TWO-YEAR STUDY ON THE INHERITANCE OF REDUCED SATURATED FATTY ACID CONTENT IN SUNFLOWER SEED

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SUMMARY

RS1 and RS2 are two genetic stocks that have low saturated fatty acid (SFA) contents. In this two-year study, we report on the inheritance of palmitic and stearic acids of RS1 and RS2 in the F_1 and F_2 generations of crosses with HA 382, an inbred line with a fatty acid composition similar to many commercial sunflower hybrids. The expression of the reduced palmitic and stearic acid traits is most important in the ${\rm F}_2$ generation because it is the ${\rm F}_2$ seed that is harvested by the farmer, sold, and crushed for the oil. Our experiments showed that the F_2 seed of reciprocal crosses of RS1 or RS2 with HA 382 was intermediate in SFA content between the two hybrid parents. The inheritance and expression of the reduced palmitic and stearic acid traits of RS1 and RS2 were controlled by more than one gene. In addition, there were significant relative changes in palmitic and stearic acid contents between the growing seasons of 2002 and 2003, suggesting that environment played a significant role in the relative proportions of these fatty acids. The total SFA content was also influenced by environment, producing opposite effects with RS1 (higher total SFA in 2003) and RS2 (lower total SFA in 2003). The data demonstrate that it will be necessary to incorporate the reduced SFA trait into both parents of a hybrid in order to achieve the lowest concentration of SFAs in the F_2 seed.

Key words: F₁ hybrid, *Helianthus annuus*, maternal effects, multigenic inheritance, palmitic acid, stearic acid

INTRODUCTION

High levels of saturated fat consumption are correlated with increased risk of coronary heart disease. Traditional sunflower oil has about 130 g kg⁻¹ saturated fatty acid (SFA) content, which is considered low compared with most vegetable oils. However, canola oil is lower, with about 70 g kg⁻¹ SFA, and remains a major

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competitor to sunflower oil in the marketplace. For sunflower oil to compete with canola oil and other vegetable oils with low SFA content, it is desirable to decrease its saturated fat concentration. To address this consumer preference, the USDA-ARS Sunflower Research Unit has recently released genetic stocks with reduced palmitic and stearic acids, the major SFAs of sunflower oil (Miller and Vick, 1999; Vick *et al.*, 2003a).

Two genetic stocks, RS1 and RS2, were released in 2001 (Vick *et al.*, 2003a). These stocks were derived from a cultivated sunflower line, PI 250542, collected in Egypt by Paul Knowles in 1958. RS1 has black seeds with gray stripes, while RS2 has light gray seeds that usually bleach to white in the sun. Both have a total SFA content (C16 to C24) of about 75 g kg⁻¹. In an earlier study (Vick *et al.*, 2002) we reported on the inheritance of reduced SFA content in crosses of RS1 or RS2 with HA 821, which typically has a total SFA content of about 115 g kg⁻¹. However, after analysis of the inheritance data, we discovered that HA 821 was naturally low in palmitic acid, similar to RS1 and RS2, making it difficult to draw conclusions about the segregation of reduced palmitic acid in the progeny of crosses with RS1 or RS2. Therefore, we conducted a second inheritance study using inbred line HA 382 to cross with RS1 or RS2. HA 382 is a high yielding, high oil inbred line released in 1992 (Miller and Gulya, 1994), which has a palmitic, stearic, oleic, and linoleic acid content similar to commercial hybrids and other sunflower inbred lines.

In this study we report the results obtained from two years of field study on the inheritance of reduced palmitic and stearic acids in crosses between HA 382 and RS1 or RS2. A comparison of the data from the two years shows similarities in inheritance patterns between years, but also some variations, apparently caused by environmental differences. Preliminary results of this study were reported at annual meetings of the U.S. National Sunflower Association's Sunflower Research Workshop (Vick *et al.*, 2003b; Vick *et al.*, 2004).

MATERIALS AND METHODS

Plant material. RS1 (PI 616494) and RS2 (PI 616495) were used as sources of reduced palmitic and stearic acids in crosses with HA 382 (PI 578871). HA 382 is an inbred line which has an SFA content (C16 to C24) of approximately 130 g kg⁻¹, that is typical of commercial sunflower. Seed was derived from reciprocal crosses of both RS1 and RS2 with HA 382. The F₁ seed of the four crosses [RS1/HA 382, HA 382/RS1, RS2/HA 382, and HA 382/RS2] was planted in the field in Fargo, ND, in 2002 and 2003, and the heads bagged for self-pollination to obtain F₂ seed. F₁ seed of the same four crosses was produced in the same field plot each year by emasculation and hand pollination. Heads of the parental lines, RS1, RS2, and HA 382, were bagged prior to flowering to obtain parental seed grown in the same environment as the F₁ and F₂ seed. The parental (N=50), F₁ (N=65-100), and F₂ (N=85-100) seed was harvested in September and was analyzed for fatty acid com-

position by gas chromatography. Seed was typically from one head, but in a few crosses with poor seed set, seed from as many as 13 heads was analyzed. Weather data, including sunflower growing degree days at the Fargo field plots, were obtained from the North Dakota Agricultural Weather Network (http://ndawn.ndsu.nodak.edu/application/sunflower-degree-days-form.html).

Fatty acid composition analysis. Parental, F_1 , and F_2 seeds were cut in half and the distal end (not containing the embryo) was placed in a 1.5-ml gas chromatograph autosampler vial. After crushing the half seed with a blunt instrument, 1.5 ml of a solution of hexane-chloroform-0.5 mol dm⁻³ sodium methoxide in methanol (75:20:5, v/v) (sodium methoxide solution purchased from Sigma, St. Louis, MO^{**}) was added to the vial and the vial capped. The sample was injected into a Hewlett-Packard 5890 gas chromatograph containing a DB-23 capillary column (30 m × 0.25 mm, J&W Scientific), which was held at 190EC for 5 min, then increased to 220EC at 10EC/min, held at 220EC for 1 min, then increased to 240EC at 20EC/ min, and finally held at 240EC for 1.5 min, for a total run time of 11.5 min. Fatty acid concentrations are expressed as percent by weight of the total fatty acids. The total SFA composition is expressed as weight of the C16 to C24 SFAs per kg of all fatty acids.

RESULTS

Inheritance of the reduced SFA trait in RS1

Parental lines (RS1 and HA 382). RS1 had significantly lower SFA content than HA 382 in the field experiments of both 2002 and 2003. RS1 had an average total SFA concentration of 73-74 g kg⁻¹ compared with 128-129 g kg⁻¹ for HA 382 (Table 1). However, the relative proportions of palmitic and stearic acid changed between years. For RS1, the palmitic acid concentration was higher in 2003 (45 g kg⁻¹) compared with 2002 (39 g kg⁻¹). Conversely, the stearic acid concentration was lower in 2003 (20 g kg⁻¹) than in 2002 (26 g kg⁻¹). Thus, the total concentration of SFA remained about the same for both years. For HA 382, the concentrations of both palmitic and stearic acids were relatively stable over years. The palmitic acid concentration was 63 g kg⁻¹ in 2003 compared with 69 g kg⁻¹ in 2002, while the stearic acid concentration was 50 g kg⁻¹ in 2003 and 46 g kg⁻¹ in 2002.

 F_1 seed. The F₁ seed from the reciprocal crosses of RS1 and HA 382 was intermediate between the parents for palmitic acid content in both 2002 and 2003 (Figures 1B, B' and 1C, C'). The cross RS1/HA 382 produced a similar distribution of palmitic and stearic acids in both years. However, in the reciprocal cross, HA 382/RS1, there was an apparent environmental effect on palmitic acid concentration. In this cross there was a shift from 45 g kg⁻¹ palmitic acid in 2002 to 57 g kg⁻¹ in 2003

^{**} Names of products are included for the benefit of the reader and do not imply endorsement or preferential treatment by the United States Department of Agriculture.

Table 1: Two-year study (2002 and 2003) on the fatty acid composition in the seed oil of RS1, HA 382, and the F_1 and F_2 progeny of reciprocal crosses of RS1 and HA 382	tudy (2002 prosses of F	Two-year study (2002 and 2003) on reciprocal crosses of RS1 and HA 382	on the fatty 382	acid compo	osition in th	le seed oil o	f RS1, HA 3	382, and th	e F ₁ and F ₂	progeny of
		Palmitic	Stearic	Oleic	Linoleic	Arachidic	Gondoic	Behenic	Lignoceric	Total
Sample	Year	16:0	18:0	18:1	18:2	20:0	20:1	22:0	24:0	Satd FA
		g kg ⁻¹	g kg ⁻¹	g kg ⁻¹	g kg ⁻¹	g kg ⁻¹	g kg ⁻¹	g kg ⁻¹	g kg ⁻¹	g kg ⁻¹
RS1†	2002	39±2	26±3	401±36	518±39	2.0 ± 0.2	2.1±0.1	5.9±2.7	1.9±0.3	74±4
RS1†	2003	45±2	20±3	317±29	596±30	1.9 ± 0.9	2.0±0.1	5.0±1.0	1.8 ± 0.5	73±5
HA 382†	2002	69 ± 4	46±8	234±30	626 ± 34	3.6 ± 0.5	1.7 ± 0.1	7.0±1.3	1.8 ± 0.8	128±8
HA 382†	2003	63±6	50±8	223±35	635±33	4.1 ± 0.6	1.8±0.1	8.8±1.4	3.3 ± 0.6	129±10
RS1/HA 382 (F ₁)†	2002	49±3	26±3	341±39	559±39	2.2±0.3	2.2±0.5	7.0±1.1	2.4 ± 0.4	87±4
RS1/HA 382 (F ₁)‡	2003	51 ± 4	24±3	303±39	600±36	2.3 ± 0.3	2.1 ± 0.2	6.5±0.8	2.7±0.4	86±11
HA 382/RS1 (F ₁)†	2002	45±2	47±5	489±28	397±29	3.7 ± 0.4	2.0±0.1	9.5±1.0	2.8±0.4	108 ± 6
HA 382/RS1 (F ₁)‡	2003	57±5	44±7	290±32	585±31	3.6 ± 0.6	1.8±0.2	8.2±1.5	2.8±0.7	113±6
RS1/HA 382 (F ₂)‡	2002	50±3	33±6	382±37	518±37	2.6 ± 0.4	1.9±0.2	7.2±1.0	1.4 ± 0.5	95±7
RS1/HA 382 (F ₂)‡	2003	56±4	36±8	232±27	648 ± 24	2.7 ± 0.5	1.8±0.2	6.9±0.9	2.2 ± 0.3	104 ± 10
HA 382/RS1 (F ₂)‡	2002	45±3	36±6	373±48	525 ± 49	2.6 ± 0.4	2.0 ± 0.2	8.4±1.2	2.3 ± 0.9	94±12
HA 382/RS1 (F ₂)‡	2003	60±4	26±7	275±27	616±27	2.4 ± 0.6	2.0±0.2	6.5±1.4	2.3±0.7	97±9
† N=50, ‡ N=100										

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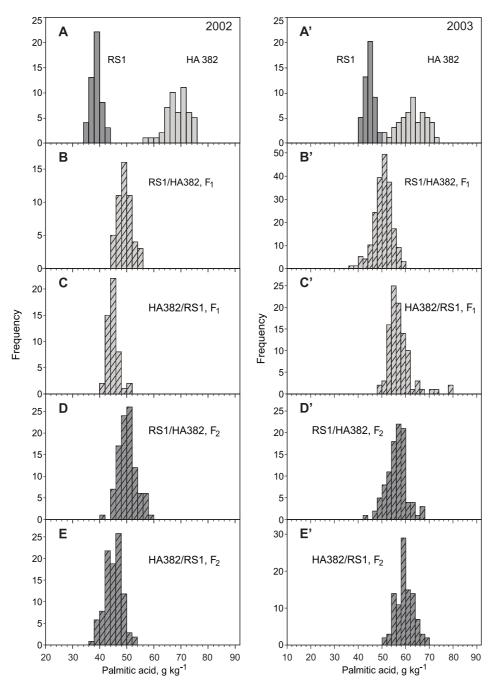


Figure 1: Frequency distribution of palmitic acid in the parents (A, A'), F_1 (B, B', C, C'), and F_2 (D, D', E, E') progeny of the reciprocal crosses between RS1 and HA 382 grown in the field in 2002 and 2003

(Table 1; and compare Figures 1C and 1C'). In addition, there appeared to be a maternal effect on total SFA content of the F_1 seed. During both 2002 and 2003, the cross HA 382/RS1 produced F_1 seed with 21 to 27 g kg⁻¹ higher SFA content than the RS1/HA 382 cross. The maternal effect was largely due to the consistently lower stearic acid in the RS1/HA 382 cross (Figures 2G, G' and 2H, H').

 F_2 seed. In both 2002 and 2003, the F_2 seed of both crosses was intermediate in total SFA content, compared with the parents (Table 1). However, the palmitic acid concentration tended to be higher in 2003 than 2002, especially in the HA 382/ RS1 cross (Table 1; Figure 1E, E'), suggesting that palmitic acid synthesis in this cross is influenced by environment more than in its reciprocal cross. The effect on total SFA in the HA 382/RS1 cross was tempered by a corresponding reduction in stearic acid content during the 2003 growing season (Table 1; Figure 2J, J'), so that the total SFA composition was similar in both years.

The palmitic acid content of the F_2 seed did not segregate over the range of the palmitic acid content of the two parents (Figures 1D, D' and 1E, E'). Instead, the F_2 seed had a somewhat narrow range of intermediate palmitic acid content. The stearic acid content of the F_2 seed segregated more broadly than did palmitic acid, and the maternal influence observed in the F_1 seed was less obvious in the F_2 generation (Figures 2I, I' and 2J, J'). For the RS1/HA 382 cross, the mean was intermediate between the parents in both years; however, the reciprocal cross HA 382/RS1 showed a reduced stearic acid content in 2003. This was probably an environmental effect, because the reduced stearic acid was not observed in the F_2 seed of this cross in 2002.

Inheritance of the reduced SFA trait in RS2

Parental lines (RS2 and HA 382). In both 2002 and 2003, RS2 was significantly lower in SFA content than HA 382. The average total SFA content of RS2 was 80-88 g kg⁻¹ compared with 127-129 g kg⁻¹ for HA 382 (Table 2). Both of the predominant SFA, palmitic and stearic, were lower in RS2 than in HA 382. But unlike RS1, the proportion of palmitic acid in RS2 did not increase in 2003; however, the concentration of stearic acid in RS2 decreased in 2003. Thus, the total concentration of SFA was slightly less in 2003, largely due to the lower stearic acid content. For HA 382, the concentrations of palmitic and stearic acids were similar between the years. The palmitic acid concentration was 63 g kg⁻¹ in 2003 compared with 69 g kg⁻¹ in 2002, while the stearic acid concentration was 50 g kg⁻¹ in 2003 and 46 g kg⁻¹ in 2002.

F₁ seed. In both 2002 and 2003, the total SFA content in the F₁ seed (85 and 102 g kg⁻¹, respectively) of the RS2 by HA 382 cross was intermediate between the parents, RS2 (80-88 g kg⁻¹) and HA 382 (127-129 g kg⁻¹) (Table 2). The reciprocal cross HA 382/RS2 produced F₁ seed with 106-112 g kg⁻¹ total SFA, also intermediate between RS2 and HA 382.

Table 2: Two-year study (2002 and 2003) on the fatty acid composition in the seed oil of RS2, HA 382, and the F_1 and F_2 progeny of reciprocal crosses of RS2 and HA 382	study (200 crosses of	Two-year study (2002 and 2003) on reciprocal crosses of RS2 and HA 382	on the faty 382	<i>y</i> acid comp	osition in tl	he seed oil o	of RS2, HA	382, and th	e F_1 and F_2	progeny of
		Palmitic	Stearic	Oleic	Linoleic	Arachidic	Gondoic	Behenic	Lignoceric	Total
sample	Year	16:0	18:0	18:1	18:2	20:0	20:1	22:0	24:0	Satd FA
		g kg ⁻¹	g kg ⁻¹	g kg ⁻¹	g kg ⁻¹	g kg ⁻¹	g kg ⁻¹	g kg ⁻¹	g kg ⁻¹	g kg ⁻¹
RS2 †	2002	45±3	32±5	429±56	477±58	2.4±0.2	1.8±0.1	7.3±0.7	1.8±0.2	88±6
RS2‡	2003	44 ± 4	26±4	340 ± 69	560±59	2.1±0.4	1.8±0.2	6.3 ± 1.2	2.0 ± 0.4	80±7
HA 382†	2002	69 ± 4	46±8	234±30	626±34	3.6±0.5	1.7 ± 0.1	7.0±1.3	1.8±0.8	127±8
HA 382†	2003	63 ± 6	50±8	223±35	635±33	4.1±0.6	1.8±0.1	8.8±1.4	3.3±0.6	129±10
RS2/HA 382 (F ₁)†	2002	45±3	29±3	437±43	468±43	2.2±0.2	1.9 ± 0.1	6.5 ± 0.9	1.8±0.3	85±3
RS2/HA 382 (F ₁) [§]	2003	62±6	29±4	285±36	602±33	2.5±0.4	1.8±0.2	5.9 ± 1.0	2.4 ± 0.5	102 ± 9
HA 382/RS2 (F ₁)†	2002	46±2	44±5	474±26	415±28	3.6±0.3	2.0±0.1	9.8±0.8	3.0 ± 0.3	106 ± 7
HA 382/RS2 (F ₁)‡	2003	55±8	42±7	270±35	606±31	3.5±0.6	1.9±0.8	8.7±1.5	3.3 ± 0.6	112±12
RS2/HA 382 (F ₂)‡	2002	47±5	45±8	450±72	439±73	3.3 ± 0.5	1.8±0.1	9.1 ± 1.5	1.5 ± 0.5	105 ± 9
RS2/HA 382 (F ₂)‡	2003	56±5	34±8	339±51	550 ± 53	2.6±0.5	1.9 ± 0.2	7.4±1.3	1.8 ± 0.5	102 ± 9
HA 382/RS2 (F ₂)‡	2002	49±3	43±7	410 ± 42	478±43	3.3 ± 0.5	1.9 ± 0.8	9.6±1.5	2.4±0.4	107 ± 8
HA 382/RS2 (F ₂)*	2003	50±4	37±6	370 ± 59	519±59	3.0±0.7	1.9±0.2	7.9±1.3	2.2±0.5	100 ± 2
† N=50 ± N=100										
[§] N=65										
* N=85										

HELIA, 27, Nr. 41, p.p. 25-40, (2004)

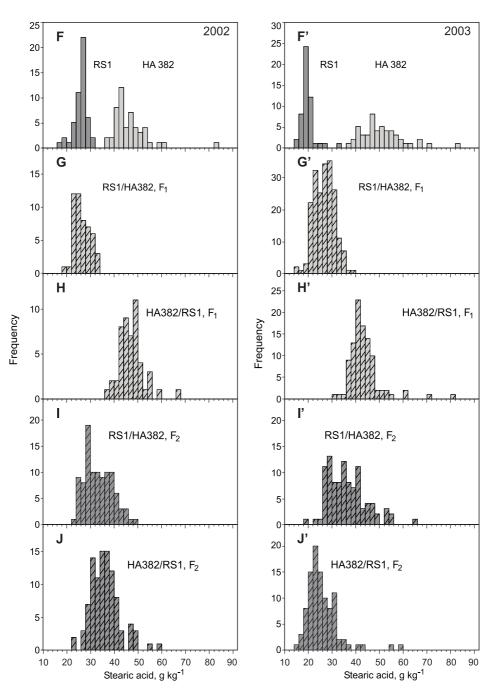


Figure 2: Frequency distribution of stearic acid in the parents (F, F'), F_1 (G, G', H, H'), and F_2 (I, I', J, J') progeny of the reciprocal crosses between RS1 and HA 382 grown in the field in 2002 and 2003

The distribution of palmitic and stearic acids in the F_1 seed from the reciprocal crosses of RS2 and HA 382 in the field in 2003 was different than that observed in 2002, suggesting that environment affected SFA synthesis. Whereas in 2002, the reciprocal crosses between RS2 and HA 382 both resulted in low palmitic acid phenotypes, in 2003 the palmitic acid concentration in the F_1 seed of the reciprocal crosses was increased. In 2003, the RS2/HA 382 cross had a higher palmitic acid content, similar to HA 382 (Table 2, Figure 3B'), while the reciprocal cross had a palmitic acid content intermediate between the two parents (Figure 3C').

The distribution of stearic acid was similar for both years (Figures 4G, G' and 4H, H'), with an apparent maternal effect for both crosses. As in 2002, the RS2/HA 382 cross resulted in a reduced stearic acid content similar to the RS2 female parent, whereas the reciprocal cross HA 382/RS2 was intermediate between the parents.

 F_2 seed. In both 2002 and 2003, the F_2 seed of both crosses was intermediate in total SFA content compared with the parents (Table 2). The palmitic acid concentration tended to be higher in 2003 (Figures D', E') than in 2002 (Figures 3D, E), an effect that was also observed in the F_2 seed of reciprocal crosses of RS1 with HA 382 (compare Figures 1D', E' with Figures 1D, E). However, the total SFA content was similar in both years because the increased palmitic acid in 2003 was balanced by a corresponding reduction in stearic acid content during the 2003 growing season.

The palmitic acid content of the F_2 seed did not segregate over the range of the parents (Figures 3D, D' and 3E, E'). Instead, the F_2 seed had a somewhat narrow range of intermediate palmitic acid content. The stearic acid content of the F_2 seed segregated more broadly than did palmitic acid (Figures 4I, I' and 4J, J'), and the maternal influence observed in the F_1 seed was less pronounced in the F_2 generation. The mean stearic acid concentration was intermediate between the parents. We did not observe the apparent environmental effect of reduced stearic acid in the F_2 seed of HA 382/RS2 that was observed in the HA 382/RS1 cross (compare Figure 4J' with Figure 2J').

DISCUSSION

The inheritance of SFA content has been studied in a number of mutant sunflower and soybean lines in which the SFA composition has been altered. Erickson *et al.* (1988) and Fehr *et al.* (1991) each developed a mutant soybean line in which the palmitic acid content was reduced from about 100-120 g kg⁻¹, typical for commercial cultivars, to about 70 g kg⁻¹. The F₂ plants of a cross between these two low-palmitic soybean lines segregated to produce some progeny that were further reduced in palmitic acid content to about 44 g kg⁻¹, indicating that two alleles at independent loci, acting additively, were responsible for the low-palmitic trait. Sim-

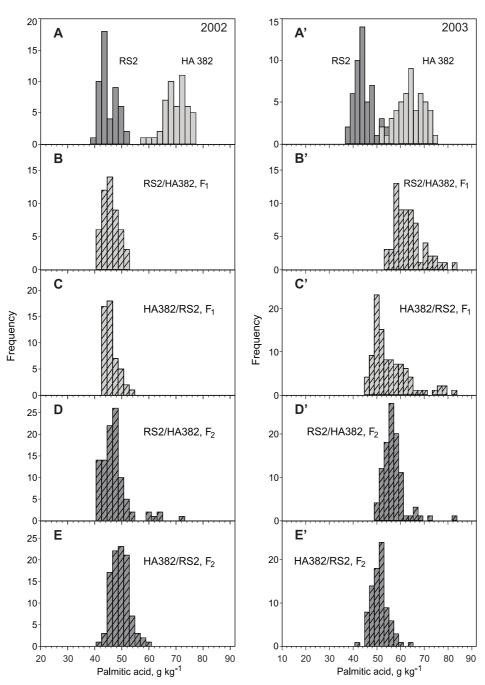


Figure 3: Frequency distribution of palmitic acid in the parents (A, A'), F_1 (B, B', C, C'), and F_2 (D, D', E, E') progeny of the reciprocal crosses between RS2 and HA 382 grown in the field in 2002 and 2003

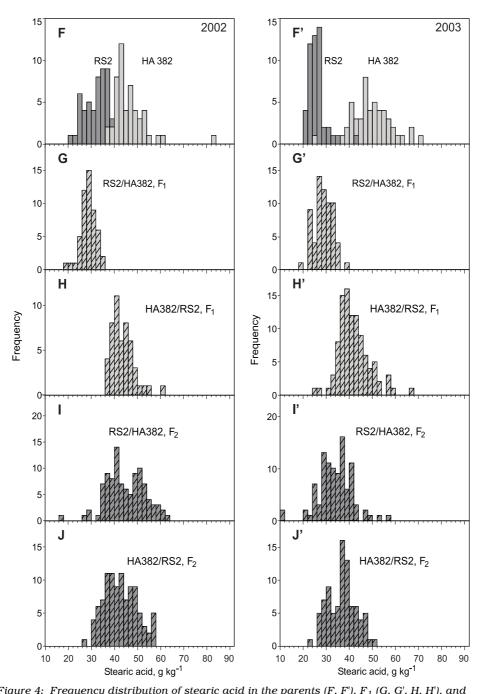


Figure 4: Frequency distribution of stearic acid in the parents (F, F'), F_1 (G, G', H, H'), and F_2 (I, I', J, J') progeny of the reciprocal crosses between RS2 and HA 382 grown in the field in 2002 and 2003

ilar results were reported by Primomo *et al.* (2002) for two new low-palmitic germplasm lines, RG3 and RG1.

Miller and Vick (1999) generated three sunflower mutant lines that were reduced in total SFAs. One mutant line was reduced in palmitic acid, and the other two mutant lines were reduced in stearic acid. Inheritance studies showed that the low-palmitic acid content was controlled by a single gene with additive gene action. The low-stearic acid content was controlled by a single gene in one mutant, and by two genes in the other mutant, both with additive gene action.

Inheritance studies have also been reported for several sunflower mutants with high saturated fatty acid content. Two high-palmitic acid sunflower mutants, CAS-5 and CAS-12, each with palmitic acid content greater than 250 g kg⁻¹, were described by Pérez-Vich et al. (1999a, 2002a). The high palmitic acid content of CAS-5 was concluded to result from the presence of a mutant recessive allele, p1, and one of two other recessive alleles, p2 or p3, already present in the line prior to mutation. The inheritance of high palmitic acid in CAS-12 was similar, being controlled by the partially recessive alleles, p1, p2, and p3. Similarly, the genetic control of a high-stearic acid (about 250 g kg⁻¹) sunflower mutant, CAS-3, has been investigated (Pérez-Vich et al., 1999b). Two recessive genes, es1 and es2, at two independent loci were proposed to be responsible for the high-stearic acid trait of CAS-3, having the genotype eslesles2es2. In another sunflower mutant, CAS-4 (medium-stearic), the stearic acid content (about 130 g kg⁻¹), was found to be controlled by the same es2 gene and a second gene, different from es1, termed es1^b (Pérez-Vich et al., 2002b). The genotype of CAS-4 was proposed to be es1^bes1^bes2es2.

In a practical sense, the expression of the reduced palmitic and stearic acid traits is most important in the F_2 seed, which is harvested commercially and crushed for oil. The F_2 generation is also important for plant breeders because it is the segregating generation where selections can be made by companies using RS1 and RS2 to lower the SFA content of their parental lines. Our experiments showed that the inheritance and expression of the reduced palmitic and stearic acid contents of RS1 and RS2 are complex and apparently multigenic. The greatest reduction (~25 %) in SFAs in the F_2 seed was achieved when RS1 was a parent.

The relative changes in palmitic and stearic acid contents between the two growing seasons suggest that environmental conditions play a significant role in the proportions of palmitic and stearic acids. While we do not know the specific environmental parameter(s) responsible for the altered proportions of palmitic and stearic acids, temperature may have contributed to the differences between years. Figure 5 shows thermal availability (daily sunflower growing degree days) during the seed development period in 2002 and 2003. The weather data for the field plots showed that there was a substantial difference in timing of heat input between years. In 2002, the period of early seed development (~August 15-21) was cool, followed by a warmer period (August 22 - September 8). In 2003, the temperature

trend during seed development was reversed, with a warm early period (~August 15-21) and cooler later (August 22 - September 8). Linoleic acid content increases with cooler temperatures, and our data show a significantly higher linoleic acid concentration in the seed harvested in 2003 (Tables 1 and 2). Therefore, it may be that differences in fatty acid composition of the seeds were largely determined by temperature during the active seed filling period from August 22 to September 8, when temperatures were cooler in 2003 than in 2002. The total SFA content was also influenced by environment, having opposite effects with RS1 (higher total SFA in 2003) and RS2 (lower total SFA in 2003).

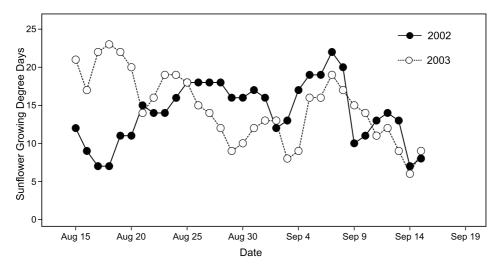


Figure 5: Daily sunflower growing degrees from August 15 through September 15, 2002 and 2003, measured in the field plots at Fargo, ND (data from North Dakota Agricultural Weather Network)

CONCLUSIONS

The reduced SFA phenotype in the F_2 generation is largely due to inheritance of the trait from RS1 and RS2. Environment also plays a role in achieving reduced content of SFA. Thus, it can be expected that the level of SFAs in cultivated hybrid sunflower with genes from RS1 or RS2 in one of the parents will vary from year to year. Therefore, to achieve the lowest possible concentration of SFAs in the F_2 seed, it will be necessary to incorporate the reduced SFA trait into both parents of a hybrid.

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ESTUDIO BIANUAL DE HERENCIA DEL CONTENIDO REDUCIDO DE ÁCIDOS GRASOS SATURADOS EN LA SEMILLA DE GIRASOL

RESUMEN

RS1 y RS2 son dos bases genéticas que contienen bajo contenido de ácidos grasos saturados. Aquí hacemos la relación de los resultados de bienio sobre la herencia de los ácidos palmítico y esteárico en RS1 y RS2, en la generación de cruzamiento F_1 y F_2 con HA 382, la línea consanguínea (inbred), cuyo contenido de ácidos grasos es semejante a muchos híbridos comerciales de girasol. La expresión del contenido reducido de los ácidos palmítico y esteárico es la más importante en la generación F_2 porque la semilla F_2 es la que el granjero siega y vende y de la cual se produce aceite. Nuestros ensayos han demostrado que la semilla F_2 de los cruzamientos recíprocos de RS1 ó RS2 con HA 382 era intermediaria en el contenido de los aceites grasos saturados entre dos progenitores híbridos. La herencia y la expresión de los ácidos palmítico y esteárico reducidos en RS1 y RS2 eran bajo control de más de un gen. Además, los cambios relativos significantes en contenido de los ácidos palmítico y esteárico entre los períodos vegetativos en los anos 2002 y 2003, indican que el entorno tenía un papel importante en las relaciones relativas entre los ácidos grasos. También, el total contenido de los ácidos grasos era bajo la influencia del entorno, generando efectos contradictorios en RS1 (incrementado el total contenido de ácidos grasos en el ano 2003) y RS2 (reducido el total contenido de ácidos grasos en el ano 2003). Esos datos indican que es necesario introducir el reducido contenido de ácidos grasos en ambos progenitores del híbrido, para lograr la más baja concentración de los ácidos grasos saturados en la semilla F_2 .

ÉTUDE DE DEUX ANS PORTANT SUR LA TRANSMISSION DU CONTENU RÉDUIT D'ACIDES GRAS SATURÉS DANS LA GRAINE DE TOURNESOL

RÉSUMÉ

RS1 et RS2 sont deux bases génétiques qui possèdent un faible contenu d'acides gras saturés. Nous faisons part ici des résultats d'une étude de deux ans sur la transmission des acides palmitique et stéarique de RS1 et RS2 dans les générations du croisement F1 et F2 avec HA 382, une lignée ayant une composition d'acides gras semblable à celle de plusieurs hybrides de tournesol commerciaux. L'expression du contenu réduit d'acides palmitique et stéarique est la plus importante dans la génération F2 parce que c'est celle qui est cultivée par les fermiers, vendue et pressée pour l'extraction d'huile. Nos expériences ont démontré que la graine F2 des croisements réciproques de RS1 ou RS avec HA 382 était intermédiaire pour le contenu d'acides gras saturés entre les deux parents hybrides. La transmission et l'expression des caractéristiques palmitique et stéarique de RS1 et RS2 ont été contrôlées par plus d'un gène. De plus, des changements relatifs importants dans les contenus d'acide palmitique et stéarique entre les saisons de végétation de 2002 et 2003 donnent à penser que l'environnement a un rôle important dans les proportions relatives de ces acides gras. Le contenu total d'acides gras saturés a aussi été influencé par l'environnement, produisant des effets opposés avec RS1 (total d'acides gras saturés augmenté en 2003) et RS2 (total d'acides gras saturés diminué en 2003). Ces données démontrent qu'il est nécessaire d'incorporer la caractéristique d'acides gras saturés réduite dans les deux parents d'un hybride de manière à obtenir la plus basse concentration d'acides gras saturés dans la graine F₂.

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