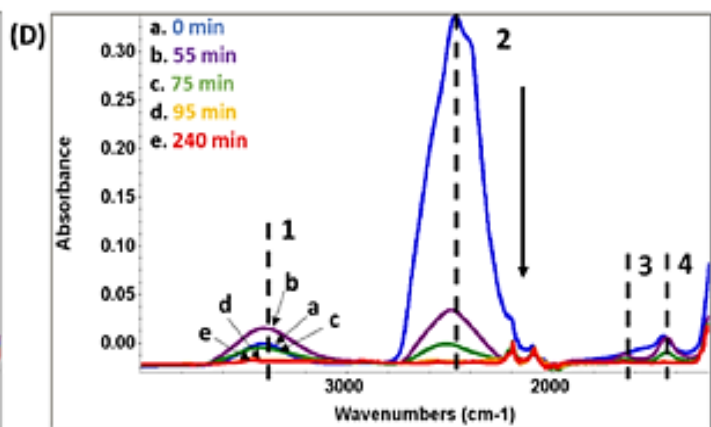
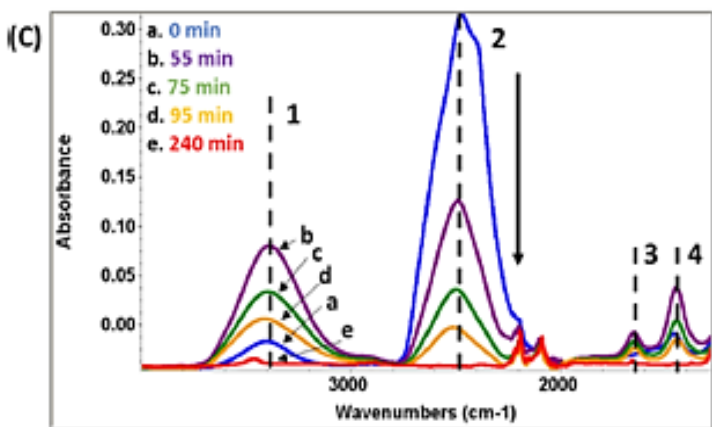
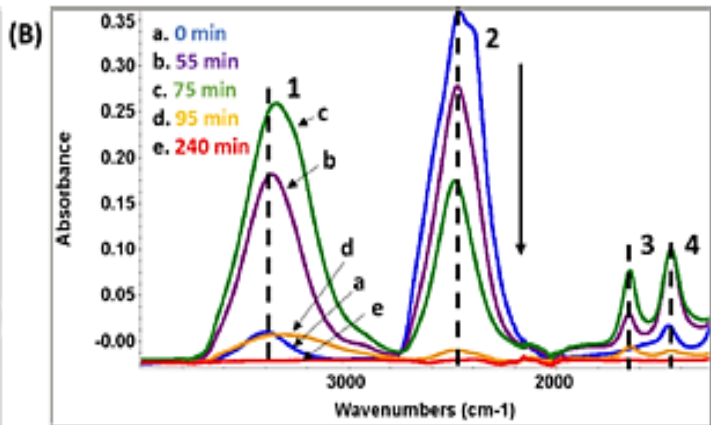
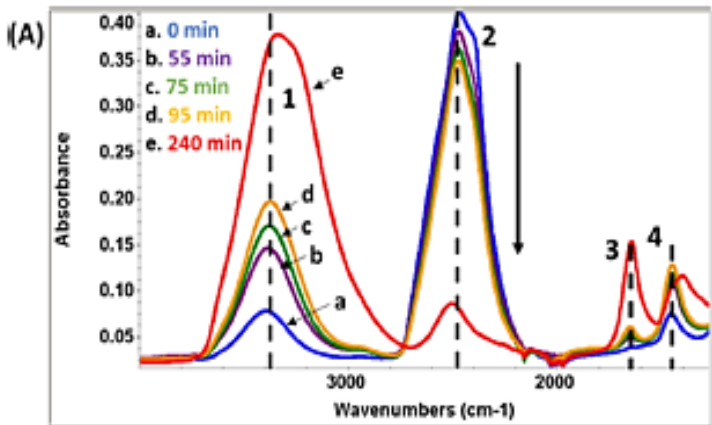
Table 2. Contact Angle Data of Non-Keratinized SCS and Keratinized SCS with and without 10 wt % SDS

	surfactant (SDS)	
	0 mM (deg)	385 mM (deg)
control (mica)	<1	17.8
non-keratinized SCS layers	12.9	19.2
keratinized SCS layers	22.3	32.4

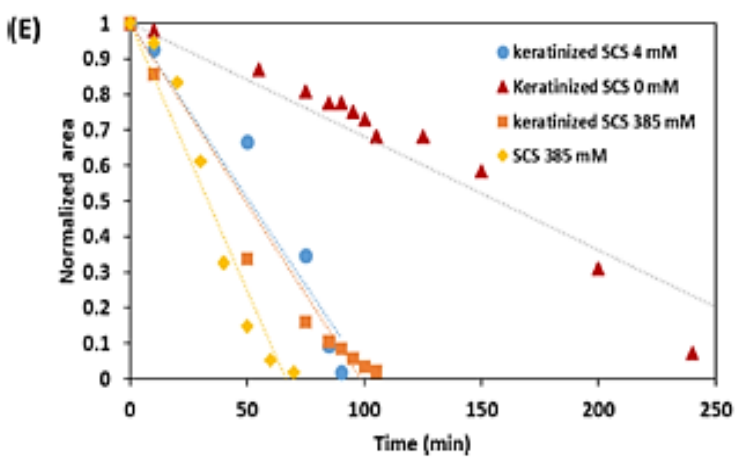
Molecular Interactions between SCS and the Surfactant.



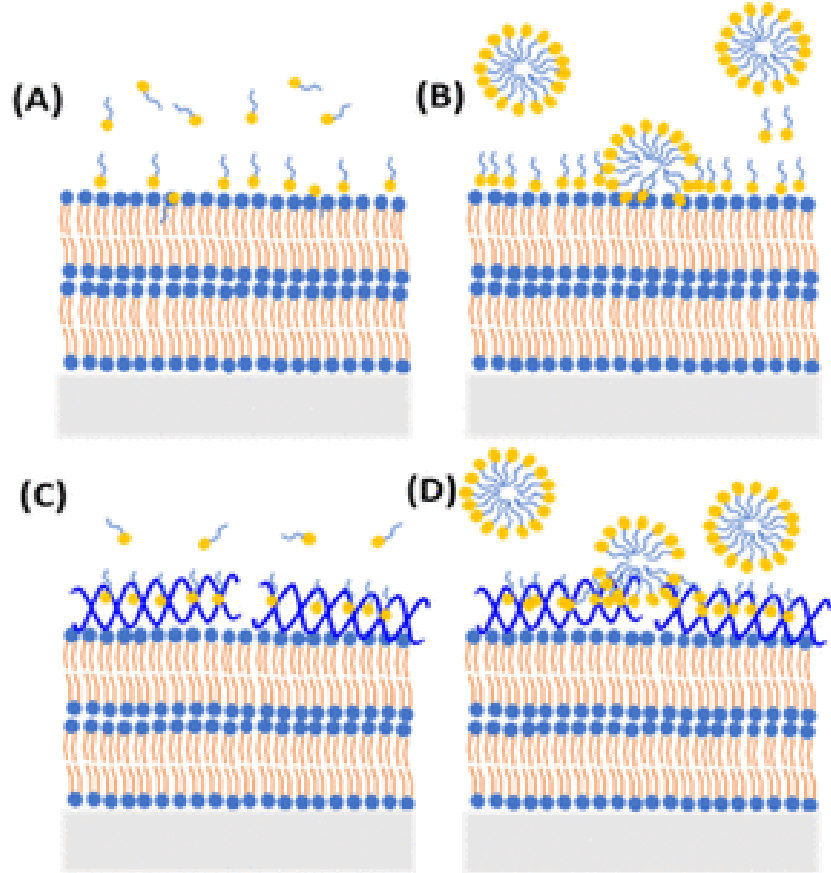
(A) Molecular structure, (B) 1H proton, and (C) 2D NOESY spectra of the SDS/keratin hydrolysate mixture.



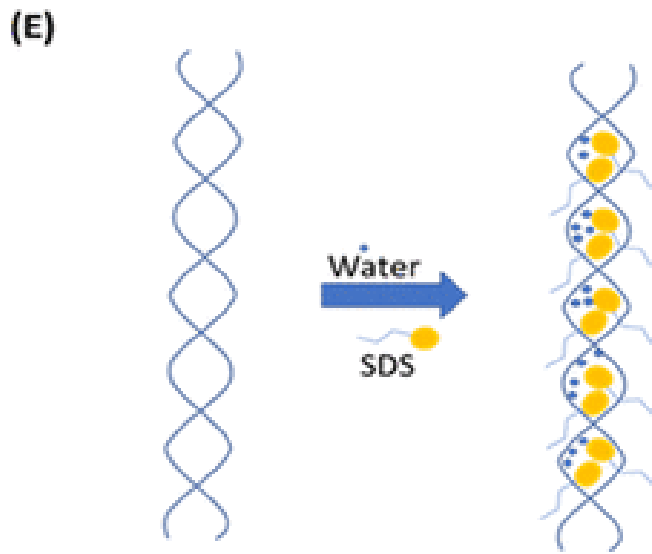
- (1) Hydroxyl (3401 cm^{-1}) peak from the SCS sample,
- (2) D_2O (2466 cm^{-1}) peak,
- (3) amide I (1648 cm^{-1}) peak, and
- (4) $-\text{CH}_3$ (1458 cm^{-1}) peak



ATR-FTIR spectral evolution during evaporation of (A) keratinized SCS and D_2O ; (B) keratinized SCS and 4 mM deuterated SDS (SDS-d25); (C) keratinized SCS and 385 mM SDS-d25; and (D) non-keratinized SCS and 385 mM SDS-d25, in D_2O . are denoted in all conditions. (E) Normalized area under the D_2O peak as a function of time.



Schematics of (A) non-keratinized SCS/4 mM SDS interaction, (B) non-keratinized SCS/385 mM SDS interaction, (C) keratinized SCS/4 mM SDS interaction, (D) keratinized SCS/385 mM SDS interaction, and (E) keratin hydrolysates and SDS interaction in an enlarged view.



Conclusion

- The keratinized SC model has shown the importance of keratin hydrolysates on regulating the dehydration process for both short term and long term.
- SDS interacts differently with keratin hydrolysates than with lipid bilayers.