Experimental Characterization of a 5 N Hydrogen Peroxide Monopropellant Thruster Prototype

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In the framework of the LET-SME program funded by the European Space Agency, ALTA S.p.A. (Italy) and DELTACAT Ltd. (United Kingdom) jointly investigated the use of advanced catalytic beds on ceramic supports as a cost-effective alternative to metal screen reactors for the decomposition of high-concentration hydrogen peroxide in small monopropellant rockets. To this purpose ALTA S.p.A. designed and realized a reconfigurable test bench for the characterization of the operation and propulsive performance of small rocket thrusters. The present paper illustrates the experimental campaign carried out on a 5 N thruster prototype operating with two platinum catalysts on γ–alumina supporting spheres, especially developed by ALTA in collaboration with the Chemistry and Industrial Chemistry Department of Pisa University, Italy. The results indicated that Pt/Al_2O_3 is an effective catalyst combination for the decomposition of 87.5% propellant grade hydrogen peroxide, with good stability and performance comparable to silver screen beds of equal geometric envelope and operational conditions. Incomplete hydrogen peroxide decomposition and the onset of flow oscillations in the reactor were observed at the tested levels of bed loading, residence time and flow pressure. Thermal stresses due to the large temperature gradients occurring during the decomposition of high grade hydrogen peroxide (87.5% by weight) caused the ceramic pellets to break and the progressive occlusion of the bed. Based on the analysis of the test results, several ways to overcome these problems in future investigations have been tentatively identified, together with the necessary modifications to the present experimental set-up.

Nomenclature

A_t	=	throat area
C_F	=	thrust coefficient
c^*	=	characteristic velocity
F	=	thrust
g_o	=	sea level gravity acceleration

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I_{sp} = specific impulse

$\dot{m}$ = propellant mass flow rate

p_a = ambient pressure

p_c = combustion chamber pressure

R = gas constant of the exhaust gases

T_{ad} = adiabatic decomposition temperature

T_{amb} = ambient temperature

T_c = combustion chamber temperature

T_{exp} = decomposition temperature (experimentally measured)

γ = specific heat ratio of the exhaust gases

η_{c^*} = characteristic velocity efficiency

$\eta_{\Delta T}$ = temperature efficiency

I. Introduction

IN the last decade there has been a renewed interest in hydrogen peroxide engines, generated mainly by the need for less toxic and dangerous propellants. As a green propellant, hydrogen peroxide allows for a drastic simplification of the health and safety protection procedures needed in its production, storage and handling. As a result, attractive savings can be obtained in low or medium thrust rocket engines for small missions, where the overall cost of propulsion systems with traditional propellants does not scale down proportionally to the rocket size.

The nominal propulsive performance of hydrogen peroxide as a monopropellant is about 20% lower than hydrazine, but the volume specific impulse achievable with 90% H_2O_2 is higher than for most other propellants due to its high density. This is particularly useful for systems with significant aerodynamic drag losses or stringent volume constraints, as is often the case for small satellites. Finally, when used in bipropellant and hybrid rocket engines, hydrogen peroxide yields specific impulses comparable to other liquid oxidizers like dinitrogen tetroxide, nitric acid and even liquid oxygen (Wernimont and Muellens [1], Wernimont and Garboden [2]). Furthermore, with respect to other high-energy green propellants, like ADN, HAN and HNF (Wucherer et al.[3], Schoyer et al.[4]), hydrogen peroxide has the significant advantage that its decomposition temperature does not require the use of extremely expensive materials and manufacturing processes for the thrust chamber.

The earliest research on hydrogen peroxide-based rockets was conducted by Walter[5] in Germany during the 1930s. He pioneered the idea of using a catalytic bed to generate a hot stream of oxygen and steam, which could either be directed to an exhaust nozzle or used to auto-ignite a fuel. These early engines were used in the ME163 fighter plane and to drive the V2 turbine pump. Interest in hydrogen peroxide in the post-war years up to about 1955 was confined largely to the United Kingdom, although both the United States and the former Soviet Union developed the Walter concept for different applications. The US X-1 and X-15 space planes, together with the early Mercury and Gemini manned spacecrafts, used hydrogen peroxide in their reaction control systems. Following these early developments, a significant amount of work on hydrogen peroxide decomposition and its application to monopropellant rockets was carried out in the 1960s at NASA laboratories[6,7], but this effort was subsequently abandoned with the advent of effective hydrazine catalysts and engines. On the other hand, the Russian Soyuz launch vehicle, which has been in active service for over forty years, continues to rely on hydrogen peroxide in its gas generator to drive the main turbine pump and in the RCS thrusters used for the descent phase. Much of the British work on hydrogen peroxide (Rothmund & Harlow[8]) was concerned with rocket-assisted take-off engines for rapid ascent and short runways. This led to several production engines including the mono-propellant De Havilland Sprite and the Screamer, dating back to 1956. Between 1953 and 1957 a British company, Saunders Roe, developed a high performance bipropellant peroxide engine, known as the Spectre. This was used in test flights of the British jet fighter SR-53. The highly reliable Gamma 301 engine, developed by Bristol Siddeley in the 1950s, was used for both the first and second stages of the Black Arrow satellite launcher, which flew from Woomera, Australia in 1971. This engine provided a vacuum specific impulse of 265 seconds and used hydrogen peroxide at a concentration of 86%.

The most significant technology challenge for the realization of hydrogen peroxide monopropellant thrusters is the development of effective, reliable, long-lived catalytic beds, providing fast and repeatable performance, insensitivity to poisoning by the stabilizers and impurities contained in the propellant, capable of sustaining the large

number of thermal cycles imposed by typical mission profiles and not requiring (if possible) pre-heating for efficient operation. Nowadays the most used catalyst materials for H_2O_2 are metallic silver (Runckel, Willis, and Salters [6]; Morlan et al.[9]), permanganates of alkali metals as in Musker[10] and manganese oxides (typically MnO_2 and Mn_2O_3). Some experience is also available with alumina-deposited platinum, ruthenium dioxide, divanadium pentoxide and lead oxide, as in Rusek[11]. None of these solutions is free from drawbacks, the most important being temperature limitations and poisoning for metallic silver (Wernimont and Mullens[12]; Ventura and Wernimont[13]), powdering and thermomechanical resistance for ceramic-deposed catalysts, excessive flow resistance for pellet beds, and flow stratification for channel matrix support catalysts (Beutien et al.[14]).

Along with the application of hydrogen as a monopropellant, there was also an evolution of the methods used for catalyst decomposition and development. The work-horse catalyst for 90% hydrogen peroxide has long been the silver screen pack[13]. This catalyst proved to be a robust choice, but its relatively low melting temperature limits its use to hydrogen peroxide concentrations lower than 92% ca. The main reason for using higher peroxide concentrations is the increase of the ratio between the adiabatic decomposition temperature and the average molecular weight of the products, which is beneficial in attaining superior propulsive performance. In particular, raising the H_2O_2 concentration to 98% can increase the monopropellant specific impulse by approximately 10 to 20 seconds. On the other hand, 98% H_2O_2-kerosene rocket engines can start to rival conventional storable propellants, and this feature has prompted various programs, such as Future-X and SMV, to consider using H_2O_2 and particularly 98% H_2O_2 as a "green" oxidizer.

Earlier 98% HP catalytic beds, realized in the 1950s and 1960s, were based on the use of screens made of different materials, like high melting point silver alloys (silver palladium), platinum, palladium, iridium, ruthenium, manganese dioxide, cobalt. However, no one of these candidate materials provided really adequate performance. Alternative techniques for decomposing hydrogen peroxide greater than 90% are necessary in order to fully exploit the higher performance offered by 98% H_2O_2. Research for the development of advanced catalytic beds has focused on the characterization of catalytic materials in batch reactors and the evaluation of the overall performance of the catalytic beds in monopropellant thruster prototypes. Ceramic-deposed catalysts have mainly been developed because of their cost-effectiveness and potentially favorable thermo-mechanical properties.

Kappenstein et al.[15] have recently investigated the thermal decomposition and the hydrothermal reduction of different permanganate precursors of manganese oxide-based catalysts, finding that higher reaction surface area and activity are exhibited when using potassium permanganate rather than sodium permanganate. An extensive experimental study carried out by Rusek[11] indicated that catalysts based on MnO_2 or Mn_2O_3 on different ceramic pellets lead to an activity about one order of magnitude higher than obtained with silver. Other catalysts, like ruthenium dioxide, displayed activities about three times higher than manganese oxides. Moreover, a series of thermal tests in the same study showed that the activity of platinum on alumina is one order of magnitude higher than exhibited by manganese oxides-based catalysts. These findings, however, are not fully consistent with those obtained by Pirault-Roy et al.[16] who investigated the activity of platinum supported on silica, silver, iridium and platinum-tin or manganese oxides supported on alumina, observing that silver on alumina yielded the highest activity, followed by manganese oxides on alumina and by platinum on silica. Another experimental activity was carried out by Eloirdi et al.[17] using a constant-volume batch reactor. Two catalysts, manganese oxide and silver supported on alumina, were tested. The manganese oxide catalyst showed a better activity with a good repeatability after several firings, while the supported silver sample was less active and showed a slight loss of activity after the first firings.

Tian et al.[18] have investigated the performance of Ir/γ-Al_2O_3 catalyst for the decomposition of high concentration hydrogen peroxide in a monopropellant thruster, finding that catalyst oxidation and surface Sn poisoning are the main reasons for catalyst deactivation. Beutien at al.[14] have illustrated the evaluation of cordierite-based catalytic beds for 98% hydrogen peroxide. The most interesting characteristic of cordierite as a catalyst supporting material, together with its relatively good mechanical strength, is that it does not melt or break when exposed to 98% hydrogen peroxide decomposition. Furthermore, high-channel density catalytic beds tend to result in higher temperatures and a more complete decomposition with respect to low pore density ones (Long and Rusek[19]).

As a direct consequence of the renewed interest in the use of hydrogen peroxide by the rocket propulsion community, ALTA S.p.A. (Italy) and DELTACAT Ltd. (United Kingdom) jointly investigated the use of advanced catalytic beds for the development of hydrogen peroxide monopropellant rocket thrusters. This activity has been funded by the European Space Agency in the framework of the LET-SME program, supporting innovative projects carried out by small and medium European companies. The objective of the activity was the design and realization of two prototype thrusters (a 5 N and a 25 N one), where catalytic beds made of different catalyst materials and substrates could be installed and validated. The present paper illustrates the performance of Pt/γ-Al_2O_3 catalysts in the 5 N thruster prototype.

II. Test Apparatus

A. The Engine Test Bench and the Propellant Feed System

The experimental characterization of the 5N thruster prototype has been carried out using ALTA's Hydrogen Peroxide Thrusters Test Facility (HPTTF). It consists of a custom-made test bench and a propellant feed system, designed for providing hydrogen peroxide to the thruster prototypes.

A three-dimensional view and a picture of the test bench are shown in Fig. 1. The mobile part of the one degree-of-freedom dynamometric force balance for the measurement of the engine's axial thrust consists of an L-shaped cradle, obtained by bolting together an horizontal and a vertical plate by means of two lateral triangular ribs. The load cell for the measurement of the axial thrust is installed on the vertical plate. The engine prototype is mounted in the cradle, which has suitable provisions for the hydrogen peroxide feed line and for accommodating thrusters of different sizes. The engine cradle is connected to the upper supporting plate of the thrust balance by means of two flexures realized out of 0.05 mm thick stainless steel sheets. These flexures are designed for transmitting to the supporting structure all of the off-axis engine loads, while being extremely flexible in the axial direction in order to minimize their interference with the thrust measurement. For bigger thrusters it is possible to easily reconfigure the balance over a relatively wide range of thrust levels by simply mounting a cell with suitable load capacity and by changing the thickness of the flexures. The thrust balance is suspended to a stiff cantilever beam supported by a second vertical I-beam, which also mounts the stop plate and screw used as mating elements for adjusting the compression preload of the measuring cell. The test bench has been designed paying particular attention to the reduction of spurious additional forces interfering with the thrust measurement. The calibration showed that the relative error on the axial thrust introduced by the force balance (i.e. by hydrogen peroxide supply tubes, electrical connections of transducers, flexures etc.) is lower than 2%.
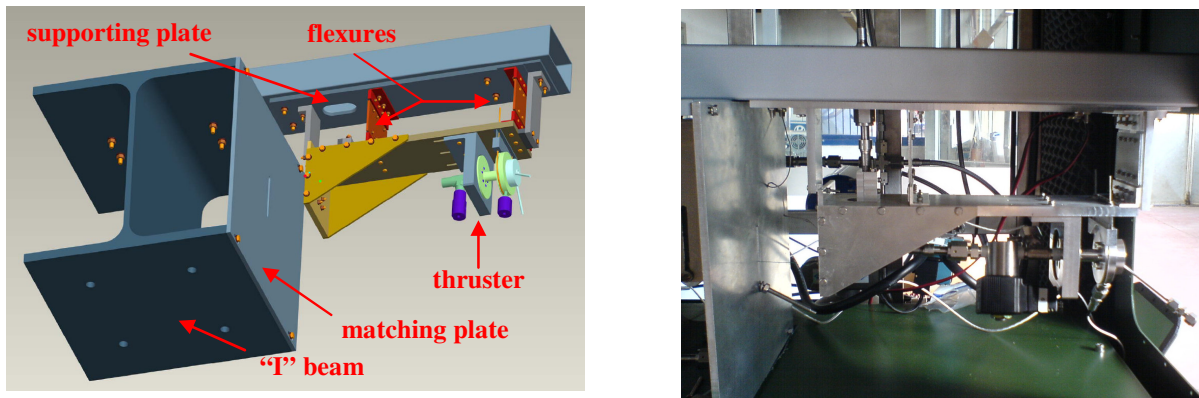

Figure 1. 3-D-drawing (left) and a picture of the test bench assembly (right).

The propellant feed system has been specifically designed for providing hydrogen peroxide to the prototype thrusters. This system is intended as a multi-purpose one, which can easily be adapted and reconfigured for testing thrusters of different operational characteristics (like bipropellants or hybrids) and target performance. A schematic of the propellant feed system is shown in Fig. 2. High-concentration hydrogen peroxide is stored at up to 40 bar pressure in the main 2.5 liters tank, made of stainless steel and internally coated with Teflon. If necessary, its storing capability can be increased by connecting it to a larger tank. In order to avoid dangerous overpressures in the case of uncontrolled hydrogen peroxide decomposition, the tank is connected to a manually operated safety valve, a burst-disk and a non-return valve. The physical conditions of the propellant stored in the tank are monitored by means of a thermocouple (6 mm diameter J-type mineral-insulated thermocouple, produced by Watlow) and a pressure transducer (PTU model produced by Swagelok with a 0-40 bar pressure range and an accuracy of 0.43% FSO). A check valve, a cavitating venturi (0.254 mm throat diameter) and a Coriolis flowmeter (Optimass MFS 7100 S04 produced by Krohne with a maximum operating pressure of 150 bar, a maximum flow rate of 100 kg/hr and an accuracy of 0.1% of the measured flow rate) have been placed along the hydrogen peroxide feeding line in order to, respectively, prevent flow reversals, regulate the flow by only adjusting the upstream pressure in the tank, and monitor the propellant mass flux with the required accuracy. For ease of reconfiguration, standard (1/4") PTFE-lined, stainless steel-braided hoses have been used for interfacing the main components of the feed system. Remotely-operated electro-pneumatic valves have been employed for operating the various fluid lines of the facility. A commercial solenoid valve with a low response time (<20 ms) has been chosen as firing valve. Gaseous nitrogen

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from a 200 bar bottle has been fed at different regulated pressures to the three lines used for propellant tank pressurization (40 bar), pneumatic valve actuation (7-8 bar), and for purging to a discharge tank and venting all of the hydrogen peroxide lines.

In order to prevent accidents, the detailed test procedure has been specified in written step-by-step instructions and ALTA's team has been equipped with the personal protective equipment (PPE) prescribed when handling high-grade hydrogen peroxide.
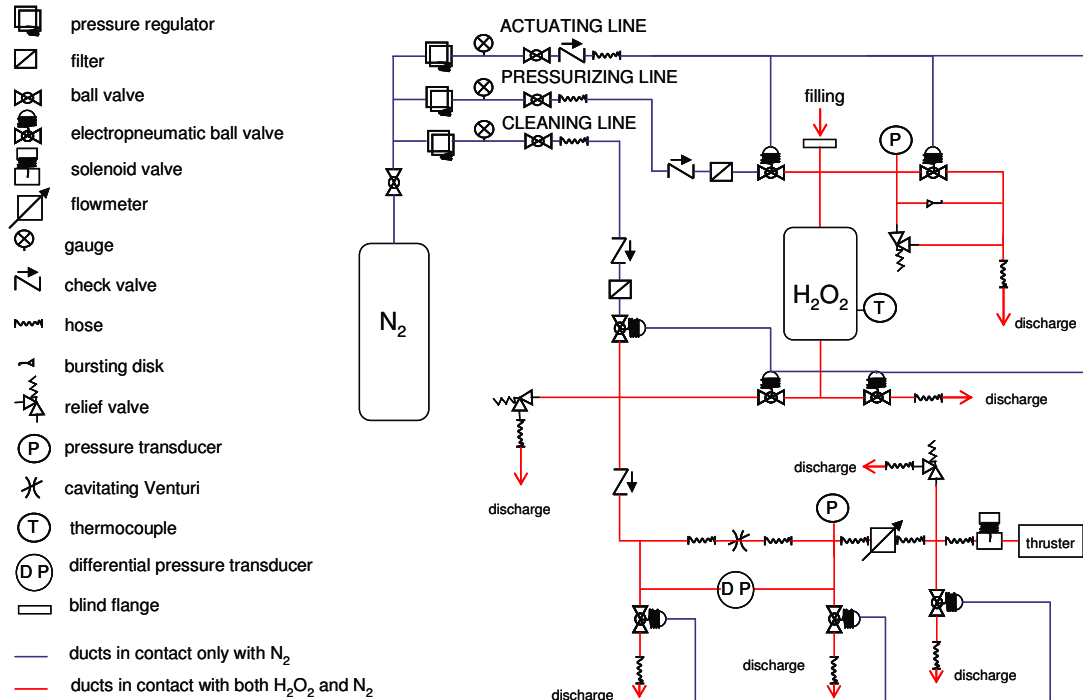
Figure 2. Schematic of the hydrogen peroxide supply facility

B. The 5 N Thruster Prototype

The thruster has been designed in a modular manner (Cervone et al.[21]) and consists of five main components in AISI 316L stainless steel (see Fig. 3): the catalytic bed, the nozzle, the connecting flange, the injection plate and the distribution plate. Sealing is obtained by means of copper-coated, inconel C–rings and two AISI 304 stainless steel screens (37x37 mesh size and 0.2 mm wire diameter) are used for retaining the catalyst pellets.

Sizing of the engine prototype and its components has been carried out using the standard simplified isentropic 1D relations presented in Ref. 21. The chamber pressure, the residence time of the propellant in the catalytic bed and the catalytic bed loading have been chosen based on the typical values reported in the open literature for similar applications. Originally the conical nozzle was designed for an exhaust pressure of 13800 Pa. Since the tests reported in this paper would have been carried out at atmospheric external pressure, the original nozzle has been shortened in order to avoid flow separation due to excessive over-expansion. Table 1 summarizes the main features of the 5 N tests thruster.

Table 1. Main characteristics of the test thruster.

|		Units	Values
Propellant	H2O2 content	g/100 g	87.5
	Tin content [ppm]	ppm	5-9
	phosphate content (as PO4)	ppm	<=0.5
Catalytic bed	inner diameter	mm	8
	length/diameter ratio	-	4
Nozzle	throat diameter	mm	2
	exhaust diameter	mm	3.73
	exhaust area ratio	-	3.48
	conical nozzle half angle	deg	15
	nozzle length	mm	3.23
Inj./dist. plates	diameter	mm	8
	diameter of holes	mm	1.5
	open area/total area ratio	-	0.5
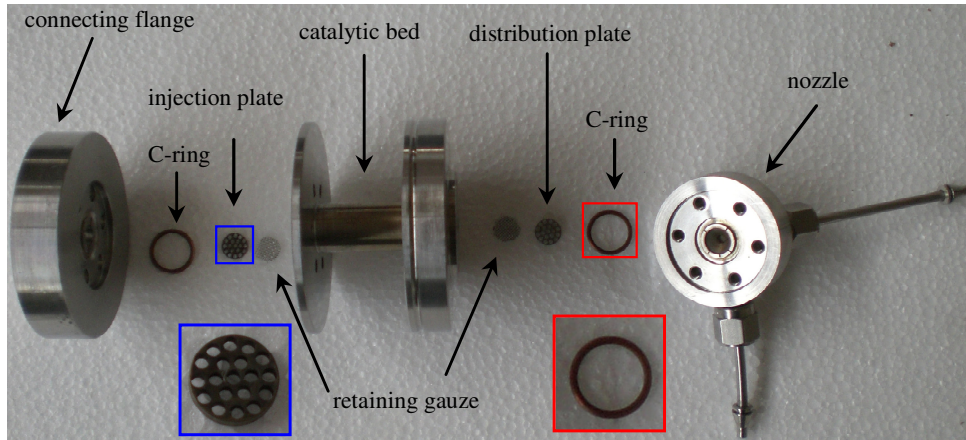
Figure 3. Main parts of the thruster prototype.

C. Measurements and Data Acquisition System

The measurements taken during a typical test can be divided into two different categories: measurements for monitoring the operation of the propellant supply system and measurements for evaluating the performance of the thruster prototype.

With reference to Fig. 2, three pressure transducers, one thermocouple and one flowmeter have been installed along the hydrogen peroxide line, in order to carry out the following measurements:

− the tank delivery pressure, which is directly related to the propellant mass flow rate as long as cavitating conditions are established in the venturi;
− the tank temperature, whose abnormal increase is usually an indication of incipient decomposition of the stored hydrogen peroxide;
− the venturi differential pressure, whose value is indicative of correct operation under cavitating conditions where the hydrogen peroxide mass flow rate only depends on the tank delivery pressure;
− the venturi outlet pressure, which gives information on the state of the fluid entering the flow meter and indirectly on the venturi inlet pressure;
− the propellant flow rate.

Figure 4 shows a schematic of the 5 N thruster prototype and the transducers arrangement as described in the previous subsection. In order to monitor the performance of the thruster prototype the following measurements are taken:

− the thruster inlet pressure, by means of a Kulite pressure transducer model XTM-190M-17 bar A, with a maximum combined error (non-linearity, hysteresis and repeatability) of 1% FSO, mounted using an adapter on the "T" junction placed just in front of the injection plate;
− the chamber pressure, by means of a second Kulite pressure transducer model XTM-190M-17 bar A mounted by means of a suitable Swagelok connector to the tube brazed on the converging portion of the nozzle;
− the chamber temperature, by means of a 0.5 mm diameter K-type mineral-insulated thermocouple inserted in the dedicated temperature tap;
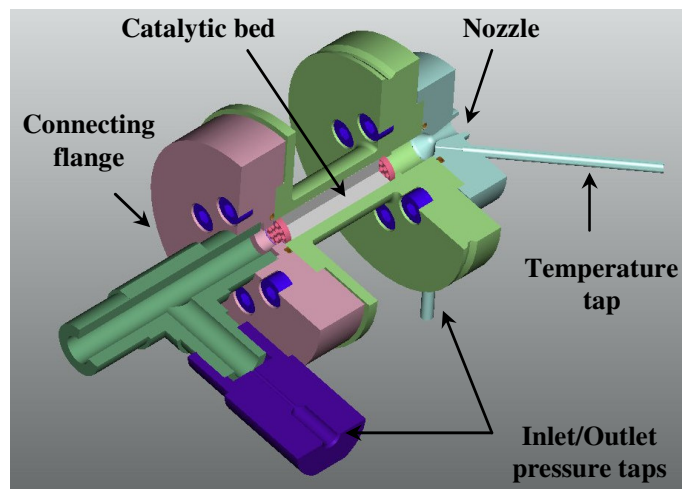− the thrust, by means of a subminiature compression load cell (Sensotec model 13)

Figure 4. Cut-off assembly drawing of the thruster prototype

with 1 kgf FS and a maximum combined error (non-linearity, hysteresis and repeatability) of 0.9% FSO.

A DC source, caable of supplying different output voltages, provides the transducers with the required excitations. The data coming from the sensors and transducers installed in the facility are acquired and transferred to a personal computer by means of a National Instruments acquisition board, capable of acquiring 32 analogical and 48 digital channels at a maximum sample rate of 1.25 MS/sec. The acquisition board is connected to:

- a SCXI 1520 module, for conditioning and filtering the signals coming from the pressure transducers, the load cell and the flow meter;
- a SCXI 1125 module, for conditioning and filtering the thermocouple signals:
- a SCXI 1163 module, for providing the digital outputs for the remote-control of the electro-pneumatic valves.

In particular, the flow meter current output (4 – 20 mA) has been changed into a voltage output by means of a calibrated 200 Ω resistance (0.8 – 4 V) in order to acquire the signal using the same SCXI 1520 module.

A LabVIEW® data acquisition and control program has been implemented. The operator can actuate the valves both manually (by clicking on the corresponding display switch) and automatically (by starting the predefined automatic procedures), having direct knowledge of the state of the valves. The acquired data are recorded and real-time displayed on the front panel. Separate windows also allow for in-line displaying the time-histories of all of the main measurements acquired during the test. This allows for the operator to fully monitor the entire system and make the proper decisions. Before the program is run it is also necessary to choose the sample acquisition rate compatibly with the maximum sampling rate of the acquisition board and the CPU performance. A value of 10 sps has been selected for the tests reported in this paper.

III.   Catalysts

The core of the hydrogen peroxide monopropellant thruster is represented by the catalyst. In collaboration with the Chemistry and Industrial Chemistry Department of Pisa University, ALTA S.p.A has developed a series of advanced catalysts on ceramic supporting spheres based on novel implantation techniques. The chemical activity of these catalysts has been preliminary assessed using 30% hydrogen peroxide solution by weight in a dedicated test bench[22,23] in order to select the most promising catalysts to be integrated in the thruster prototype. Two platinum-based catalysts, obtained from two different precursors (marked 1st and 2nd type in Table 2) using the same γ-alumina carrier and deposition procedure, have been selected, as they proved to be very active and relatively insensitive to poisoning problems. The last column in Table 2 reports the catalyst load expressed in percent atomic content (At%), which refers to the average SEM reading over the optical framing window (200 × 180 µm), down to the typical penetration depth of the electron beam below the catalyst surface (about 200 atomic layers).

Figure 5. LR-59 catalytic spheres.

Table 2. Main characteristics of the catalytic systems used in the thruster experiments.

ALTA's code	Catalyst	Carrier	Deposition procedure	SEM At%
LR-57	Platinum 1st type	*SASOL 06/170 γ-alumina spheres* Diameter: 0.6 mm Surface area: 170 m^2/g Pore volume: 0.53 ml/g Al$_2$O$_3$: 96.1%	1- Impregnation phase (Pt precursor and solvent) 2- Drying phase (mechanical depressurization at 10^{-4} atm and room temperature) 3- Reduction phase (under hydrogen atmosphere at room temperature)	0.39
LR-59	Platinum 2nd type			0.47

IV. Experimental Results and Discussion

Two catalytic beds made with the LR-57 and LR-59 platinum-deposed ceramic spheres of Table 2 have been tested in the mono-propellant thruster prototype. Both catalysts have decomposed hydrogen peroxide at concentration of 87.5%. For one of them, experimental data on the operation with 70% hydrogen peroxide are also available. For ease of assessment, the performance of these catalytic beds has been compared to that of a conventional silver screen catalytic bed of equal geometric envelope (diameter and length).

A. Performance Definition and Theoretical Evaluation

These tests have been mainly used to evaluate the behavior of the catalytic beds. In propulsive applications, the characteristic velocity efficiency ("C-Star efficiency") represents a significant parameter closely related to the capability of the catalytic bed of effectively decomposing the propellant for generating thrust. This parameter is based on the quasi-1-D theory of ideal rocket performance (perfect gas with constant composition, quasi 1-D steady isentropic frictionless flow, critical throat and negligible velocity in the combustion chamber) and compares the characteristic velocity obtained from the measurements of the propellant mass flow rate and chamber pressure with the theoretical value computed for a chamber temperature and composition corresponding to complete adiabatic decomposition of the propellant:

$$\eta_{c^*_{T_{ad}}} = \frac{c^*_{exp}}{c^*_{T_{ad}}} = \frac{\dfrac{p_{c_{exp}} A_t}{\dot{m}_{exp}}}{\sqrt{\dfrac{R_{T_{ad}} T_{ad}}{\gamma_{T_{ad}}}} \left(\dfrac{\gamma_{T_{ad}}+1}{2}\right)^{\frac{\gamma_{T_{ad}}+1}{2(\gamma_{T_{ad}}-1)}}}$$

(1)

A second significant parameter is the "temperature efficiency", which expresses how close the measured chamber temperature is to the adiabatic temperature corresponding to complete decomposition of the propellant:

$$\eta_{\Delta T} = \frac{T_{exp} - T_{amb}}{T_{ad} - T_{amb}}$$

(2)

In order to evaluate the deviation of the real behavior of the propellant gas in the thrust chamber from the theoretical one, it has been decided to employ the above definition of the "C-Star efficiency" where the reference characteristic velocity is computed from the experimental values of the chamber temperature and composition:

$$\eta_{c^*_{T_{exp}}} = \frac{c^*_{exp}}{c^*_{T_{exp}}} = \frac{\dfrac{p_{c_{exp}} A_t}{\dot{m}_{exp}}}{\sqrt{\dfrac{R_{T_{exp}} T_{exp}}{\gamma_{T_{exp}}}} \left(\dfrac{\gamma_{T_{exp}}+1}{2}\right)^{\frac{\gamma_{T_{exp}}+1}{2(\gamma_{T_{exp}}-1)}}}$$

(3)

This efficiency takes into account viscous effects and the other sources of losses and inefficiencies that are not considered in the quasi-1-D model of ideal rocket performance.

Finally, in space propulsion applications the pressure drop across the catalytic bed is another important operational parameter because of its obvious repercussions on the propellant pressurization, and indirectly on the design of the entire feed system. Bed pressure losses depend on the flow configuration and operating conditions (bed porosity, length, loading, flow pressure and temperature) and in general must be reduced in order to minimize the mass of the propellant feed system. Later results compare the pressure drop measured at the same operational conditions in catalytic beds realized by means of spherical pellets with those obtained in silver screen reactors of equal geometric envelope (diameter and length).

In the following, in order to characterize the propulsive performance of the engine prototype, the measured time-evolution of the thrust and the chamber pressure have been compared with the theoretical values obtained from the quasi-1-D model of ideal rockets. Based on the experimental results, it has also been possible to evaluate the thrust coefficient and the specific impulse using the well-known expressions:

$$C_{F_{exp}} = \frac{F_{exp}}{p_{c_{exp}} A_t} ; \qquad I_{sp_{exp}} = \frac{c^*_{exp} C_{F_{exp}}}{g_o}$$

(4)

B. 70% H₂O₂ Firing

In the first test, the tank has been filled with 450 ml of 70% hydrogen peroxide and a single, relatively long firing has been carried out up to complete depletion of the tank. The mass flow rate of hydrogen peroxide has been controlled using the cavitating venturi, previously calibrated with bi-distilled water in order to identify its effective throat area. The use of the cavitating venturi also allowed for the reliable measurement of the mass flow rate, in spite of a temporary malfunction of the Coriolis flowmeter.

The firing comprised an initial transient phase (lasting about 12 s), a longer steady-state operation (lasting about 75 s), followed by the progressive occlusion of the catalytic bed due to the thermal rupture of the ceramic pellets. Measurements of the thrust chamber parameters indicated the occurrence of large-amplitude oscillations about their mean values due to the onset of a flow instability. Since the data have been acquired at 10 sps without analog filtering, the unstable frequency, which was higher than the Nyquist frequency corresponding to the acquisition rate[20], could not be resolved by Fourier analysis of the acquired signals. A moving average has therefore been used to smooth out short-term fluctuations, in order to digitally simulate low-pass filtering of the acquired data. The theoretical and the experimental values of the thrust and the chamber pressure are respectively shown in Fig. 6 and Fig. 7, while their steady state values are summarized in Table 3 together with the other main operational parameters of the catalytic bed. The bed loading and chamber pressure were 76.6 kg/s m² and 6 bars, respectively. The low values of the decomposition temperature, and consequently of the C-Star and temperature efficiencies that characterize the catalyst behavior, clearly indicate the incomplete decomposition of hydrogen peroxide as the main reason for the attainment of relatively poor propulsive performance. On the other hand, during the steady state phase the pressure drop across the catalytic bed reached 4 bar, a relatively high value. Later in the run (at the relative time 5908 s), the pressure losses started to increase even further, due to the progressive occlusion of the catalytic bed.
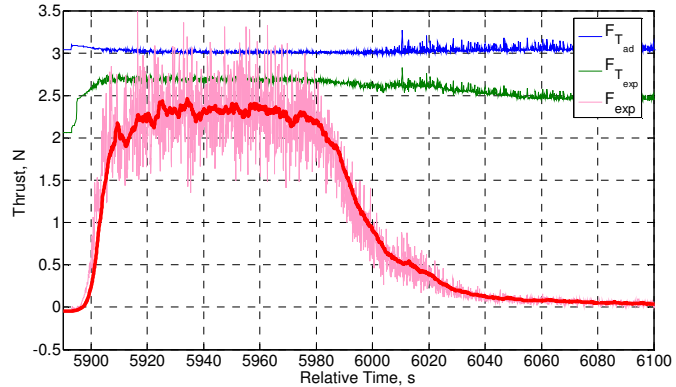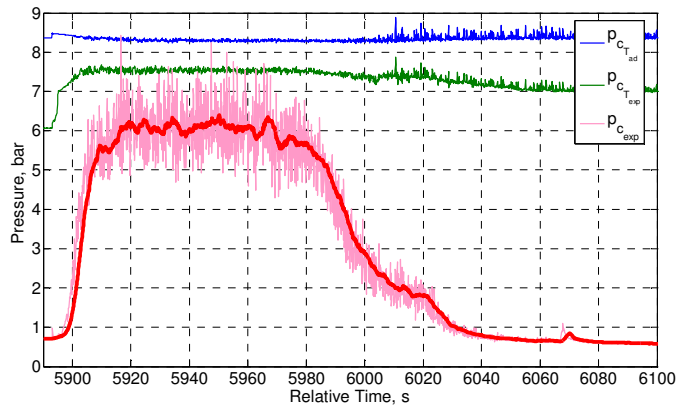

Figure 6. Thrust (LR-59, 70% H₂O₂).

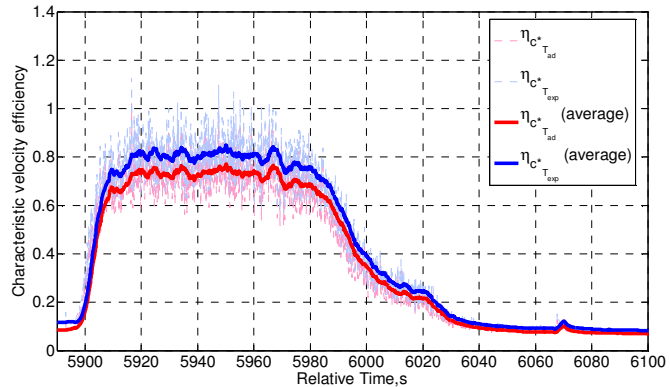**Figure 7. Chamber pressure (LR-59, 70% H₂O₂).**

Figure 8. C-Star efficiency (LR-59, 70% H₂O₂).
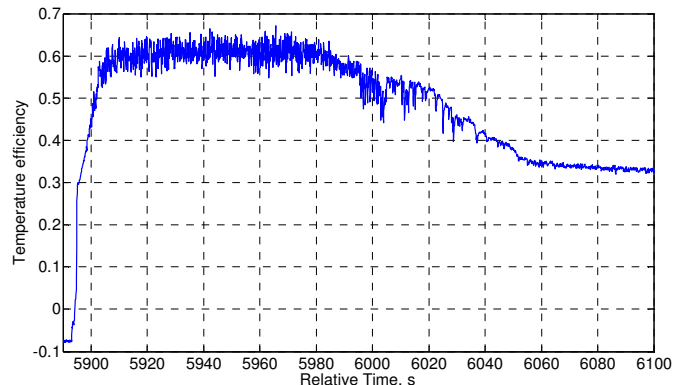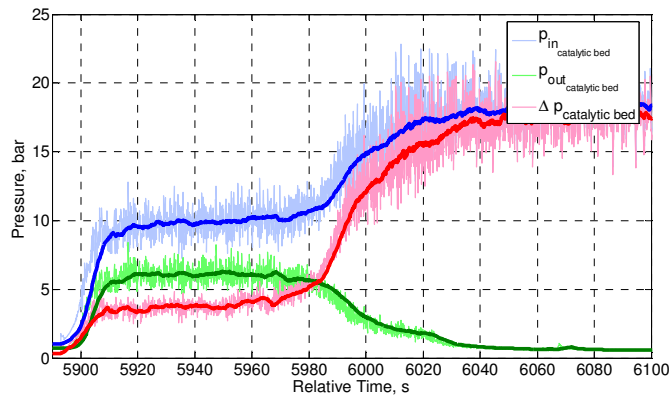

Figure 9. Temperature efficiency (LR-59, 70% H₂O₂).

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Figure 10. Catalytic bed pressure drop (LR-59, 70% H_2O_2).

Table 3. Steady-state performance with LR-59 catalyst and 70% H_2O_2.

Performance	5 N Thruster Pt/ Al_2O_3 2nd type (LR-59)	Ideal Adiabatic Decomposition
Thrust	2.3 N	3.0 N
Chamber Pressure	6 bar	8.5 bar
Chamber Temperature	433 K	514 K
Mass Flow Rate	3.85 g/s	3.85 g/s
Characteristic velocity	500 m/s	680 m/s
Thrust Coefficient	1.16	1.16
Specific Impulse	60 s	80 s
Bed Loading	76.6 kg/s m^2	76.59 kg/s m^2
$c*$ efficiency	0.73	-
ΔT efficiency	0.60	-
Catalytic bed pressure drop	4 bar	-

C. 87.5% H_2O_2 Firings

The Pt/ Al_2O_3 2nd type (LR-59) catalyst has also been tested in the engine prototype with the same procedure using higher grade 87.5% hydrogen peroxide. During the first firing the catalytic bed started to occlude just a few seconds after the reaching steady-state operation and the firing had to be terminated. Later attempts to restart the thruster have been unsuccessful.

All experimental data have been low-pass filtered by means of a 10 Hz cut-off frequency analog Butterworth filter and acquired at 10 sps, thereby eliminating the need to use the moving average in the reduction of the data. Figures 11 and 12 indicate that the initial transient phase of the test run lasted for about 10 seconds, while steady state

Figure 11. Thrust (LR-59, 87.5% H_2O_2).

operation has only been attained for a few seconds. However, even this short period of time has been sufficient for the evaluation of the steady state conditions, summarized in Table 4.

The operating parameters of the catalytic bed were very similar to the previous firing using 70% hydrogen peroxide. In particular, the chamber pressure was 5.5 bars and the bed loading 73.2 kg/s m^2. Not surprisingly, the chamber pressure and thrust display quite similar trends. Also in this case, their relatively low values are due to the
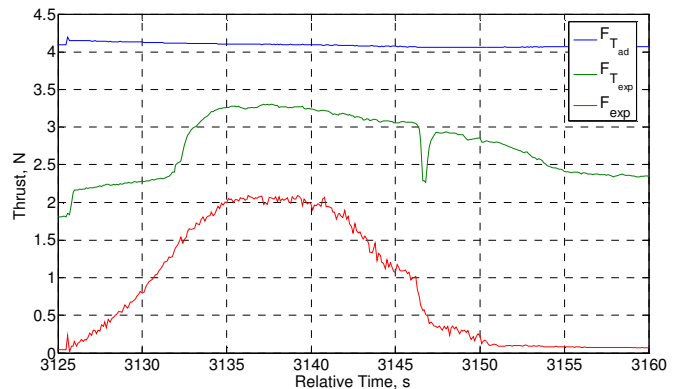
incomplete decomposition of hydrogen peroxide, as indicated by the temperature efficiency, 62%, and the characteristic velocity efficiency, 52%.
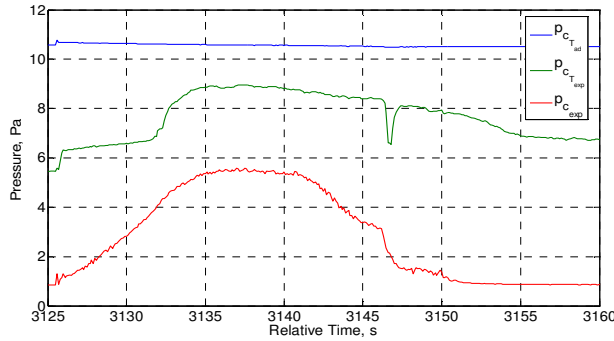

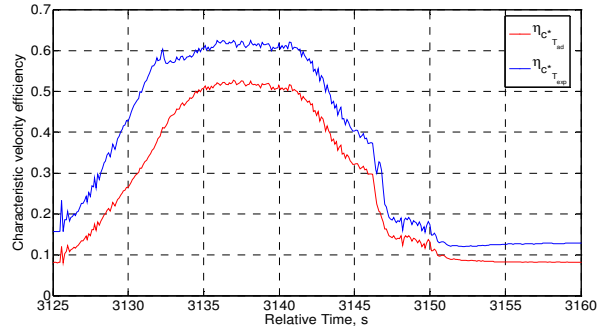**Figure 12. Chamber pressure (LR-59, 87.5% H₂O₂)**

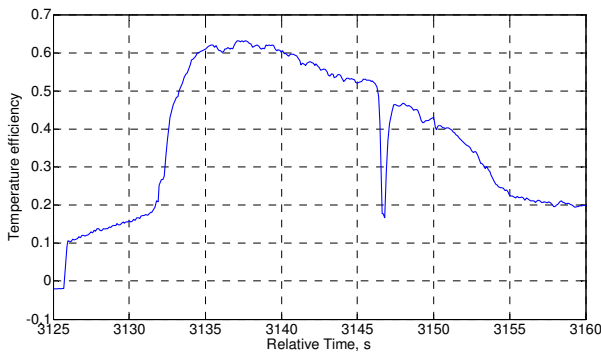**Figure 13. Characteristic velocity efficiency (LR-59, 87.5% H₂O₂)**

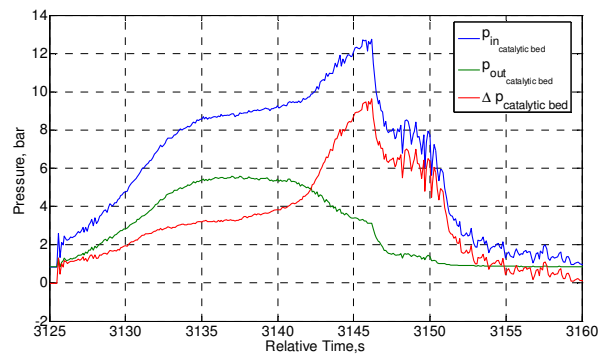**Figure 14. Temperature efficiency (LR-59, 87.5% H₂O₂)**

Figure 15. Catalytic bed pressure drop (LR-59, 87.5% H₂O₂)

Table 4. Steady-state performance with LR-59 catalyst and 87.5% H₂O₂.

Performance	5 N Thruster Pt/ Al₂O₃ 2nd type	Ideal Adiabatic Decomposition
Thrust	2.05 N	4.1 N
Chamber Pressure	5.5 bar	10.5 bar
Chamber Temperature	703.15 K	952.85 K
Mass Flow Rate	3.68 g/s	3.68 g/s
Characteristic velocity	470 m/s	904 m/s
Thrust Coefficient	1.20	1.23
Specific Impulse	57 s	113 s
Bed Loading	73.21 kg/(s m²)	73.21 kg/(s m²)
$c*$ efficiency	0.52	-
ΔT efficiency	0.62	-
Catalytic bed pressure drop	3.2 bar	-

The starting time of the occlusion of the catalytic bed can be easily identified from the diagram in Fig. 15 as the inflection point where the pressure drop starts to increase suddenly. As a consequence of bed occlusion when using the Pt/ Al₂O₃ 2nd type catalyst (LR-59), several tests with high grade hydrogen peroxide have been carried out in ALTA's catalytic activity facility[22,23] in order to identify the most promising substitute. The Pt/ Al₂O₃ 1st type catalyst (LR-57) proved to be the least sensitive to pellet rupture, and therefore has been tested in the engine prototype under the same operating conditions as the LR-59 bed reactor.

A total of 275 ml of 87.5% H_2O_2 have been used, acquiring the experimental data at 10 sps without filtering. Also in this case, a moving average has been employed to smooth out short-term fluctuations. The data of Table 5 indicate that the performance of the LR-57 catalyst bed turned out to be slightly better than that of the LR-59 catalyst. In particular, the characteristic velocity efficiency (Fig. 18) improved because the chamber pressure had increased (Fig. 17). Unfortunately, the occlusion of catalytic bed still occurred. The steady-state operation lasted almost twice as much as in the previous case (10 seconds), but the pressure drop across the bed also dramatically increased.
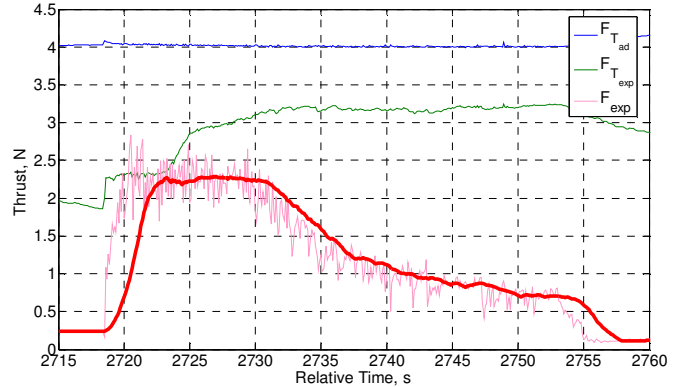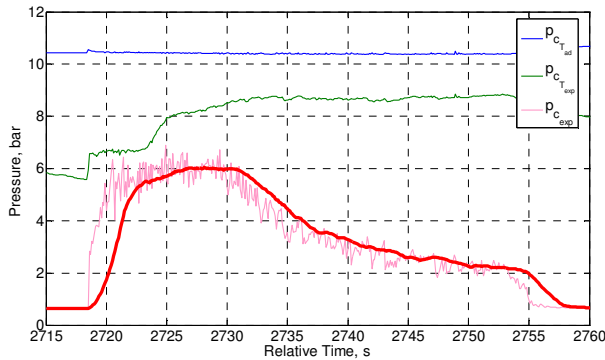

Figure 16. Thrust (LR-57, 87.5% H_2O_2).
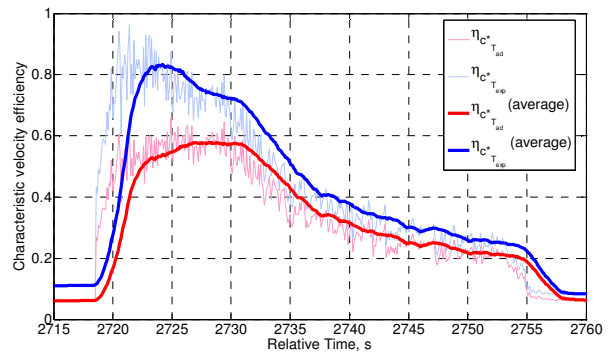

Figure 17. Chamber pressure (LR-57, 87.5% H_2O_2).

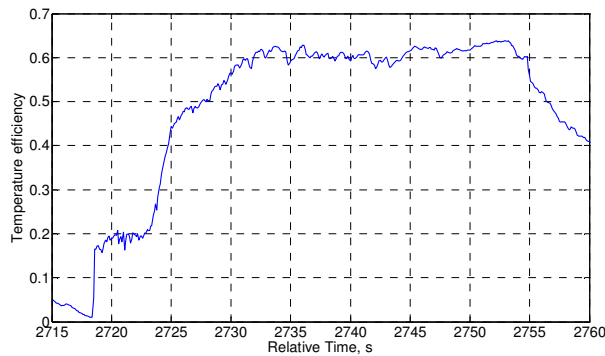**Figure 18. Characteristic velocity efficiency (LR-57, 87.5% H_2O_2).**

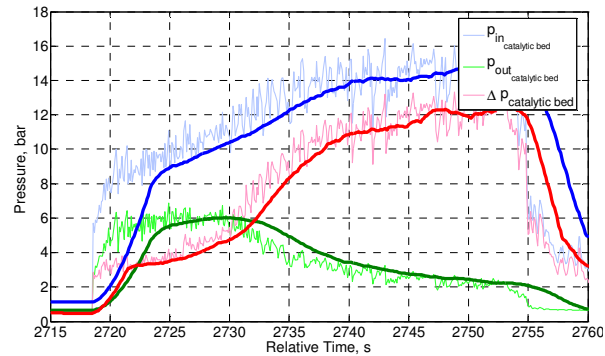

Figure 19. Temperature efficiency (LR-57, 87.5% H_2O_2).

Figure 20. Catalytic bed pressure drop (LR-57, 87.5% H_2O_2).

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Table 5. Steady-state performance with LR-57 catalyst and 87.5% H_2O_2.

Performance	5 N Thruster Pt/ Al_2O_3 1st type	Ideal Adiabatic Decomposition
Thrust	2.3 N	4.1 N
Chamber Pressure	6 bar	10.5 bar
Chamber Temperature	703.15 K	952.85 K
Mass Flow Rate	3.68 g/s	3.68 g/s
Characteristic velocity	520 m/s	904 m/s
Thrust Coefficient	1.20	1.23
Specific Impulse	65 s	113 s
Bed Loading	73.2 kg/s m^2	73.2 kg/s m^2
$c*$ efficiency	0.58	-
ΔT efficiency	0.62	-
Catalytic bed pressure drop	6 bar	-

Finally, Table 6 summarizes the steady state performance of a silver screen catalytic bed with the same envelope dimensions (diameter and length) and tested at equal operating conditions as the previous pellet reactors. 140 silver screens (99.9% purity) with a 80x80 in^{-2} mesh size and a wire diameter of 0.115 mm have been used as catalyst. The decomposition efficiency, the characteristic velocity efficiency and the temperature efficiency were significantly higher than for the previous pellet reactors, but still far from their theoretical limits, most likely as a consequence of insufficient bed length and possibly because of the occurrence of flow channeling phenomena. The increase in the decomposition temperature allowed for a higher value of the chamber pressure, and consequently a higher thrust. Furthermore, the pressure drop through the bed turned out to be lower than for the previous ceramic pellet reactors.

Table 6. Steady-state performance with silver screen catalyst and 87.5% H_2O_2.

Performance	5 N Thruster Silver Grids	Ideal Adiabatic Decomposition
Thrust	3 N	4.0 N
Chamber Pressure	7 bar	10.5 bar
Chamber Temperature	773.15 K	952.85 K
Mass Flow Rate	3.62 g/s	3.62 g/s
Characteristic velocity	724 m/s	904 m/s
Thrust Coefficient	1.22	1.23
Specific Impulse	80 s	113 s
Bed Loading	72.00 kg/s m^2	72.00 kg/s m^2
$c*$ efficiency	0.80	-
ΔT efficiency	0.72	-
Catalytic bed pressure drop	2 bar	-

V. Conclusions

The current design of HP catalytic reactors based on novel catalyst implantation techniques on γ-alunima ceramic supports has led to unsatisfactory levels of performance. The main problems encountered can be summarized as follows:

- rupture of the pellets and consequent occlusion of the catalytic bed;
- low c* and temperature efficiencies;
- onset of catalyst bed flow instabilities;
- high pressure drop across the catalytic bed.

Preliminary analyses of the relevant physical phenomena by means of reduced order models suggested that the main causes of these problems can be tentatively identified as follows:

- Rupture and powdering of the catalyst pellets due to excessive thermal stresses induced by the rapid and large change of their surface temperature during the initial phase of hydrogen peroxide decomposition. The occurrence of either one of these cracking modes seems to depend on the intensity of the superficial heat release and the ensuing different evolution of the radial temperature profile in the catalyst pellets.
- Progressive occlusion of the bed by the debris generated by the thermal cracking of the catalyst pellets.
- Mass diffusion limitations of the decomposition rate due to the large velocity difference between the incoming liquid reactant and the outgoing gaseous products, which effectively reduce the c* and temperature efficiencies below the theoretical values.
- Flow channeling, whose occurrence is suggested by the observed occurrence of fluid dynamic instabilities in the reactor. Flow channeling is known to interfere with the spatial distribution and intensity of hydrogen peroxide decomposition, with the consequence that the pressure drop in the reactor becomes very sensitive to changes of the H_2O_2 flowrate. Any accidental decrease of the decomposition rate results in a temporary increase of the liquid hydrogen peroxide concentration and flow velocity, due to the parallel decrease of the pressure losses. As soon as the local accumulation of hydrogen peroxide eventually succeeds in increasing the decomposition rate, the liquid is replaced by faster gaseous products and the flow velocity increases, preparing the conditions for the next oscillation cycle.

The identification of the physical causes of the problems affecting the catalytic reactor provides some guidelines in finding possible solutions. Stratified or multiple beds with increasing decomposition activity in the flow direction can conceivably reduce thermal stresses down to sustainable levels, at the expense, however, of larger bed volumes and higher pressure losses. Mechanical analyses of thermal loads in the catalyst pellets suggests that the cracking of the catalyst support, and the consequent occlusion of the reactor, might be avoided by using ceramic materials with higher thermal shock resistance. Four ways of pursuing this objective have been identified:

- reducing the product of the thermal expansion coefficient times the Young's modulus of the supporting material;
- increasing the ultimate stress of the supporting material;
- increasing the thermal conductivity of the supporting material;
- decreasing the typical thickness of the the supporting material.

If susceptible of effective catalyst deposition, cordierite, titania, zirconia, silicon nitride and silicon carbide are some potentially interesting ceramic materials for catalyst support. Besides, the intensity and distribution of thermal stresses is a strong function of the pellet Biot number hD/k when its value is of order unity. Since the superficial heat release due to hydrogen peroxide decomposition can be expressed in terms of an effective heat transfer coefficient, also the reduction of the pellet size can effectively contribute to mitigate the thermal stress problem of the catalyst support.

Several actions can be undertaken to improve the decomposition activity of the catalytic bed:

- increasing the active area of the catalyst by means of longer beds or smaller pellet size, accepting the associated pressure loss penalty;
- increasing the catalyst deposition on the support;
- increasing the reactant concentration by raising the flow pressure.

More recent preliminary experiments suggest that the reactor performance is not greatly affected by higher catalyst depositions on the same ceramic pellets. On the other hand, the results of 1-D steady analyses of the two-phase homogeneous reacting flow through the catalytic bed indicate that the last alternative might be particularly attractive because of its positive combined influence on the decomposition rate of hydrogen peroxide in gaseous phase, on the propulsive efficiency of the engine and on the pressure losses through the reactor. Finally, flow instabilities can probably be suppressed by the introduction in the catalytic bed of flow redistribution elements capable of negatively interfering with channeling phenomena, even though at the expense of increased pressure losses.

The experience gained during the present investigation showed that catalytic activity tests under atmospheric conditions are quite helpful during the early stages of development in selecting the most promising catalysts and provide useful indications on their relative susceptibility to thermal cracking, especially when carried out with high grade hydrogen peroxide (87.5% by weight). However, the effective investigation of the above critical aspects can only be carried in out in engine prototype tests, and implies the need of independently adjusting the bed loading, the residence time in the reactor and the chamber pressure over wider ranges than can be afforded in the present set-up.

Therefore, future redesigns of the engine prototype will need to accommodate catalytic beds of different cross-section, length, flow distribution and management devices, catalyst retention systems, etc., and incorporate easily reconfigurable thrust chambers realized in three modular segments in order to separately modify the chamber diameter, the nozzle cross-section and the nozzle exhaust area as required by the specific experiment.
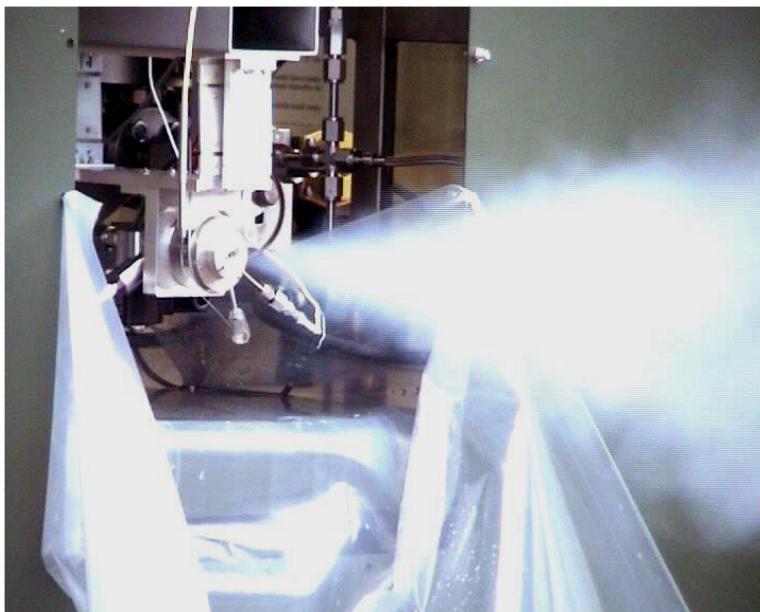

Figure 21. LR-59 70% H_2O_2 firing.

Acknowledgments

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