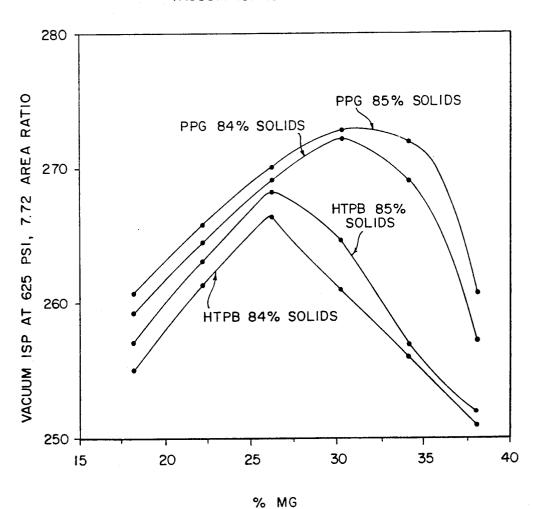
United States Patent [19] 5,076,868 [11] Patent Number: Doll et al. Date of Patent: Dec. 31, 1991 [54] HIGH PERFORMANCE, LOW COST SOLID 3,198,677 8/1965 Thomas 149/19.6 3,680,483 Staudacher et al. 149/19.6 8/1972 **PROPELLANT COMPOSITIONS** PRODUCING HALOGEN FREE EXHAUST 3,720,553 3/1973 Henderson 149/19.6 3,956,890 5/1976 Davis 149/19.4 [75] Inventors: Daniel W. Doll, N. Ogden; Gary K. 4,111,728 9/1978 Ramnarace 149/19.6 8/1979 Lund, Ogden, both of Utah 4,165,247 Brew et al. 149/19.6 4,358,327 11/1982 Reed et al. 149/19.4 [73] Assignee: Thiokol Corporation, Ogden, Utah 4,799,980 1/1989 Reed 149/19.4 4,925,909 5/1990 Kubota et al. 149/19.6 [21] Appl. No.: 531,728 Primary Examiner—Edward A. Miller [22] Filed: Jun. 1, 1990 Attorney, Agent, or Firm-Lawrence I. Field; Ronald L. Lyons [52] U.S. Cl. 149/19.4; 149/19.6; [57] **ABSTRACT** 149/20; 149/111 [58] Field of Search 149/19.6, 19.4, 20, High performance solid propellant compositions pro-149/111 ducing halogen-free exhaust products comprised of Ammonium Nitrate and powdered magnesium and op-[56] References Cited

U.S. PATENT DOCUMENTS

VACUUM ISP vs PERCENT MG

binder.

tionally containing polyoxypropylene glycol as a



VACUUM ISP vs PERCENT MG

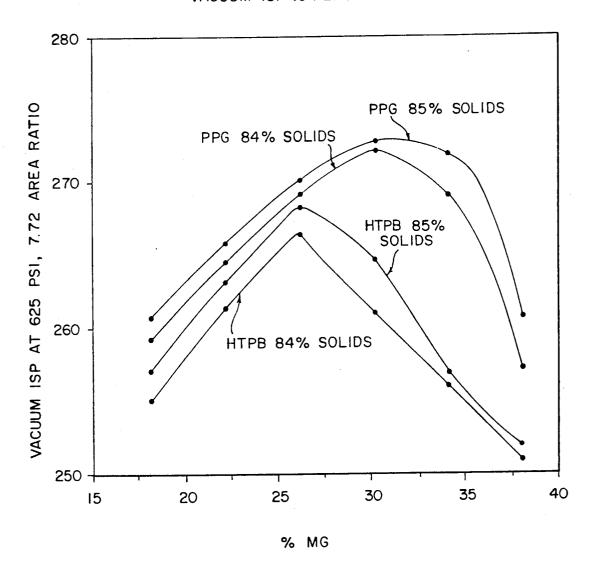


FIG. I

FIG. 2

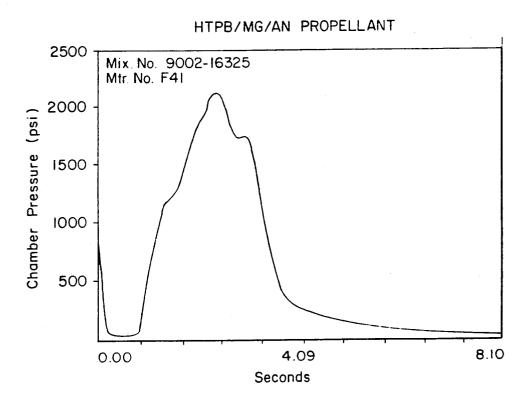
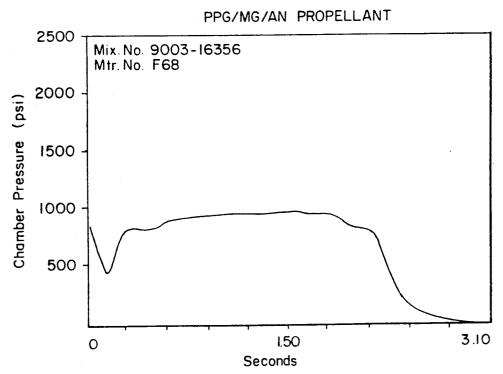


FIG. 3



HIGH PERFORMANCE, LOW COST SOLID PROPELLANT COMPOSITIONS PRODUCING HALOGEN FREE EXHAUST

This invention relates to high performance, low cost solid propellant compositions producing halogen free exhaust. More particularly, it relates to solid propellant compositions which are free of chlorine containing constituents and which therefore produce an exhaust 10 which is free from any chlorine or other halogen either as the element or as a halogen containing compound.

In general, it has been the experience of the propellant industry as a whole, that use of ammonium nitrate as a solid propellant oxidizer in the absence of substan- 15 present a safety hazard due to ignition sensitivity to tial amounts of ammonium perchlorate (or other similar solid oxidizers) produces unsatisfactory combustion when formulated with aluminum powder. Consequently, propellant performance is poor and addition of amounts of nitrate esters or use of energetic polymers is required to achieve adequate combustion temperatures to ignite the aluminum powder. These additives are expensive and often increase the explosive sensitivity of complexity.

One object of the invention is to provide a low cost propellant composition in which ammonium nitrate is the sole oxidizer, which burns without leaving any solid or liquid residue and which does not require the pres- 30 ence of energetic polymers or other additives to obtain such complete combustion.

Another object is to provide a propellant composition which does not include any halogen containing

These and other objects are achieved by a composition in which metallic magnesium is the fuel and ammonium nitrate is the sole oxidizer and which may contain polyoxypropylene glycol as a binder.

The invention will be more fully understood from the 40 description which follows taken in conjunction with the drawings in which:

FIG. 1 shows graphs depicting theoretical Isp for various percentages of Mg in an Mg/AN propellant for two different binders; and

FIGS. 2 and 3 are graphs depicting chamber pressure vs time for two Mg/AN propellants.

AMMONIUM NITRATE

Ordinary fertilizer grade ammonium nitrate is satis- 50 factory in formulating the compositions of this invention, provided it contains less than 0.1% of water, by weight.

For certain applications requiring AN propellants to be exposed to temperatures exceeding 120° F., it is pre- 55 formulated to equivalent oxidizer to fuel ratios.

ferred to use AN that contains phase stabilizers (eg. KNO₃, ZnO, NiO, MgO, etc.). Usually two particle sizes of AN are used in the propellant compositions of this invention namely: a coarse fraction (200- to 2000micron) and a fine fraction (20- to 200-micron). The coarse fraction preferably has rounded edges, e.g., a prill. The fine fraction can be ground from the coarse AN. The preferred fine particle size is 40- to 100micron.

MAGNESIUM

Any Mg powder coarser than 50-micron and finer than 800-micron has been found to be suitable. Smaller sizes (<50-micron) can be used. However, these often electrostatic energy and thus are to be avoided. Spherical or ellipsoidal particles are preferred although not required.

It has been found that formulations utilizing ammocombustion improving ingredients such as large 20 nium nitrate as the sole oxidizer ignite and combust completely with little or no slag formation without the addition of high energy ingredients when magnesium powder is employed instead of aluminum powder. Table I compares the ballistic behavior of a series of the composition, greatly increasing propellant costs and 25 ammonium nitrate propellants utilizing various binders with combinations of aluminum and magnesium all formulated to equivalent oxidizer to fuel ratios.

BINDER

The AN/Mg propellant compositions may contain a binder. A preferred binder is polyoxypropylene glycol (PPG).

OTHER INGREDIENTS

Other ingredients commonly used in formulating propellant compositions and which may be present in the compositions of this invention include: burn rate catalysts, plasticizers, phase stabilization agents, bonding agents, and the like. Any or all of these may be used, provided they do not contain a halogen such as chlorine.

The propellant ingredients are typically blended in a 1-pint Baker-Perkins vertical mixer. Propellant is vacuum cast into 1.5×2.5 inch center perforated motors for 45 ballistic testing and JANNAF Class C uniaxial tensile specimens for mechanical property testing.

It has been found that formulations utilizing ammonium nitrate as the sole oxidizer ignite and combust completely with little or no slag formation without the addition of high energy ingredients when magnesium powder is employed instead of aluminum powder. Table I compares the ballistic behavior of a series of ammonium nitrate propellants utilizing various binders with combinations of aluminum and magnesium all

TABLE I

		85 PE		N WITH Mg AND AICENT SOLIDS, 15% BINDER					
	16372	16373	16374	28883-1	28883-2	28883-3	28886-1	28886-2	28886-3
Binder	НТРВ	НТРВ	НТРВ	PPG	PPG	PPG	GAP	GAP	GAP
Α1 20μ	_	11.40	21.0	_	13.50	25.0	_	14.50	27.00
Mg Hart 160µ	25.0	11.40	_	30.0	13.50	_	31.50	14.50	
ΑΝ 600μ	30.0	31.1	32.0	38.0	40.0	41.5	36.75	38.50	39.90
AN 35μ	30.0	31.1	32.0	16.0	17.0	17.50	15.75	16.50	17.10
Viscosity	82	37	39	53	19	23	143	112	>160
(kP)									
Rb (in./sec)	0.104	0.102	would	0.140	would	would	0.260	0.206	0.174
Slope	0.31	0.20	not	0.26	not	not	0.36	0.44	0.96

TABLE I-continued

AN WITH Mg AND Al								
85 PERCENT SOLIDS, 15% BINDER								
16372	16373	16374	28883-1	28883-2	28883-3	28886-1	28886-2	28886-3
		ignite		ignite	ignite			

Rb is propellant burning rate at 1000 psi in inches/second

Use of the energetic binder, GAP(Glycidyl Azide Polymer) resulted in sufficient combustion of either 10 magnesium or aluminum fuel to obtain measurable burning rates, whereas formulations prepared with the non-energetic binders, HTPB(hydroxy terminated polybutadiene) and PPG(polyosypropylene glycol) taining aluminum in all cases. In the present invention

(HTPB)/Mg/AN propellants tend to display pressure versus time traces that are indicative of erratic combustion. Test firings of PPG/Mg/AN propellants display pressure versus time traces that are indicative of stable combustion.

Table II is a comparison of the ballistic and mechanigave very poor or no combustion in formulations con- 15 cal properties of both PPG and HTPB based Mg/AN propellants.

TABLE II

Mg/AN Propellant Comparison						
Binder	% Solids	% Ground AN	% Mg (160μ)	Rb	n	
НТРВ	85	18	25	0.102	0.11	Kp = 50-100
		21		0.121	0.59	Kp = > 100
		24		0.114	0.42	•
		27		0.118	0.33	
		30		0.115	0.42	
		30 (1% Al ₂ O ₃)		0.085	0.23	
		30 (1% Pyrocat)		0.103	0.69	
PPG/DOA	85	15	25	0.134	0.134	Kp = 50-100
		18		0.127	0.120	•
		21		0.127	0.092	
		24		0.133	0.120	
PPG/DOA	84	16	30	0.134	0.169	Kp = 50-100
		19				Kp = > 100
		22		0.135	0.232	-
		25		0.146	0.244	
		28		0.130	0.350	
		34		0.148	0.253	
		37		0.145	0.33	

	MECHANICAL PROPERTIES FOR Mg/AN PROPELLANTS				
	PPG Binder	HTPB Binder			
	85 Solids, 25 Mg, 18% Ground	85 Solids, 25 Mg, 30% Ground			
E ^{2.6} (psi)	348	480-1780			
$\epsilon_m^c(\%)$	15	9-12			
€f(%)	19	12-22			
σ_{m}^{c} (psi)	49	40-127			
Shore A	50	48-73			

Rb is propellant burning rate at 1000 psi

high cost GAP is not required and lower cost binders may be used.

tages over the use of hydroxy terminated polybutadiene (HTPB) as it permits substantially higher metal loading than does HTPB, possibly because of the higher oxygen content of PPG. Consequently higher performance (Isp) is achievable with PPG binders than with HTPB 55 binders at the same weight % solids loading.

FIG. 1 compares metal loadings with PPG as the binder vs HTPB as the binder and it will be seen that the former permits higher metal loadings, with conse-HTPB as a binder.

FIGS. 2 and 3 are pressure vs time curves obtained in small motor tests for comparing the combustion behavior of Mg/AN propellants containing PPG and HTPB binders. The pressure versus time trace for Mg/AN 65 lations without peril provided use temperature requirepropellants, tested in 1.5-×2.5-inch motors, serves to illustrate the improved combustion of PPG binders compared to HTPB binders. Test firings of R-45M

The overall costs of the propellants is lowest with The use of polyosypropylene glycol offers advan- 50 PPG binder formulations. The low viscosity and low hydroxyl reactivity of PPG combine to allow room temperature processing and cure of the formulations using highly reactive cure catalysts such as dibutyltindilaurate. PPG/Mg/AN propellant formulations have been found to achieve a full state of cure at ambient temperature in a similar time as required for conventional propellants which are cured at elevated (120°-135° F.) temperatures.

The ability to process and cure at room temperature quently higher performance (Isp) than is achieved with 60 is particularly important for ammonium nitrate propellants since ammonium nitrate undergoes volume expansion due to crystalline phase changes above about 100° F. Thus, very inexpensive, non-phase stabilized grades of ammonium nitrate may be employed in these formuments do not exceed the phase transition temperatures.

> In summary the compositions of this invention comprise the following in percent by weight:

n is ballistic pressure exponent E²⁶ is propellant modulus (psi)

 $[\]epsilon_m^c$ is propellant strain corrected maximum stress (%)

 $[\]epsilon /$ is propellant strain at failure (%)

 $[\]sigma_m^c$ is corrected maximum propellant stress (psi)

AN (oxidizer)	40 –70
Mg (fuel)	16-36
Binder (PPG)	10-25 (12-18 preferred)

As indicated above, other additives commonly used in propellant compositions may be included in the compositions provided they do not include any halogen or halogen containing compounds.

One specific example of a preferred propellant formulation shown below, contains a binder (which is also a fuel) that is typically composed of a PPG polymer, curative, plasticizer, and a cure catalyst. The main fuel 15 is Mg metal (160-micron) and the non-chlorine oxidizer is solely comprised of AN (600-micron and 35-micron).

Typical properties of the sample composition are: burn rate (ips) at 1000 psi=0.14, burn rate pressure 20 and wherein the combustible binder is a polyoxypropylexponent = 0.26, strain (%) = 15, and stress = 50 psi.

Ingredient	% by weight
Binder	
PPG Polymer	11.89
Isophorone Diisocyanate (Curative)	1.10
Dioctyl Adipate (Plasticizer)	2.00
Dibutyltin Diacetate (Catalyst)	0.01
Fuel	30.00
Mg Metal	
Oxidizer	55.00

-con	tin	ıue	d

% by weight

Having now described a preferred embodiment of the invention it is not intended that it be limited except as may be required by the appended claims, we claim:

1. A high-performance, low-cost, solid propellant composition in which ammonium nitrate is the sole oxidizer and which consists essentially of the following in weight percent:

Ammonium Nitrate (coarse and fine particles)	40-7 0
Magnesium Particles (coarser than 50 microns and	16-36
finer than 800 microns)	
Combustible Binder	10-25

ene glycol cured with an aliphatic diisocyanate.

2. The composition of claim 1 wherein teh coarse particles are 200-2000 microns and the fine particles are 20–200 microns.

3. The composition of claim 2 wherein the fine particles are 40-100 microns.

4. The composition of claim 1 including a phase stabilizer for the ammonium nitrate.

5. The composition of claim 1 in which the propor-30 tions are approximately 55% NH₄NO₃, 30% Mg powder and 15% binder.

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40

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50

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60