Propylenated fatty acids as emulsifiers

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RESUMEN

Acidos grasos propilenados como emulsionantes.

Se prepararon estearato, palmitato, laurato, oleato y linoleato de hidroxipropilenilos mediante reacción de óxido de propileno con ácido graso a 160 °C durante cinco horas de agitación en presencia de hidróxido potásico como catalizador. Se determinaron las propiedades físico-químicas de los cinco productos, en cuanto a su uso como emulsionantes.

PALABRAS-CLAVE: *Acido graso propilenado - Emulsionante alimentario - Propiedades físicas - Propiedades químicas.*

SUMMARY

Propylenated fatty acids as emulsifiers.

Hydroxy propylenyl stearate, palmitate, laurate, oleate and linoleate were prepared by reaction of propylene oxide with fatty acid at 160 °C for five hours stirring in presence of potassium hydroxide as a catalyst. Physico-chemical properties of the five products, regarding their use as emulsifiers, were determined.

KEY-WORDS: *Chemical properties - Food emulsifier - Physical properties - Propylenated fatty acid.*

1. INTRODUCTION

In general, for a surfactant to act as an emulsifier, it must show good surface activity and produce a low interfacial tension in the particular system in which it is to be used. This means that it must have a tendency to migrate to the interface, rather than to remain dissolved in either one of the bulk phases (1). The emulsifiers were used to improve bread volume and texture, to extend shelf-life, and to control cookers spread factor (2). Various food products, such as butter, low fat spreads, shortening, chocolate, and margarine are water- in - oil (W/O) emulsions and require emulsifiers to prevent separation (3-5). The degree of solubility of surfactants in water often suggests approximate hydrophilic lipophilic balance (HLB) values are indicators of behavioral characteristics and do not indicate emulsifying efficiency. The selection of a suitable emulsifier for an O/W or a W/O emulsion must be determined by experimentation. However a

low HLB value (3-6) will promote or stabilize W/O emulsion, an intermediate HLB value (8-13) will stabilize O/W emulsion, and a high HLB value (15-18) will act as a solubilizer (6).

2. MATERIAL AND METHOD

- Hydroxy propylenyl fatty acids were prepared by adding propylene oxide slowly to fatty acid at 160 °C while stirring for five hours.
- pH was measured with a pH meter (Corian model SA 520).
- Surface tension was measured in distilled water at room temperature (15 °C) with a (Sc-Du Nouy Interfacial Tensiometer Model 70545) (7).
- HLB was determined as described in (9).
- Viscosity and shear stress were determined with Bohlin Visco 88 B v according to (10).
- Solubility was determined in different solvents, as water, ether, petroleum ether (40/60), n-Hexane, oil and ethanol.
- Emulsifying power was determined according to Titus (11).
- Infrared (IR) spectra were determined with Backman according to (12).

3. RESULTS AND DISCUSSION

The solubility of the mentioned emulsifiers in different solvents are recorded in table I where it can be seen that all emulsifiers are soluble in both oil and ethanol at 75 °C. Generally, hydroxy propenyl stearate and palmitate are insoluble with different degrees in all solvents at 25 °C while they are soluble in oil and ethanol only at 75 °C and insoluble in the rest of solvents at this temperature.

Water seems not to be the suitable solvent for solubility of these mentioned emulsifiers. The insolubility of these emulsifiers in water suggests a non polar nature, while the solubility of some emulsifiers in oil indicate that these emulsifiers can be blended with lipids during formation of food system without employing organic solvents (13). Figure 1 shows that pH values of the five emulsifiers are alkaline in character which agree with those of

sorbitol fatty acids reported by (14). Melting point, as shown in Figure 2, indicates that the value concerning hydroxy propylenyl stearate is more or less double of those of laurate and palmitate, while oleate and linoleate are liquid emulsifiers. Surface tension of the five emulsifiers are drawn in Figure 3. Those of stearate, linoleate and laurate are more or less the same as the value of hydroxy propylenyl palmitate while hydroxy propylenyl oleate shows the lowest value. Those results are in close agreement with sorbitol fatty acids reported by (14).

Table I
Solubility of some synthetic emulsifiers

Solvent	Temp. (°C)	Compounds				
		(1)	(2)	(3)	(4)	(5)
Water	25	I	SC	SC	SS	SS
	75	SS	C	C	PS	PS
Ether	25	SS	SS	S	S	S
	75	PS	PS	S	S	S
P. ether (40/60)	25	SS	PS	S	S	S
	75	PS	S	S	S	S
n-Hexane	25	SS	SS	S	S	S
	75	PS	PS	S	S	S
Cotton seed oil	25	I	I	S	S	S
	75	S	S	S	S	S
Ethanol	25	PS	PS	S	S	S
	75	S	S	S	S	S

(1) Hydroxy propylenyl stearate
(2) Hydroxy propylenyl palmitate
(3) Hydroxy propylenyl laureate
(4) Hydroxy propylenyl oleate
(5) Hydroxy propylenyl linoleate

I = Insoluble
PS = Partially soluble
SS = Springly soluble
S = Soluble

1 Hydroxy propylenyl stearate
2 Hydroxy propylenyl palmitate
3 Hydroxy propylenyl laureate
4 Hydroxy propylenyl oleate
5 Hydroxy propylenyl linoleate

Figure 1
pH values of emulsifiers

Figure 2
Melting points (°C) of emulsifiers
(See Figure 1 for emulsifier types)

Figure 3
Surface tension (dyn/cm²) of emulsifiers
(See Figure 1 for emulsifier types)

Viscosity of hydroxy propylenyl stearate is more or less double the value of those of linoleate and oleate. Hydroxy propylenyl laureate has the lowest viscosity value and that of palmitate is much higher than that of laurate (Figure 4). Shear stress of hydroxy propylenyl stearate is about three times of that of linoleate and oleate and about four times of that of laurate and palmitate as refere in Figure 5. Emulsifying power values are given in Figure 6, which shows that all emulsifiers tested have more or less the same value. On the other side the HLB range from 3.4 HLB to 6.2 HLB (Figure 7) so these emulsifiers promote or stabilize W/O emulsions (15,16). The relation between both shear stress to viscosity and emulsifying power to HLB are more or less parallel to each other as shown in Figures 8 and 9. From Figures (10-14), concerning the infrared spectra, it can be stated that:

- Broad band appears for stretching vibration of O-H group absorbed near $3600-3400\text{ cm}^{-1}$.
- Forked band for stretching vibration band for C-H group absorbed near $3000-2800\text{ cm}^{-1}$ for methylene and methyl groups. The symmetrical bending vibration for (CH_3) occurs near 1373 cm^{-1} and assymmetrical bending vibration near 1450 cm^{-1} .

- O-H bending vibration in region of $1420-1300\text{ cm}^{-1}$ in primary and secondary alcohol.
- H bending couple with C-H wagging vibration to produce band near 1420 and secondary 1330 cm^{-1} . C-O stretching vibration in alcohols

produces strong band in $1260-1000\text{ cm}^{-1}$. C-O stretching coupled with the adjacent C-C stretching vibration. This in primary alcohol gives better vibration. Stretching vibration of C = O of acid and ester absorbed near 1715 cm^{-1} .

Figure 4
Viscosity (mp) of emulsifiers
(See Figure 1 for emulsifier types)

Figure 7
Hydrophilic - lipophile balance (HLB) of emulsifiers
(See Figure 1 for emulsifier types)

Figure 5
Shear stress (Pa) of emulsifiers
(See Figure 1 for emulsifier types)

Figure 8
The relationship between the Shear Stress and Viscosity

Figure 6
Emulsifying power (%) of emulsifiers
(See Figure 1 for emulsifier types)

Figure 9
The relationship between Emulsifying Power Percentages and Hydrophilic Lipophilic Balance

Figure 10
Infra red spectrum of hydroxy propylenyl stearate

Figure 11
Infra red spectrum of hydroxy propylenyl palmitate

Figure 12
Infra red spectrum of hydroxy propylenyl laureate

Figure 13
Infra red spectrum of hydroxy propylenyl oleate

Figure 14
Infra red spectrum of hydroxy propylenyl linoleate

REFERENCES

1. Akoh, C.C. and Chigozie, V.N. (1992).—«Emulsification properties of polyesters and sucrose ester blends II: Alkyl glycoside polyesters».—*Journal of the American Oil Chemists' Society* **69** (1), 14-19.
2. Akoh, C.C. (1992).—«Emulsification properties of polyesters and sucrose ester blends I: Carbohydrate fatty acid polyesters».—*Journal of the American Oil Chemists' Society* **69** (1) 9-13.
3. Breyer, L.M. and Walker, C.E. (1983).—«Comparative effects of various sucrose-fatty acid esters upon bread and cookies».—*Journal of Food Science* **48**, 955-958.
4. Ebeler, S.E. and Walker, C.E. (1983).—«*Cereal Chem.*».—**60**, 270.
5. Pomeranz, Y. (1981).—«*Ibid.*».—**58**, 190.
6. Akoh, C.C. and Swanson, B.G. (1989).—«Synthesis and properties of alkyl glycoside and stachyose fatty acid polyesters».—*Journal of the American Oil Chemists' Society* **66**, 1295-1301.
7. Alexander, A.E. and Haytex, J.B. (1989).—«Physical Methods of Chemistry».—John Wiley and Sons, New York, 1971, pp. 501-655.
8. Lutton, E.S. (1950).—«Review of the polymorphism of saturated even glycerides».—*Journal of the American Oil Chemists' Society* **27**, 276-281.
9. Othmer, K. (1965).—«*Encyclopedia of Chemical Technol.*».
10. Tobias, J. and Tracy, P.H. (1950).—«Basic viscosity of ice cream mixes ice cream».—*Tr. J.* **46**, 183.
11. Titus, T.C., Wiancko, N.N., Barbour, H.F. and Mickle, J.B. (1968).—«Emulsifier efficiency in model systems of milk fat or soybean oil and water».—*Food Technology* **22**, 115-117.
12. Akoh, C.C. and Swanson, B.G. (1987).—«One-stage synthesis of raffinose fatty acid polyesters».—*Journal of Food Science* **52** (6) 1570-1576.
13. Feuge, R.O., Zeringue, H.J., Weiss, T.J. and Brown, M. (1970).—«Preparation of sucrose esters by interesterification».—*Journal of the American Oil Chemists' Society* **47**, 56-60.
14. El-Shattory, Y., Saadia, M. Aly and Megahed, M.G. (1995).—«Performance characteristics of some synthetic emulsifiers».—*SOFW Journal* **121** (10) 762.
15. Nawar, W.W. (1985).—«*Food Chemistry*».—Edited by O.R. Fennema, Marcel Dekker, Inc., New York, pp. 139-244.
16. Gupta, R.K., James, K.A. and Smith, F.J. (1983).—«Sucrose esters and sucrose ester/glyceride blends as emulsifiers».—*Journal of the American Oil Chemists' Society* **60**, 862-869.

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