

FIELD EXPOSURE OF TWO UPLAND COVER TYPES TO SOLID ROCKET MOTOR FUEL EMISSIONS¹

by

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ABSTRACT

*A series of replicated field experiments in which two important upland cover types were exposed to solid rocket motor fuel emissions did not reveal any immediate impact on the vegetation. Soil pH and Cl^- concentrations remained unchanged. Root biomass one year after exposures was not significantly different among the treatments. Likewise, twig growth of two browse species, dwarf live oak (*Quercus minima*) and myrtle oak (*Q. myrtifolia*), was unimpaired. Flower and fruit formation by the oaks, saw palmetto (*Serenoa repens*), and blueberry (*Vaccinium myrsinites*) appeared normal one year after exposure. Litter fractions (twigs, leaves, and fine material) were not significantly different among treatment groups.*

The ecological effects and environmental fate of solid rocket motor (SRM) exhaust in and around the Kennedy Space Center (KSC) represents an area of concern to the National Aeronautics and Space Administration (NASA). Over the next several years (1978-1981) an operational Space Shuttle which employs SRM's is anticipated to be launched at KSC on a routine basis (reviewed by von Braun 1972). This aerospace activity may be expected to continue for perhaps several decades. Solid rocket motor emission products, therefore, will continue for many years to enter the various ecosystems making up Merritt Island and the Kennedy Space Center.

The purpose of this research was to determine if single, short-term exposure (10 min.) of selected ecosystems to SRM fuel emissions would result in demonstrable changes in certain response parameters. Field studies commenced in July 1973 when permanent study areas were selected in undisturbed plant communities representative of pine flatwoods and scrubby flatwoods ecosystems and when baseline data on soil variables and plant abundance were gathered. Exposures of the intact plant communities were carried out in August 1974. Field observations and final data gathering was completed July 1975.

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MATERIALS AND METHODS

Study Areas

The Kennedy Space Center occupies most of Merritt Island on Florida's east central coast. Merritt Island National Wildlife Refuge shares a common boundary with KSC. The

¹ Contribution No. 3 of the Merritt Island Ecosystems Studies

newly created Canaveral National Seashore is contiguous with and to the North of the refuge. The Eastern Test Range, under control of the U. S. Air Force is on the southern portion of Cape Canaveral.

Merritt Island is subtropical in its climate. Summer temperatures range from daily highs in excess of 32C to night time lows of 21-26C. Winter temperatures average 18.3C during the day and freezing conditions are seldom realized. Rainfall averages 127-139 cm per year, with an uneven distribution throughout the year (Bufson and Prine 1968). Rain is concentrated in the period of June-September while the winter months may be quite dry.

Pine Flatwoods—This ecosystem is a counterpart of a community type found widely over peninsular Florida (Laessle 1942). The typical overstory of longleaf pine (*Pinus palustris*) is not present in the stands studied on northern Merritt Island, but a typically developed understorey of broadleaf evergreen shrubs is present. Study of 40 systematically located 1-m² plots revealed 9 dominant woody plants on the study plots. Dwarf live oak occurred at a density of 20.2/m². Saw palmetto was very conspicuous but less dense (0.8/m²). Shiny lyonia (*Lyonia lucida*) and shiny blueberry were most common among the remaining woody plants.

The pine flatwoods study plots are found on Immokalle sand soil type (Huckle et al. 1974). The soil is dark gray and highly acid (pH 4.5). Drainage is poor and standing water may be present for short periods during the summer months.

Scrubby Flatwoods—This ecosystem is very similar to the sandpine scrub communities that occupy well drained sands in many parts of Florida (Laessle 1958). The coastal examples of these communities on Merritt Island lack the sandpine (*P. clausa*) overstorey. Our sampling (40 1-m² plots) revealed 10 common scrub species. Three oaks were dominant elements based on density. Myrtle oak was most prevalent (10.1/m²) and Chapman (*Q. chapmanii*) and dwarf live oak were less important, 3.9 and 4.3/m². Three *Lyonia* species, *L. fruticosa*, *L. ferruginea*, and *L. lucida*, were present. Saw palmetto occurred at the same density (0.8/m²) as in the pine flatwoods study area.

The scrubby flatwoods study plots were on the Pomello sand soil type (Huckle et al. 1974). The soil is light gray and has a pH of 5.0. Permeability is rapid and the communities are not subject to flooding during the rainy season.

Research Design and Exposures

Experimental design for the pine flatwoods and scrubby flatwoods communities was identical. Six plots were exposed to SRM emission products in each community type during August 1974. Three treatments of 5, 50, and 100 ppm HCl based on theoretical concentration in SRM fuel exhaust and a control were replicated in each community. Plots were identified in the field and assigned to treatment levels according to a randomized complete block design (Steel and Torrie 1960).

A 10-m² field enclosure constructed of four side panels and two hinged roof panels with 6 mil polyethylene transparent plastic walls rested over the vegetation and confined the exhaust gases for a period of 10 minutes.

Exhaust components (Table 1) from SRM fuel were generated by open burning of pre-weighed quantities of SRM fuel. Fuel was placed on a sand substratum within an open-topped metal cylinder which served to shield the plants from thermal damage. Ignition was accomplished electronically after the fuel was attached and nichrome wire (0.254 mm diam.) to lead wires from a battery box located outside the enclosure.

An impinger gas sampling system was used to determine dosage. All components of the exhaust could not be independently ascertained; therefore, HCl, the most toxic component, was monitored with a system Madsen (1974) developed for laboratory and field use. Air from the enclosure atmosphere was continuously drawn by a vacuum pump thru a gas dispersion tube and trapped in deionized water. The chloride content of the deionized water was then related to the HCl content of the enclosure atmosphere. Routine calculation of HCl concentration was carried out by direct potentiometric determination of chloride. A calibration curve was prepared based on standard chloride solutions and results expressed in ppm HCl.

Table 1. Major theoretical combustion products of solid rocket motor fuel and the weight of these products during the launch of a Titan III-C booster.

Product	Form	Mole Fraction Vickers 1974	Kg per Launch	
			NASA Anon Undated	Cesta and McLouth 1969
H ₂	Gas	0.32008	7,582	7,774
CO	Gas	0.25761	84,898	98,899
HCl	Gas	0.14744	63,106	75,871
H ₂ O	Gas	0.09663	20,657	38,772
N ₂	Gas	0.07801	2,724	3,795
Al ₂ O ₃	Solid	0.07708	95,616	116,878
CO ₂	Gas	0.01407	8,535	11,332
H	Gas	0.00506	—	—
Cl	Gas	0.00170	—	—
FeCl ₂	Solid	—	1,226	—

Response Parameters

Studies on soil variables were limited to monitoring pH and Cl⁻ concentration. Multiple soil samples were collected from the study sites prior to and 10 days after exposure to SRM fuel exhaust. Soil pH was determined by the 1:1 soil:water ratio method. The pH of the resulting suspension was measured with a previously calibrated Beckman Expandomatic pH meter operated in the standard range. Silver-silver chloride indicator and reference electrodes were employed to determine chloride content of the soil suspensions.

Structural conditions of leaves of plants growing within the study plots were documented with line drawings, notes, and color photographs prior to exposure. Four to six woody plants were identified and three branches (top, middle, and bottom of the canopy) were tagged for sequential observation of three terminal leaves per branch.

Possible delayed effects of SRM exhaust on the vegetation were studied in July 1975. Treatment effects on root biomass were evaluated by taking ten 1- x 25-cm soil cores from each plot using objective procedures. Cores from a given plot were pooled and the root material separated from the soil by use of a No. 20 sieve. Root biomass was oven dried 40 hours at 58C and weighed to the nearest 0.1 g.

Twig growth was studied by clipping current growth and determining dry weight. Within the pine flatwoods area only *Q. minima* was suitable for this measurement. Ten twigs were collected from a single plant (judged to be typical) in each plot. The same procedures were used in sampling the scrubby flatwoods plots. Owing to their abundance, two individuals of *Q. myrtifolia* per plot were sampled. All twigs were oven dried 40 hrs. at 58C and their mass individually determined on a Mettler H542 microbalance.

Within each plot on the two study areas, two 0.1-m² quadrats were objectively located and all plant litter (leaves, twigs and fine particulate matter) was collected. Material passing through a screen with a 2- x 2-mm mesh was discarded. Woody material was handpicked and the remaining fraction sorted into fine material with a screen of 6- x 6-mm mesh. Material not passing through the screen was defined as leaves. All fractions were oven dried for 40 hrs. at 58C and the mass determined.

RESULTS

Theoretical and atmospheric concentrations of HCl in the field enclosure are given in Table 2 according to cover type. All fuel was ignited in each exposure; therefore, the expected concentrations were probably reached for some brief period. Differences between theoretical and observed values may be explained by absorption of the emission products (including HCl) on the foliage and enclosure walls. The data suggest that some vertical stratification in the enclosure occurred with concentration increasing with height.

Table 2. Theoretical and atmospheric concentrations of HCl (ppm) as measured in 16.7-m³ field enclosure after open burning of SRM fuel. Three impingers were located at 0.4-m and two at 1.0-m.

Theoretical Concentration of HCl (ppm)	Mean \pm SE concentration of HCl (ppm)			
	Pine Flatwoods		Scrubby Flatwoods	
	0.4-m (n = 6)	1.0-m (n = 4)	0.4-m (n = 6)	1.0-m (n = 4)
5 ppm	1.98 \pm 0.50	1.89 \pm 0.44	0.46 \pm 0.13	0.27 \pm 0.04
50 ppm	6.45 \pm 1.02	7.37 \pm 2.29	3.05 \pm 0.55	10.12 \pm 3.24
100 ppm	13.16 \pm 1.71	16.85 \pm 1.30	6.55 \pm 1.17	22.97 \pm 7.00

Soil

Analysis of variance of the treatments and soil pH in the pine flatwoods 10 days after exposure to SRM fuel exhaust indicated no differences among the plots (F=1.708, df 1 and 7, NS). A similar analysis of the scrubby flatwoods revealed no treatment effect (F=0.435, df 1 and 7, NS).

Mean Cl⁻ concentrations in soil samples from the replicate plots are presented in Table 3. Plots were exposed to SRM fuel exhaust 10 days prior to the September observations. Inspection of these data suggests no detectable change in soil Cl⁻ was apparent between April and September.

Impact on Leaf Structure

Visual inspection of leaves on labeled plants immediately after exposure, and at 2 weeks, and 4 weeks post-exposure revealed no discernible damage. Previous work with laboratory plants (Stout unpublished work) had indicated acid burns as the typical kind of damage to be expected. No further inspection of leaves on the plots was made after 4 weeks.

Root Biomass

The relationship between root biomass and exposure concentration on the pine flatwoods plots one year after exposure was described by the equation $y=19.83+0.02(x)$, where y is root biomass and x is the theoretical concentration of HCl in the SRM fuel

Table 3. Soil Cl⁻ concentrations on the study plots prior to and following exposure to various concentrations of SRM fuel emissions.

Community	Treatment (ppm HCl)	Mean soil Cl ⁻ concentration (ppm)	
		April 1974	September 1974 ^a
Pine Flatwoods	Control n = 2	66	51
	5 ppm n = 2	24	19.5
	50 ppm n = 2	28	26
	100 ppm n = 2	29	36
Scrubby Flatwoods	Control n = 2	20.5	26
	5 ppm n = 2	18.5	21
	50 ppm n = 2	24.5	20
	100 ppm n = 2	17	17

^aExposures were in August 10 days prior to determination.

exhaust (Fig. 1). The coefficient of determination (r^2) was 0.1031 and indicated little of the variation in root biomass was explained by the treatments. Similarly, the equation $y = 14.30 - 0.03(x)$ fit the root biomass data from the scrubby flatwoods area. Little variation in root biomass was accounted for by the treatments ($r^2 = 0.1005$).

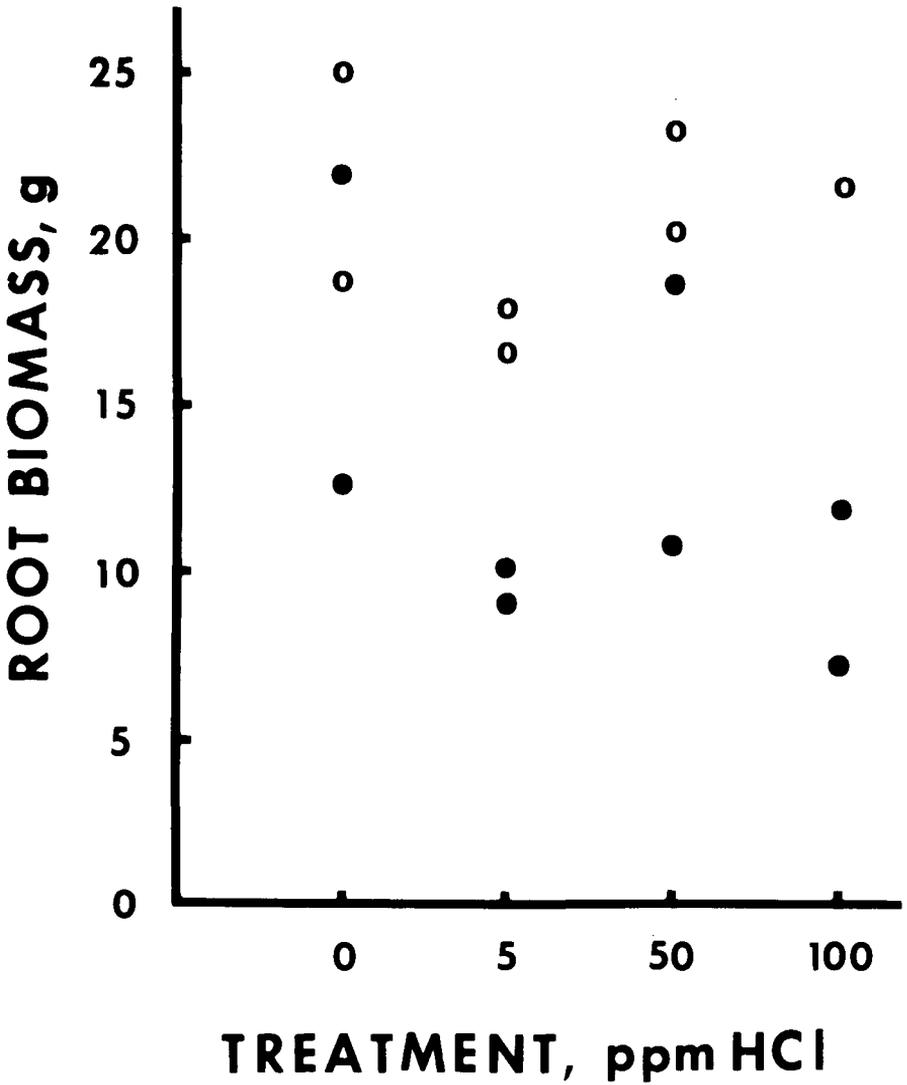


Figure 1. Relationship between root biomass and treatment with SRM fuel emissions (ppm HCl). Replicate plots in the pine flatwoods are designated by open circles and plots in scrubby flatwoods by closed circles. Exposures were completed 1 year prior to determination of root biomass.

Twig Growth

Mean dry weights of dwarf live oak twigs from the pine flatwoods revealed no treatment effects ($F = 2.628$, 3 and 3 df, NS, Table 4). In contrast, dry weights of myrtle oak twigs from scrubby flatwoods indicated heterogeneity among the means ($F = 7.097$, 3 and 12 df, $P < .05$, Table 5). The 100 ppm treatment group was significantly larger than the other groups ($p < .05$).

Flowering and Fruiting

Observations in July 1975 on the pine flatwoods and scrubby flatwoods indicated that fruiting and flowering of shrubs such as the oaks, saw palmetto, and blueberries were normal. These qualitative data are summarized in Stout (1975:64-65).

Table 4. Dry weight of early summer (July 1975) twig growth of dwarf live oak (*Quercus minima*) from a pine flatwoods community. Plots were exposed to SRM fuel emissions during August 1974.

Treatment (ppm HCl)		Mean \pm SE dry weight of twigs (g)
Control	n = 2	0.056 \pm .012
5 ppm	n = 2	0.100 \pm .008
50 ppm	n = 2	0.132 \pm .035
100 ppm	n = 1 ^a	0.042 \pm .010

^aOne plot was destroyed prior to measurement.

Table 5. Dry weight of early summer (July 1975) twig growth of myrtle oak (*Quercus myrtifolia*) from a scrubby flatwoods community. Plots were exposed to SRM fuel emissions during August 1974.

Treatment (ppm HCl)		Mean \pm SE dry weight of twigs (g)
Control	n = 4	.122 \pm .017 ^a
5 ppm	n = 4	.126 \pm .042 ^a
50 ppm	n = 4	.158 \pm .061 ^a
100 ppm	n = 4	.271 \pm .071

^aMeans followed by letter are not significantly different ($P < .05$) according to Duncan's new multiple range test.

Table 6. Summary of analysis of litter on plots in pine flatwoods. Plots were exposed to SRM fuel exhaust in August 1974 and litter samples were collected in July 1975.

Treatment (ppm HCl)	Mean \pm SE dry weight of litter components			
	Twigs	Leaves	Fine Material	Total
Control n=4	14.62 \pm 4.41	34.00 \pm 9.08	13.07 \pm 1.18	61.77 \pm 13.68
5 ppm n=4	22.87 \pm 10.06	40.05 \pm 8.66	22.50 \pm 3.95	85.42 \pm 17.59
50 ppm n=4	13.07 \pm 4.76	34.40 \pm 8.34	21.10 \pm 3.95	68.57 \pm 15.45
100 ppm n=2	9.00 \pm 0.30	24.40 \pm 1.60	19.40 \pm 2.20	52.80 \pm 0.30

Table 7. Summary of analysis of litter on plots in scrubby flatwoods. Plots were exposed to SRM fuel exhaust in August 1974 and litter samples were collected in July 1975.

Treatment	Mean \pm SE dry weight of litter component			
	Twigs	Leaves	Fine Material	Total
Control n=4	10.70 \pm 2.96	35.62 \pm 10.85	14.25 \pm 1.51	60.57 \pm 10.17
5 ppm n=4	26.62 \pm 3.64	38.22 \pm 5.82	12.87 \pm 3.48	77.72 \pm 11.75
50 ppm n=4	14.77 \pm 5.55	21.72 \pm 4.15	12.90 \pm 1.78	49.40 \pm 9.31
100 ppm n=4	18.40 \pm 3.30	34.87 \pm 4.12	13.45 \pm 1.18	74.97 \pm 6.99

Litter Standing Crop

Standing crop of litter components on the pine and scrubby flatwoods plots are given in Tables 6 and 7. Analysis of variance of treatment and litter components on the pine flatwoods was not significant ($p > .05$). A similar analysis of treatment effects on the scrubby flatwoods indicated no significant differences among plots ($p > .05$).

DISCUSSION

The objective of this work was to determine if wildlife habitat was significantly altered by single, and short-term exposure to SRM emission products. Exposure of natural vegetation to the total complex of emission products appeared to be a realistic way to help answer the question. Furthermore, we assumed that the potential impact of greatest concern to long-term stability of the habitats was the disruption of plant growth and reproduction, i. e., primary production (Woodwell 1970).

Indirect but subtle changes in soil chemistry could influence plant growth. Exposure to SRM fuel emission products did not result in changes in soil pH and Cl⁻ concentration of the study plots. A reduced soil pH could have made soluble aluminum more available and led to toxic conditions for wildlife food plants (Clarkson 1969). Input of Al₂O₃ was not likely to cause any alteration in growing conditions because of its stability and abundance in the soil. Buckman and Brady (1969:24) state that from 4.5 to 13.14 percent of soil is Al₂O₃.

No structural damage in the form of acid burns, chlorosis, margin curl or pitting was observed on leaves of plants exposed in the field. Heck et al. (1970) reviewed the scanty literature on hydrogen chloride as a phytotoxin. They list, among other plants, maple (*Acer* sp.), cherry (*Prunus* sp.), larch (*Larix* sp.), and viburnum (*Viburnum* sp.) as sensitive to HCl; whereas, oaks, spruce (*Picea* sp.), and some maples are resistant. Lind and London (1971) found marigold (*Tagetes* sp.) exhibited no visible effect when exposed to 95 ppm HCl. A spectrum of sensitivities might be expected in a natural community, comprised of numerous species. The lack of evidence of sensitivity to SRM fuel emissions in our studies is probably best explained by natural resistance. All the plants are adapted to grow in xeric environments with nutrient deficient soils. Evergreenness and thick, leathery leaves are morphological features which further aid in provision of resistance to SRM fuel emissions (Seddon 1974:71).

The importance of oaks and other scrub species as cover and food for wildlife in general and deer (*Odocoileus virginianus*) specifically (Harlow and Jones 1965:100) prompted a re-examination of the study areas a year after the exposures. Laboratory studies of HCl gas (Godish 1970) have shown it to affect photosynthesis and respiration of tomato plants. Thus lack of overt damage to our study plants did not preclude delayed responses in terms

of diminished growth or flower and fruit formation. However, no differences in root biomass, twig growth of oaks, flower and fruit formation, or litter production were found.

A major deficiency of our study is a lack of replication in other seasons. However, based on our experience with two upland cover types, single and short-term exposure to SRM fuel emission products did not have an influence on the variables under study after one year.

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