common type non-ionics while on cotton the efficiency seems to be poorer at the lower concentrations but at use concentrations again to be comparable with other type non-ionics.

The sudsing characteristics of the wax acid nonionics were observed in launderometer tests as well as in many of the empirical shaking tests. In most instances the amount of suds formed was small and not too lasting.

## Conclusion

Non-ionic synthetic detergents having detergent properties similar to those of alkyl phenol polyethylene glycol ethers and alkyl mercapto polyethylene glycol ethers can be prepared by reacting the optimum quantity of ethylene oxide with the composite crude fatty acids made by the air oxidation of low melting chemical grade paraffin wax.

## Testing of Drying Oils. II. Evaluation of Natural and Synthetic Oils\*

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In a previous paper Bolley and Gallagher (1) have described in detail a method for the preliminary examination of drying oils. This method has been used by the National Lead Company's Research Laboratory for the evaluation of a large number of oils. It is the purpose of this paper to present some of the data obtained during the evaluation program.

Since this is a preliminary evaluation which is to be completed in a relatively short time, the scheme does not include exposure tests. The object is to characterize the oils to determine their probable application; final evaluation must await exposure tests. Accelerated exposure tests were not included since these have been found to be unreliable in many cases.

One test has been added to the evaluation program as previously described. This is the determination of flash point according to the standard A.S.T.M. Method D92-33 and was included when it was noticed that some synthetic oils had relatively low flash points. A comparison of the data presented in this paper with the previously published method will disclose that not all of the tests have been reported. Although all the tests were actually run on the oils, it seemed desirable to eliminate the results of lesser interest and significance in order to keep the already lengthy tables of data to a minimum.

The following oils, referred to in the abreviated form used in the tables, were tested:

G Bodied Linseed—An alkali refined, bleached, and refrigerated linseed oil heat bodied to a G viscosity.

Conjugated Linseed—A refined linseed oil catalytically treated to contain 8.1% conjugated linoleic isomer and 0.5% conjugated linolenic isomer.

Linseed Penta.—Technical pentaerythritol esterified with nearly equivalent amounts of linseed oil fatty acids.

Dehydrated Castor-Regular commercial unbodied dehydrated castor oil.

Fatty Tall Oil—Tall oil processed to remove a large amount of the rosin acids and the fatty acid concentrate esterified with technical pentaerythritol.

G Bodied Soybean—Refined soybean oil heat bodied to a G viscosity.

Conjugated Soybean—A refined soybean oil catalytically treated to contain 10.1% conjugated linoleic isomer and 1.2% conjugated linolenic isomer.

Soybean Penta.—Technical pentaerythritol esterified with nearly equivalent amounts of soybean oil fatty acids.

Soybean Mannitol—One mol. of mannitol esterified with four mols. of soybean oil fatty acids.

Soybean Sorbitol—One mol. of sorbitol esterified with four mols, of soybean oil fatty acids.

Q Bodied Linseed—An alkali refined, bleached, and refrigerated linseed oil heat bodied to a Q viscosity.

Linseed Polypenta.—Technical polypentaerythritol esterified with nearly equivalent amounts of linseed oil fatty acids.

Linseed Mannitol—One mol. of mannitol esterified with four mols. of linseed oil fatty acids.

Linseed Sorbitol—One mol. of sorbitol esterified with four mols. of linseed oil fatty acids.

TABLE I Chemical Constants

	Visc.	Color	Appear- ance	Odor	Acid Value	Sap. Value	Acetyl Value	Iodine Value	% Unsap.	% Ash	Ref. Index	Specific Gravity
G Bodied Linseed	G	6	Clear	Normal	2.1	189.0	5.4	165.0	1.43	0.000	1.4841	0.9430
	F	7	Clear	Bodied Oil	3.3	190.7	7.4	154.7	1.20	0.000	1.4838	0.9431
	I—	12—	Clear	Bodied Oil	4.0	180.9	28.4	155.6	1.99	0.000	1.4850	0.9524
	II+	6	Clear	Normal	8.5	198.3	1.6	137.3	0.96	0.000	1.4820	0.9377
	J—	10	Clear	Fatty	6.7	170.8	3.6	133.1	4.80	0.055	1.4861	0.9490
G Bodied Soybean	F	5+	Clear	Normal	2.2	189.6	5.0	119.0	1.15	0.000	1.4765	0.9358
	G	6	Clear	Normal	3.2	192.3	3.4	118.5	0.92	0.000	1.4771	0.9356
	H+	12	Clear	Normal	3.3	185.9	52.2	125.1	1.00	0.000	1.4796	0.9475
	G	7-	Clear	Normal	29.5	183.2	3.1	112.1	0.71	0.000	1.4774	0.9402
	F	6	Clear	Normal	16.5	185.1	7.8	119.6	0.80	0.020	1.4775	0.9463
Q Bodied Linseed	U QTPP	5	Clear	Bodied Oil	5.9	192.2	6.9	141.4	1.06	0.000	1.4858	0.9585
Linseed Polypenta		11+	Clear	Bodied Oil	1.6	181.2	45.3	154.3	1.54	0.006	1.4852	0.9592
Linseed Mannitol		10	Clear	Normal	31.3	179.7	7.3	131.8	2.05	0.000	1.4862	0.9654
Linseed Sorbitol		9	Clear	Bodied Oil	18.2	185.0	8.0	146.8	1.63	0.018	1.4857	0.9606
Soybean Polypenta		10	Clear	Bodied Oil	1.2	177.0	37.4	129.8	0.89	0.013	1.4819	0.9553
Soybean Maleic Glyc	Q	8	Clear	Normal	6.3	212.6	37.5	120.1	0.14	0.000	1.4800	0.9592
	S-	9	Clear	Normal	14.5	221.0	24.1	120.5	1.63	0.005	1.4806	0.9614
	Z2	6	Clear	Bodied Oil	6.6	194.0	4.8	123.8	1.54	0.000	1.4890	0.9664
	Z1+	7-	Clear	Acrid	7.9	211.8	23.4	138.0	1.05	0.014	1.4879	0.9763
	Z2+	7	Clear	Acrid	11.8	212.8	24.8	155.5	0.98	0.000	1.4872	0.9731

<sup>\*</sup> Presented at 22nd annual fall meeting, American Oil Chemists' Society, Nov. 15-17, 1948, New York City.

TABLE II
Oil and Oil Film Tests

	В	odying Tes	st	Dryi	ng Test	Sward		Resin Compati- bility		Reactivity (zinc oxide)		
	Minutes	at 307°C.	Color	Set to Touch	Dryness	Hard.	Point °F.	Urea Formal-	Mod.	Orig. Visc.	<u>'</u>	crease
	<b>Z</b> 2	Gel.	Z2	Hr.	48 Hr.			dehyde	Alkyd.	Sec.	24 Hr.	1 Week
G Bodied Linseed	72	253 199 120 111 420+	9— 8 13 8 15	3 % 2 % 2 ½ 1 % 4	9+ 9+ 9+ 9+ 9-	3 3 3 2 3	565 580 495 460	C C I	O C 1 O I	166.0 154.0 208.0 110.0 112.0	$ \begin{array}{c c} -11.5 \\ -2.6 \\ 3.9 \\ 60.0 \\ 28.3 \end{array} $	-22.9 $-26.5$ $-24.5$ $-60.0$ $-54.8$
G Bodied Soybean	$\frac{260}{120}$	480+- 450 195 275 348	11+ 11 14+ 18 18+	8 7 3¾ 4 3	8+ 8+ 9+ 8 9-	3 3 4 1 4	555 530 540 435 465	C C C	C C C I	145.0 190.0 207.0 34.0 38.0	$ \begin{array}{c c} -9.0 \\ 19.0 \\ -27.0 \\ 0 \\ 57.9 \end{array} $	$\begin{array}{c c} -33.8 \\ -29.5 \\ -45.4 \\ 0 \\ 1150.0 \end{array}$
Q Bodied Linseed. Linseed Polypenta. Linseed Mannitol Linseed Sorbitol. Soybean Polypenta.	102 48 38 45 73	221 83 120 120 123	9 13 14 18— 13	3 ½ 1 ¼ 3 2 ¼ 2 ¼	9 9 8 9 9	2 3 2 2 2 3	565 515 450 470 530	I I I I	C C I	69.6 49.4 104.0 153.0 60.0	13.5 - 1.8 70.0 Livered -10.8	44.7 - 7.9 70.0 Livered -14.0
Soybean Maleic Glyc	76 	257 130 126 58 16	11+ 13 	5 6 31/3 21/2 2	9 10— 9 10— 10—	4 5 2 2 4	480 535 530 520 470	C C C I I I I I I I I I I I I I I I I I	C C C C	87.0 55.5 110.0 52.0 109.0	$\begin{array}{c c} 77.0 \\ -3.1 \\ 81.0 \\ 104.0 \\ 292.0 \end{array}$	84.0 9.2 96.0 144.0 787.0

Soybean Polypenta.—Technical polypentaerythritol esterified with nearly equivalent amounts of soybean oil fatty acids.

Soy. Maleic Glyc.—A soybean oil adduct made with 5% maleic anhydride and esterified with a near equivalent amount of glycerine.

Soy. Maleic Penta.—A soybean oil adduct made with 5% maleic anhydride and esterified with a near equivalent amount of technical pentaerythritol.

Z2 Bodied Linseed—An alkali refined, bleached, and refrigerated linseed oil heat bodied to Z2 viscosity.

Lin. Maleic Glyc.—A linseed oil adduct made with 5% maleic anhydride and esterified with a near equivalent of glycerine.

Lin. Maleic Penta. -A linseed oil adduct made with 5% maleic anhydride and esterified with a near equivalent of technical pentaerythritol.

The bodied linseed, bodied soybean, and dehydrated easter oils are included for the purpose of providing comparison controls for the synthetic and modified oils. Only one soybean oil control was used because in most cases the synthetic oils based on soybean fatty acids have performance characteristics which approximate or surpass those of linseed oil.

The analytical constants of the various oils are listed in Table I. Upon examination it will be seen

that they merely serve to characterize the oils and do not predict performance characteristics. The first four tests-viscosity, color, appearance, and odor--are purely physical in nature and describe the outward appearance of the oils. The term "normal" as used to describe the odor of an oil refers to a faint, bland, fatty odor such as that of a good quality alkali refined linseed oil. With synthetic esters such as are being discussed in this paper, the acid value and acetyl value show the degree of completion attained in esterification. It will be noticed that the mannitol and sorbitol esters have an excess of acid while the pentaerythritol and polypentaerythritol esters and the maleic treated oils contain some unreacted hydroxyl groups. The maleic treated oils also display a characteristically high saponification value (over 210). Because a certain amount of heat bodying always occurs during esterification, the iodine value of a synthetic ester is not such a reliable indication of quality as it is with the natural glycerides. The unsaponifiable and ash content of Fatty Tall Oil is much higher than that of the other oils as might be expected. The last two constants, refractive index and specific grav-

TABLE III
Oil and Oil Film Tests

			Film So	lubility				ld Water esistance	Hot V	Vater Resistance	Alkali Resist.
i.	Wa	ster	Hexane	Acetone	Alcohol	Benzene	Time to	Time to	Time to		Time to
	Per Cent	Acid Value	Per Cent	Per Cent	Per Cent	Acid Value	Whiten Hr.	Fail Hr.	Whiten Min.	Failure	Fail Min.
G Bodied Linseed	9.4 11.2 10.8 16.7 14.9	193.9 212.1 164.0 178.8 128.0	20.2 21.8 18.1 22.4 24.4	44.7 64.6 34.4 80.4 75.0	48.8 71.2 37.2 86.5 88.4	104.2 94.0 116.4 132.0 100.4	140  18	140 24 • 24 16 42	35 9 8 5 1	Dull Broken blisters Dull, sl. white Film removed Broken blisters	32 11 8 9
G Bodied Soybean	15.9 18.4 10.2 16.7 12.8	118.4 144.2 182.0 207.8 211.8	31.1 34.2 23.8 28.0 30.1	97.7 98.2 56.2 94.0 91.0	98.2 98.7 60.8 95.5 92.3	89.7 93.7 111.9 120.5 103.3	16 	16 16 16 16 16	5 45 5 7	Soft, dull, sl. white White, dull Soft, dull, sl. white Film removed Film removed	56 100 34 9 14
Q Bodied Linseed	9.7 12.8 12.3 11.4 10.2	178.0 149.7 195.0 171.5 120.8	17.0 9.9 20.9 17.4 10.5	42.4 17.8 49.6 38.0 20.4	46.9 18.0 56.4 41.1 22.4	101.8 141.2 113.9 120.4 169.8	141 190 24 	142 190 40 16 190	36  10 7 	Dull Broken blisters Soft, white, dull White, dull, v. soft Broken blisters	34 2 5 11 2
Soybean Maleic Glyc	9.2 6.6 6.3 7.7 10.3	187.5 357.5 194.6 224.3 187.0	21.6 20.5 18.0 16.6 18.2	40.5 35.0 40.0 34.2 36.7	43.4 37.5 46.5 39.1 40.0	120.8 152.1 96.6 101.5 112.9	144 	16 Pass 192 144 40 40	7  40 40 40	Pass Pass Dull Dull Pass	14 17 35 9 5

TABLE IV
Bakelite BR 254 Varnishes

	Cooking Time	Cooking	Visc.	Acid	Kauri Red.	:	Drying Te	st	Sward Hard-	Cold Water Resist.	Hot Water Resistance	Alkali Resist.
	Min.	Loss %	VISC.	Value	Pass %	Set to Touch Min.	Dust Free Min.	Dry- ness 48 Hr.	ness	Time to Fail Hr.	Failure	Time to Fail. Hr.
G Bodied Linseed	135 86	10.0 10.0 6.1 9.2	E D D E	17.9 15.3 19.4 22.7	130 220 150 220	30 21 25 20	50 30 58 28	9+ 9 9 10	7 5 4 7	>168 >168 >192 >168 	Dull White,dull Pass Pass	46 19 114 66
G Bodied Soybean	285 150 195	13.1 12.6 6.1 15.0 15.0	D D D D D D D D D D D D D D D D D D D	13.2 13.6 15.1 22.0 17.0	140 170 230 140 140	30 25 20 23 28	50 35 65 60 45	9 9 8-+- 9+ 9	4 2 2 4 4	>192 190 >192 190 >192 , >192	Pass Soft, dull Pass Pass Pass	40 4 66 19 27
Q Bodied Linseed. Linseed Polypenta. Linseed Mannitol. Linseed Sorbitol. Soybean Polypenta.	31 120 95	8.2 3.7 12.6 9.7 4.4	D + D+ E+ D+	18.7 21.8 28.2 24.3 20.0	160 180 120 90 190	41 28 43 20 30	61 330 53 40 340	10- 10- 10 10- 9	4 7 11 14 3	>168   >192   190   >192   >192   >192	Pass Broken blisters Pass Soft, white, dull Broken blisters	288 19 40 288
Soybean Maleic Glyc Soybean Maleic Penta Z2 Bodied Linseed Linseed Maleic Glyc Linseed Maleic Penta	47 60	7.6 8.0 4.2	D D 	20.6 28.8 27.6	150 90 130 	25 75 253 	60 328 430 	10- 10- 7+ 	6 10 6 	196 >192 >168 	Soft, dull Pass Broken blisters	4 4 27 

ity, are of a purely physical nature and vary more with the viscosity of the oil than with its composition.

Table II lists some of the performance characteristics of the oils and oil films. An examination of the results of the bodying test shows that the polypentaerythritol esters are much faster bodying than the other synthetic esters. The pentaerythritol esters are next and the mannitol and sorbitol esters are the slowest bodying. Soybean Sorbitol is much slower bodying than Soybean Mannitol, but no such difference occurs with the corresponding linseed esters. The sorbitol and mannitol esters also produce much darker bodied oils than the other synthetic oils. The maleic treated oils which have been esterified with pentaerythritol are much faster bodying than those esterified with glycerine. Most of the oils have about the same flash and fire points, except for the sorbitol and mannitol esters which are much lower. The drying times of the synthetic esters follow the same order as the bodying rates and all are faster than the corresponding natural glycerides. The fastest drying time is shown by the polypentaerythritol esters, followed by pentaerythritol, sorbitol, and mannitol in that order. None of the synthetic oils dried a great deal harder than the natural glycerides and the mannitol and sorbitol esters were somewhat softer. Resin compatibility is not a particularly important test from a performance standpoint. It is intended only to discover any unusual compatibility characteristics of the oils which might be made the basis of some novel practical applications.

Oil and oil film properties are continued in Table III. The film solubility test is designed to give an insight into the film structure. The amount of low molecular weight, water soluble materials produced is approximately the same for all oils. Variations between the oils become apparent in the amounts of slightly higher molecular weight, hexane soluble products and become most apparent in the acetone and alcohol-benzene soluble fractions. The last two solvents appear to remove not only the decomposition products but also uncombined and loosely bound oil molecules. The amount of material which is soluble in acetone or alcohol-benzene varies directly with the

TABLE V
Amberol 801 Varnishes

						* (61.116114.)	· 					
	Cooking Time Min.	Cooking Loss	Visc.	Acid Value	Kauri Red. Pass	Set to Touch Min.	Drying Te	Dry- ness 48 Hr.	Sward Hard. 48 Hr.	Cold Water Resist. Time to Fail Hr.	Hot Water Resist. Failure	Alkali Resist. Time to Fail Min.
G Bodied Linseed	104 68 43	1.1 1.1 3.2 1.8	D F+ E+ F+ 	11.9 11.0 9.2 11.4	90 100 100 130	133 60 58 56	140 74 84 76	10 10 10 10 10	43 6 21 6 	>168 114 >120 190	Pass Dull Pass Soft, dull	40 20 32 15
G Bodied Soybean Conjugated Soybean Soybean Penta Soybean Mannitol Soybean Sorbitol	95 67 26	2.9 1.8 2.1 1.3 1.6	F F D D	11.7 10.0 8.8 12.3 14.4	190 140 90 70 70	61 63 75 75 75	87 81 105 105 105	9 9 9-1- 9+ 10-	6 2 12 17 17	>120 114 >112 112 112 112	Pass Soft, dull White, dull White, dull Dull	30 36 80 34 27
Q Bodied Linseed	* 75 69	1.1 * 4.2 4.3 *	D * D+ F+ *	11.9 * 16.2 13.4 *	90 * 80 40 *	133 * 62 70 *	140 * 77 100 *	10 * 10 10	43 * 17 18 *	>168 * 114 >112 *	Pass * Very dull White, dull	40 * 37 32 *
Soybean Maleic Glyc	0 58 25	3.2 1.3 1.1 1.3 3.9	D+ D- D F E	11.7 15.3 11.9 13.6 15.5	120 50 90 100 80	58 140 133 57 65	89 246 140 77 85	10 10— 10 10 10	8 10 43 9 12	>120 >192 >168 190 190	Pass Pass Pass Soft, dull Soft, dull	15 50 40 20 14

<sup>\*</sup> Incompatible.

TABLE VI Limed Rosin Varnishes

	Cooking	Cooking	•		Kauri	1	Drying Te	st	Sward	Cold Water Resist.	Hot Water	Alkali Resist.
!	Time Min.	Loss %	Visc.	Acid Value	Red. Pass %	Set to Touch Min.	Dust Free Min.	Dry- ness 48 Hr.	Hard. 48 Hr.	Time to Fail Hr.	Resistance Failure	Time to Fail Min.
G Bodied Linseed	105 75 30 75	6.1 6.9 2.9 6.8	D+ E D+ F	18.3 14.5 16.4 18.4	90 100 90 100	120 53 45 58	180 100 115 120	9+ 10- 10- 10- 	36 16 14 11	48 43 112 43	White, dull Broken blisters Blistered White, dull	25 6 62 12
G Bodied Soybean	195 75 115 115 125	8.4 6.1 5.0 11.8 13.7	E D E D+	16.8 15.3 13.7 17.1 18.7	200 120 130 90 110	56 47 75 70 75	216 100 105 100 105	8+ 9 10- 10- 10-	3 6 10 16 14	43 43 39 64 24	White, dull Film removed White, dull White, dull White, dull	7 4 41 10 13
Q Bodied Linseed	105 35 100 90 48	6.1 6.4 10.0 11.8 5.3	D+ D+ E+ F	18.3 16.6 27.2 22.8 15.4	90 90 80 60 100	120 46 52 85 87	180 338 72 115 348	9+ 10 10 10- 9	36 11 18 21 7	48 19 120 64 19	White, dull Broken blisters Dull White, dull Broken blisters	25 5 1 22 5
Soybean Maleic Glyc	67 105 *	11.6 6.1 * 8.2	D- D+ *	12.0 18.3 *	70 90 *	140 120 *	246 180 *	10 9+ *	13 36 *	>192 48 * 47	Broken blisters White, dull* Broken blisters	157 25 *

<sup>\*</sup> Incompatible.

functionality of the alcohol and the unsaturation present in the fatty acids and, with alcohol-benzene, ranges from a low of 18.0% for Linseed Polypentaerythritol to a high of 98.2% for G Bodied Soybean. It may be remarked that the hexahydric alcohols, mannitol and sorbitol, exhibit greater solubility than the tetrahydric pentaerythritol, but it must be remembered that mannitol and sorbitol both lose water with the formation of inner ethers at esterification temperatures, so that their actual functionality is usually less than four. In most cases the reactivity of the oils with zinc oxide was quite satisfactory, only the sorbitol esters and Linseed Maleic Pentaerythritol showed excessively large increases in viscosity. The polypentaerythritol esters showed the greatest resistance to cold and hot water, followed by pentaerythritol, mannitol, and sorbitol in that order. The soybean maleic oils had slightly better cold and hot water resistance than the linseed maleic oils. No

particular trend can be noticed in the results of the alkali resistance tests.

Table IV shows the characteristics of varnishes made from the various oils and a pure phenolic resin, Bakelite BR 254. The polypentaerythritol esters gave the fastest cooking times, followed by the pentaerythritol, sorbitol, and mannitol esters in that order. Soybean Maleic Pentaerythritol was faster cooking than Soybean Polypentaerythritol while Soybean Maleic Glycerine required about the same time as Soybean Pentaerythritol. No varnishes were made with the linseed maleic oils because these oils both had Z2 viscosities while the resin requires a lower viscosity oil for good results. Most of the oils had excellent Kauri reduction values, only two of them, Linseed Sorbitol and Soybean Maleic Pentaerythritol, failing as low as 100%. The various esters do not produce any great differences in the drying times of the varnishes made from them, but the varnishes made with sorbitol

TABLE VII

	Appli	cation	Consi	stency	Drying Odor	D	ry to Touc	h	Hardness			
	S. P.	М.Р.	1 Day	7 Days		Hours at 77°F.	Hours at	Min. at 230°F.	I Day		3 days	
	Z.2.		М. Р.	М. Р.	М.Р.	M.P.	М.Р.	М.Р.	S.P.	М.Р.	S.P.	M.P.
G Bodied Linseed	Good Good	Good V. good Good Good Good	183 195 219 260 204	180 198 220 260 208	Mod. Mod. Mod. Mod. Mod.	6 ½ 6 ¼ 4 ½ 3 ½ 8+	42 50 26 26 *	6 ½ 14 7 ½ 10 25	V. soft Soft Soft Firm Soft	Soft Firm Firm Hard Soft	Hard Soft Firm Hard Firm	Hard Hard V. hard V. hard Hard
G Bodied Soybean	V. good Good	Good Good Good Poor Fair	186 197 205 700+ 700+	184 207 212 700+ 700+	Mod. Mod. Mod. Mod. Mod.	8+ 8+ 4½ 7¼ 7¾	* 42 42 42	19 15 10 21 17	Soft V. soft V. soft V. soft Soft	Soft Soft Firm Firm Firm	Soft Soft Firm Soft Hard	Firm Firm Hard Hard Hard
Q Bodied Linseed. Linseed Polypenta. Linseed Mannitol. Linseed Sorbitol. Soybean Polypenta.	Good Fair Poor	Good Fair Poor Fair Fair	381 230 700+ 408 230	400 220 700+ 401 225	Slight Strong Mod. Slight Strong	6 3½ 4¾ 5¾ 3½	26 18 26 26 18	6 6 12 6½ 6	Soft Firm Soft V. soft Hard	Soft Firm Soft Soft Hard	Hard Hard Hard Firm Hard	Hard Hard Firm Hard V. hard
Soybean Maleic Glyc	Good V. good Good	Fair Fair V. good V. good V. good	310 275 205 288 248	339 302 215 309 268	Mod. Mod. Slight Mod. Mod.	5 1/4 5 1/2 4 1/2 3 1/4 2 3/4	50 42 42 42 26	6 ½ 7 7 4 3 ½	Soft Soft Soft Firm Firm	Firm Firm Hard Hard Hard	Firm Hard Firm Hard V. hard	Hard Hard Hard V. hard Hard

<sup>\*</sup> Didn't dry.

esters produced the hardest films, followed by mannitol, polypentaerythritol, and pentaerythritol. Both of the soybean maleic oils produced harder films than the soybean esters. The cold water and hot water resistances of the varnishes were generally good. Both of the polypentaerythritol ester varnishes failed in hot water resistance because of broken blisters but showed no sign of whitening. The alkali resistance of the polypentaerythritol ester varnishes was outstanding, followed by pentaerythritol, sorbitol, and mannitol in decreasing order. The soybean maleic varnishes had less alkali resistance than any of the varnishes made with soybean esters.

Table V shows the characteristics of varnishes made from various oils and a rosin-maleic resin, Amberol 801. All the oils were bodied to Z2 viscosity before cooking these varnishes. For this reason it was only necessary to prepare one varnish for the three bodied linseed oils. The maleic treated oils were the fastest cooking group. Most of the esters required approximately the same cooking times, the exceptions being the polypentaerythritols, which were incompatible, and Soybean Mannitol, which had a much shorter cooking time than the other esters. All of the varnishes made with synthetic oils dried rapidly and produced hard films. The pentaerythritol esters produced about the best results, but were not greatly superior to the other esters. Soybean Maleic Pentaerythritol produced a varnish that remained slightly tacky but was as hard as those made with the other soybean base synthetic oils. The soybean maleic oils and Linseed Pentaerythritol produced varnishes having excellent cold water and hot water resistances. The Soybean Pentaerythritol and Soybean Maleic Pentaerythritol varnishes had excellent alkali resistance. The other synthetic oils produced varnishes whose alkali resistances were not quite as good as that of the Z2 Bodied Linseed varnish.

Table VI shows the characteristics of varnishes made from the various oils and 4% Limed Rosin. All the oils were bodied to Z2 viscosity before cooking these varnishes and, as with the Amberol 801 varnishes, only one varnish was necessary for the various bodied linseed oils. Linseed Maleic Glycerine and Soybean Maleic Glycerine were incompatible

with Limed Rosin. Linseed Polypentaerythritol and Linseed Pentaerythritol are much faster cooking than the other linseed base synthetic oils, and Soybean Polypentaerythritol is much faster than the other soybean base oils. The varnishes made with the synthetic oils dried faster than those made with the natural oils and those based on soybean oil also dried harder than the varnish made with G Bodied Soybean. The polypentaerythritol esters produced varnishes having less cold water but more hot water resistance than those made with the natural oils, while the other synthetic oils tended to produce greater cold water and about equal hot water resistances. The greatest alkali resistances were produced by the varnishes made with the pentaerythritol esters and Soybean Maleic Pentaerythritol.

Table VII lists some of the properties of paints made with the oils. Two paints were made with each oil, one a single pigment type using zinc sulfide as the sole pigment and the other a mixed pigment paint. The application test, which includes appearance, brushing, leveling, and gloss records a master painter's opinion of the handling qualities of the paints. Most of the paints had fair to good consistencies but those of paints made with the sorbitol and mannitol esters are excessively high. The drying times of the paints corresponded to the drying times of the oils from which they were made. The paints made with polypentaerythritol esters were the fastest drying, followed by those made with pentaerythritol, sorbitol, and mannitol esters in that order. All of the paints made with synthetic oils dried faster than those made with the corresponding natural oils.

Table VIII lists some further characteristics of the paints. The results of the after yellowing, water permeability, and flexibility tests do not show any pronounced trends. In general, the films of paints made with the synthetic oils had less elongation than those of the paints made with natural oils. The two maleic glycerine oils seem to have a tendency to promote brittleness. All of the linseed base synthetic oils produced greater tensile strength than bodied linseed oil of the same viscosities. The two linseed maleic oils produced greater tensile strength in one formulation and less in another. In the mixed pig-

TABLE VIII
Paint Tests

	Water Permea- bility M. P.	Flexibil- ity M.P.	Elongation Stripped Film		Adhesion S. P.	Tensile Strength of of Stripped Film		Taber Abrasion M. P.	Shear Hard- ness M.P.	60° Gloss M.P.	Daylight Reflect. M.P.	After Yellowing M. P.
			S.P.	M.P.		S. P.	М. Р.	1				
G Bodied Linseed. Conjugated Linseed. Linseed Penta. Dehydrated Castor. Fatty Tall Oil.	0.57	28+	34.0	8.0	Fair	5,0	20.0	339	500	35.5	81.0	Considerable
	0.63	28+	34.0	6.0	V. poor	8.0	27.0	344	405	45.5	80.3	Slight
	0.59	28+	18.0	5.0	Poor	20,0	37.0	200	583	26.0	81.0	V. Slight
	0.64	28+	25.5	9.0	Poor	14.7	38.7	264	428	36.2	81.0	V. Slight
	0.54	28+	25.5	3.1	Fair	11.0	16.3	310	637	29.3	80.0	Slight
G Bodied Soybean	0.52	28+	84.0	8,0	Fair	0,0	9.0	724	234	30.5	81.7	Slight
Conjugated Soybean	0.76	28+	80.0	7,0	Fair	0,0	0.0	930	182	34.0	80.6	V. Slight
Soybean Penta	0.94	16.0	12.5	5,0	Fair	11.0	30.0	399	373	14.5	83.0	Slight
Soybean Mannitol	0.55	18.0	60.0	3,0	Fair	0.0	10.0	395	278	18.3	80.8	None
Soybean Sorbitol	0.53	19.0	39.0	3,0	Good	8.0	18.0	331	396	17.7	82.9	V. Slight
Q Bodied Linseed	0.57	28+	24.0	7.0	Good	13.0	32.0	254	145	69.0	80.2	V. Slight
	0.54	20.0	6.0	1.5	Good	33.3	58.7	215	625	27.0	82.3	Considerable
	0.50	28+	36.0	4.0	Fair	12.0	30.0	216	595	27.0	81.0	V. slight
	0.49	14.0	21.0	4.0	Good	18.0	48.0	199	524	32.5	78.8	V. slight
	0.56	25.0	10.5	1.5*	Good	25.5	36.9	227	645	27.5	83.3	Slight
Soybean Maleic Glyc	0.63	14.0	17.0	3.0*	V. poor	0.0	25.0	370	362	46.0	78.5	None
	0.67	28+	12.5	6.0	Poor	30.8	40.9	320	1000	30.4	81.2	V. Slight
	0.45	16.0	32.0	2.0	Good	19.0	29.0	236	536	13.0	81.5	None
	0.52	13.0	18.0	2.0	Poor	14.0	38.0	218	597	15.7	82.0	Slight
	0.53	12.0	16.0	0.0*	Poor	28.0	22.0	250	730	12.5	82.1	Slight

<sup>\*</sup> Brittle.

ment paints, the soybean synthetics produced greater tensile strength than G Bodied Soybean, while in the single pigment paints there was in several cases no difference. The paints made with Soybean Maleic Pentaerythritol and the two polypentaerythritol esters had tensile strengths which were much superior to those of the paints made with natural oils. Conjugated Soybean did not have any tensile strength in either formulation. Abrasion resistance was not determined on the single pigment paints because they were too soft. Abrasion resistance of paints made with the various oils showed only that most of the synthetics were better than the corresponding natural oils. Conjugated Soybean, however, had less abrasion resistance than G Bodied Soybean. In most cases the shear and Sward hardnesses of the paints made with synthetic oils were greater than those of the paints made with the natural oils. Soybean Maleic Pen-

taerythritol and the two polypentaerythritol esters produced particularly hard films. Gloss measurements on the paint films show no particular trends.

Daylight reflectance is a measure of the whiteness of the paints and also shows no particular trends.

In a following paper it is intended to present a method whereby the experimental data gathered under this evaluation scheme may be condensed and correlated for convenience in reference and comparison.

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## REFERENCE

1. Bolley, Don S., and Gallagher, E. C., J. Am. Oil Chem. Soc. 24, 146 (1947).



CHEMICAL TREATMENTS OF SEEDS TO PREVENT HEAT-ING AND DETERIORATION DURING STORAGE. A. M. Altschul (So. Reg. Res. Lab., New Orleans, La.). Cotton Gin & Oil Mill Press, Jan. 8, 1949. Forty-eight chemicals which inhibit heating and deterioration of seeds during storage are recorded. Mixtures of 2 chemicals were more efficient than either one used alone.

Solvent extraction of soybean oil by mixtures OF TRICHLOROETHYLENE AND ETHYLENE ALCOHOL. S. G. Measmer, O. R. Sweeney, and L. K. Arnold. Proc. Iowa Acad. Sci. 54, 189-97 (1947). A solvent consisting of denatured alcohol mixed with trichloroethylene to give a specific gravity of 0.910 can be used successfully in a continuous extraction plant to remove oil from soybeans. The extraction is carried out at 70° and the miscella cooled to cause it to separate into two phases. The lower phase can be separated and stripped to remove the solvent while the upper phase can be returned to the system without evaporation to extract more oil. The moisture content in the solvent will not build up if beans having less than 6% moisture are used and the drying is done with 10-lb. per square inch steam pressure on the drier. (Chem. Abs. 43, 1999.)

Pyrolysis of Palm oil waste. R. Francois. Oleagineux 3, 602-6(1948). The possibility of manufacture of coke, tar, gas, fertilizer, oil, etc. from palm oil cake and pulp is discussed from laboratory tests on the subject.

Alcohols by sodium reduction. A staff-industry COLLABORATIVE REPORT. M. L. Kastens and H. Peddicord. Ind. & Eng. Chem. 41, 438-46(1949).

Unsaturated long-chain aliphatic esters. Cata-LYTIC OXIDATION. T. M. Patrick and W. S. Emerson (Monsanto Chem. Co., Dayton, Ohio). Ind. Eng. Chem. 41, 636-41(1949). The liquid-phase air oxidation of the Me esters of tall oil fatty acids is described. In order to develop a technic for the method, oxidation of Et oleate and of oleic acid was studied in some detail. Oxidative scission was expected to take place at or near the double bonds to yield mono- and dicarboxylic acids. Large amounts of high molecular weight by-products, in addition to the expected scission products, were formed in these oxidations. The identified products comprised most of the members of the homologous series from caproic to palmitic acid and from suberic to undecanedioic acid, inclusive.

THE NITROGENOUS CONSTITUENTS OF THE TISSUE LIPIDES. I. THE EXTRACTION, PURIFICATION, AND HY-DROLYSIS OF TISSUE LIPIDES. J. M. McKibbin and W. E. Taylor (Syracuse Univ. College Med.). J. Biol. Chem. 178, 17-27(1949). A technic for the determination of total lipide nitrogen is described. The amounts of nitrogen, phosphorus, and choline present in lipide extracts of 9 tissues from several dogs are presented.

THE NITROGENOUS CONSTITUENTS OF THE TISSUE LIPIDES. II. THE DETERMINATION OF SPHINGOSINE IN TISSUE LIPIDE EXTRACTS. J. M. McKibbin and W. E. Taylor (Syracuse Univ.). J. Biol. Chem. 178, 29-35 (1949). A method is described for an approximate determination of sphingosine in whole lipide extracts which have been freed of non-lipide impurities. The method is based on a chloroform extraction of the aqueous lipide hydrolysate which is both specific and relatively quantitative for sphingosine. Sphingosine is then determined on the chloroform extract as nitrogen.

TESTING EMULSIONS OF VEGETABLE OILS BY MEANS of ultrasonic waves. A. Audouin and G. Levavasseur. Oleagineux 4, 95-100(1949). The authors treat the influence of the frequency of sound waves in the production and rupture of water-in-oil and oil-inwater emulsions.

Oxidation of esters of linoleic acid by oxygen. W. O. Lundberg (Hormel Inst., Austin, Minnesota). Oleagineux 4, 86-93(1949).

AN OXIDATION-REDUCTION CYCLE IN EMULSION POLY-MERIZATION SYSTEMS. F. T. Wall and T. J. Swoboda (Univ. Illinois). J. Am. Chem. Soc. 71, 919-24(1949).