Gas Phase Detonations: Effective Pressures Acting on the Walls of the Enclosures and Probability of Deflagration-to-Detonation Transition in Pipes, Vessels and Packings

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Overview

- Brief background info: deflagrative and detonative explosions in gaseous mixtures
- Motivation
- Detonations in pipes
- Detonations in empty vessels (i.e. no turbulence enhancing elements inside)
- Detonations in vessels filled with dry packings
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Gas Phase Explosion (Self-sustaining flame front) V_{flame} < V_{sound} V_{flame} > V_{sound}

V_{flame} < V_{sound}
(typically: 0.5 – 10 m/s)

V_{flame} > V_{sound} (typically: 1600 – 2800 m/s)

Deflagration

Detonation

occurrence of shock wave:

no

yes

- mechanisms for triggering the reaction in the unburnt mixture
- transfer of heat from flame front to unburnt mixture
 diffusion of radicals from flamefront into unburnt
- adiabatic compression by shock wave heats up gas mixture to T >> T_{autoignition} (flame front is coupled to shock front)

propagation speed of pressure v_{pressure}:

 $V_{pressure} = V_{sound}$

 $V_{pressure} = V_{flame} > V_{sound}$

pressure venting:

possible

not possible $(v_{flame} > v_{sound}!!)$

explosion pressure ratio $r = p_{ex}/p_{initial}$:

r ≤ 25

much larger than for deflagrations

influence of geometry of enclosure on $r = p_{ex}/p_{initial}$:

no

yes, substantial

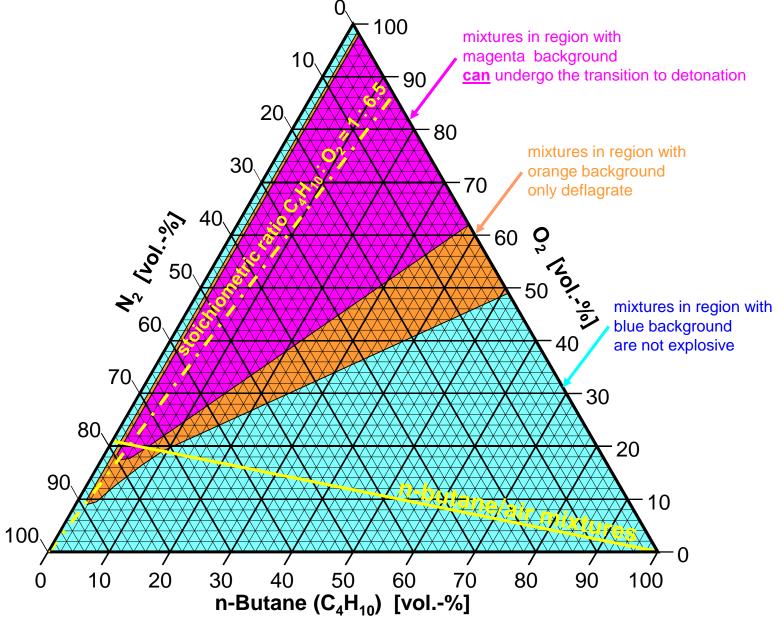
spatial pressure distribution:

same pressure at any location (⇒ no net force on containment)

substantial differences between pressures at different locations (huge net forces on containment)

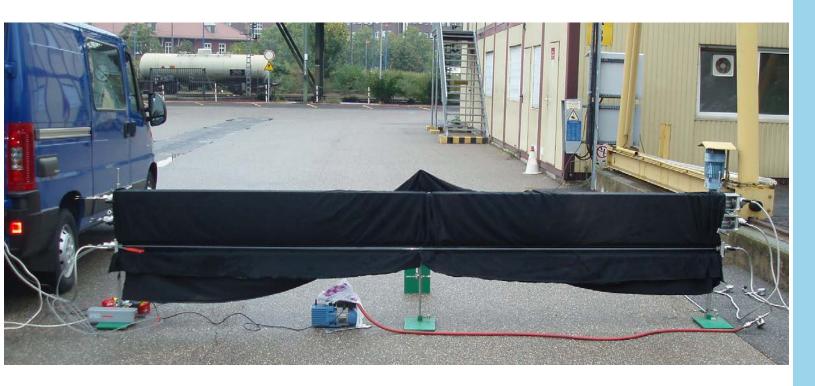


Example: Deflagrative and potentially detonative explosion regime of n-Butane/O₂/N₂ at 1 bar abs, 20 °C





Practical demonstration: Transition from deflagrative to detonative explosion of a gaseous mixture in a long pipe



Deflagration to Detonation transition in almost stoichiometric propane/air-mixture at about $90*\phi_i$ distance to point of ignition (=1.8 m), 3 bar abs, 15°C, pipelength = 4 m, ϕ_i = 20 mm

4.9 vol.-% C₃H₈, Versuch 9, slow motion

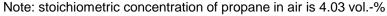
4.9 vol.-% C₃H₈, Versuch 9, real time

4.03 vol.-% C₃H₈, Versuch 7, slow motion

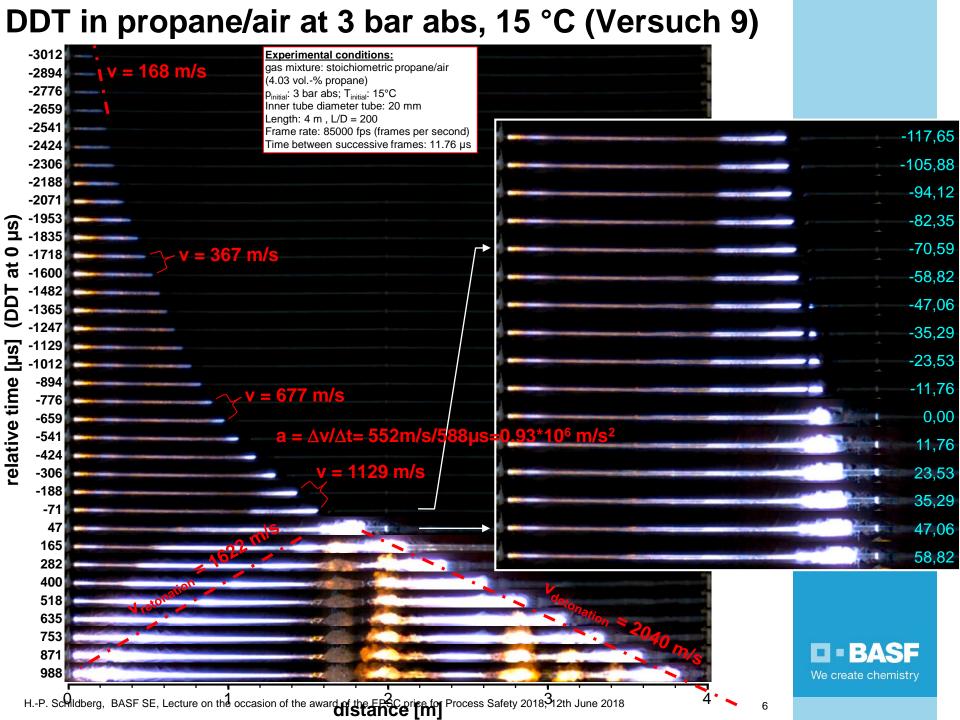
4.03 vol.-% C₃H₈, Versuch 7, real time











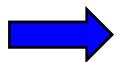
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Motivation

- In chemical process plants <u>detonable gas mixtures do occur</u> and effective ignition sources can, in general, not be ruled out with certainty
- The sole safety concept in this case is <u>explosion pressure proof design</u> of the affected plant components
- Worldwide there are <u>no guidelines published</u> by standardization organizations or interest groups (ISO, NFPA, ASME, CGA, CEN, EIGA, BSI, DIN, VDI) for explosion pressure proof design against the load generated by gas phase detonations
- Scientific literature:
 - Focussed mainly on the explosive mixture itself, not on the interaction mixture-enclosure
 - pressure/space/time profiles only understood for the two most simple detonative pressure scenarios with lowest pressure generation. No systematic classification of the remaining scenarios, not to mention their pressure/space/time profiles



BASF started research in 2008 aimed at developing a guideline for detonation pressure proof pipe design.



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Main results of the work on detonations in pipes

- Detonations in pipes can be described by <u>8 distinctly different pressure scenarios</u>:
 - 4 Scenarios in "long" pipes
 - 4 Scenarios in "short" pipes
- 6 scenarios are design-relevant
- An experimental method ("pipe wall deformation method") was established to directly determine the "static equivalent pressures" of each detonative scenario
- Once the static equivalent pressures are know, the <u>classical pressure vessel</u> formulae, which can only cope with static loads, can be applied for detonation pressure resistant design
- Results can be generalized to apply to any combustible/O₂/N₂ mixture by a parameter R, whose typical variation over the entire explosion triangle is provided.



Publications with Experimental Results

- [1] H.P. Schildberg, J. Smeulers, G. Pape, Experimental determination of the static equivalent pressure of gas phase detonations in pipes and comparison with numerical models, Proceedings of ASME 2013 Pressure Vessels and Piping Conference, Proc. ASME. 55690, Volume 5: High-Pressure Technology, ISBN: 978-0-7918-5569-0; doi: 10.1115/PVP2013-97677 (Paper No. PVP2013-97677, pp. V005T05A020; 15 pages; conference from July 14-18, 2013, Paris, France)
- [2] H.P. Schildberg, Experimental determination of the static equivalent pressure of detonative decompositions of acetylene in long pipes and Chapman-Jouguet pressure ratio, Proceedings of ASME 2014 Pressure Vessels and Piping Conference, Proc. ASME. 46025, Volume 5: High-Pressure Technology, ISBN: 978-0-7918-4602-5; doi: 10.1115/PVP2014-28197 (Paper No. PVP2014-28197, pp. V005T05A018; 13 pages; conference from July 20-24, 2014, Anaheim, California, USA
- [3] H.P. Schildberg, Experimental Determination of the Static Equivalent Pressures of detonative Explosions of Stoichiometric H₂/O₂/N₂-Mixtures in Long and Short pipes, Proceedings of the ASME 2015 Pressure Vessels and Piping Conference, Proc. ASME. 56987; Volume 5: High-Pressure Technology; Rudy Scavuzzo Student Paper Competition and 23rd Annual Student Paper Competition; ASME NDE Division, V005T05A015.July 19, 2015, PVP2015-45286, doi: 10.1115/PVP2015-45286 (Paper No. PVP2015-45286, 13 pages, conference from July 19-23, 2015, Boston, Massachusetts, USA)
- [4] H.P. Schildberg, Experimental Determination of the Static Equivalent Pressures of Detonative Explosions of Stoichiometric CH4/O2/N2-Mixtures and of CH4/O2-Mixtures in Long Pipes, Proceedings of the ASME 2016 Pressure Vessels and Piping Conference, (Paper No. PVP2016-63223, 14 pages, conference from July 17 - 21, 2016, Vancouver, BC, Canada)
- [5] H.P. Schildberg, Experimental Determination of the Static Equivalent Pressures of Detonative Explosions of Stoichiometric C2H4/O2/N2-Mixtures and of C2H4/O2-Mixtures in Long Pipes and of stoichiometric C6H12/O2/N2 Mixtures in long and short pipes (to be published in Proceedings of the ASME 2018 Pressure Vessels and Piping Conference)

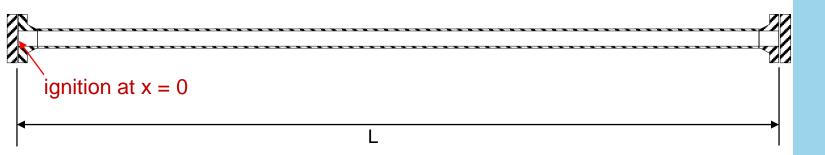


Publications giving a General Overview

- [7] H.P. Schildberg, Gas phase detonations in pipes: the 8 possible different pressure scenarios and their static equivalent pressures determined by the pipe wall deformation method. (part 1), Chemical Engineering Transactions, 48, 241-246; DOI:10.3303/CET1648041 (Proceedings of the 15th International Symposium on Loss Prevention and Safety Promotion in the Process Industries, 5.6.-8.6.2016, Freiburg, Germany. Guest Editors: Eddy de Rademaeker, Peter Schmelzer, Copyright © 2016, AIDIC Servizi S.r.l., ISBN 978-88-95608-39-6; ISSN 2283-9216)
- [8] H.P. Schildberg, Gas phase detonations in pipes: the 8 possible different pressure scenarios and their static equivalent pressures determined by the pipe wall deformation method. (part 2), Chemical Engineering Transactions, 48, 247-252; DOI:10.3303/CET1648042 (Proceedings of the 15th International Symposium on Loss Prevention and Safety Promotion in the Process Industries, 5.6.-8.6.2016, Freiburg, Germany. Guest Editors: Eddy de Rademaeker, Peter Schmelzer Copyright © 2016, AIDIC Servizi S.r.I., ISBN 978-88-95608-39-6; ISSN 2283-9216)
- [9] Technische Regel für Gefahrstoffe 407 (TRGS 407), Tätigkeiten mit Gasen Gefährdungsbeurteilung, Gemeinsames Ministerialblatt Nr. 12-17 (26.04.2016), p. 328 – 364, ISSN 0939-4729 Note 1: In the attachment A4 (page 48 – 56 of TRGS 407) the pressure scenarios in long pipes and their static equivalent pressure are for the first time mentioned in a guideline (here only related to detonative decompositions of acetylene).
 - Note 2: The TRGS 407 is published by German Bundesministerium für Arbeit und Soziales (Federal Ministry for Work and Social Affairs).
 - Note 3: Das Gemeinsame Ministerialblatt (GMBI) ist das amtliche Publikationsorgan der Bundesregierung und wird vom Bundesministerium des Innern seit 1950 herausgegeben. Hier veröffentlichen nahezu alle Bundesministerien die von ihnen erlassenen oder ergänzten Allgemeinen Verwaltungsvorschriften, Verordnungen, Richtlinien, Erlasse, Anordnungen, Rundschreiben und Bekanntmachungen von allgemeiner Bedeutung sowie Stellenausschreibungen einschließlich ihres nachgeordneten Bereichs.



Help to visualize the different detonative pressure scenarios

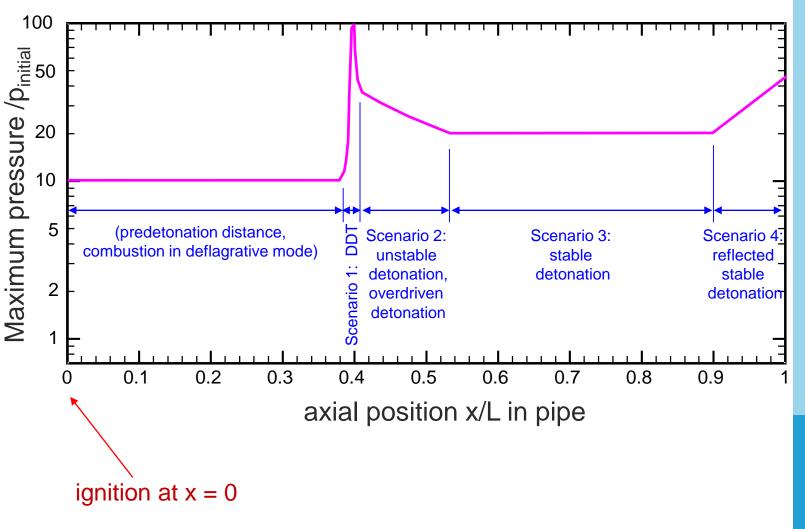


1st step: trigger an explosion with transition to detonation inside a pipe

2nd **step:** record the maximum pressure ratios found in the pipe at any axial position during the course of the explosion

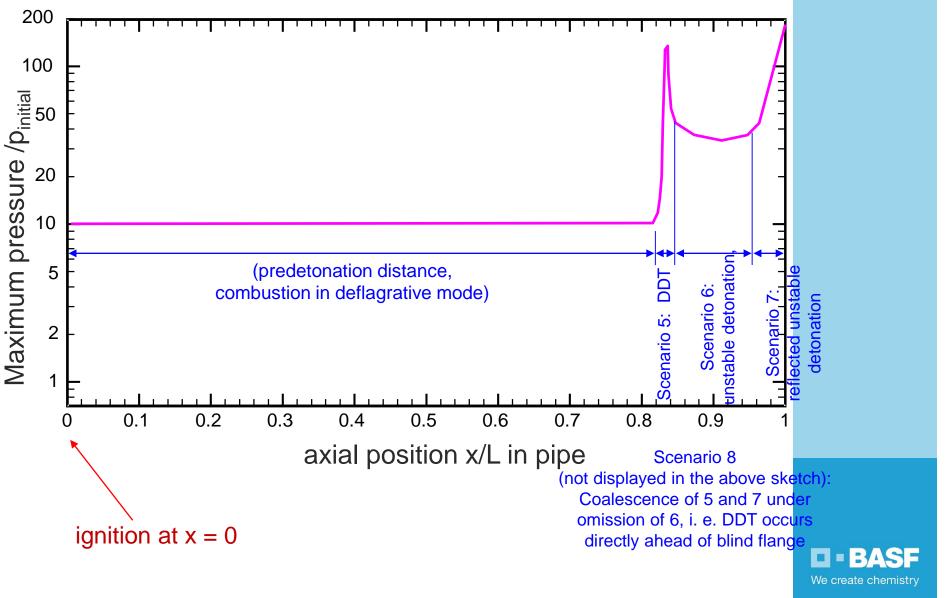


Maximum pressure ratios found in a long pipe at different axial positions in the course of an explosion involving a transition from deflagration to detonation (schematic)

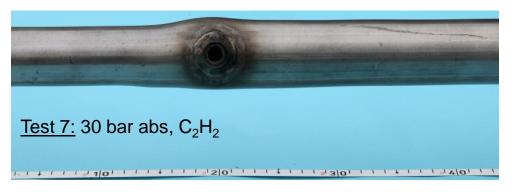


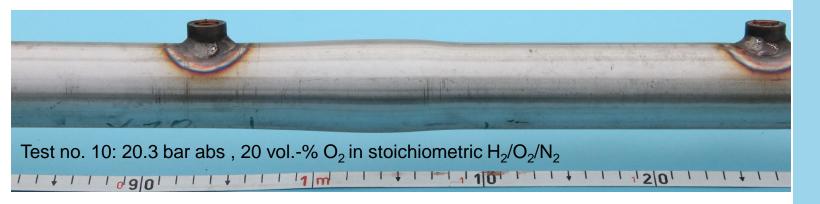


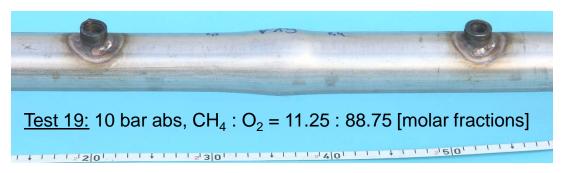
Maximum pressure ratios found in a <u>short</u> pipe at different axial positions in the course of an explosion involving a transition from deflagration to detonation (schematic)



Example: bulging in long pipes at DDT (scenario 1)

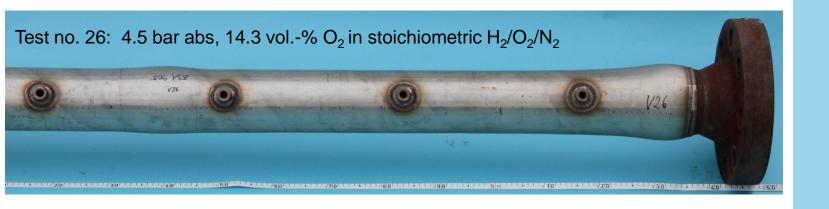


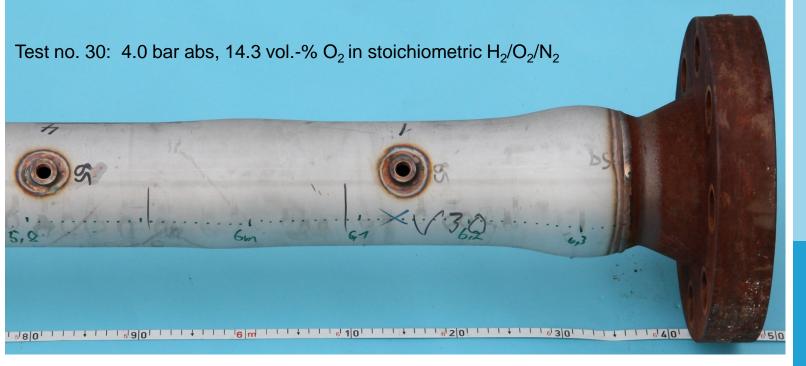






Example: bulging in short pipes (scenario 5 and 7)







Example: bulging in short pipes (scenario 8)

<u>Test no. 28:</u>

4.5 bar abs, 14.05 vol.-% O₂ in stoichiometric H₂/O₂/N₂



Test no. 29:

4.63 bar abs, 14.175 vol.-% O_2 in stoichiometric $H_2/O_2/N_2$





Static equivalent pressures for the 8 detonative pressure scenarios in pipes

7	Гуре с	of pressure scenario	Static equivalent pressures for any detonable
no.	pipe	name	gas mixture
1	pipe	DDT	$p_{\text{stat_DDT_long}} = R \bullet p_{\text{stat_stable}}$
2		unstable detonation	(irrelevant for pipe design)
3	long	stable detonation	$p_{\text{stat_stable}} = \alpha \cdot p_{\text{CJ_r}} \cdot p_{\text{initial}}$
4	<u> </u>	reflected stable detonation	$p_{\text{stat_reflected_stable}} = 2.4 \bullet p_{\text{stat_stable}}$
5		DDT	$p_{\text{stat_DDT_short}} = 1.5 \bullet p_{\text{stat_DDT_long}}$
			= 1.5 • R • p _{stat_stable}
6	short pipe	unstable detonation	(irrelevant for pipe design)
7		reflected unstable detonation	$p_{\text{stat_reflected_instable}} = 1.5 \bullet 2 \bullet p_{\text{stat_reflected_stable}}$
			$= 1.5 \cdot 2 \cdot 2.4 \cdot p_{stat_stable}$
8	S	coincidence of DDT and	$p_{\text{stat_coincidence_DDT_reflection}} = 2.4 \bullet p_{\text{stat_DDT_short}}$
		reflection	= 2.4 • 1.5 • p _{stat_DDT_long}
			= 2.4 • 1.5 • R • p _{stat_stable}

Factor **R** depends on reactivity of gas mixture

Formulae for p_{stat} are valid for any other explosive gas mixture at any p_{initial} and T_{initial}.

Short pipe scenario can be predicted based on long pipe scenarios 1 and 3

Note:

- $\alpha = 0.7$ (valid in general)
- p_{C_v} of the mixture can be calculated (based on combustion enthalpy, mean molar mass and c_p/c_v -values)
- 2.4 applies for reflection of the stable detonation and is assumed to also apply for reflection of instable detonations and DDT's
- R must be determined experimentally (ratio between effective load at DDT and effective load for stable deto.)
- justification for using factors 1.5 and 2 -> next slide

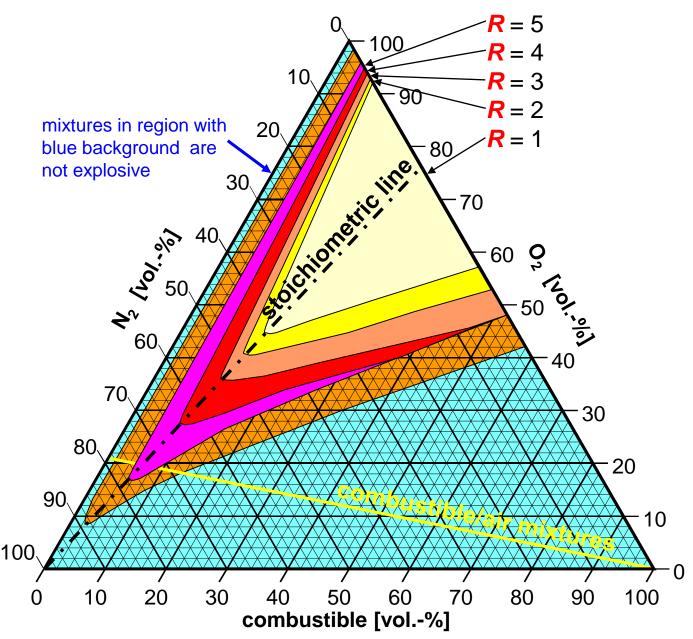


Example: static equivalent pressures <u>measured</u> for detonations of stoichiometric Ethylene/air mixtures at 15°C

	Туре	of pressure scenario	p _{stat}
no.	pipe	name	(expressed as multiple of p _{initial})
1	(D)	DDT	64.6
2	long pipe	unstable detonation	-
3		stable detonation	13.2
4)	reflected stable detonation	33
5	short pipe	DDT	81.8
6		unstable detonation	-
7		reflected unstable detonation	128
8	S	coincidence of DDT and reflection	232



Variation of R over the explosive range of a ternary mixture of type combustible/ O_2/N_2 (tentative)



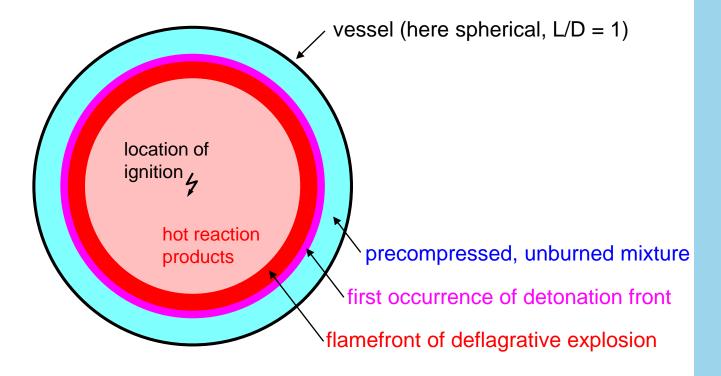


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Fundamental problem when deflagration to detonation transitions occur in empty vessels

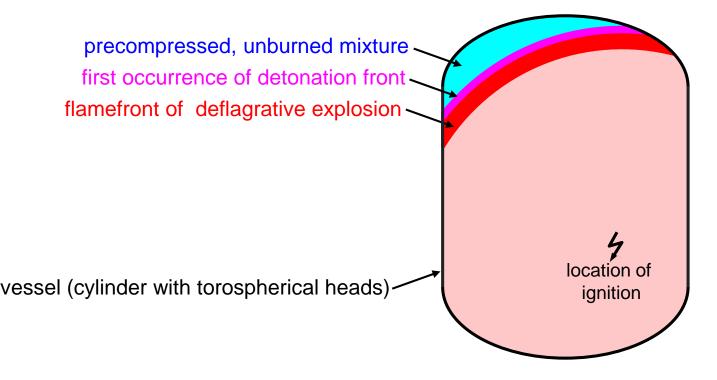


If there is a transition from deflagration to detonation in the vessel, precompression will almost always occur, because the diameter is usually not much larger than the predetonation distance. The precompression factor may attain the highest possible value, i. e. the deflagration pressure ratio.

Note that the detonation propagates faster than the speed of sound in the reaction gases, i.e. pressure relief into the central section of the vessel occurs <u>after</u> the wall has "seen" the detonative pressure peak



Schematic sketch of an explosion with DDT inside a vessel



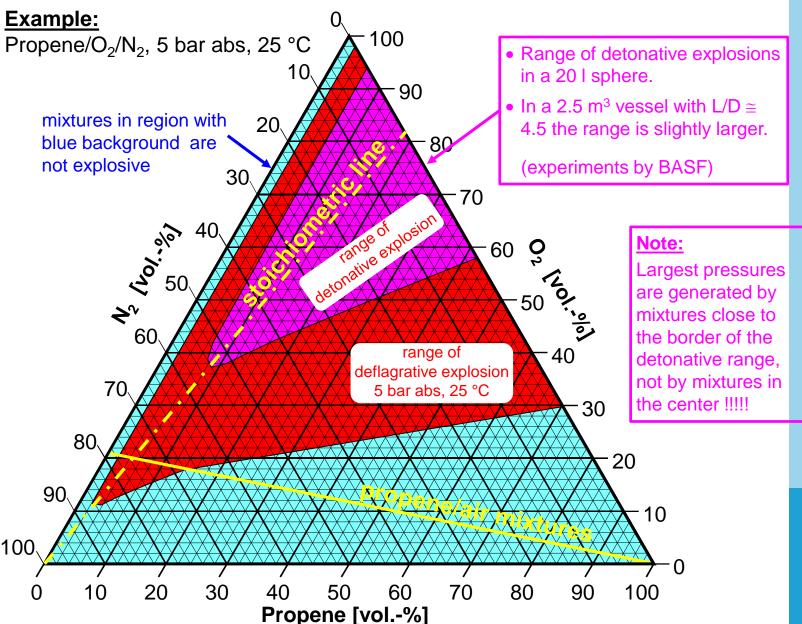
If a DDT occurs at this time instant, maximum pressure loads can be expected, because the unreacted mixture is almost precompressed to r·p_{initial}

(r denotes the deflagration pressure ratio, p_{intial} the pressure in the vessel at the moment of ignition)

Note that the detonation propagates faster than the speed of sound in the reaction gases, i.e. pressure relief into the lower section of the vessel occurs <u>after</u> the wall in the upper left section has "seen" the detonative pressure peak

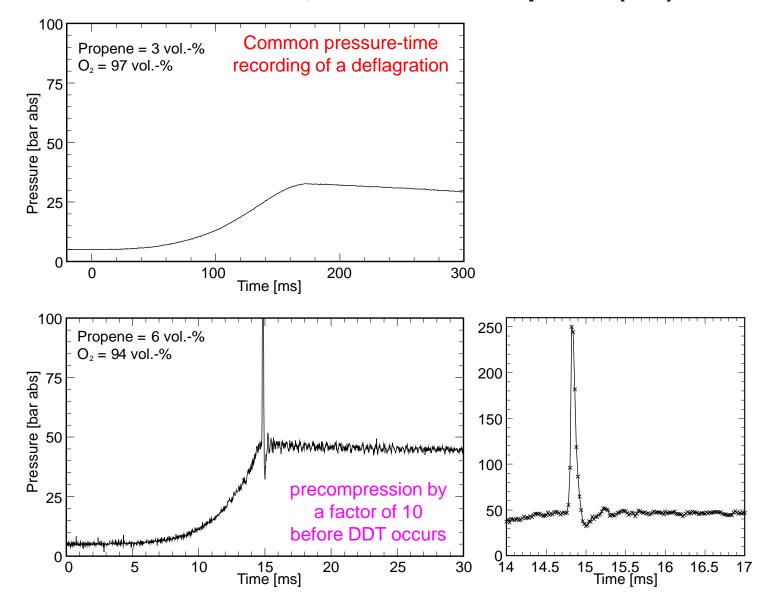


Mixtures which undergo a DDT inside vessels





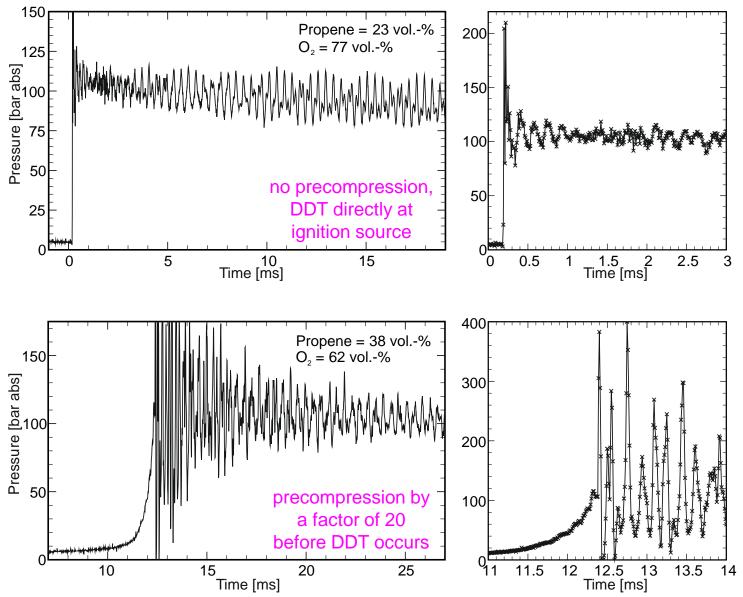
Pressure/Time recordings of explosions of Propene/O₂-mixtures at 5 bar abs, 20°C in a 20 I sphere (1/2)





Reference: "The course of the explosions of combustible/O₂/N₂ mixtures in vessel-like geometry", H.-P. Schildberg, Forschung im Ingenieurwesen (2009) 73, 33-65, DOI 10.1007/s10010-009-0091-6

Pressure/Time recordings of explosions of Propene/O₂-mixtures at 5 bar abs, 20°C in a 20 I sphere (2/2)





Fundamental questions when quantifying hazards associated with potential detonations in vessels

- Under what conditions (mixture composition, p_{initial}, T_{initial}, volume, L/D) do we have to assume that a DDT occurs?
- What is the largest conceivable value of the static equivalent pressure p_{stat}? (it should be larger than p_{stat} of the pipe scenario no. 8).

Hint: In vessels having dimensions as used in the process industry the width of the detonative peaks hitting the wall will be much less than half of the cycle time of the fundamental oscillation modes. This helps to reduce p_{stat} . But excitation is usually unsymmetric and thereby triggers all higher harmonics, which have much smaller cycle times.

How to cope with the massive net forces acting on the vessel, which cause the vessel to be displaced?



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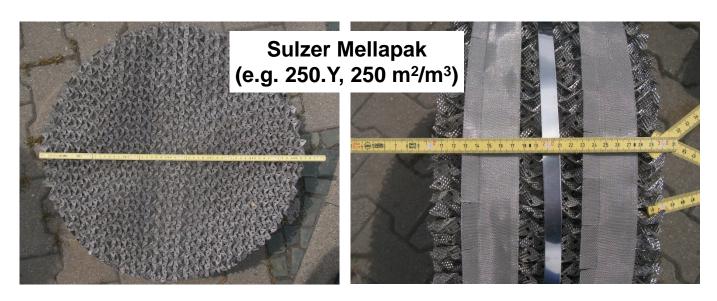
Typical packings used in the process industry

Raschig Rings, L/D = 1, typical: 15 mm \leq L \leq 50 mm



Pall Rings, L/D = 1, typical: 15 mm \leq L \leq 50 mm







Typical packings used in small-scale equipment





Typical applications of vessels filled with dry packings

- Vessels acting as demisters
- Vessel acting as flame arrestors
 (e.g. for self decomposable gases like Acetylene or Ethylene Oxide)
- Vessels acting as static mixers



Fundamental questions pertaining to the course of explosions in vessels with dry packings

Can mixtures, which explode in deflagrative manner in empty vessels, transition to detonation in packed vessels?

Yes! Example: hydrocarbon/air mixtures do not transition to detonation in empty vessels, but will do in a packing, if the packing diameter is larger than 1/3 of the detonation cell size of the mixture.

If a DDT occurs:

- Can we specify a <u>predetonation distance</u> as function of the dimensions of the packing elements? Yes! Example: stoichiometric hydrocarbon/air mixtures at 20 °C and 1 bar abs < p_{initial} < 5 bar abs have a predetonation distance of about 100·φ_i in straight pipes (φ_i is the inner pipe diameter). In packings this distance is much smaller (φ_i denotes the packing diameter).
- How large is the maximum <u>precompression factor</u> relative to the maximum precompression factor in the empty vessel? (specific BASF know how)
- How large is the <u>static equivalent pressure</u> acting on the wall of the vessel compared to a detonation in an empty vessel? (specific BASF know how)

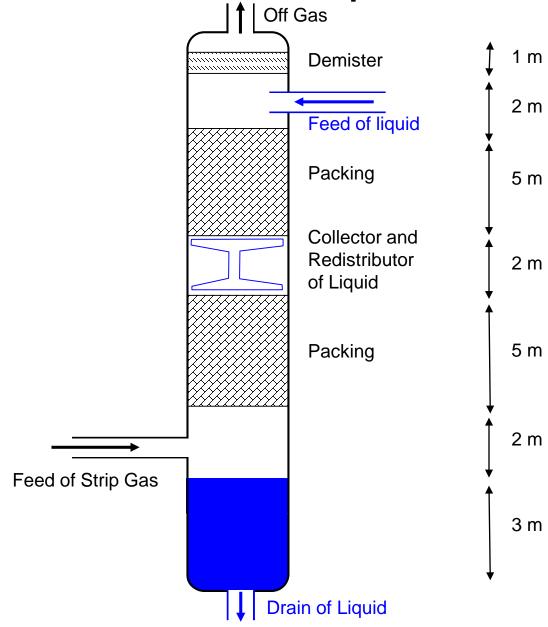


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General application of an irrigated packing: distillation column or desorption tower





Fundamental questions pertaining to the course of explosions in vessels with irrigated packings

Can a DDT, which would happen inside the dry packing, be suppressed by the irrigation?

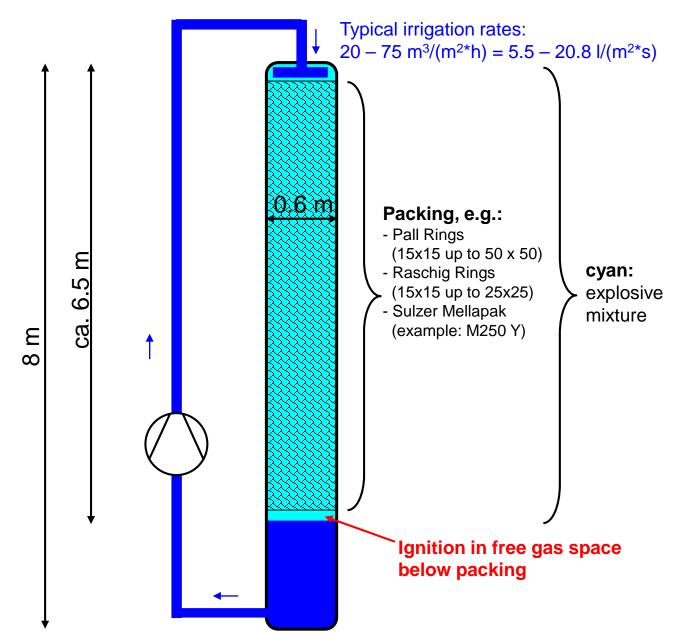
- If the DDT can be suppressed:
 - What is the required irrigation rate as function of p_{initial}?
 - What is the required irrigation rate as function of the diameter of the packing elements?

Answers

- Answers do very much depend on the process conditions of the system under investigation
- Number of investigated systems is still too small to generalize the results



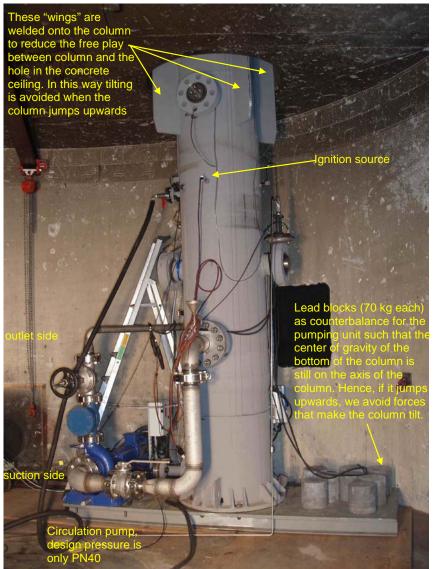
Vessel geometry used by BASF for experiments





Example: PN250 column, 8 m long, installed in bunker K348 for testing suppression of DDT by irrigation

Lower part of the column in the test room of the bunker K348





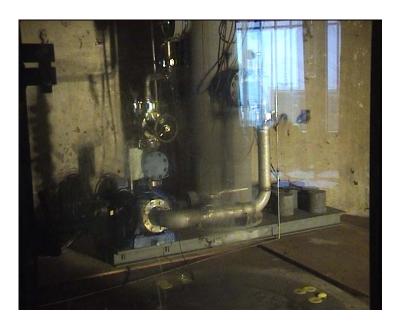
View on the sparger for water mounted under the upper blind flange



(here with rust, picture taken after test no. 4)



Example: Tests without and with detonation in PN250 column filled with 50x50 Pall Rings



Acetylene:

 $p_{initial} = 8 \text{ bar abs}$

irrigation rate:

Pall rings: 50mm x 50mm

no DDT!!!



Acetylene:

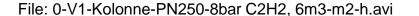
 $p_{initial} = 12 \text{ bar abs}$

irrigation rate:

Pall rings:

50mm x 50mm

DDT !!!



0-V2-Kolonne-PN250-12bar C2H2, 6m3-m2-h.AVI



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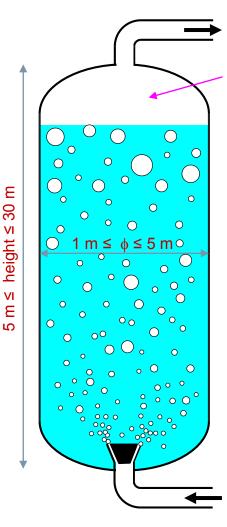
Schematic sketch of a gas-liquid partial oxidation or vinylation process (injection from bottom)

Characteristics of process:

- Large hold-up of organic liquid
- Injection of a gaseous oxidant or gas mixture containing the oxidant
- 5 to 20 % of the liquid are taken by gas bubbles
- Large fraction of the bubbles can be in the <u>explosive range</u> due to vapour of the organic liquid (in case of reaction breakdown all bubbles can become explosive)
- Gas space in reactor dome is usually in <u>explosive range</u> due to organic vapour and/or organic mist
- Ignition sources can not be excluded (mostly: chemical ignitors)

Note:

Explosive range encompasses the purely deflagrative regime and the potentially detonative regime



off gas pipe

Upper reactor dome: Usually with detonable gas phase (oxidant, vapours, mists, inert gases)

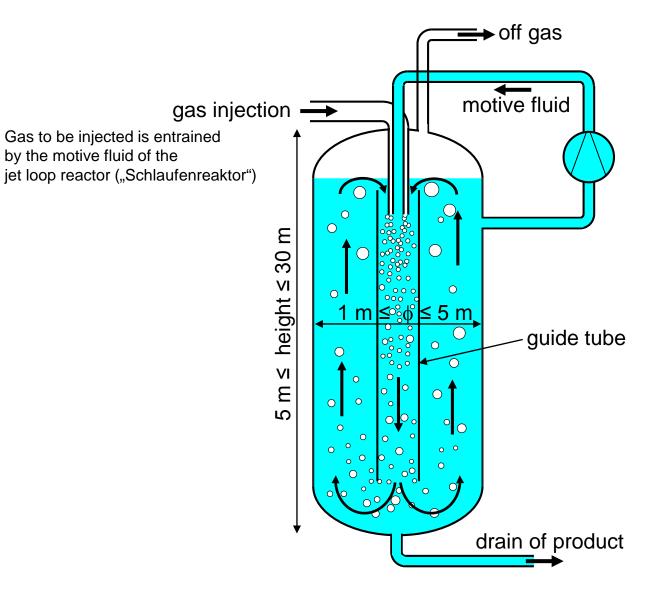
Black: Reactor, $20^{\circ}C \le T \le 300^{\circ}C$, $6 \text{ barg} \le p_{\text{design}} \le 325 \text{ barg}$

Blue: Liquid to be partially oxidized or vinylated, $20^{\circ}C \le T \le 300^{\circ}C$, $0 \text{ barg} \le p_{\text{operation}} \le 30 \text{ barg}$ $5 \text{ m}^3 \le V \le 500 \text{ m}^3$

Feed gas pipe for oxidant containing gas mixture or acetylene: $O_2/N_2/CO_2/CO/H_2O, C_2H_2$ 0 barg $\leq p_{operation} \leq 30$ barg $1000 \text{ Nm}^3/h \leq V \leq 50000 \text{ Nm}^3/h$



Schematic sketch of a gas-liquid partial oxidation or vinylation process (injection from top with motive fluid)





Typical applications with the risk of bubble swarm detonations triggered by detonation in head space of reactor

- Partial oxidation reactions of organic liquids with air,
 O₂ or N₂O
- Vinylation reactions of organic liquids



Fundamental questions pertaining to the course of bubble swarm detonation triggered by detonation in the head space of the reactor

- Can the bubbles be ignited by adiabatic compression generated by the shockwave in the liquid?
 (Assumption: shock wave is triggered by gas phase detonation in head space)
- If ignition occurs:
 - Can there really be a DDT inside the bubbles?
 - How large is the <u>static equivalent pressure</u> acting on the wall?
 - Will all conceivable courses of the explosion in the bubbles (homogeneous runaway reaction, deflagration, detonation) lead to the same static equivalent pressure acting on the wall?



Research data not suited to assess bubble swarm detonation triggered by detonation in head space

No.	Parameter	Reactors used in	
		published research	process industry
geometric data			
1	dimensions of reactors	L ≤ 1 m;	L ≤ 15 m; φ _i ≤ 6 m
2	volumetric gas fraction in liquid	very small (mostly single bubbles)	1 to 20 vol%
3	bubble diameter	≤ 20 mm	≤ 200 mm
experimental results			
4	ignitability of bubbles containing explosive mixture	yes	(unknown)
5	pressure at wall caused by exploding bubbles	negligible	(unknown)
6	water hammer	negligible	(conceivable)

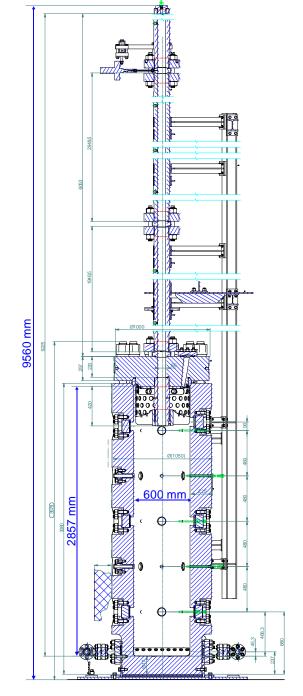
Due to large differences in (1), (2) and (3), the results from lab scale tests are **not suited** to assess real scale scenarios!!

(e.g. liquid in real reactors has much higher compressibility (due to higher gas fraction) and will be accelerated to much higher speed (combined effect of higher gas fraction and larger length of reactor)



Experimental setup of BASF to investigate bubble swarm detonation

First results to be expected by the end of 2018



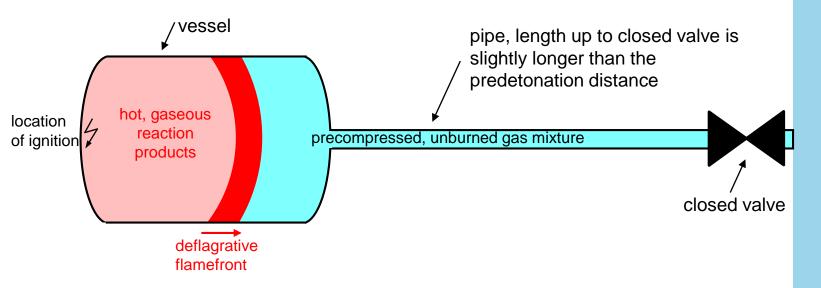


Overview

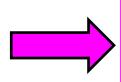
- Brief background info: deflagrative and detonative explosions in gaseous mixtures
- Motivation
- Detonations in pipes
- Detonations in empty vessels (i.e. no turbulence enhancing elements inside)
- Detonations in vessels filled with dry packings
- Detonations in vessels filled with irrigated packing
- Detonations in bubble swarms rising upwards in a liquid
- Particularly dangerous geometries in context with detonations
 - Short pipe connected to large vessel
 - Long pipe connected to large vessel
- Present status of adopting the pipe results in regulation, guidelines, standards etc.



Problem with short pipe connected to large vessel



- Many gas mixtures only deflagrate in vessel-geometry but undergo transition to detonation in pipe geometry
- Usually: volume of vessel >>> volume of short pipe



At the moment when the deflagrative flame front reaches the point where the pipe is tied in, the unburned mixture in the pipe is precompressed by a factor equal to the deflagration pressure ratio (typically 4 to 25).

When, upon further propagation, the flame transitions to detonation, the <u>resulting pressures are extremely large</u>

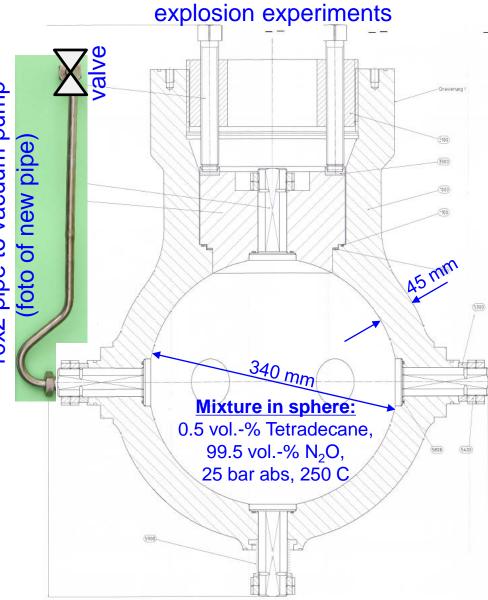


Example 1: 20 I sphere with 45 cm pipe (ϕ_i = 6 mm)

10x2 pipe to vacuum pump

10x2 pipe ruptured at location of DDT, thereafter the curved part molt.

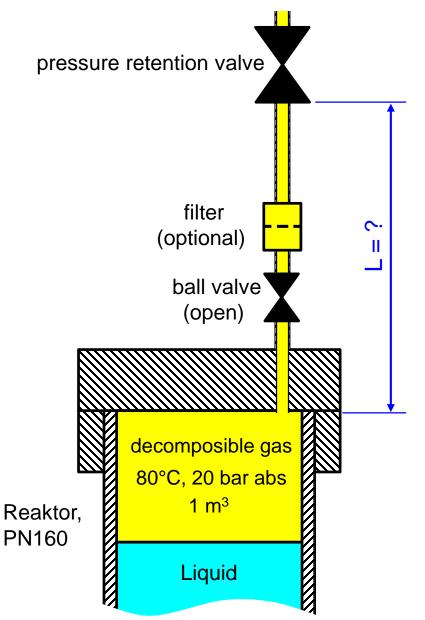
(hydraulic burst pressure of pipe is about 3480 bar; pipe material: 1.4541; deflagration pressure ratio of the mixture was only ca. 5 at T_{initial} = 250 C)



20 I sphere used for



Example 2: 1 m³ vessel with DN80 pipe

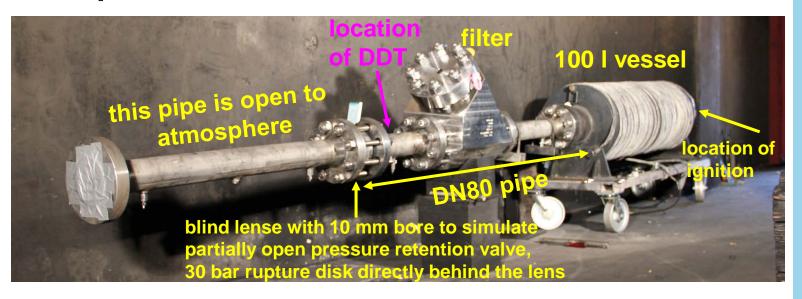


Question:

What is the maximum permissible length L of the DN80 pipe such that a deflagrative decomposition starting in the headspace of the reactor does not transition to detonation in the pipe?



Example 2: result of a real-scale test

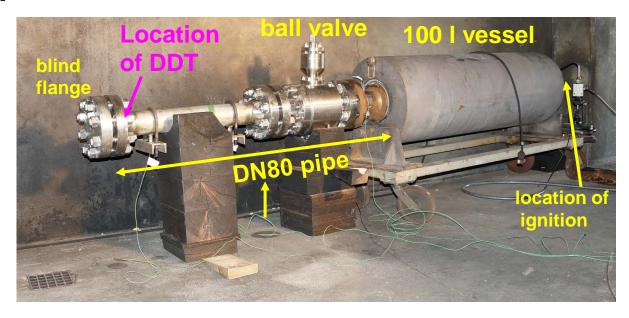








Example 2: result of a real-scale test







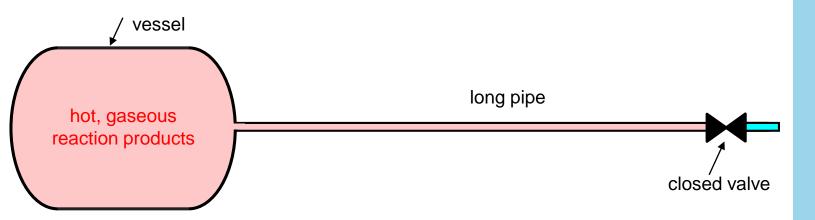


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Problem with long pipe connected to a large vessel



- Explosive mixture present in vessel and pipe explodes in deflagrative or detonative manner. The resulting pressures are sustained by the equipment.
- Usually: volume of vessel >>> volume of long pipe

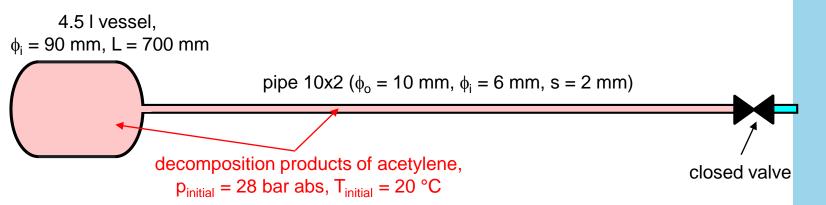
The cooling rates in the pipe are much faster than in the vessel.

- ⇒ Hot reaction gases (2400 K to 3000 K) flow from the vessel into the pipe.
- ⇒ Excessive heating of pipe at point where connected to vessel
- ⇒ Rupture of pipe at that point because yield strength R_{p0.2} drops to very low values

Note: Let n denote the number of moles in the pipe directly after the explosion has terminated. Then the pipe will typically receive 9*n hot moles from the vessel within a short time span after completion of the explosion.



Example 1: vessel with long pipe



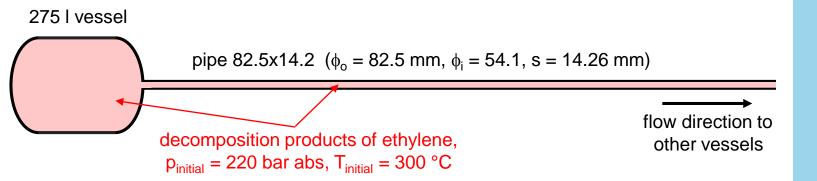
Case 1: pipe length is 1.5 m: no rupture

Case 2: pipe length is 4.3 m: rupture





Example 2: 275 I vessel with 82.5x14.2 pipe



- Ignition occurred in vessel (decomposition reaction is slow)
- About 60 s after ignition: pipe ruptured at flange where it was connected to the vessel due to excessive heating



Example 3: laboratory setups with small Swagelock pipes and Whitey-bombs

<u>Test conditions:</u> decomposition of acetylene, $p_{initial} = 28$ bar abs, $T_{initial} = 20$ °C

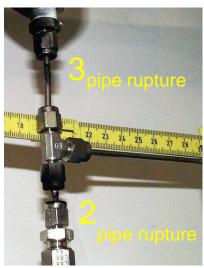
Pipes: 3.16 x 0.5, 6 x 1

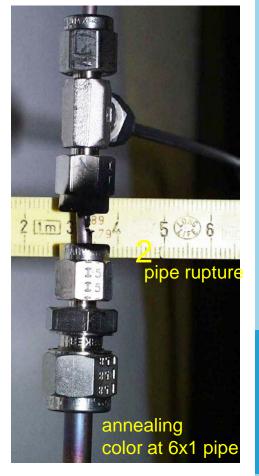
Vessels: Whitey bombs with 0.1 to 1 l, design pressures ca. 200 to 300 bar

After the decomposition reaction has terminated, flow of hot reaction products sets in from somewhere to somewhere and wall sections under high thermal load rupture.











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State of adopting the pipe results in regulation, guidelines, standards etc

- Germany: TRGS 407, Anhang 4
 - long pipe scenarios for C₂H₂-Detonations included
 - a pipe is considered as detonation pressure resistant only on basis of the wall thickness, not on basis of its official design pressure.
- ASME: In 2015 a working group was established to develop a new ASME code case on detonation pressure resistant pipe design. Work has been postponed so far due to work overload.
- NFPA: NFPA 67 "Guide on Explosion Protection for Gaseous Mixtures in Pipe Systems" is going to be revised shortly, in particular chapters 5 to 8 dealing with principles of detonations in pipes. Members of BASF Corporation take part.



- Still much work to be done to make the pipe results "penetrate" the existing guidelines.
- Possibly: Extra chapter to be included in DIN EN 13480

 Metallic industrial piping Part 3: Design and calculation





The Chemical Company