



Pressure Peaking Phenomena for Unignited Liquid Hydrogen Releases: Experiments and Analysis

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ABSTRACT

Liquid hydrogen (LH₂) has been identified as a carbon-free fuel in the transition to Net Zero. The transport sector has a particular interest in LH₂ due to its increased density (71 kg/m³ at -253°C, 0.1 MPa) compared to pressurised gaseous hydrogen (24 kg/m³ at 20°C, 35 MPa). The Health and Safety Executive's (HSE) Science Division has been researching and assessing LH₂ transport hazards for over a decade with supporting UK Safe Net Zero as one of the organisation's key objectives.

Pressure peaking is a phenomenon unique to lighter-than-air gases, where an overpressure develops due to a release of unignited gas, which can be high enough to cause damage to structures. This work investigates the extent to which pressure peaking can occur in the event of significant unignited LH₂ releases inside a ventilated enclosure. The experiments presented are intended to be representative of a full-bore LH₂ hose failure inside a transfer connection space.

LH₂ releases of 180 g/s into a 1 m³ enclosure with a vent area varying from 0.0416 to 0.0104 m² were studied. It was found that pressure peaking can occur, with a maximum peak overpressure of 3.5 kPa observed for the smallest vent area. Compared to a modelled gaseous release of the same mass flow rate, the peak overpressure of the LH₂ experiment is significantly lower.

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For hydrogen to be used safely as a zero-carbon fuel, the associated hazards must be fully understood and characterised. One novel hazard associated with hydrogen compared to other fuels is the ‘pressure peaking phenomenon’, where an overpressure can occur in a vented enclosure due to a leak of hydrogen. These overpressures may be capable of causing damage to structures even without ignition.

This phenomenon has been observed experimentally with helium and gaseous hydrogen (GH₂) leaks (Lach, Gaathaug and Vaagsaether, 2020; Shentsov, Kuznetsov and Molkov, 2015) and modelled using numerical methods (Brennan and Molkov, 2013) and computational fluid dynamics (Cirrone et al., 2021).

The aim of this work was to investigate whether liquid hydrogen (LH₂) leaks can result in pressure peaking, the extent of the overpressure and the effect of varying ventilation area. The potential damage from the overpressure is analysed, and the results are compared to modelled gaseous releases of an equivalent size.

GH₂ PRESSURE PEAKING

Experiments have shown that releases of buoyant gas (helium and hydrogen) into a vented enclosure can cause overpressure before decaying to a steady-state pressure (Shentsov, Kuznetsov and Molkov, 2015). This effect is not observed for releases of gases heavier than air. Figure 1 shows the largest hydrogen release rate test performed in the experimental program of Shentsov, Kuznetsov and Molkov.

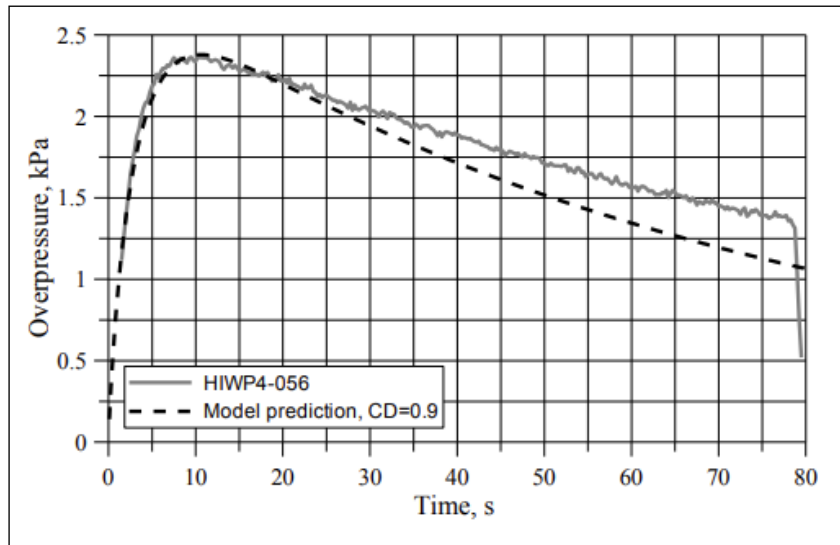


Figure 1 Pressure peaking test (1.086 g/s gaseous hydrogen leak in a 1 m³ enclosure with a 16.5-mm-diameter vent). The grey line shows experimental overpressure; the black dashed line shows the modelled overpressure (Shentsov, Kuznetsov and Molkov, 2015).

Larger-scale experiments conducted by Lach, Gaathaug and Vaagsaether (2020) released gaseous hydrogen into a 14.9 m³ chamber at flow rates of 1.9–10.10 g/s with vent areas of 0.0006–0.0020 m². This achieved peak overpressures between 0.42 and 8.05 kPa. It was found that the larger enclosure did not have a significant effect on the peak overpressure achieved, but that the peak occurred later. Decreasing the ventilation area also caused the peak pressure to occur later.

A phenomenological model of hydrogen pressure peaking in a vented enclosure was developed by Brennan and Molkov (2013). The steady-state behaviour is found analytically by modelling the inlet flow $\dot{m}_{nozz}$ using Bernoulli’s equation and modelling the vent flow $\dot{m}_{vent}$ as a subsonic orifice and equating the two. This results in an expression for the steady-state vent mass flow rate (1) and steady-state overpressure (2):

$$\dot{m}_{vent} = CA \left(\frac{2\gamma}{\gamma-1} \right) P_{atm} \rho_{vent} \left[\left(\frac{P_{atm}}{P_{encl}} \right)^{\frac{2}{\gamma}} - \left(\frac{P_{atm}}{P_{encl}} \right)^{\frac{\gamma-1}{\gamma}} \right]^{\frac{1}{2}} \quad (1)$$

$$P_{encl} - P_{atm} = \frac{\rho_{vent} \left(\frac{\dot{m}_{nozz}}{\rho_{vent} A} \right)^2}{2} \quad (2)$$

$\dot{m}_{vent}$: vent flow rate (kg/s); C : coefficient of discharge; A : vent area (m^2); γ : specific heat ratio of venting gas; P_{atm} : atmospheric pressure (Pa); ρ_{vent} : density of gas at the vent (kg/m^3); P_{encl} : enclosure pressure (Pa); $\dot{m}_{nozz}$: nozzle mass flow rate (kg/s).

Equation (2) shows that the steady-state overpressure is inversely proportional to the density of the fluid at the vent. This is important for the liquid tests because the density of vaporised cryogenic hydrogen can be many times higher than the density of ambient hydrogen gas.

To determine the transient pressure increase in the enclosure, a system of equations (enclosure mass, enclosure moles, ideal gas equation and vent flow rate) is solved numerically. As there is no spatial dimension in this approach, perfect mixing of hydrogen and air is assumed. If this approach were to be applied to LH_2 leaks, a homogeneous temperature distribution would also have to be assumed.

METHODOLOGY

LH_2 was released from a tanker through a 1" ID pipe into a $1 m^3$ ($1 m \times 1 m \times 1 m$) enclosure with 3-mm-thick walls, shown in Figure 2. The enclosure had four passive vents, shown in Figure 2, to which the vent covers (bottom of Figure 3) were attached. These were swapped for blank flanges to reduce the vent area. The enclosure also had a larger foil burst panel to protect the enclosure in case of overpressurisation. The dimensions of the enclosure and vent geometry are shown in Figure 3.

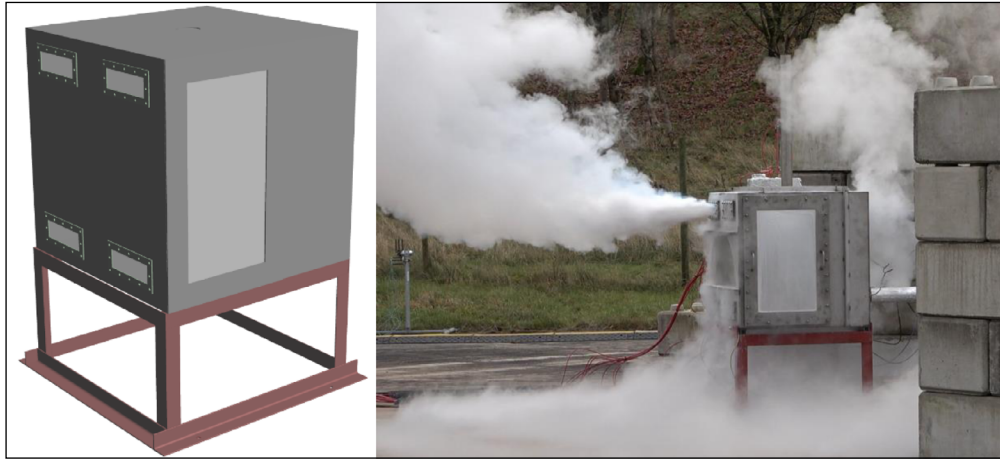


Figure 2 Left: A CAD model of the $1 m^3$ enclosure with four passive vents and a burst panel on a frame. Right: Image during Test 3 with one open vent.

Three tests were conducted with vent areas of 0.0416, 0.0208 and 0.0104 m^2 .

A sensing array inside the enclosure measured temperature and hydrogen concentration at various points. Gas in the enclosure was sampled and piped away from the experiment to the hydrogen sensors with a total sampling flow rate of approximately 60 L/min. Two pressure transducers (PT1, PT2) were attached to the box on the right side of the enclosure (Figure 3) at two points to measure the increase in pressure in the enclosure. Mass flow rate was not measured directly, but using the pressure and pipe diameter and correlating to previous experimental LH_2 releases conducted at HSE, the mass flow was estimated to be 180 g/s. Details of experimental instrumentation are given in Table 1.

RESULTS

TEST 1

For the first test, all four vents were open, resulting in a total vented area of 0.0416 m^2 . LH_2 was released into the enclosure at 330 kPa. Figure 4 shows the pressure measured at

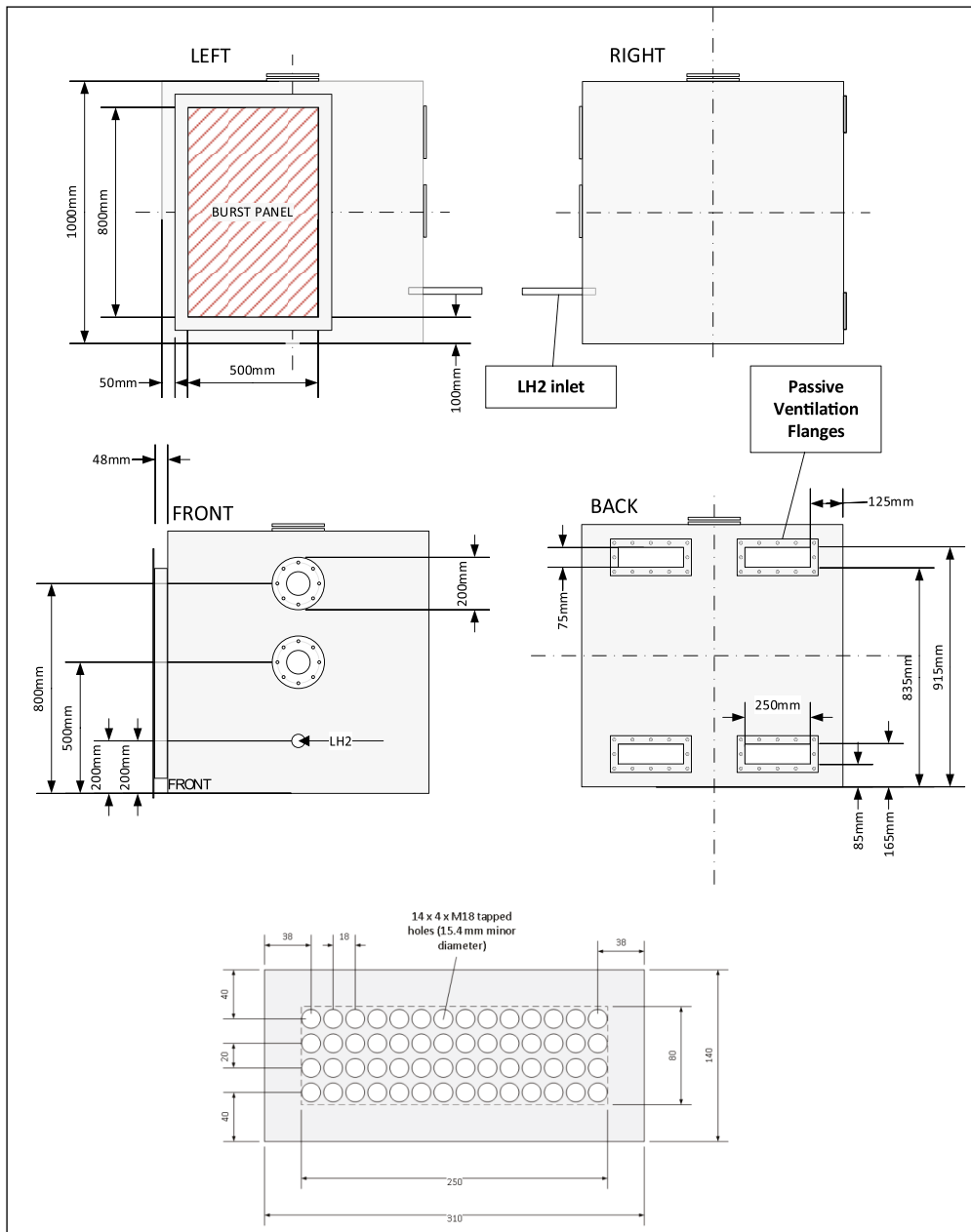


Figure 3 Top: Left, right, front and back dimensioned views of the 1 m³ enclosure. Note that for these tests, the two flanges on the front side of the enclosure were blanked. Bottom: Dimensions of plates attached to open flanges. The total ventilation area of each plate is 0.0104 m².

MEASUREMENT	INSTRUMENT	RANGE	ACCURACY	UNCERTAINTY
Release pressure	ESI IS 4200EXH0016AB Pressure Transducer	0.1–1.1 MPa	±0.25%	±0.002 MPa
Release temperature	Type T thermocouple, 2 m × 1.5 mm	4.2–400 K	±8 K (@ 20 K)	±8 K
Enclosure internal temperature	Type T thermocouple, 2 m × 1.5 mm	77–400 K	±1 K	±1 K
Enclosure wall temperature	Type T thermocouple, 2 m × 1.5 mm	77–400 K	±1 K	±1 K
Enclosure internal overpressure (PT1, PT2)	WIKA A-10 pressure transducer	0–10 kPa	±0.5%	±0.03 kPa

two locations on the wall of the test rig. When LH₂ was released into the test rig at 153 s, a sharp increase in overpressure was measured by both pressure transducers. PT1 measured a maximum overpressure of 0.30 kPa, and PT2 measured a maximum overpressure of 0.55 kPa. The burst panel, which was rated to fail at an overpressure around 12 kPa, did not rupture. The overpressure in the test rig then decayed over 26 s to a steady state of 0.00 kPa.

The peak overpressure was achieved 0.15 s after the start of the release. The logging rate of the pressure transducers was 20 Hz, which may not have been fast enough to capture the true peak pressure.

Table 1 Instrumentation used in experiments.

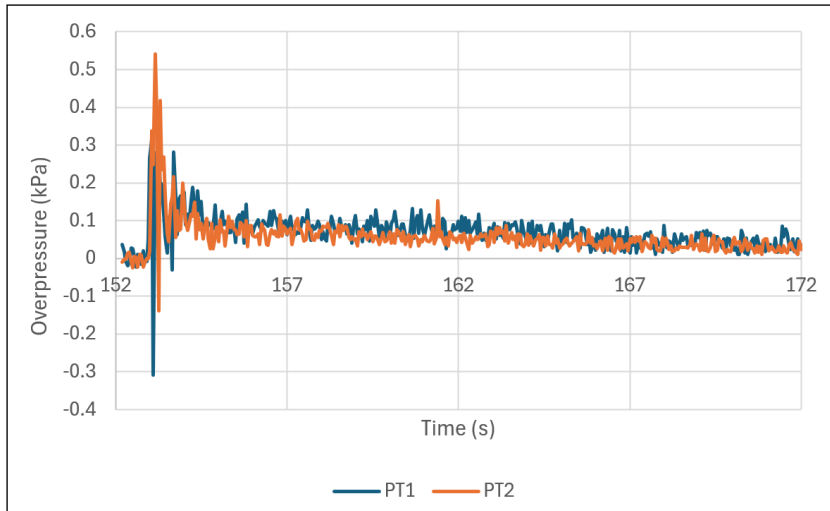


Figure 4 Test 1 enclosure overpressure.

TEST 2

For the second test, the bottom two vent flanges were blanked off, and the two open vents had a total vent area of 0.0208 m².

LH₂ was released into the enclosure at 330 kPa. Figure 5 shows the pressure measured in the test rig. Again, a sharp increase in pressure was observed when the LH₂ release started at 167 s. PT1 measured a maximum overpressure of 1.10 kPa, and PT2 measured a maximum overpressure of 1.20 kPa. The burst panel did not rupture. The overpressure in the test rig then decayed over 63 s to a steady state of 0.05 kPa.

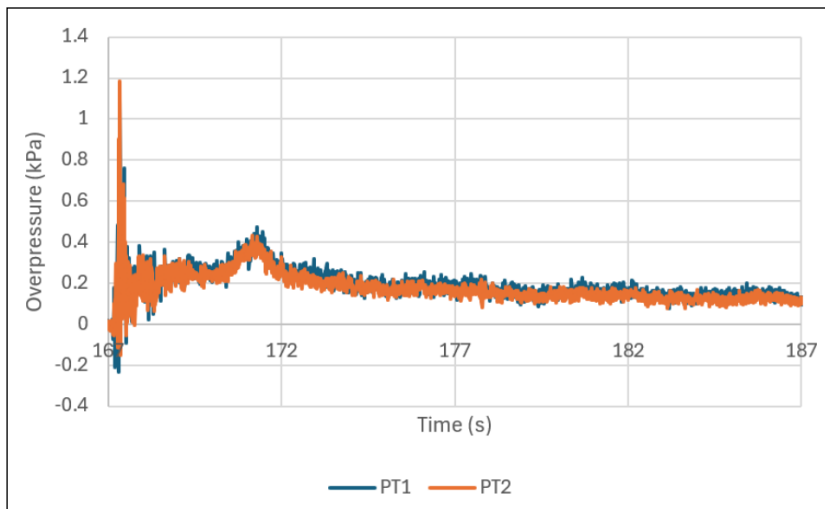


Figure 5 Test 2 enclosure overpressure.

The peak overpressure was achieved 0.2 s after the start of the release. It should be noted that the peak pressure was achieved before the end of the pressure drop due to opening the release valve (i.e. before the maximum LH₂ flow rate is achieved).

Interestingly, during this test, there was a secondary peak in overpressure of 0.40 kPa which occurred 3.9 s after the primary overpressure peak. The reason for this is currently unclear.

TEST 3

For the third test, the bottom two vent flanges and the top left vent flange were blanked off so that only the top right vent was open with a total vent area of 0.0104 m².

The pressure at the release rig prior to opening the release valve was 340 kPa. Figure 6 shows the pressure measured in the cube. Again, a sharp increase in pressure was observed when the LH₂ release started at 168 s. PT1 measured a maximum overpressure of 3.30 kPa, and PT2 measured a maximum overpressure of 3.35 kPa. Maximum overpressure was achieved 0.3 s

after the start of the release. Pressure in the cube then decayed rapidly in 3.7 s to 0.60 kPa, where it plateaued for 6.8 s. The overpressure in the cube then decayed further over 56 s to a steady state of 0.20 kPa. The burst panel did not rupture.

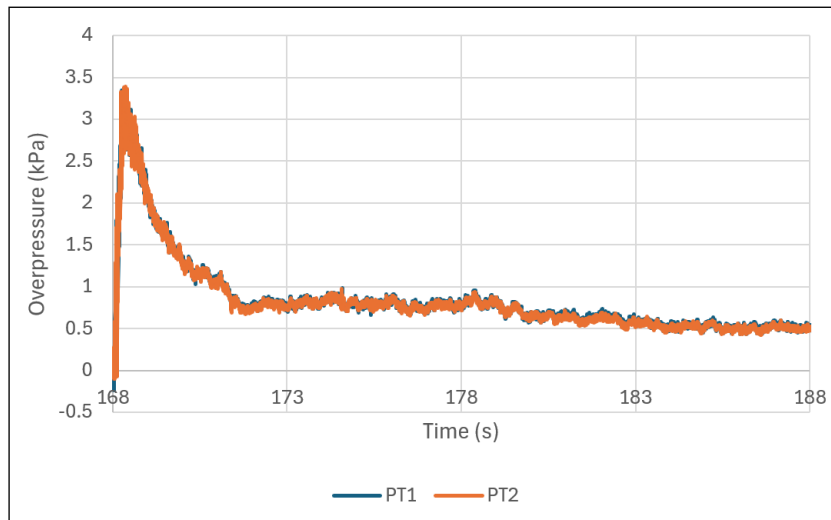


Figure 6 Test 3 enclosure overpressure.

Peak pressure was achieved 0.33 s after the start of the release. As in the previous test, the peak pressure was achieved before the end of the pressure drop due to opening the release valve (i.e. before the maximum LH_2 flow rate is achieved). The maximum rate of pressure increase was 33.3 kPa/s.

DISCUSSION

The time-to-peak pressure from the start of the release increases as a vent area increases for the LH_2 experimental releases. This is notable because the opposite phenomenon is observed for gaseous releases. In Tests 2 and 3, plateaus in overpressure are observed for a few seconds before the overpressure reaches steady state, which is not typically observed during gaseous releases. This discontinuity in gradient may be indicative of LH_2 -specific pressure peaking behaviour because it is not observed in GH_2 pressure peaking.

To model transient pressure peaking using a numerical model, it was assumed that the mixture in the enclosure was homogeneous with respect to hydrogen concentration and temperature. [Figure 7](#) shows the measured temperature distribution in the enclosure for Test 3 at 168.5 s, just after the peak overpressure occurs. Temperature in the enclosure ranges from 107 to 275 K, with colder temperatures being restricted to the jet axis. It should also be noted that the temperature probes will have non-insignificant thermal inertia, so it is possible for this temperature range to be even wider in reality.

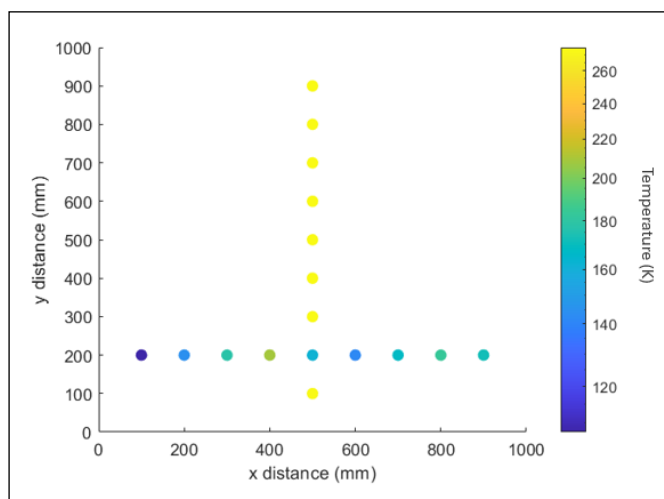


Figure 7 Temperature distribution in the enclosure during Test 3 at 168.5 s. All temperature probes are in the plane of the LH_2 inlet, with the inlet at (0.200).

Peak overpressures achieved by the tests are comparable to unconfined, uncongested hydrogen-air deflagrations (flash fires); however, the overpressure behaviour is significantly different because the positive phase duration is much longer than for explosive overpressures. Therefore, it is important to consider the impulse of the overpressure rather than only comparing peak overpressures.

Because an overpressure is observed throughout the continuous LH_2 release, the associated impulse and phase duration are dependent on the duration of the LH_2 release (the total impulse of the overpressure will keep increasing as the LH_2 release continues, and the positive phase duration is essentially the same as the LH_2 release duration). Overpressure reaches a steady state asymptotically, and therefore in order to determine the duration of the overpressure, the trace is truncated to the point where 95% of the steady state is achieved; the overpressure before this time is taken to be the positive phase, as shown in Figure 8.

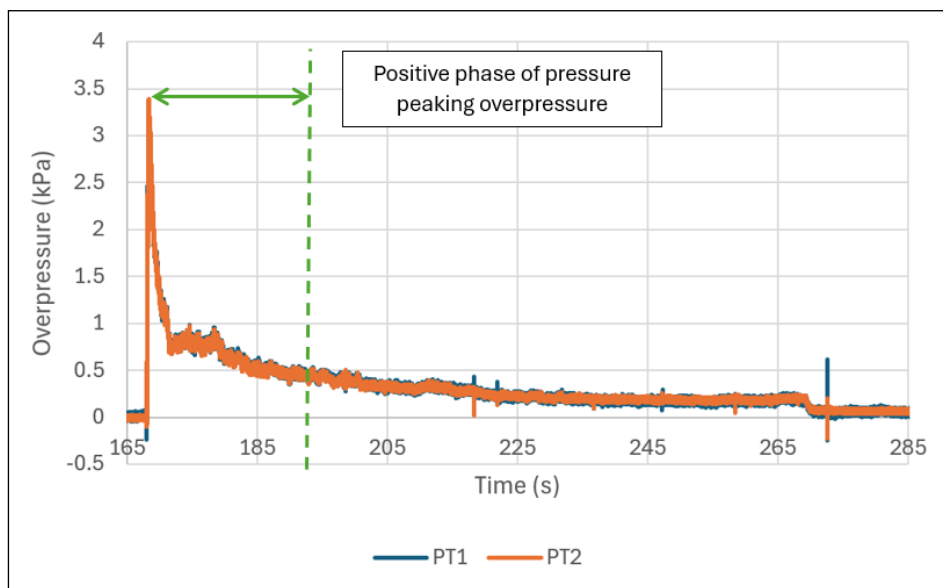


Figure 8 Graph of overpressure during Test 3 showing the positive phase cutoff.

Using this assumption, the positive phase duration is taken to be 32 s. This is much longer than the positive phase duration for explosive overpressures, which are on the order of 10 ms. The impulse is the integral of overpressure with respect to time, which is 22,500 Pa-s, much higher than for an explosion (on the order of 100s Pa-s) due to the extended length of the positive phase duration. Even considering only the sharp peak in the period 0–4 s, the impulse is 6,000 Pa-s, an order of magnitude larger than for an explosion. Because of the high impulse and relatively low rate of pressure rise, a pressure wave from this unignited pressure peaking will act on a structure quasi-statically, and the structure will not experience an increase in strength due to high strain-rate behaviour (an apparent increase that should be taken into account when designing structures that may be exposed to pressure peaking phenomena).

However, the overpressures observed in these tests are unlikely to be high enough to cause significant damage to even very light wooden structures. Higher overpressure could be caused by increased mass flow or decreased vent area.

COMPARISON WITH GH_2 PRESSURE PEAKING

The pressure peaking phenomena for gaseous releases are modelled using the *pressure peaking phenomenon for unignited releases* tool created by Ulster University (Shentsov, Makarov and Molkov, 2019). Only the third pressure peaking test was analysed as it had a high enough logging rate to capture transient phenomena.

The gaseous release was modelled using the parameters provided in Table 2.

PARAMETER	SYMBOL	VALUE	UNIT
Atmospheric pressure measured in experiment	P_{atm}	96,900	Pa
Enclosure temperature	T_{encl}	280	K
Enclosure volume	V_{encl}	1	m ³
Vent height	H_{vent}	0.1	m
Vent width	W_{vent}	0.104	m
Hydrogen mass flow rate	$\dot{m}_{H_2}$	0.18	kg/s
Coefficient of discharge	C_d	0.6	–
Integration time step	Δt	0.002	s
No. of time steps	n_{max}	1000	–

Table 2 Input parameters into the Ulster University GH₂ pressure peaking model.

The vent area is aggregated into a single rectangular opening with the same total area as the LH₂ experiment.

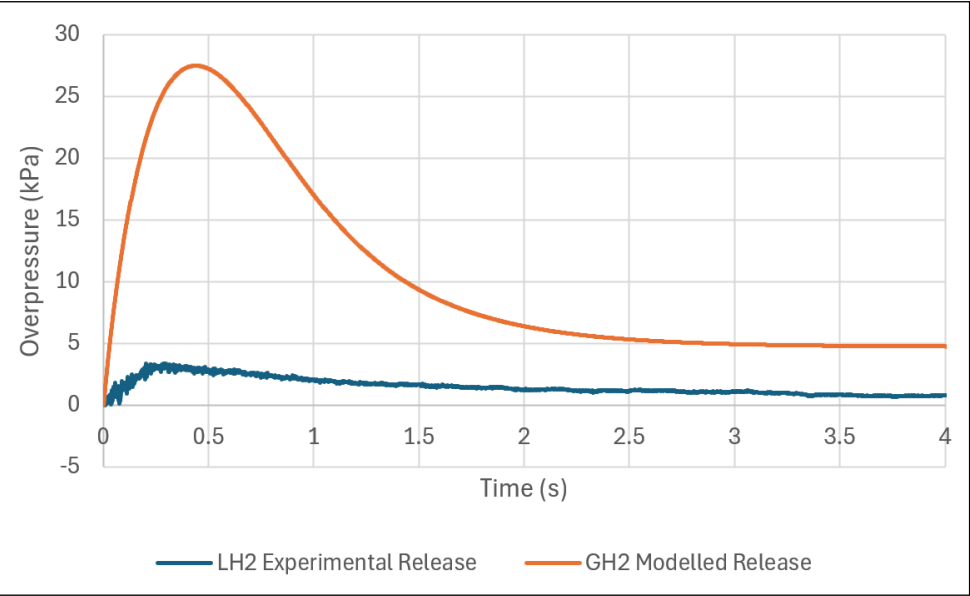


Figure 9 Graph of enclosure overpressure for LH₂ experimental release (Test 3) and GH₂ modelled release using Ulster University pressure peaking model.

A modelled gaseous release of 180 g/s achieves a maximum overpressure of 27.6 kPa in 0.43 s before decaying to a steady state of 4.6 kPa. The maximum rate of pressure increase is 149 kPa/s and the impulse (applying the same method as for the LH₂ experiment) is calculated to be 21,600 Pa-s.

Figure 9 shows that for an equivalent mass flow in the same volume enclosure, the LH₂ experiment was found to achieve peak pressure approximately 8.3 times lower than the GH₂ modelled release. The peak pressure was achieved 0.11 s faster for the LH₂ experiment. The steady-state overpressure in the enclosure is also lower for the LH₂ experiment. However, the LH₂ experiment takes much longer to reach steady state; hence, the LH₂ experiment and GH₂ model have similar impulses despite the GH₂ model having a much higher peak pressure. It is known that pressure peaking does not occur for releases of heavier-than-air gases, so it is assumed that the largest contributor to the decrease in peak overpressure for the LH₂ experiment is the increased density of the venting fluid (approximately 3.7 times greater for cryogenic hydrogen).

By setting the inlet mass flow equal to the vent mass flow, which must be true for the system to be at steady state, equations (1) and (2) can be combined to produce an equation for the discharge coefficient, which is dependent on the ambient and enclosure pressures (at steady state) and the specific heat ratio of the venting gas:

$$C = \sqrt{\frac{\gamma}{\gamma - 1} \frac{P_{atm} \left\{ \left(\frac{P_{atm}}{P_{encl}} \right)^{\frac{2}{\gamma}} - \left(\frac{P_{atm}}{P_{encl}} \right)^{\frac{\gamma+1}{\gamma}} \right\}}{P_{encl} - P_{atm}}} \quad (3)$$

This yields a discharge coefficient of 0.998 for the LH₂ pressure peaking experiment. This discharge coefficient is much larger than those found experimentally by Shentsov, Kuznetsov and Molkov (2015) or by Lach, Gaathaug and Vaagsaether (2020), which range between 0.65 and 0.75. One explanation for this is that the enclosure was not fully sealed at the LH₂ inlet or other flanges, and therefore the vent area would be larger than 0.0104 m², with the increased vent area being accounted for by a larger discharge coefficient. The very low pressure differential across the vent may also result in a high discharge coefficient.

Equations (1) and (2) are used to model the steady-state behaviour of Test 3. In Test 3, the steady-state vent temperature was measured to be 59 K. Assuming that the concentration at the vent was 100% H₂, the density of the venting hydrogen is 0.412 kg/m³ (Leachman et al. 2009). The specific heat ratio at this temperature is 1.67.

Using equation (1) with a density of 0.412 kg/m³, an overpressure of 0.2 kPa, a vent area of 0.0104 m², a specific heat ratio of 1.67 and a coefficient of discharge of 0.998, the steady-state vent flow rate can be calculated to be 177 g/s, which is close to the nozzle flow rate of 180 g/s.

Using equation (2) with a density of 0.412 kg/m³, nozzle flow rate of 0.18 kg/s and vent area of 0.0104 m², a steady-state enclosure overpressure of 0.36 kPa is predicted, which is double the experimental steady-state overpressure of 0.2 kPa.

The release was modelled again using the same parameters as in Table 2 except for the enclosure temperature being equal to the steady-state vent temperature, 59 K, and the discharge coefficient being equal to the calculated value, 0.998. This results in a peak overpressure of 4.1 kPa and a steady-state overpressure of 0.38 kPa as shown in Figure 10.

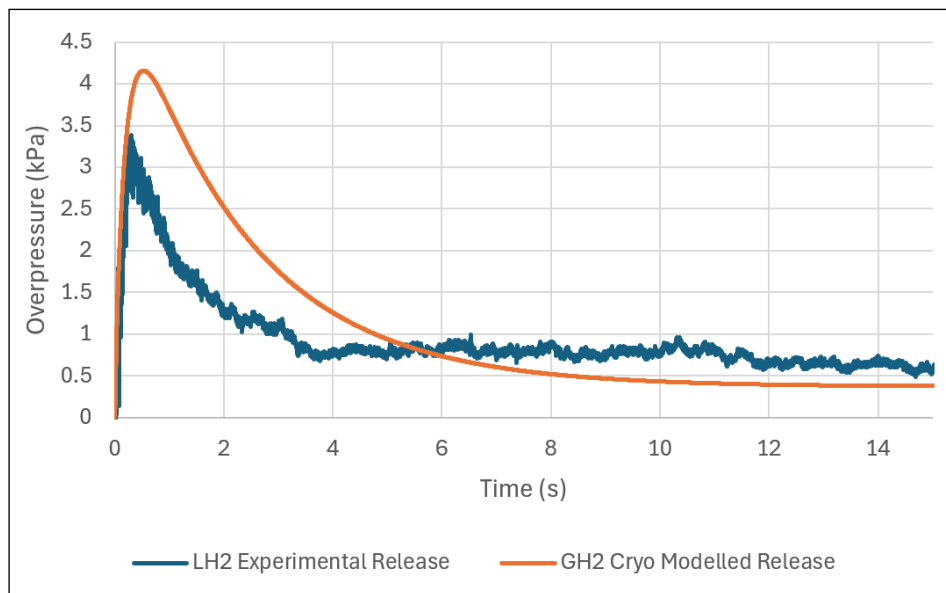


Figure 10 Graph showing enclosure overpressure for LH₂ experimental release (Test 3) and GH₂ modelled release using Ulster University model with cold enclosure (59 K) and increased Cd (0.998).

Agreement between the model and the experiment is good in the transient period (below 4 s) and then deviates beyond that. The decay in overpressure is slower for the modelled release but falls below the experimental release at 6 s, around the point where the experimental overpressure first plateaus. This may indicate that there are additional phenomena occurring, causing the plateau and slower decay, which are not captured in the model.

The impulses of the experiment and model are similar (12,000 Pa-s for the modelled release, 16,900 Pa-s for the experimental release).

This approach was also applied to Test 2, which had a steady-state vent temperature of 60 K. The modelled overpressure is compared to the test overpressure in Figure 11.

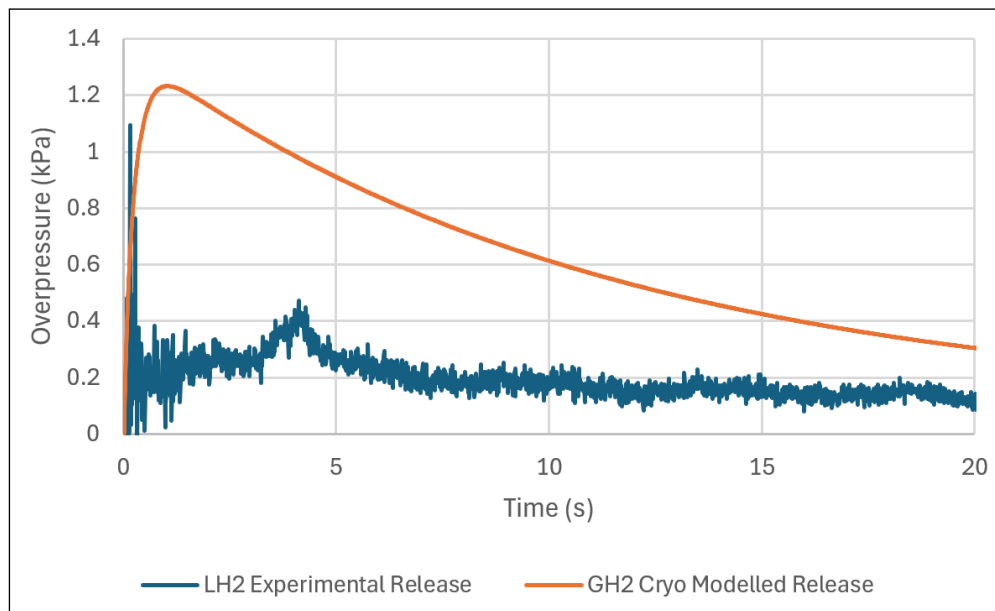


Figure 11 Graph showing enclosure overpressure for LH₂ experimental release (Test 2) and GH₂ modelled release using the Ulster University model with a cold enclosure (60 K) and increased Cd (0.998).

Agreement between the modelled overpressure and Test 2 experimental overpressure is worse, especially for the overpressure decay, though the model produces a similar peak overpressure to the experimental data.

Notably, these analytical models do not consider the cooling and condensation of air, which will cause a reduction in overpressure, nor do they account for the phase change of LH₂ as it vaporises to a cryogenic gas. The model has also not been validated for liquid or cryogenic hydrogen releases. It is worth restating that the LH₂ mass flow rate is estimated, and any error in the estimation will have a significant impact on the modelled overpressure. As such, comparisons between the modelled and experimental results should be taken to be indicative rather than definitive. However, the modification of parameters based on experimental data may serve to advise changes to the model to account for cryogenic pressure peaking phenomena as well as provide validation data for future models.

CONCLUSION

Liquid hydrogen was released into an enclosure with a variable vent area to assess the extent of pressure peaking for LH₂ leaks. It was found that pressure peaking did occur and that the peak overpressure increased as the vent area decreased. Decreasing the vent area had the effect of increasing the time-to-peak overpressure.

The overpressure was analysed, and it was found that the pressure peaking from even the smallest vent area in the LH₂ experiment conducted was unlikely to cause significant damage to a structure. The experimental results for the third test were compared to modelled GH₂ releases of the same mass flow rate, enclosure volume and vent area, and it was found that the peak overpressure due to LH₂ pressure peaking in one of the experiments was 8.3 times lower than the overpressure of a modelled GH₂ release of the same mass flow rate, which is attributed to the higher density of cryogenic hydrogen.

The results of these experiments may aid in the development of models for cryogenic pressure peaking phenomena and the design of enclosures and vent sizes to mitigate damage arising from pressure peaking. The results show that, generally, pressure peaking from LH₂ releases is not a significant hazard compared to pressure peaking from GH₂ releases.

Future work may include the development of analytical or numerical models that take into account the contribution of air cooling and condensation to pressure peaking overpressure. As LH₂ technologies develop and the size and structure of typical transfer connection spaces become clearer, it may be useful to conduct pressure peaking experiments on a more realistic transfer connection space. The effect of increased vaporisation rate (due to increased surface area or heating) on the pressure peaking of LH₂ releases, particularly for subcooled LH₂ releases, would contribute to a better understanding of this phenomenon.

The data from these experiments can be found under the pressure peaking section of the following dataset: <https://dataverse.no/dataset.xhtml?persistentId=doi:10.18710/JXJP0H>.

Additional sensor information can be found in the following dataset: <https://dataverse.no/file.xhtml?persistentId=doi:10.18710/JXJP0H/SUAIAR>.

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DISCLAIMER

This paper and the work it describes were undertaken by the Health and Safety Executive (HSE). Its contents, including any opinions and/or conclusions expressed or recommendations made, do not supersede current HSE policy or guidance.

COMPETING INTERESTS

Janni Vizma is a reviewer for Hydrogen Safety journal. Authors have no other competing interests.

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