

# Green Monopropellants

Subjects: Energy & Fuels | Others

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## Definition

Green propellants are usually defined as low-hazard, low-toxicity, environmentally friendly propellants that are considered safe during various phases of spacecraft development, launch, and operations. Such propellants provide safe handling and storability when compared to conventional toxic propellants such as hydrazine and its derivatives that require special handling protocols and adhering to strict safety measurements that, in addition to others, include using Self-Contained Atmospheric Protective Ensemble (SCAPE) suits. Due to their favorable characteristics, green propellants demonstrate higher commercial value by being able to cut costs related to transportation, storage, handling, and further reduces ground operations time. Recently, a more specified definition has been noted by Mayer et al. (2018) [3], based on the Acute Toxicity Classification (ATC) by the Global Harmonized System of classification and labeling of chemicals (GHS) [6], which denotes that propellants possessing ATC levels of three and safer are considered as green propellants. ATC levels are typically categorized on a 1:5 scale where level one denotes the most toxic class and level five is considered the least toxic class.

Moreover, a controversial topic arises when referring to some modern green propellants, whether to address them by the term “monopropellants” or by more specific terms including (premixed propellants, fuel blends, or mixtures). Monopropellants are fundamentally defined as propellants consisting of chemical compounds (for example  $N_2H_4$ ), which release energy through exothermic chemical decomposition. Since the evolution of liquid gun propellants based on HAN compound and other nitrate salts aqueous solutions, the term “monopropellants” was used to describe such premixed formulations. As widely used in literature and industry, some modern green propellants, for instance the Energetic Ionic Liquids (EILs), are undoubtedly classified and described as “monopropellants.”

Basically, it can be interpreted from the previous that a "green monopropellant" may be defined as: *"A low-hazard, low-toxicity, and safe-to handle propellant that is stored in a single tank and is able to decompose from its storage state by the help of a catalyst or other ignition method, such as thermal or electric ignition, can be considered a "green monopropellant" as long as it does not require another separately stored propellant for decomposition."*

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## 1. Introduction

The current trend in the rocket propulsion field is directed towards greenifying the use of propellants. Monopropellant hydrazine was classically widely used and favored for thrusters and gas generators due to its high performance, system's simpler design, and “clean” relatively cool exhaust products as compared to bipropellant systems at that time [1]. European Chemicals Agency (ECHA) in (REACH) Registration, Evaluation, Authorization and restriction of Chemicals has included hydrazine on the list of Substances of Very High Concern (SVHC) for authorization, thus opening a process that will eventually lead to a ban on the use of hydrazine and its derivatives as space propellant in European countries [2]. Moreover, transportability and handling of hydrazine and similar hazardous propellants extend an economic burden on the space industry. Accordingly, greener alternatives that would compensate for these drawbacks are being studied and developed rapidly nowadays [3][4]. Different global entities were involved in accelerating such research activities through various projects and missions such as Green Advanced Space Propulsion (GRASP), Pulsed Chemical Rocket with Green High Performance Propellants (PulCheR),

and Replacement of hydrazine for orbital and launcher propulsion systems (RHEFORM) European projects and Green Propellant Infusion Mission (GPIM) technology demonstrator project by NASA. Since the beginning of their development, modern green propellants have shown high favorability not only in terms of operability, cost efficiency, and environmental safety but also in performance, and physicochemical properties [5].

As widely interpreted, green propellants are usually defined as low-hazard, low-toxicity, environmentally friendly propellants that are considered safe during various phases of spacecraft development, launch, and operations. Such propellants provide safe handling and storability when compared to conventional toxic propellants such as hydrazine and its derivatives that require special handling protocols and adhering to strict safety measurements that, in addition to others, include using Self-Contained Atmospheric Protective Ensemble (SCAPE) suits. Due to their favorable characteristics, green propellants demonstrate higher commercial value by being able to cut costs related to transportation, storage, handling, and further reduces ground operations time. Recently, a more specified definition has been noted by Mayer et al. [3], based on the Acute Toxicity Classification (ATC) by the Global Harmonized System of classification and labeling of chemicals (GHS) [6], which denotes that propellants possessing ATC levels of three and safer are considered as green propellants. ATC levels are typically categorized on a 1:5 scale where level one denotes the most toxic class and level five is considered the least toxic class.

Before discussing each green monopropellant family in detail, relatively different classifications in literature were proposed for green alternatives of hydrazine monopropellant, one of them considered mainly three types: Energetic Ionic Liquids, Hydrogen Peroxide, and Nitrous Oxide by Batonneau et al. [7] as cited in [8]. However, Mayer et al. [3] further classified propellants including nitrogen compounds with oxygen into two groups: Oxides of Nitrogen subcategory and Nitro Compounds subcategory. The former included mono and dinitrogen oxides ( $\text{NO}$ ,  $\text{NO}_2$ ,  $\text{N}_2\text{O}$ ,  $\text{N}_2\text{O}_3$ ,  $\text{N}_2\text{O}_4$ ,  $\text{N}_2\text{O}_5$ ), which were evaluated as potential oxidizers for bipropellant systems; among which the only compound that was considered as potential green propellant was the nitrous oxide ( $\text{N}_2\text{O}$ ) due to its relative nontoxicity (GHS [6] class 5) and being liquid within a wide part of the typically requested temperature-pressure envelope of  $[-30, +80]^\circ\text{C}$  and  $[0.1, 3]$  MPa, respectively. The latter, Nitro Compounds group, was described as organic substances containing dinitrogen monoxide (i.e., hydrocarbons and nitrous oxide) group, such as (mono)nitromethane ( $\text{CH}_3\text{NO}_2$  or NM), which was also considered as a promising candidate green propellant being relatively nontoxic (GHS class 4) [3]. It was found that the current available green monopropellants that are suitable for space propulsion, either in primary or auxiliary (secondary) propulsion systems, can be classified into three more collective major categories:

1. Energetic Ionic Liquids (EILs) green monopropellants (or premixed oxidizer/fuel ionic aqueous solutions).
2. Liquid  $\text{NO}_x$  Monopropellants (either in binary compound, nitro compound, or premixed/blend form).
3. Hydrogen Peroxide Aqueous Solutions (HPAS).

## 2. Green Monopropellants Classification

A controversial topic arises when referring to some modern green propellants, whether to address them by the term “monopropellants” or by more specific terms including (premixed propellants, fuel blends, or mixtures). Monopropellants are commonly defined as propellants consisting of chemical compounds (for example  $\text{N}_2\text{H}_4$ ), which release energy through exothermic chemical decomposition. Since the evolution of liquid gun propellants based on HAN compound and other nitrate salts aqueous solutions—discussed in the next section, the term “monopropellants” was used to describe such premixed formulations. As widely used in literature and industry, some modern green propellants, for instance the EILs, are undoubtedly classified and described as “monopropellants.” Basically, it can be interpreted from the previous example that a propellant that is stored in a single tank and is able to decompose from its storage form by the help of a catalyst or other ignition method, such as thermal or electric ignition, can be considered a

“monopropellant” as long as it does not require another separately stored propellant for decomposition. Nitrous Oxide Fuel Blends (NOFBs)—discussed in [Section 2.2](#)—were mostly described as green monopropellants as well, maintaining the above-mentioned unique conditions for storage and decomposition of monopropellants.

This section handles each of the three classes proposed for the state-of-the-art green monopropellants. History and origins of development are entitled along with technical data and characteristics including the chemical formulations and constituents of each monopropellant. Thermodynamic and thermochemical properties are gathered from various literature sources—especially resources solely focused on studying propellants thermochemistry. Flight heritage of mature propellants is mentioned, and promising lab test results of other promising propellants are highlighted when possible. Ignition techniques are another important property sought by propulsion systems designers that assist in giving insights about spacecraft mass and volume preliminary requirements as well as electric power needs, thus is noted and discussed deliberately whenever reliable data were available. Finally, physical properties and performance parameters of each propellant are tabulated, and relevant group comparisons are made for the reader’s convenience.

## 2.1. Energetic Ionic Liquids (EILs)

Energetic Ionic Liquids (or premixed oxidizer/fuel ionic propellant blends) consist of oxidizer salts dissolved in aqueous solutions, called Ionic Liquids (ILs), mixed with Ionic Fuel (IF) or Molecular Fuel (MF), refer to [Table 1](#), forming a premixed propellant (i.e., Energetic Ionic Liquid monopropellant as widely referred to among the rocket propulsion community [\[8\]](#)). Addition of the fuel component increases the performance of the propellant blend by reducing the high adiabatic temperature of the ionic liquid binary aqueous solution and further stabilizing the combustion process. Typically, methanol is used to control the burning rate of the monopropellant while the ammonium nitrate (AN) is used as a stabilizer [\[9\]](#) beside other stabilizing additives; a new article by M. Claßen et al. [\[10\]](#) introduced novel additive promoters of new azido esters and suggested it would improve the total energy and performance of ionic liquid propellants. As an example, the maximum specific impulse of 78 wt% ADN in water mixture (Ionic Liquid) is 192 s when used as monopropellant, while the specific impulse rises to 252 s when methanol (molecular fuel) is added to the mixture, as in the case of LMP-103S (63.4 wt% ADN, 25.4 wt% water and 11.2 wt% methanol) at a nozzle area expansion ratio of 50 [\[11\]](#) as cited in [\[3\]](#). In the next paragraphs different EILs will be reviewed (i.e., HAN, HAN/HN, HNF, ADN based green monopropellants) emphasizing on their composition, physical properties, performance, stability of storage and handling, toxicity, material compatibility, ignition methods, and in-flight heritage or proposed missions.

HAN-based monopropellants origins can be traced back to the development of liquid gun propellants (LGPs) in the U.S. Army [\[12\]](#). Three formulations of LGPs were addressed, namely LP1846, LP1845, and LP1898 [\[5\]](#) and their properties are listed in [Table 2](#). The first two of these aqueous solutions are HAN/TEAN-based (tri-ethanol-ammonium nitrate) while the third is HAN/DEHAN-based (di-ethyl hydroxyl ammonium nitrate). The unsuitability of these propellants for rocket’s relatively low combustion pressure [\[13\]](#), as well as the high combustion temperature (2500 K [\[5\]](#)) eventually lead to the development of the state-of-the-art AF-M315E (Air Force Monopropellant 315E) HAN-based green monopropellant for space propulsion formulated by the U.S. Air Force Research Laboratory (AFRL) [\[14\]](#).

**Table 1.** Energetic ionic liquids: oxidizers and fuels thermochemical properties [\[8\]](#)[\[15\]](#)[\[16\]](#)[\[17\]](#)[\[18\]](#).

Ionic Oxidizer		Molecular Weight (g mol <sup>-1</sup> )	Standard Heat of Formation (kJ mol <sup>-1</sup> )
HAN, hydroxyl ammonium nitrate	[NH <sub>3</sub> OH] <sup>+</sup> [NO <sub>3</sub> ] <sup>-</sup>	96.04	-338.97 <a href="#">[19]</a>
ADN, ammonium dinitramide	[NH <sub>4</sub> ] <sup>+</sup> [N(NO <sub>2</sub> ) <sub>2</sub> ] <sup>-</sup>	124.06	-134.6 <a href="#">[20]</a> as cited in <a href="#">[21]</a>

Ionic Oxidizer		Molecular Weight (g mol <sup>-1</sup> )	Standard Heat of Formation (kJ mol <sup>-1</sup> )
HNF, hydrazinium nitroformate	[N <sub>2</sub> H <sub>5</sub> ] <sup>+</sup> [C(NO <sub>2</sub> ) <sub>3</sub> ] <sup>-</sup>	183.08	-72.104 <sup>[19]</sup>
AN, ammonium nitrate	[NH <sub>4</sub> ] <sup>+</sup> [NO <sub>3</sub> ] <sup>-</sup>	80.043	-365.28 <sup>[19]</sup>
HN, hydrazinium nitrate	[N <sub>2</sub> H <sub>5</sub> ] <sup>+</sup> [NO <sub>3</sub> ] <sup>-</sup>	95.06	-211.36 <sup>[19]</sup>
Ionic Fuel			
AA, ammonium azide	[NH <sub>4</sub> ] <sup>+</sup> [N <sub>3</sub> ] <sup>-</sup>	60.06	113.66 <sup>[19]</sup>
HA, hydrazinium azide	[N <sub>2</sub> H <sub>5</sub> ] <sup>+</sup> [N <sub>3</sub> ] <sup>-</sup>	75.07	228.53 <sup>[19]</sup>
HEHN, 2-hydroxyethyl-hydrazinium nitrate	[HO-C <sub>2</sub> H <sub>4</sub> -N <sub>2</sub> H <sub>4</sub> ] <sup>+</sup> [NO <sub>3</sub> ] <sup>-</sup>	139.11 <sup>[22]</sup>	-388.69 <sup>[23]</sup>
Molecular Fuel			
MMF, mono-methylformamide	CH <sub>3</sub> HNCHO	59.067	-247.4 <sup>[21]</sup>
DMF, di-methylformamide	(CH <sub>3</sub> ) <sub>2</sub> NCHO	73.094	-239.3 <sup>[24]</sup> as cited in <sup>[21]</sup>
Methanol	CH <sub>3</sub> OH	32.04	-238.77 <sup>[19]</sup>
Ethanol	CH <sub>3</sub> CH <sub>2</sub> OH	46.07	-277.755 <sup>[19]</sup>
Glycerol	(CH <sub>2</sub> OH) <sub>2</sub> CHOH	92.094	-669.6 <sup>[25]</sup>
Glycine	NH <sub>2</sub> CH <sub>2</sub> COOH	75.07	-528.0 <sup>[26]</sup>
Urea	CO(NH <sub>2</sub> ) <sub>2</sub>	60.06	-333.43 <sup>[19]</sup>

**Table 2.** Composition of US Army liquid gun propellants (LGPs) <sup>[27][28]</sup>.

Propellant	Component, wt%			
	HAN	TEAN	DEHAN	Water
	[NH <sub>3</sub> OH] <sup>+</sup> [NO <sub>3</sub> ] <sup>-</sup>	[NH(C <sub>2</sub> H <sub>4</sub> OH) <sub>3</sub> ] <sup>+</sup> [NO <sub>3</sub> ] <sup>-</sup>	[(CH <sub>3</sub> CH <sub>2</sub> )HNOH] <sup>+</sup> [NO <sub>3</sub> ] <sup>-</sup>	H <sub>2</sub> O
LP1846	60.8%	19.2%	0.0%	20.0%
LP1845	63.2%	20.0%	0.0%	16.8%
LP1898	60.7%	0.0%	19.3%	20.0%

AF-M315E when decomposed produces an adiabatic flame temperature around 2100 K, which is much higher than that of hydrazine (nearly 1200 K). AF-M315E offers 13% increase of specific impulse and 63% increase in density over hydrazine <sup>[12]</sup>, which makes it superior in the miniaturization of propulsion systems over the latter. The propellant possesses high solubility and negligible vapor-pressure of all its solution constituents, thus promoting low toxicity hazards and high mixture stability at various temperature levels, which makes exposure in open environments have no safety issues <sup>[29]</sup>. An advantage AF-M315E possesses over current state-of-the-art green propellants is its maturity. Thorough development has taken place to reach this product and be able to test in space on 1 N and 22 N thrusters through the Green Propellant Infusion Mission (GPIM) launched in 2019 <sup>[30]</sup>. However, a disadvantage over the latest state-of-the-art green propellants rises from the relatively high flame temperature, which makes it difficult to rapidly manufacture an economic and simpler design of thrusters especially for the micro/nano satellites industry. It is worthy of mentioning that current advancements, especially related to rapid prototyping, in low-cost thrusters of small spacecrafts would benefit greatly from using additive manufacturing techniques such as metal 3D printing. Such techniques facilitate the design process and reduce the build time, they typically use metal alloys such as Ti-6Al-4V (Ti64) and Inconel<sup>®</sup>-625 (nickel-

chromium superalloy) with melting points of approximately 1900 and 1570 K, respectively [31][32]. Catalytic decomposition of AF-M315E requires higher preheating temperature, compared to hydrazine, where it typically consumes up to 15 kJ of energy [33] and the catalyst bed preheating nominal start temperature is 315 °C [34], while Busek Co. Inc researchers reported successful ignition at 400 °C preheating temperature [12].

SHP163 is another very interesting HAN-based green propellant, which was being developed since the year 2000 at the Institute of Space and Astronautical Science (ISAS)/(JAXA). SHP163 is composed of 73.6 wt% HAN, 3.9 wt% AN, 16.3 wt% methanol, and 6.2 wt% water [35]. This propellant has density of 1.4 g cm<sup>-3</sup> yet achieves high volumetric specific impulse  $\rho_{isp}$  of 396 g s cm<sup>-3</sup>, which is higher than AF-315E (at 0.7 MPa chamber pressure and 50:1 nozzle expansion ratio at frozen conditions) [35]. The flame temperature is considered very high, as it records about 2400 K [35][36]. As SHP163 shows to be one of the most energetic propellants for use in a thruster, it demonstrates operational stability and shows enough safety levels to be accepted as a green and safe liquid propellant [37]. SHP163 is only ignitable using a preheated catalyst-bed under 1.0 MPa [38][39]. Finally, SHP163 was tested in space in the Green Propellant Reaction Control System (GPRCS) utilizing a 1 N class thruster in the RAPIS-1 satellite launched in 2019 by JAXA.

HNP-family (High-performance Non-detonating Propellant) are HAN/HN-based green propellants that have been under development for over 10 years by IHI Aerospace co. in Japan. This green monopropellant family include HNP209, HNP221, and HNP225, and they are formulated from HAN, HN, methanol, and water [32]. They all possess volume specific impulse ( $\rho_{isp}$ ) superior to hydrazine, but what characterizes them most is their relatively low adiabatic flame temperature compared to other energetic ionic liquid monopropellants such as AF-M315E and SHP163. HNP209 typically has a theoretical specific impulse around 260 s with the highest combustion temperature (~1900 K), while HNP221 and HNP225 have specific impulse of 241 and 213 s, respectively (at chamber pressure of 1.0 MPa and expansion ratio of 100:1) [40][41][42], as shown in Table 3.

**Table 3.** Physical properties and performance (@ 1 MPa chamber pressure, 100:1 expansion ratio, and vacuum conditions, using NASA Chemical Equilibrium Analysis (CEA) [31] and verified from [4][29][32][35][36][43][44][45]).

Properties	Hydrazine	AF-M315E	SHP163	HAN/HN-Based	
				HNP221	HNP225
Theoretical Specific Impulse $I_{sp}$ (s)	239	260-270	276	241	213
Density $\rho$ (g cm <sup>-3</sup> ) (@ 20 °C)	1.0	1.47	1.4	1.22	1.16
Volumetric Specific Impulse $\rho_{isp}$ (g s cm <sup>-3</sup> )	239	~390	386	294	247
Adiabatic Flame Temperature (K)	1170	2166	2401	1394	990
Freezing Point (°C)	1.5	<-80	≤-30	≤0	≤-10

HNP225 is the one among the family with the least adiabatic flame temperature around 1000 K (even less than hydrazine ~1200 K), while HNP221 is approximately 1400 K [32][42]. The low temperature combustion gasses allowed IHI Aerospace co. to develop low-cost thrusters since the need for high heat resistant materials or complex cooling for the thruster's combustion chamber is no longer required. The HNP2xx family of propellants are ignited using catalytic decomposition. Igarashi et al. 2017 [32] performed tests for HNP221 and HNP225 with newly developed catalysts showing excellent response and combustion pressure stability compared to hydrazine, either in continuous mode or pulsed mode operation, with preheating temperatures starting from 200 and 300 °C for HNP221 and HNP225, respectively.

GEM or the Green Electrical Monopropellant is a novel HAN-based energetic ionic liquid composed of HAN, AN, (2,2'-dipyridyl), (1,2,4-triazole), 1H-pyrazol, and water [46]. GEM is a proprietary of Digital Solid-State Propulsion company (DSSP) [46] and is developed as a superior replacement for AF-M315E [45]. This propellant is demonstrated on a lab-scale to be capable of taking place in a multi-mode propulsion system. A "multi-mode" system is where a propulsion system in a satellite can operate as two or more separate modes (e.g., chemical high-thrust mode and electric high-specific impulse mode) under a condition of using a shared propellant tank [47] as cited in [48]. The most appealing properties in GEM is that it can also be electrically ignited without the use of any heavy catalytic beds, and it possesses a significantly large volumetric specific impulse as compared to AF-M315E and the ADN-based LMP-103S green monopropellants [45], refer to Table 4.

**Table 4.** Performance and physical properties of Green Electric Monopropellant (GEM) compared to state-of-the-art Green Monopropellants (@ 2.0 MPa chamber pressure, 50:1 expansion ratio, and vacuum conditions) [45].

Properties	Hydrazine	LMP-103S	AF-M315E	GEM
Theoretical Specific Impulse $I_{sp}$ (s)	236	252	266	283
Density $\rho$ (g cm <sup>-3</sup> ) (@ 20 °C)	1.0	1.24	1.47	1.51
Volumetric Specific Impulse $\rho I_{sp}$ (g s cm <sup>-3</sup> )	236	312.48	391	427
Vapor Pressure $P_v$ (kPa) (@ 25 °C)	1.91	15.1	1.4	<1
Toxicity	High	Moderate	Low	Low

ADN (ammonium dinitramide)-based green propellants development started at the Swedish Defense Research Agency (FOI) in Europe in 1997 [21][49][50]. The ADN-based monopropellants family mainly consists of FLP-103, 105, 106, 107 and LMP-103S, where the latter was developed by Bradford ECAPS Co. LMP-103S and FLP-106 are the most mature, and the former was qualified by the European Space Agency (ESA) and in-space demonstrated through the High Performance Green Propulsion system (HPGP) on the Mango-PRISMA satellite launched in June 2010 [51][52][53]. Different fuels were used within this energetic ionic liquid mixture such as methanol, monomethyl-formamide MMF and dimethyl-formamide DMF. However, methanol was found incompatible with ADN unless by addition of ammonia (NH<sub>3</sub>) in order to increase the pH of the mixture [21][54]. Composition of some ADN-based monopropellants are shown in Table 5 where the performance of the FLP-family is shown to be higher than LMP-103S. However, all ADN-based monopropellants mentioned in this study possess volumetric specific impulse lower than that of AF-M315E (391 g s cm<sup>-3</sup>). ADN-based green monopropellants are not only ignited by preheated catalytic beds, same as all monopropellants, but can also be ignited electrically or using thermal ignition. Larsson et al. [55] found that ADN-based propellants can be ignited using resistive heating by conducting electrical current through the propellants, and very rapid ignition was obtained (less than 2 ms); moreover, the least amount of electric energy utilized for successful ignition was in terms of (20 J). While Wilhelm et al. [52] found that glow-plug ignition was successful for LMP-103S and FLP-106 with satisfying ignition behavior and decomposition. Advantages of the LMP-103S and FLP-family over AF-M315E include, but are not limited to, lower combustion temperature, which allows using materials with lower melting point, and simpler designs for thruster development. Moreover, flexibility in using different ignition techniques and not just being restricted to catalytic decomposition of ADN-based green monopropellants would allow for development of novel designs of monopropellant thrusters.

**Table 5.** ADN-based monopropellants properties [21][56][57] (ideal vacuum  $I_{sp}$  by [56] using NASA CEA @ 2.0 MPa chamber pressure, 50:1 expansion ratio assuming frozen condition [52]).



Propellant.	Formulation	Theoretical $I_{sp}$ (s)	Density $\rho$ (g cm <sup>-3</sup> ) *	$\rho I_{sp}$ (g s cm <sup>-3</sup> )	$T_c$ (°C)
LMP-103S	(1) 63.0% (2) 18.4% (6) 18.6%	252	1.24	312.48	1630
FLP-103	(1) 63.4% (2) 11.2% (5) 25.4%	254	1.31	332.74	1760
FLP-106	(1) 64.6% (3) 11.5% (5) 23.9%	255	1.357	344.6	1814
FLP-107	(1) 65.4% (4) 9.3% (5) 25.3%	258	1.351	348.5	1869

(1) ADN. (2) Methanol. (3) MMF. (4) DMF. (5) Water. (6) Ammonia (aq. 25% concentration). \* @ 20 °C.

## 2.2. Liquid NOx Monopropellants

In this section green monopropellants of N<sub>2</sub>O, nitro compounds, and premixed N<sub>2</sub>O with hydrocarbons (i.e., nitrous oxide fuel blends NOFB [43][58]) will be introduced, emphasizing on relevant properties needed for propulsion system design either for small satellites or high-thrust in-space propulsion up to nearly 600 N.

Generally, the binary compounds of nitrogen and oxygen (e.g., NO, NO<sub>2</sub>, N<sub>2</sub>O, N<sub>2</sub>O<sub>3</sub>, N<sub>2</sub>O<sub>4</sub>) have been considered as oxidizers in hybrid rocket engines and bipropellant systems [59]. Nitric oxide NO is gaseous in the typically requested temperature-pressure range of [−30, +80] °C and [0.1, 3] MPa, which does not suit the propulsion applications of concern, since the sought propellant is required to be in liquid phase and easily storable in lighter weight tanks within the given operation conditions. Although NO<sub>2</sub>, N<sub>2</sub>O<sub>3</sub>, and N<sub>2</sub>O<sub>4</sub> form an equilibrium mixture within the mentioned temperature–pressure range, they are still highly toxic and considered as GHS class 1 acute-toxicity [6]. Nitrous oxide N<sub>2</sub>O is the only compound in this group that falls under green propellant umbrella since it is considered GHS class 5 relatively nontoxic, moreover, the critical point stands at 36.4 °C and 7.24 MPa [60] and is liquid in a wide part of the pressure–temperature range mentioned above. N<sub>2</sub>O possesses good storability characteristics at room temperature especially for long term storage since it does not have decomposition or boiling problems when compared to H<sub>2</sub>O<sub>2</sub> or cryogenic LOX as examples. At 20 °C the saturated vapor pressure of nitrous oxide is ~5.2 MPa, which is high compared to EILs and HPAS, but still considered favorable when considering this propellant for self-pressurizing feed systems.

N<sub>2</sub>O (liquid) storage density is ~0.745 g cm<sup>-3</sup> at 20 °C and ~5.2 MPa vapor pressure [61]. Thus, with suitable pressure vessels N<sub>2</sub>O can be kept under stable and readily-operating conditions, adding to that, its material compatibility with common tank materials including metals, plastics and composite-materials [59]. Although N<sub>2</sub>O as monopropellant has lower performance than most EILs, it has an experimental result  $I_{sp} = 206$  s ( $T_c = 1913.15$  K, @  $p_c = 0.3$  MPa, nozzle expansion 200:1 [62]), which is higher than high-test peroxide HTP ~180 s [61]. The most compelling about N<sub>2</sub>O for modern propulsion system design is that it can be used in the so called “multi-mode” propulsion system [47], where it can act as a propellant for cold-gas, monopropellant propulsion, and/or bipropellant systems while sharing the same propellant tank.

Nitromethane (NM, CH<sub>3</sub>NO<sub>2</sub>) is a promising nitro compound green monopropellant candidate for modern in-space propulsion systems for relatively low- to high-thrust range. NM is a relatively nontoxic (GHS class 4 toxicity), viscous, flammable liquid with density of 1.1371 g cm<sup>-3</sup>, and has a freezing temperature of −28.4 °C [63]. It shows good storability for in-space applications and by adding stabilizer additives (such as, ditertiarybutyl peroxide or chloral, and diacetyl [64]) it could be a highly-attractive liquid monopropellant [65]. Nitromethane can be considered in “multi-mode” systems since it can be used in both mono- and bipropellant propulsion systems, thus increase the overall system optimization. NM as a green monopropellant, possesses high performance ( $I_{sp} = 289$  s) [3], high volumetric specific impulse, and combustion temperature around 2449 K.

NOFB nitrous oxide fuel blends were studied as monopropellants since World War II by the Germans [66]; however, their utilization and development were ceased due to the availability of hydrazine since the 1960s until recently. Because of the current economic and political reforms to greenify the use of propellants, nitrous oxide fuel blends research has been revived in Europe by the European Fuel Blend Development (EUFBD) program carried between 2015–2018 [67].

NOFBX<sup>TM</sup> monopropellant was invented in the U.S. between 2005 and 2007 under the NASA Mars Advanced Technology program, and later a proprietary of Firestar company [68]. NOFBX<sup>TM</sup> was demonstrated in the 0.4 N–445 N thrust range with measured specific impulse performance around 325 s, while the theoretical value was ~345 s and chamber temperature ~3200 K at 0.7 MPa with stoichiometric O/F = 3 (or 25%fuel). It is storable as saturated liquid under a wide range of temperatures, with a critical point of 39.48 °C and 7.19 MPa [69].

HyNOx or Hydrocarbon NOx as denoted by German scientists in the German Aerospace Center (DLR) are premixed monopropellants where oxidizer and fuel are premixed in a single tank [70]. HyNOx fuel blend used nitrous oxide with ethylene (i.e., ethene IUPAC name) (N<sub>2</sub>O/C<sub>2</sub>H<sub>4</sub>) due to the similar vapor pressure of the two compounds [71] and has density of 0.879 g cm<sup>-3</sup>. The recorded theoretical vacuum specific impulse is 303 s obtained with O/F ratio = 6 (~14.29%fuel) under 230 K and 2.5 MPa temperature and pressure [43]. A high combustion temperature up to 3264 K is also recorded as a drawback besides the important need for a flashback-arrestor, by Werling et al. [72].

Nitrous oxide/ethanol (N<sub>2</sub>O/C<sub>2</sub>H<sub>5</sub>OH), another hydrocarbon NOx blend, was the mixture of choice after a tradeoff study carried by Mayer et al. [73] in the EUFBD program. Among hydrocarbons, ethanol showed better ignitability and moderate flame temperature, and further demonstrates stability and miscibility with nitrous oxide. Physical properties of the selected mixture resulted in saturated liquid density of 0.892 g cm<sup>-3</sup> with a stoichiometric O/F ratio of 5.73 (~14.86%fuel), the critical point stood at 36.45 °C and 6.3 MPa and the vapor pressure at bubble point was 2.6 MPa, while a theoretical specific impulse  $I_{sp} = 331$  s and combustion temperature 3093 K were reported [73]. A test campaign was carried out with a 600 N thruster and specific impulse of 259 s was achievable [74]. Drawbacks reported during this study are: high combustion temperature, which requires complex engine design and active cooling system, incompatibility of the nitrous oxide fuel blend with titanium was expected to exist, possibility of flammable vapors in the propellant tank, and low density at practical storage temperatures [71][73][74]. Advantages of nitrous oxide fuel blends, as most green monopropellants, are nontoxic and noncarcinogenic nature, low freezing point, higher specific impulse than hydrazine, and the most prominent advantage is self-pressurization capabilities, which allows for simple feed-system and tank-pressurization system design. Performance and properties of liquid NOx monopropellants are shown in Table 6.

**Table 6.** Ideal (vacuum) performance and physical properties of the Liquid NOx Monopropellants class (compounds and premixed fuel blends).

Propellant	Theoretical $I_{sp}$ (s)	$\rho$ (g cm <sup>-3</sup> ) <sup>a</sup>	$\rho I_{sp}$ (g s cm <sup>-3</sup> )	$T_c$ (°C)
N <sub>2</sub> O (liquid) *	206	0.745	153.5	1640
Nitromethane **	289	1.1371	328.6	2175.85
NOFBX <sup>TM</sup> ***	350	0.700	245	2926.85
HyNOx (Ethene) †	303	0.879	266.3	2990.85
NOx/Ethanol ‡	331	0.892	295.3	2819.85

<sup>a</sup> @ 20 °C. \* @  $p_c = 0.3$  MPa. \*\* @  $p_c = 1$  MPa. \*\*\* @  $p_c = 0.7$  MPa and stoichiometric O/F = 3. † @  $p_c = 2.5$  MPa and stoichiometric O/F = 6. ‡ @  $p_c = 1$  MPa and stoichiometric O/F = 5.73.



### 2.3. Hydrogen Peroxide Aqueous Solutions (HPAS)

Hydrogen peroxide ( $\text{H}_2\text{O}_2$ ) has been used as monopropellant in different aerospace applications since 1938 [75].  $\text{H}_2\text{O}_2$  is type-classified according to its concentration in aqueous solution, and grade-classified according to the concentration of stabilizers and impurities [76], as shown in Table 7. HTP (High-Test Peroxide) is a highly concentrated  $\text{H}_2\text{O}_2$ , greater than >85% weight concentration. Rocket grade HTP is used in space propulsion for low and medium thrust applications and is typically of 98% concentration [77]. The high density of 98% HTP ( $\sim 1.43 \text{ g cm}^{-3}$ ), and the nontoxic nature, makes it an interesting candidate for storage in propulsion systems in general. In monopropellant systems it can catalytically decompose reaching temperatures in terms of 1222 K [77]. Performance of HTP 98% in monopropellant systems is  $\sim 20\%$  less than hydrazine [78], with  $I_{sp} \sim 186 \text{ s}$  (at 1 MPa and 50:1 expansion conditions). However, in bipropellant systems, especially with hydrocarbons such as ethanol, it can reach  $I_{sp} > 325 \text{ s}$  (combustion temperature 2752 K) [3], which makes it a very competitive propellant for this kind of propulsion system. Another concentration of hydrogen peroxide that is widely used is the HTP 87.5% with a density of  $\sim 1.38 \text{ g cm}^{-3}$  at 20 °C and possesses a theoretical specific impulse of  $\sim 144 \text{ s}$  when evaluated at a chamber pressure of 1 MPa and  $A_e/A_t$  of 7.841 at sea level [79].  $\text{H}_2\text{O}_2$  at concentrations of 90% and 85% were simulated on RPA to give, respectively, theoretical vacuum specific impulse of 172.13 and 150.47 s, chamber temperature 1019.3 and 892.65 K at 1 MPa, and expansion ratios of 40:1 and 10:1 applying the shifting equilibrium model for the whole nozzle.

**Table 7.** Physical and chemical properties of  $\text{H}_2\text{O}_2$  propellant with different concentrations [76][80].

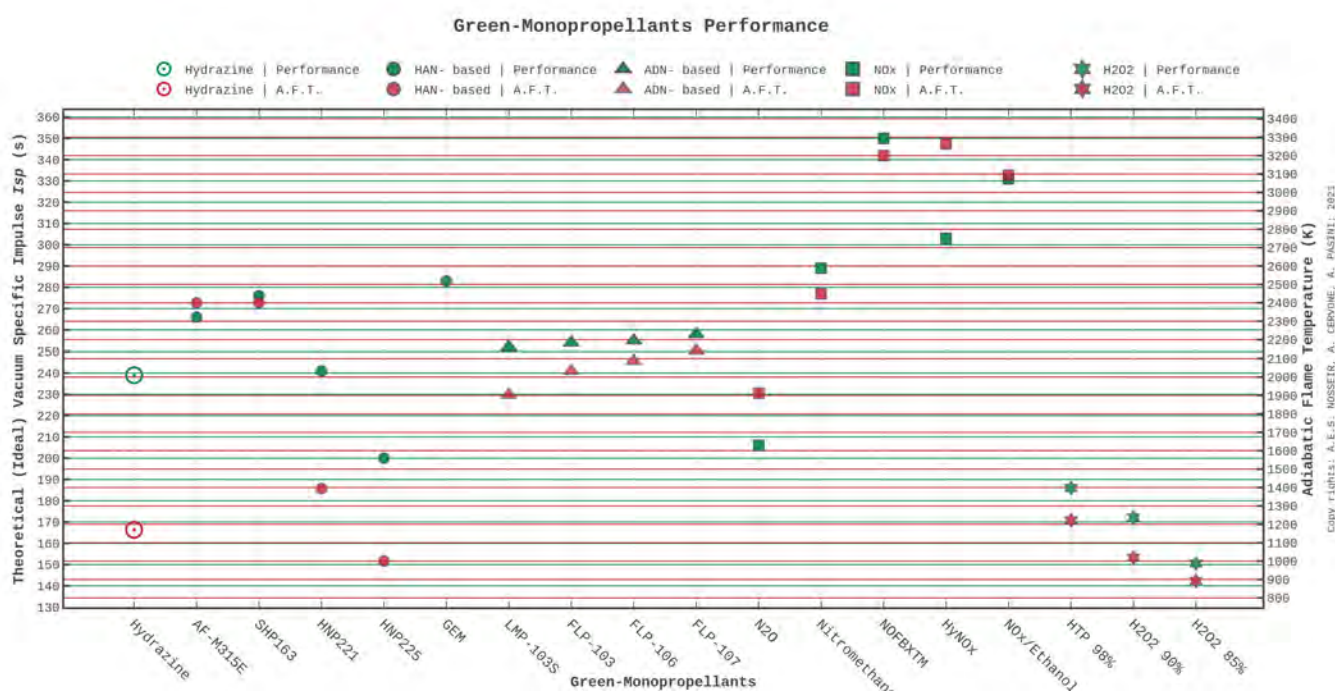
Properties	H <sub>2</sub> O <sub>2</sub> Propellant Classification				
	Type 70	Type 85	Type 90	Type 98	
	Grade ES	Grade ES	Grade ES	Grade HP	Grade HP
Concentration %	71.0–73.0	85.0–87.0	90.0–91.5		98.0–99.0
Stability *	$\leq 2\%$				
Density $\rho$ (g cm <sup>-3</sup> ) **	$\sim 1.29$	$\sim 1.34$	$\sim 1.40$		$\sim 1.43$
Freezing point (°C)	−40	−17	−12		−2
Boiling Point (°C)	125	137	140		147

ES: Extra Stabilizers; HP: High Purity; \* (24 h/100 °C) %Loss of active O<sub>2</sub>; \*\* @ 20 °C.

The Hydrogen Peroxide Aqueous Solutions (HPAS) class possesses the lowest performance values among green monopropellants; however, a unique characteristic of this family of propellants can make it of high interest from the point of view of rocket propulsion designers in terms of increasing the overall system performance and size optimization. This unique characteristic is the hypergolic ignition of hydrogen peroxide with various propellants making it a distinguishable candidate for in-space propulsion systems. Hydrogen peroxide has been experimented thoroughly for hypergolic ignition with hydrocarbons such as ethanol [81] and propyne [82]. Further, hypergolic ignition with ionic-liquid fuels such as 1-ethyl-3-methyl imidazolium cyanoborohydride ([EMIM][BH<sub>3</sub>CN]) [83] and 1-allyl-3-ethyl imidazolium cyanoborohydride ([AEIM][BH<sub>3</sub>CN]) [84] would allow for developing new generations of green propellants for effective in-space bipropellant propulsion systems, namely green hypergolic ionic liquids (HILs) [85].

Undoubtedly, the utilization of hydrogen peroxide as a well-studied and experimented propellant in modern green propulsion systems can be of great benefits to the overall system optimization and overall increase in performance, since it would allow for implementation of multi-mode [48] systems where two or more propulsion systems can rely on the same propellant tank. In a multi-mode system, hydrogen peroxide can decompose catalytically in monopropellant auxiliary propulsion (e.g., reaction control

system RCS or attitude control system ACS), as well as being used as an oxidizer in bipropellant primary propulsion system as it distinguishably can ignite through a hypergolic reaction without the need for separate ignition power source. A recent research work by Rhodes and Ronney (2019, 2020) theorized [86], investigated, and experimented [87] a first prototype of a Hydrogen Peroxide Vapor Propulsion, which can provide attitude control capability for small satellites with thrust range of millinewtons. Although hydrogen peroxide in vapor phase is extremely unstable and prone to detonation as widely known, in this novel concept it is claimed that the reactive vapor phase was utilized within a low-thrust propulsion system where the vapor was used as a propellant. Hydrogen peroxide reactive vapor was vacuum evaporated from the surface of the stored propellant in liquid phase and then passed over a catalytic bed where a chemical reaction occurs and temperature is increased; this eventually leads to a theoretical specific impulse in vacuum ( $>200$  s), which is very high when compared to conventional  $\text{H}_2\text{O}_2$  systems. This proposed novel system would add to the scope of applications of hydrogen peroxide aqueous solution propellants.



**Figure 1.** State-of-the-art green monopropellants performance chart.

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## Keywords

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green propellant;monopropellant;chemical rocket propulsion;small satellites;CubeSat;in-space propulsion;liquid propulsion system

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