

Explosion Hazards of Hydrogen-Air Mixtures

Professor John H.S. Lee
McGill University, Montreal, Canada



Hydrogen Safety Issues

- Wide spread use of hydrogen requires significant efforts to resolve safety issues
- Hydrogen is already used extensively in many industrial applications (but general public not exposed to the dangers)
- Extensive research efforts have already been devoted to hydrogen safety issues
- Post-Three Mile Island accident – information not widely disseminated

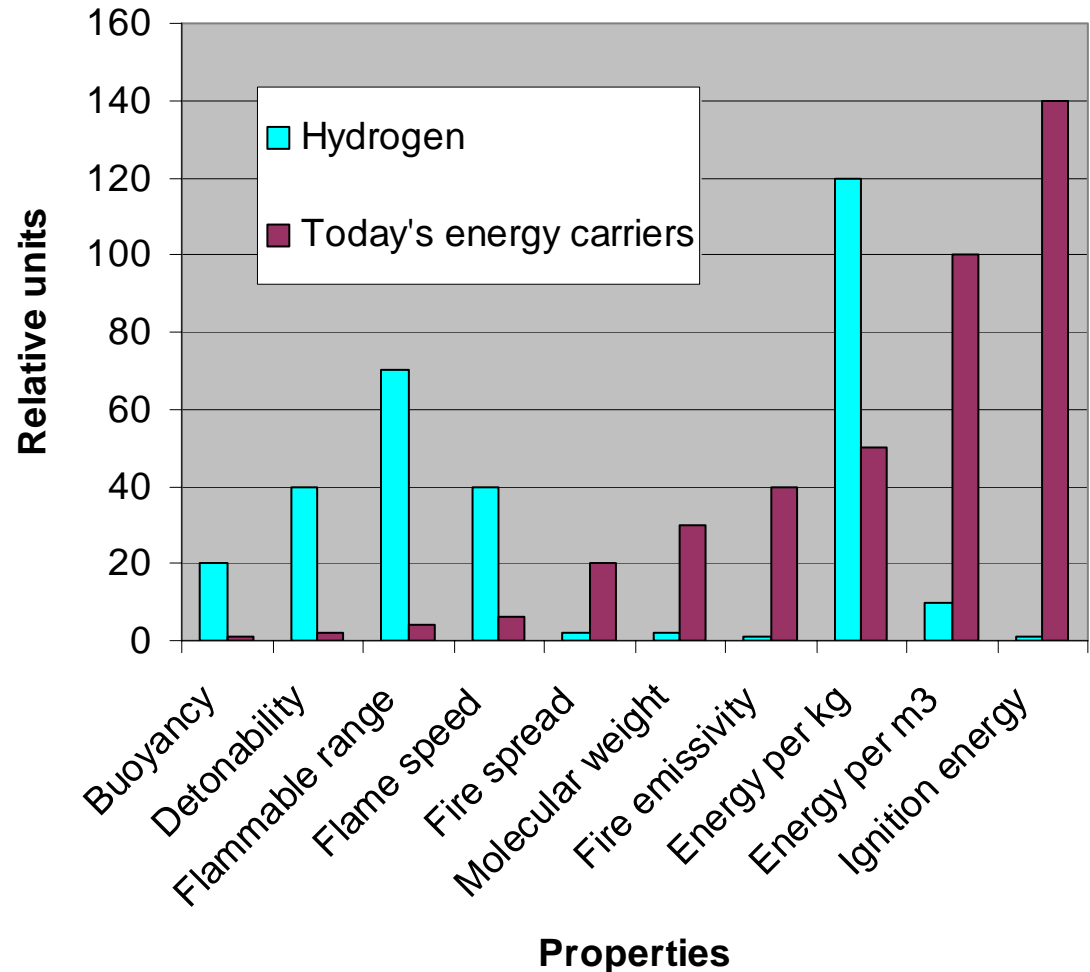
Hydrogen Safety Research

BEFORE HYDROGEN CAN BE USED AS A COMMON ENERGY CARRIER:

- Achieve public acceptance of hydrogen-technologies
- Provide at least the same level of safety, reliability, comfort as today's fossil fuels
- No solutions are available in terms of widely accepted standards, methodologies, mitigation techniques and regulations)

Hydrogen and today's fuels

- Qualitative comparison of “Safety profiles”
- Properties of hydrogen are different from today's fuels
 - H_2 is less dangerous in terms of thermal and fire hazards,
 - may be responsible for stronger pressure effects



Safety Issues

- To evaluate hydrogen safety the following set of issues should be addressed for each of the applications
- Hydrogen release, mixing, and distribution
 - Thermal, pressure, and missile effects from H₂ fires and H₂-air cloud explosions
 - Mitigation techniques for detection, dilution, and removal of hydrogen
 - Risk evaluation, both specific and in comparison with today's fossil energy carriers
 - Standardization, and regulatory issues

Objectives

- To contribute to common understanding and approaches for addressing hydrogen safety issues
- To integrate experience and knowledge on hydrogen safety
- To integrate and harmonise the fragmented research base
- To provide contributions to safety requirements, standards and codes of practice
- To contribute to an improved technical culture on handling hydrogen as an energy carrier
- To promote public acceptance of hydrogen technologies

Accident scenarios

Unconfined Explosions

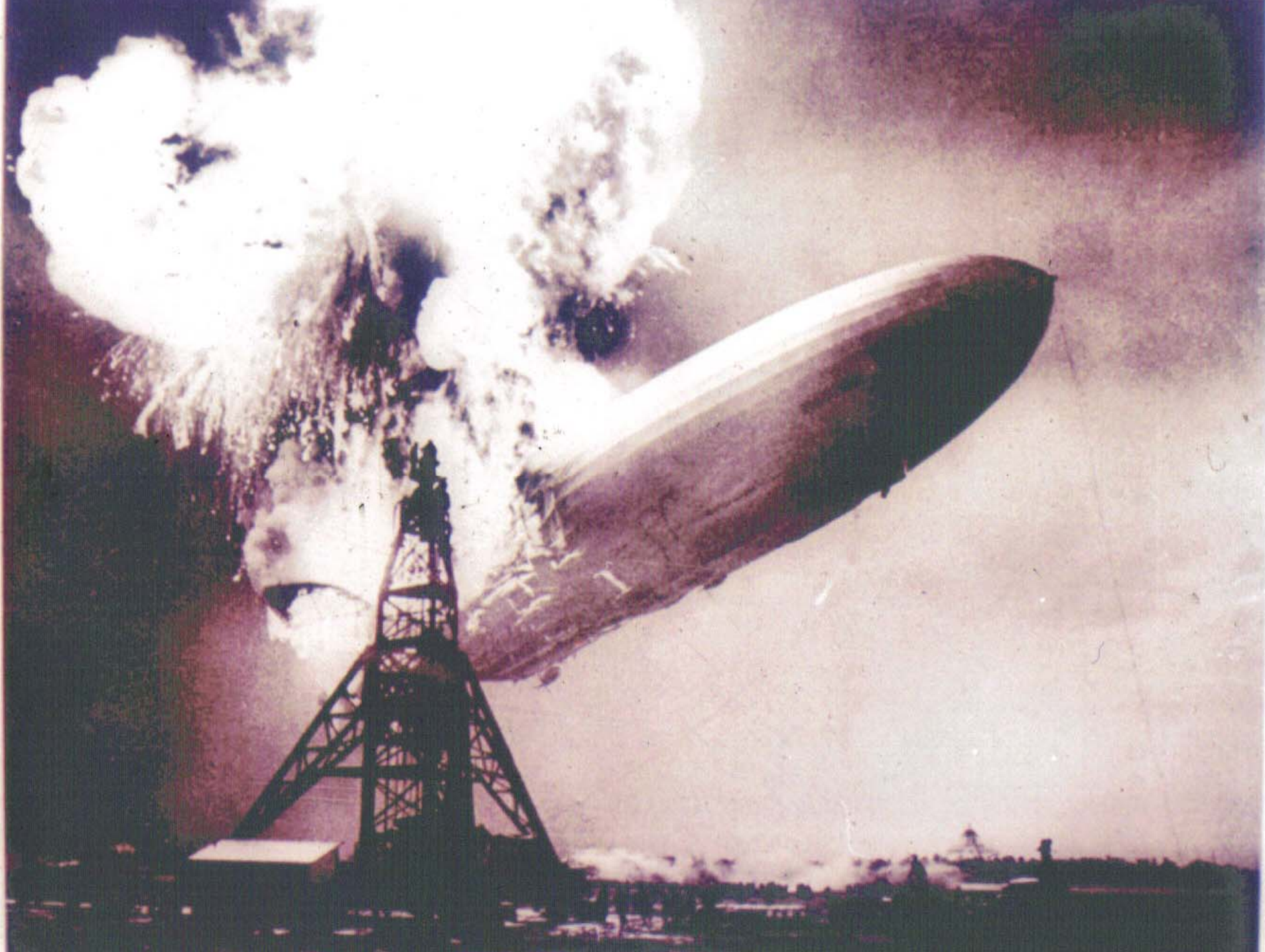
- major rapid release into the atmosphere

Confined Explosions

- leakage of H_2 into buildings
- contamination of high pressure H_2 storage facilities by air

Hindenburg (May 6, 1937)

- Lakehurst (New Jersey)
- Fired started near tail during landing
- Flame spread ~ 50 m/s
- Ship was 803 ft. ~ 245 m long
- Destruction completed in 32 seconds
- 36 lives lost





Crescent City, Illinois



Crescent City, Illinois



Jackass Flat (Nevada)

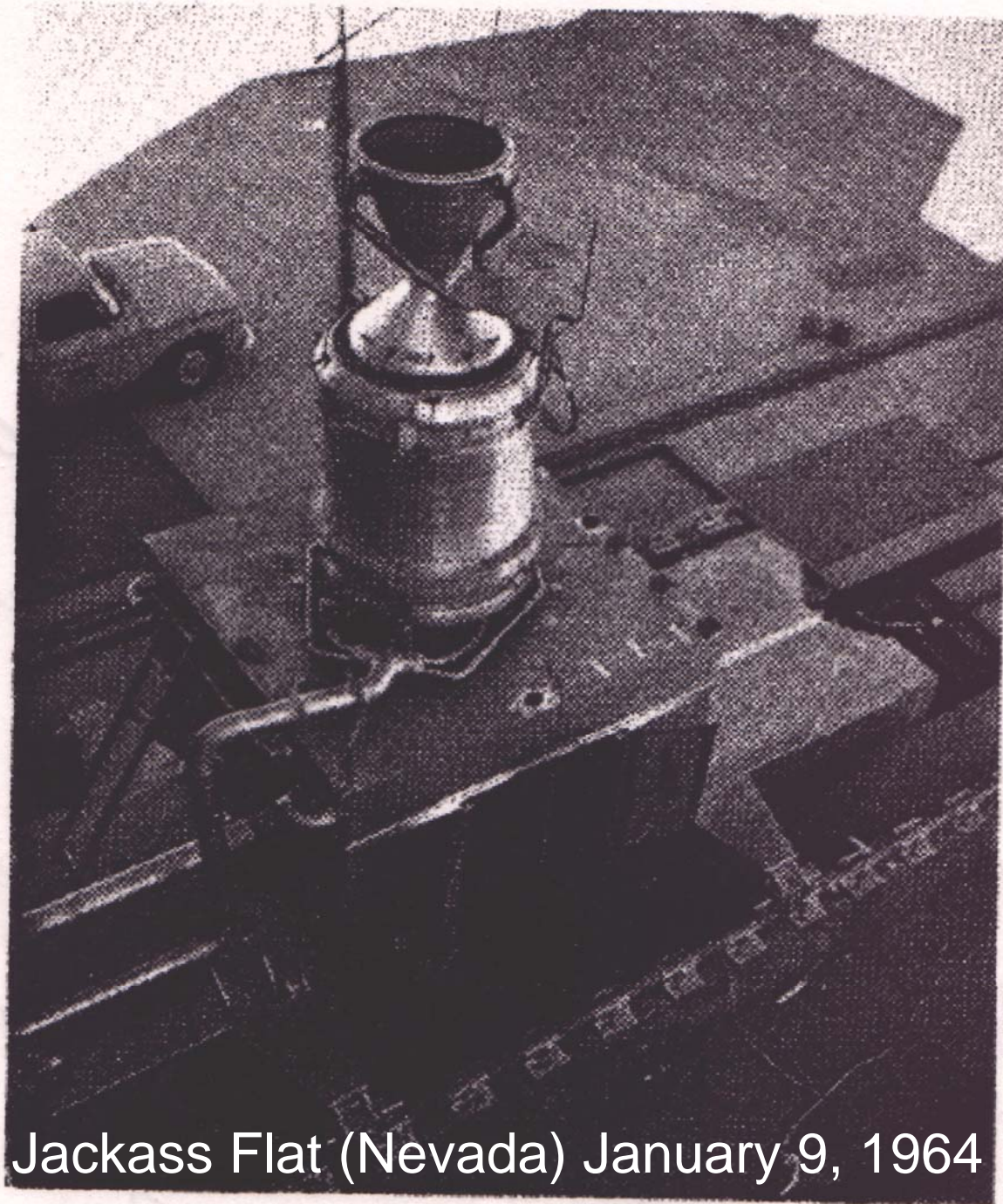
January 9, 1964

- Unconfined H₂-air explosion
- Test to measure acoustic noise due to high flow rate hydrogen
- 1000 kg H₂ discharged from vertical rocket nozzle at 23 MPa in 30 seconds
- Discharge rate uniformly increased to 55 kg/s, maintained for 10 seconds then reduced to zero
- Ignition occurs 26 seconds after discharge begins

Jackass Flat (Nevada)

January 9, 1964

- No pressure wave detected in near field less than 0.8 km
- Explosion heard 3.2 km away
- Wide spread minor damage near hydrogen discharge, but superficial
- Estimate 10 kg of H₂ involved in the explosion
- TNT equivalent of 8%



Jackass Flat (Nevada) January 9, 1964

Polysar

(April 19, 1984)

- Unconfined H₂-air explosion
- Rapid release of H₂ from a ruptured gasket of a Worthington Compressor at 600psi
- 10-20 seconds delay before ignition
- Three fatalities
- Extensive major structural damage in the near field
- Glass and minor structural damage up to 1 km
- Detonation occurred in near field
- Damage compatible to detonation of about 0.1 kg H₂-air cloud







China Light and Power Cast Peak Generating Station (August 28, 1992)

- Confined explosion
- Explosion in hydrogen receiver
- Production of hydrogen by electrolysis
- Low pressure compressor: 500 kPa
- High pressure compressor: 13.6 MPa
- Two hydrogen receivers: 8.68 m long x 1.12 m diameter
- Hydrogen plant shut down August 24 to 26
- Hydrogen plant resume to supply H₂ to receivers @ 06:30 on August 27

China Light and Power Cast Peak Generating Station (August 28, 1992)

- Pressure at receiver: 6.9 MPa
- August 28 from 00:30 to 02:00 gas from receiver supplied to generator
- Hydrogen purity in generator dropped to 85%
- Receiver disconnected from generator at 02:30; H₂ supplied from bottles
- Sampling indicated hydrogen purity in receivers about 95%
- Receiver #1 reconnected to generator to supply H₂ to generator at 09:45 on August 28

China Light and Power Cast Peak Generating Station (August 28, 1992)

- A drop in H₂ purity in generators noted immediately
- Both receivers exploded at 10:05
- Two fatalities; 18 injured by fragments
- Extensive blast damage ~ 100 m radius
- TNT equivalent 275 kg
- Conclusion: all the gas supplied to the receiver over a 20 hour period (from 06:30 on August 27 to 02:30 on August 28) was air!

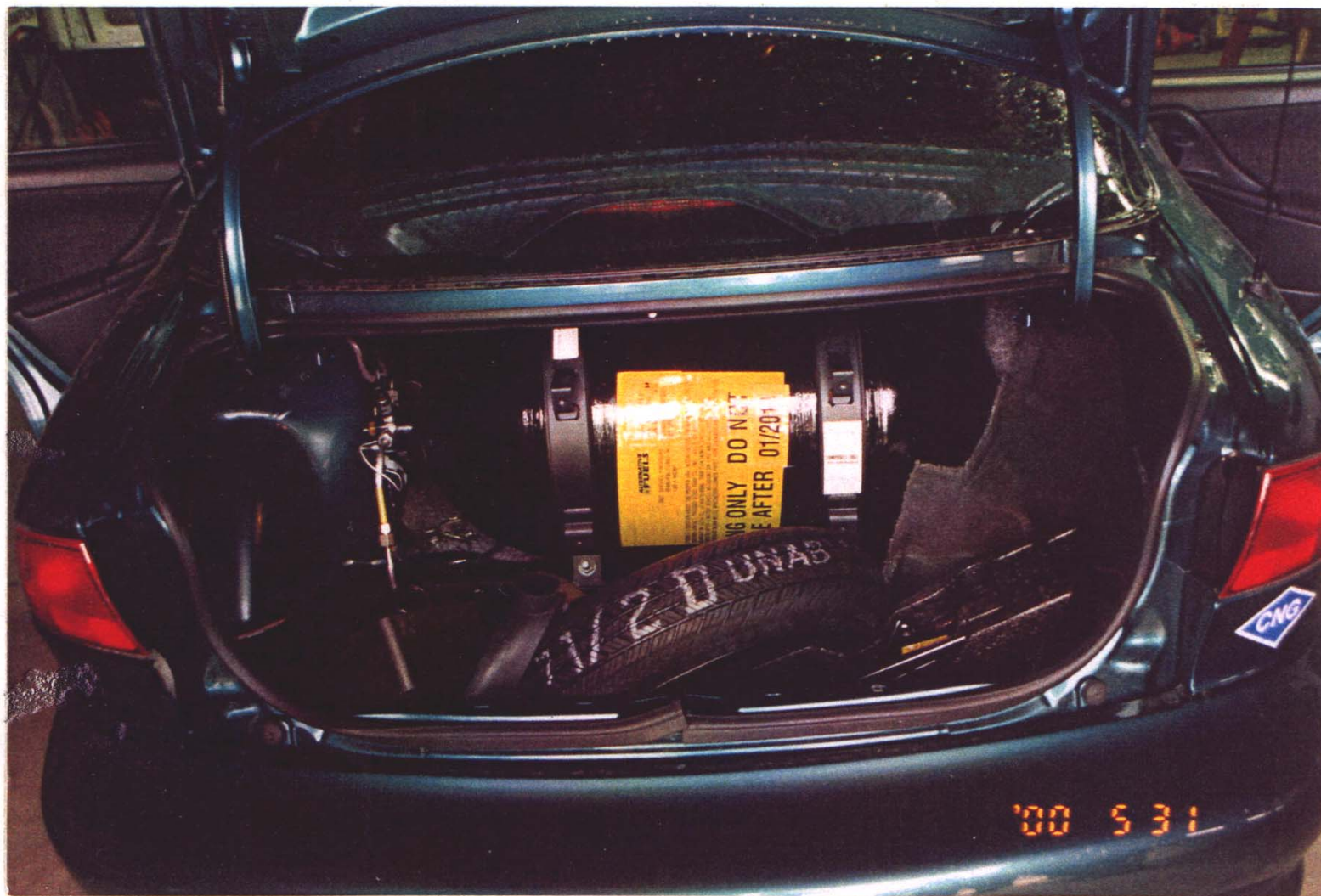


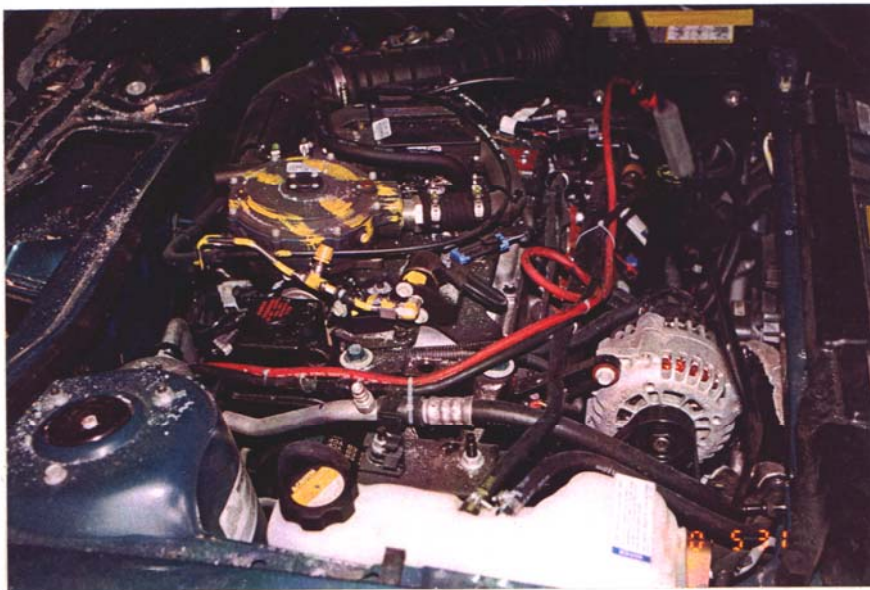


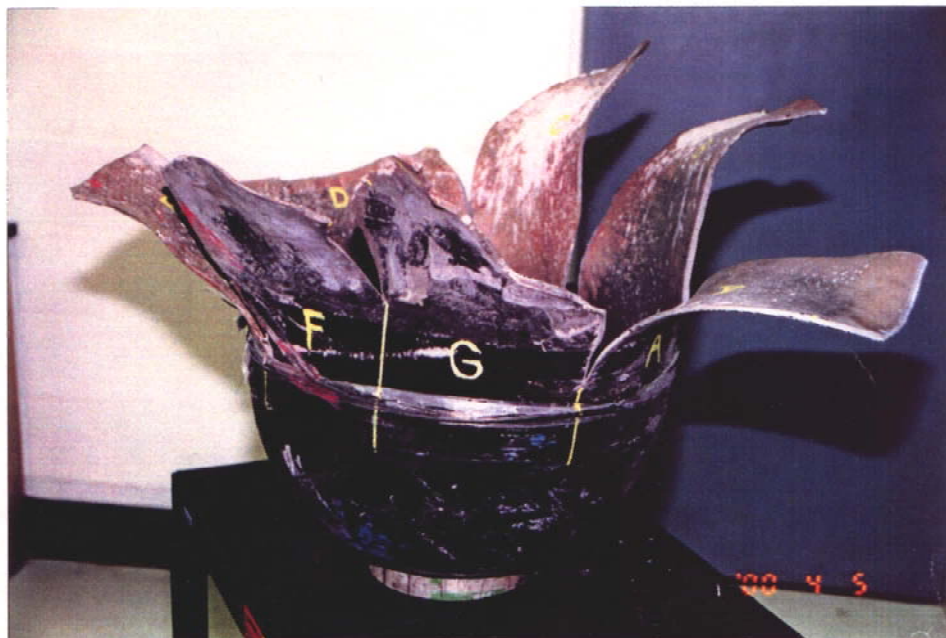


Blainville, Quebec (March, 2000)

- Confined explosion
- Motor vehicle test center
- Tank with 350 psi natural gas filled with air to 3500 psi instead of nitrogen
- Explosion occur during pressure adjustment before crash test
- Extensive damage to car and building
- 3 workers killed







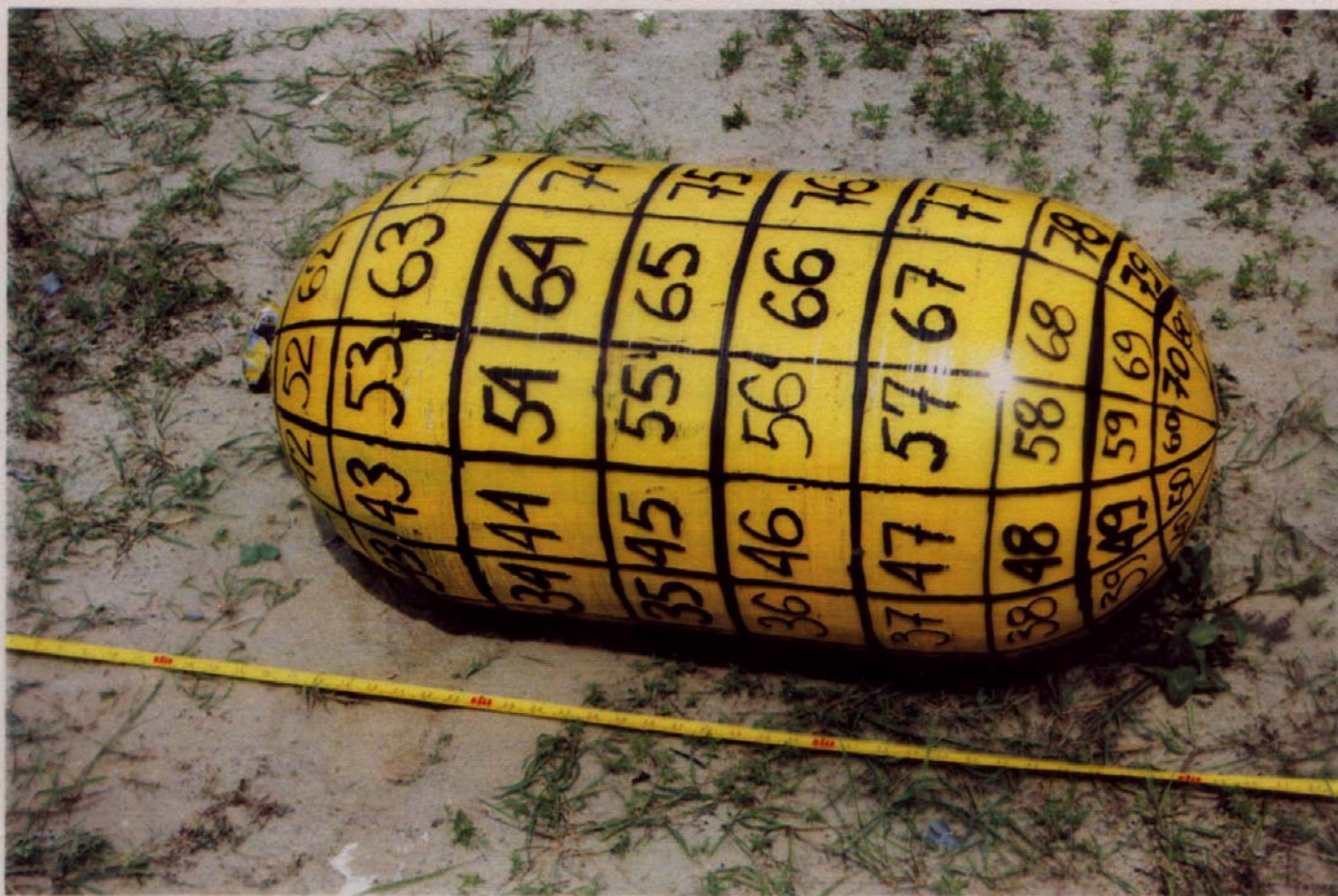


Fig. 4 The painted tank with a numbered grid for recovery of fragments

c)



d)



Conclusion from Accidents

- Rapid release in open atmosphere (Jackass Flat)
 - minor blast damages
- Rapid release in a congested area with equipment, structure etc. (Polysar)
 - severe blast damages, DDT
- Contamination of high pressure storage facility by air (China Light)
 - severe blast damages

Accident scenarios to avoid

- Rapid release in congested area (high density of equipment)
- Air contamination of high pressure hydrogen storage facilities
- Leakage of hydrogen into poorly vented enclosures

Explosion properties of hydrogen

- Equilibrium thermodynamics properties for hydrogen explosion well established
- Chemical kinetics of hydrogen oxidation sufficiently understood quantitatively (explosion limits, laminar flame propagation)
- Explosion parameters are also well established (flammability limit, ignition energy, quenching distance, etc.)

Explosion properties of hydrogen

- Detonation states are well known (Chapman-Jouguet detonation velocity, overpressure, etc.)
- Dynamic detonation parameters adequately known (initiation energy, detonability limit, critical diameter)
- Detonation sensitivity of high pressure H₂-air mixtures does not increase as other hydrocarbon fuels do
- Transition and onset of detonation (i.e. quantitative description of turbulent flame acceleration, condition for the onset of detonation) still not understood

Major unresolved problem

- Development of turbulent combustion models to describe high speed deflagrations with consideration of compressibility effects
- Quantitative theory for the onset of detonation

The Problem of the Transition from Deflagration to Detonation

Current Understanding
and Outstanding Problems

Two Modes of Combustion

Deflagration

- propagation via diffusion mechanism

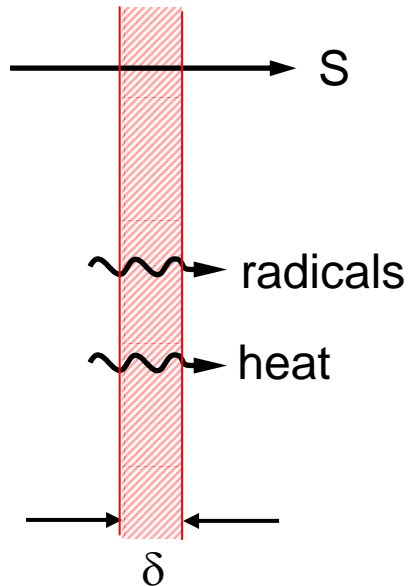
Detonation

- Propagation via shock ignition

Slowest Burning Rate

Laminar Flame

- molecular diffusion of heat and species



$$S \sim \sqrt{\frac{\alpha}{t_c}} \sim \sqrt{\frac{10^{-5}}{10^{-3}}} \approx 10^{-1} \text{ m/s}$$

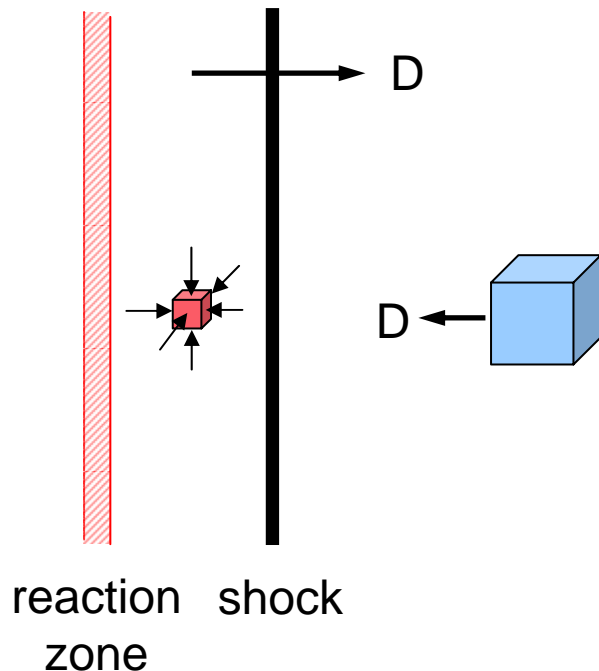
Flame Thickness:

$$\delta \sim \sqrt{\alpha t_c} \sim \sqrt{10^{-5} \cdot 10^{-3}} \approx 10^{-1} \text{ mm}$$

Fastest Burning Rate

CJ Detonation

- Ignition by adiabatic shock compression



$$D \sim c$$

$$c \sim \sqrt{e} \approx \sqrt{Q}$$

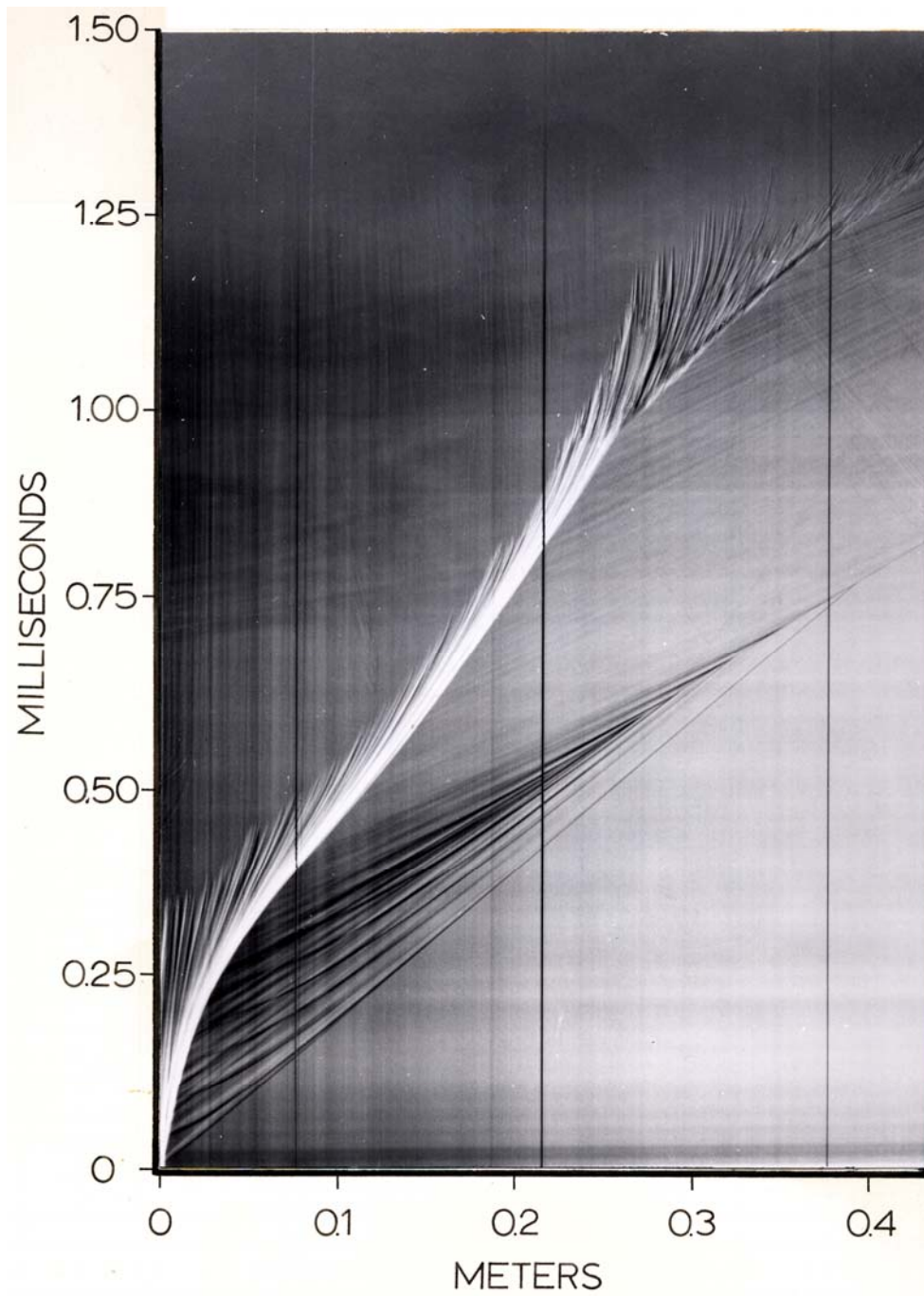
$$Q \approx 30e_o \sim 30c_o^2$$

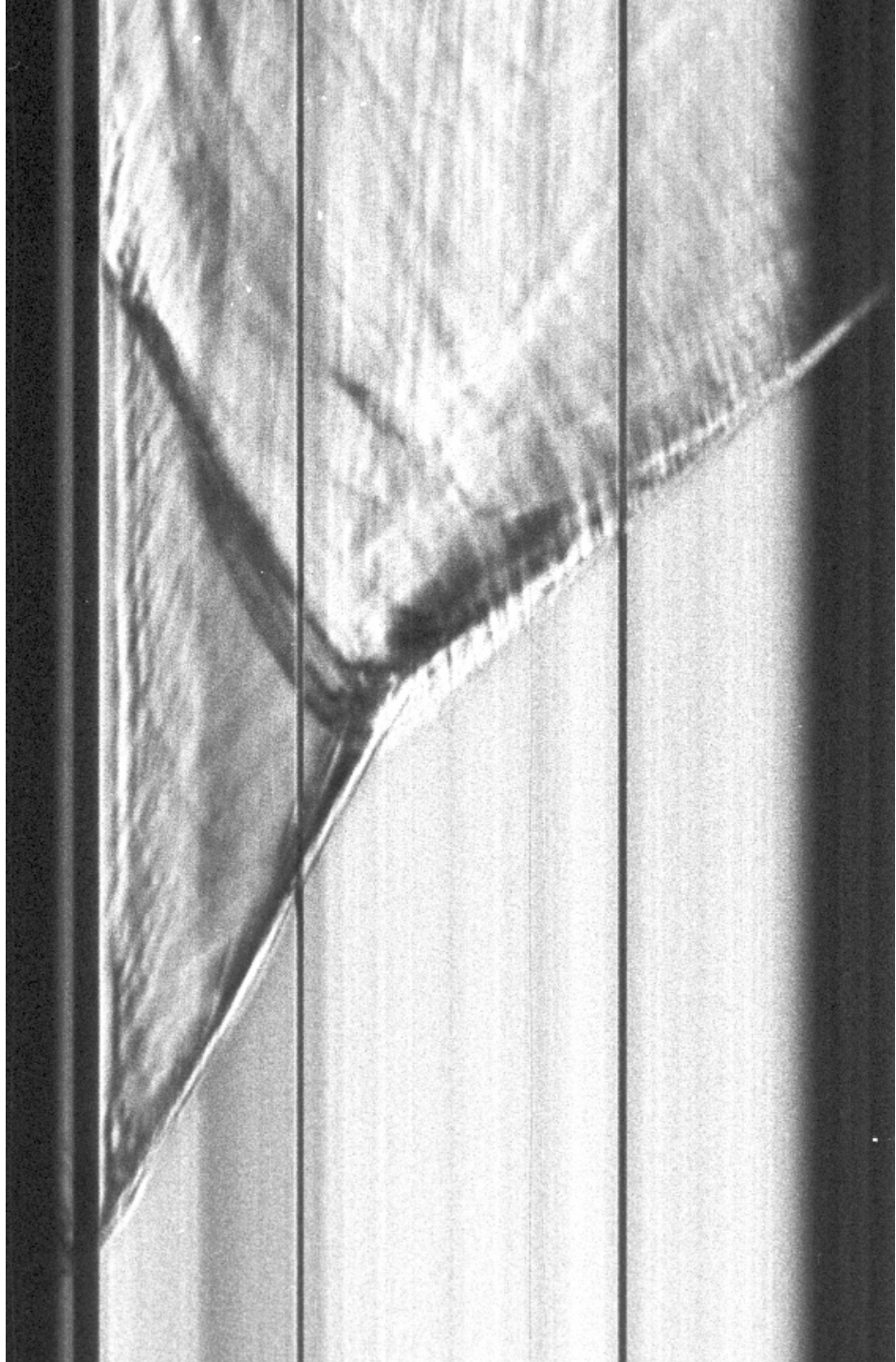
$$D \approx 5.5c_o \approx 1800\text{m/s}$$

Self-Propagating Deflagration Waves

- are unstable
- accelerate to some critical state and undergo transition to detonation waves



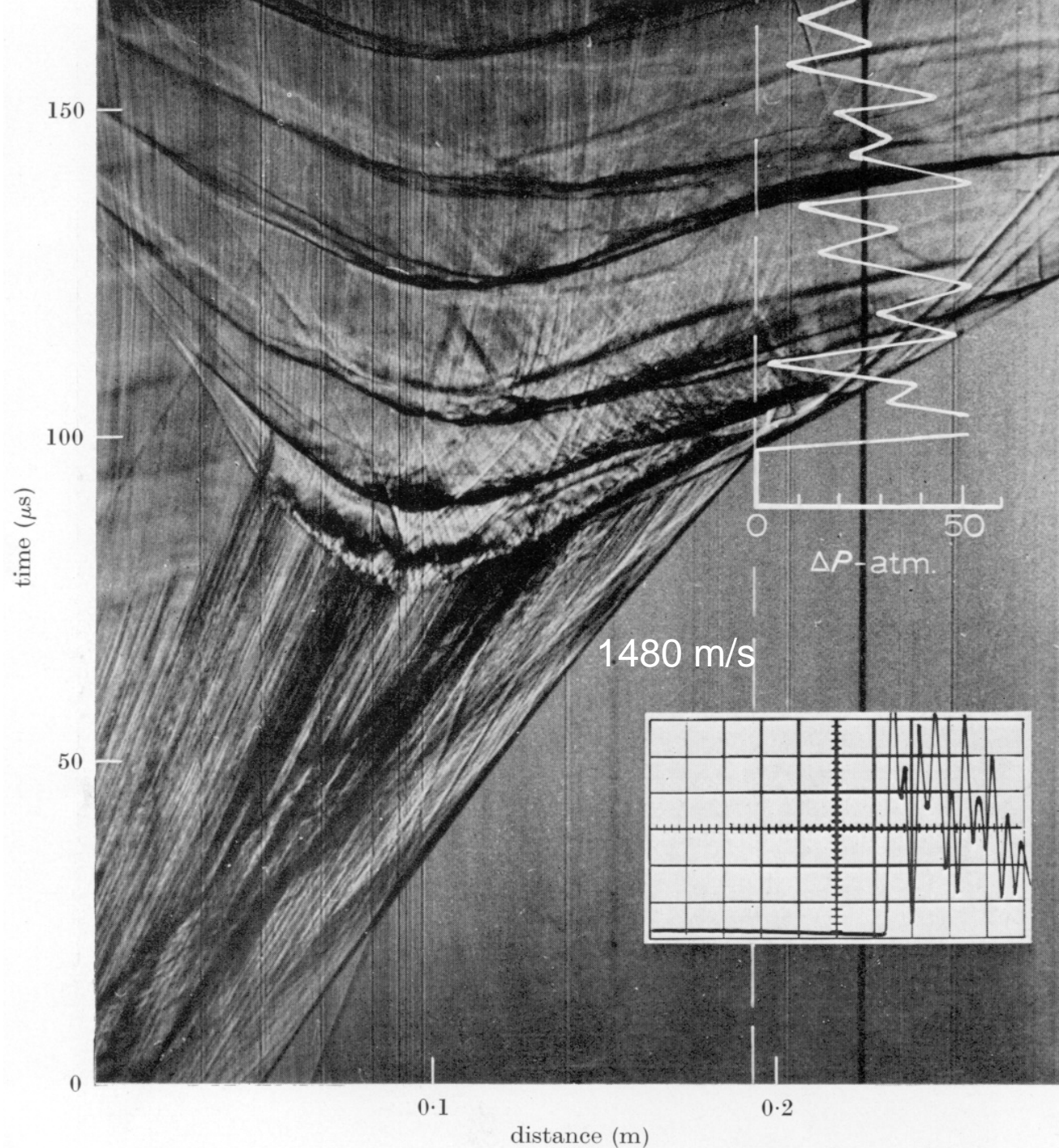


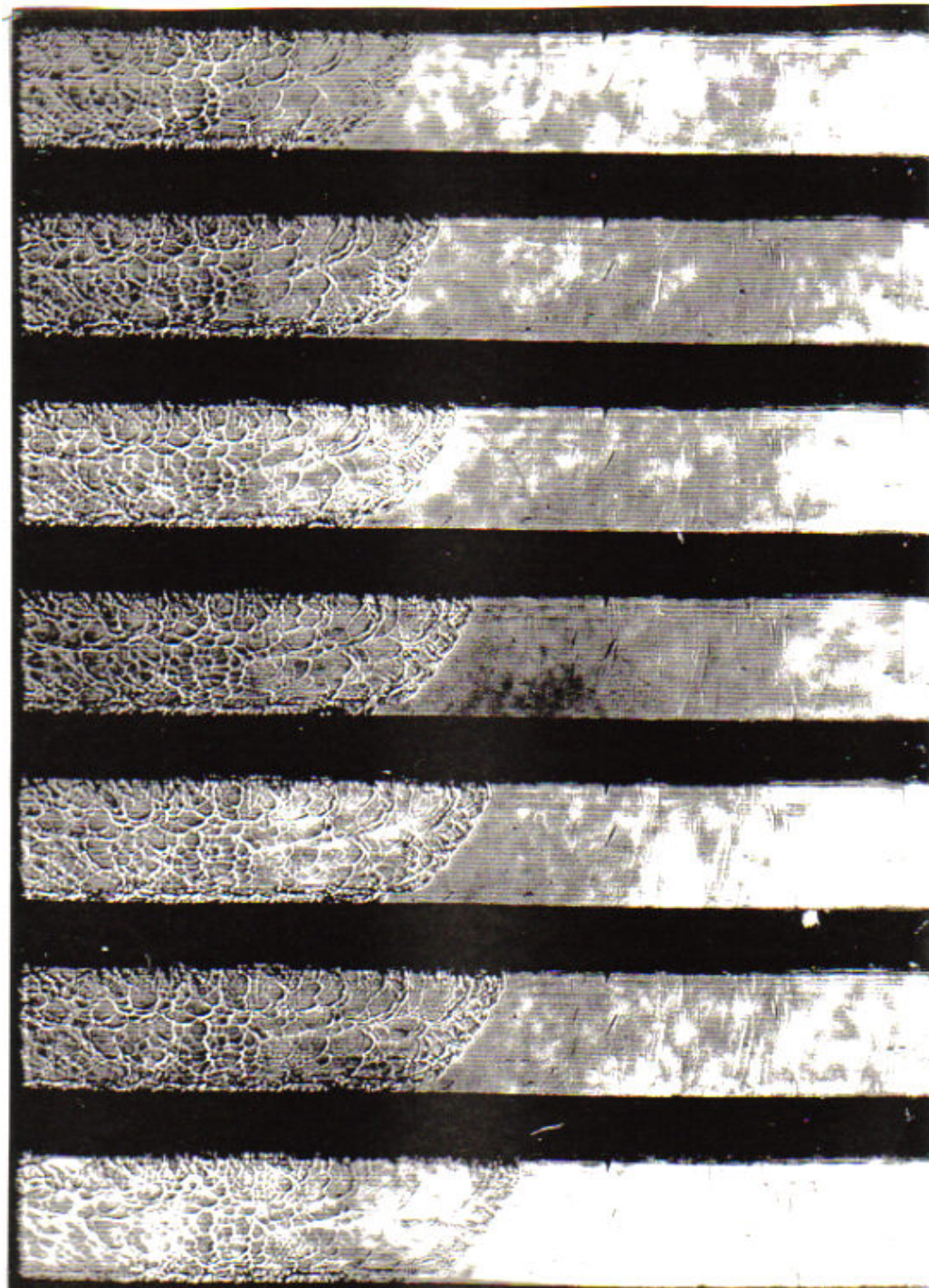


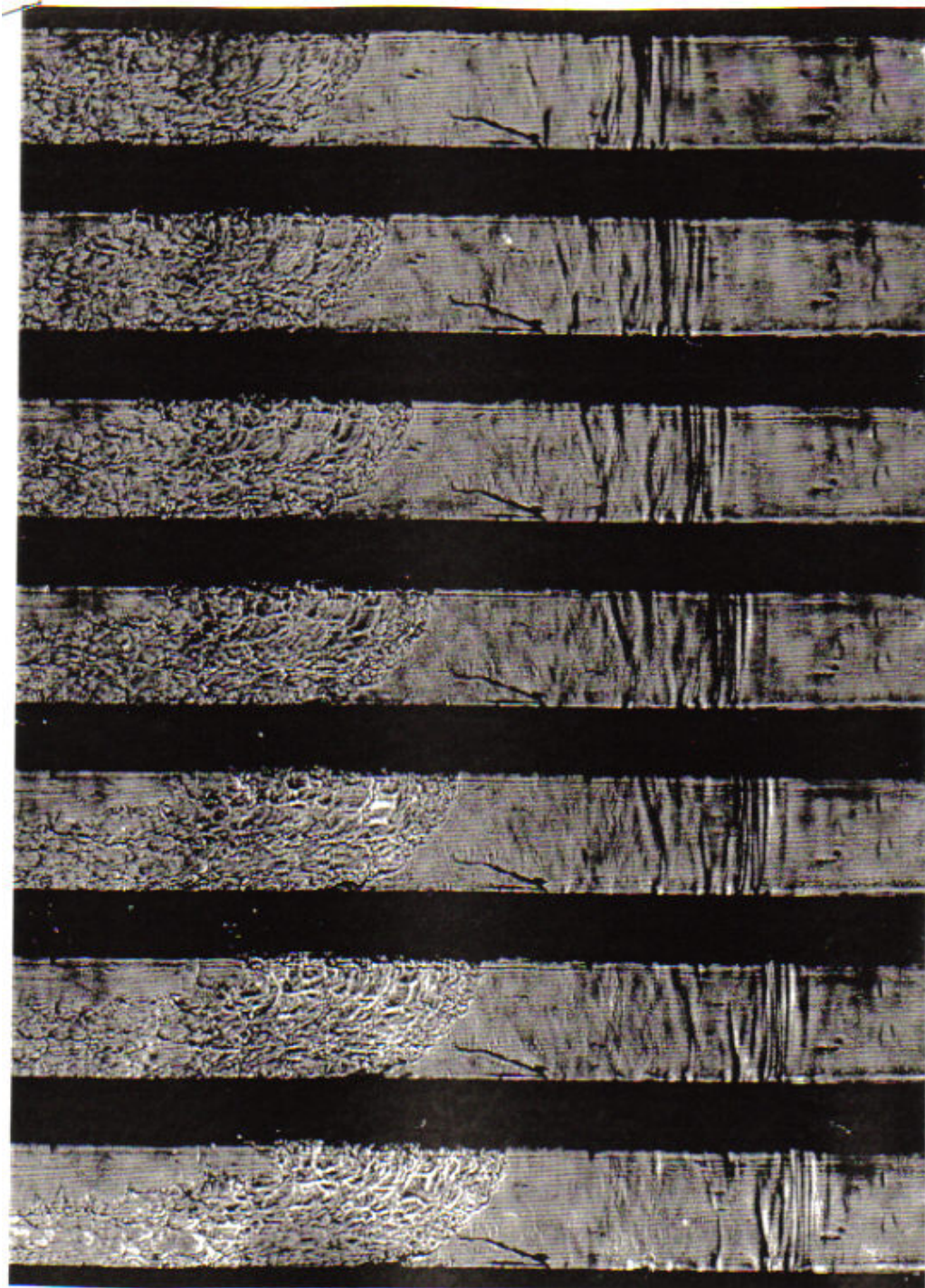
Urtiew & Oppenheim
(1966)

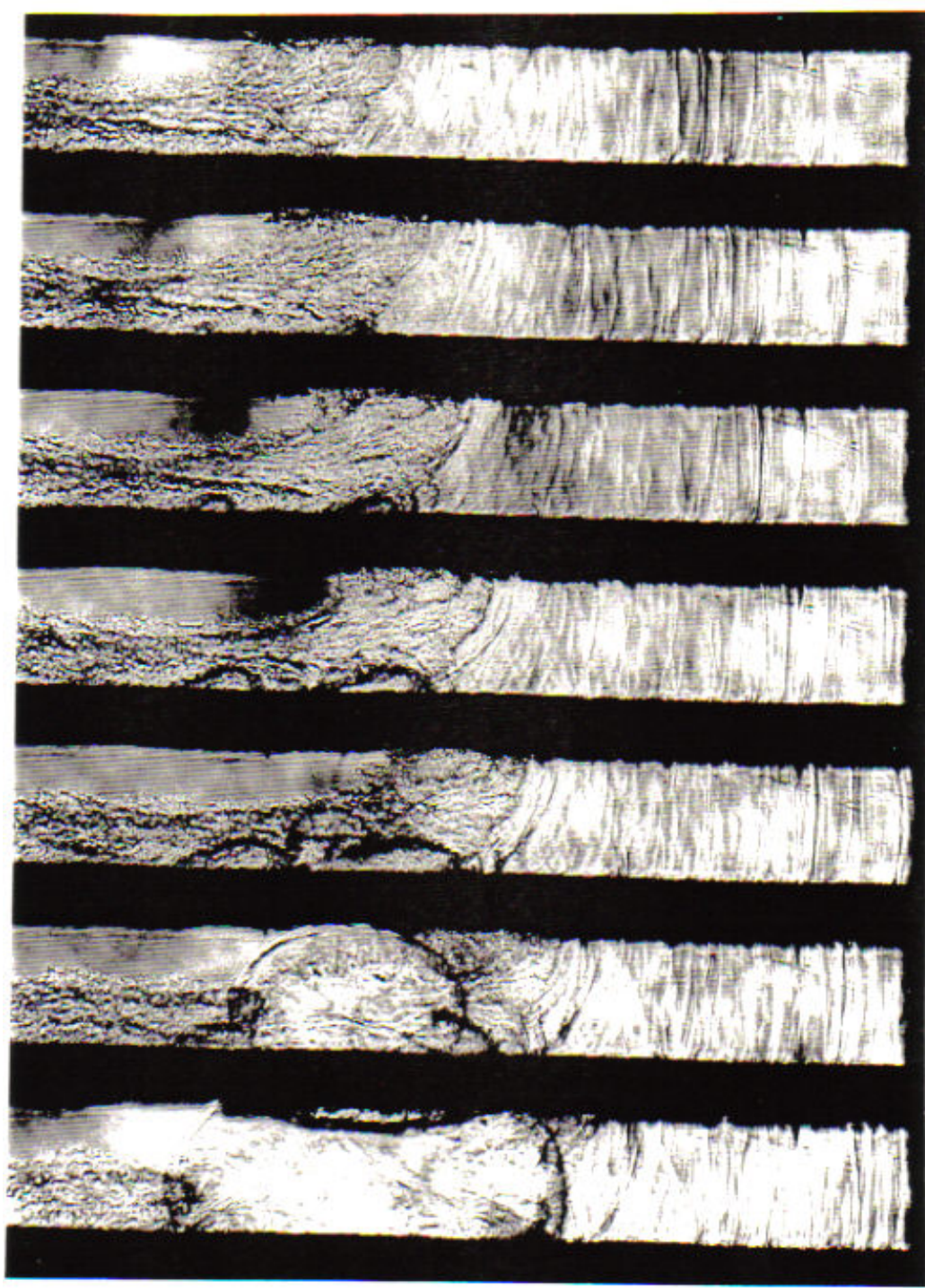
$\text{H}_2 + 0.5 \text{O}_2$
@ $P_o = 1 \text{ atm}$

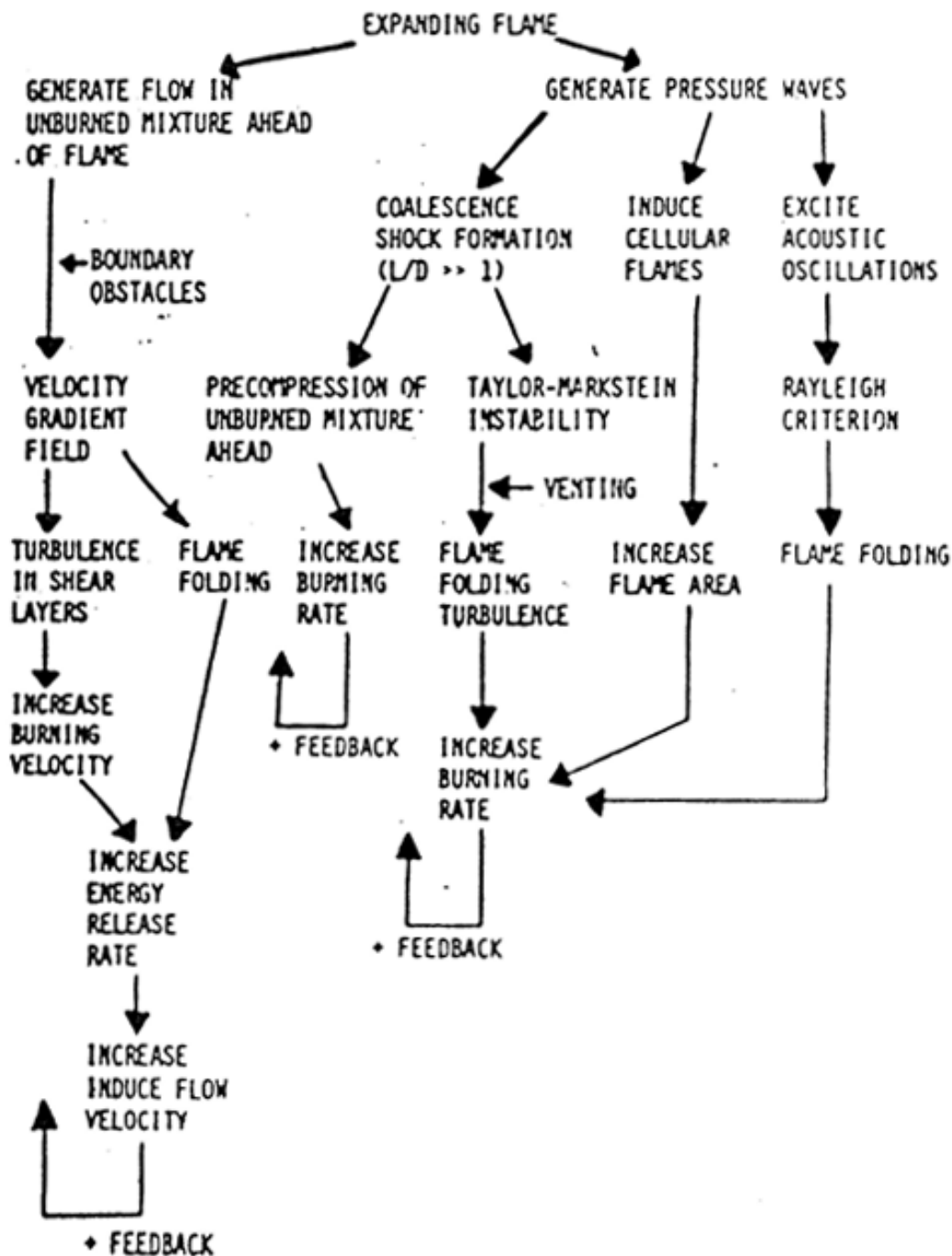
$V_{\text{CJ}} = 2837 \text{ m/s}$







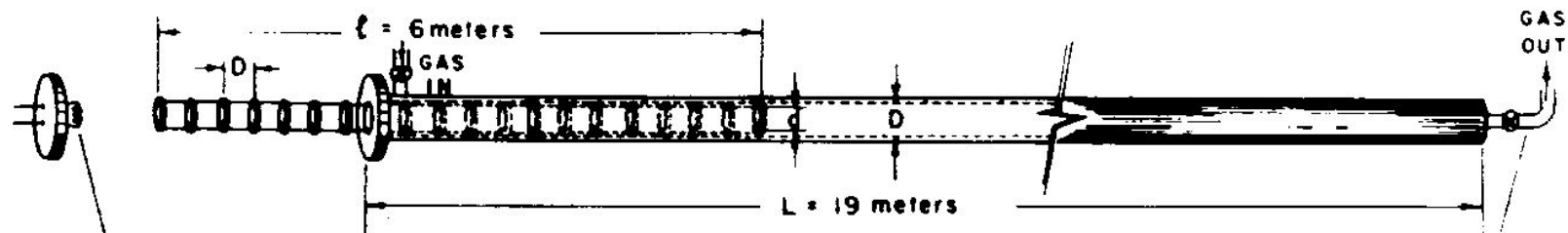




- initial phase of flame acceleration involves numerous instability mechanisms
- not possible to characterize the flame acceleration phase by a single reproducible parameter like the run-up distance

- bypass the initial phase and look at the final phase of the onset of detonation
- determine the critical deflagration speed prior to onset of detonation
- use obstacles to get to critical speed rapidly

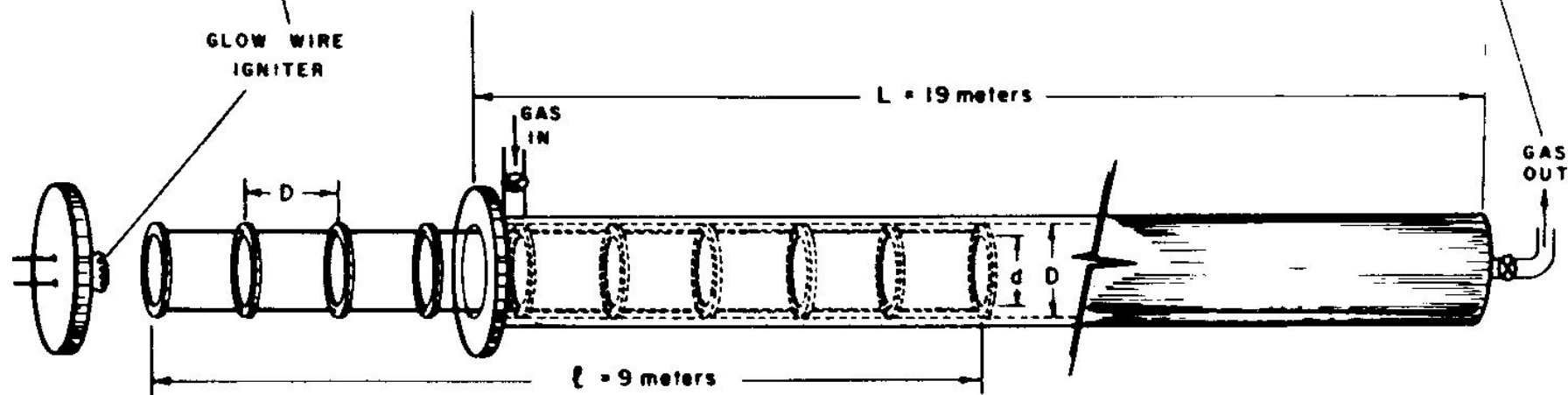
- systematic studies of DDT in rough tubes began at McGill in the late 1970's
- tubes from 5 cm to 2.5 m were used
- obstacles were in the form of orifice plates, cylindrical rods, Shchelkin spirals, etc.



TUBE DIAMETER $D = 5$ cm, OBSTACLE PITCH $P = 5$ cm

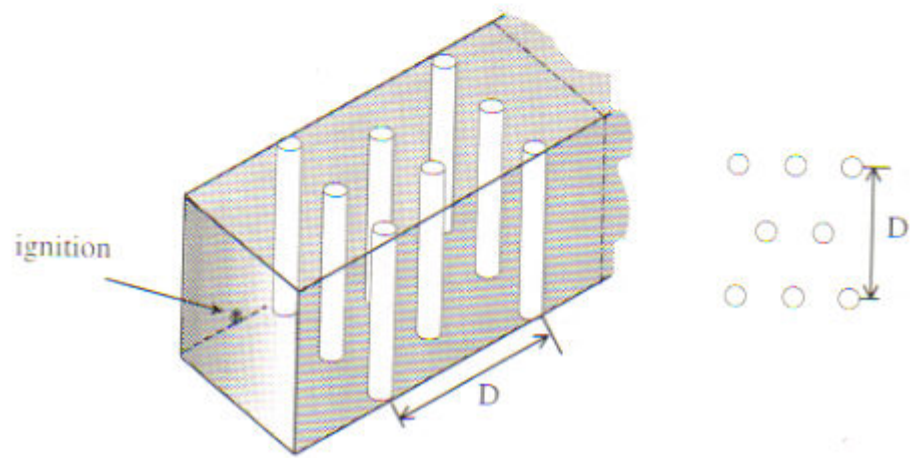
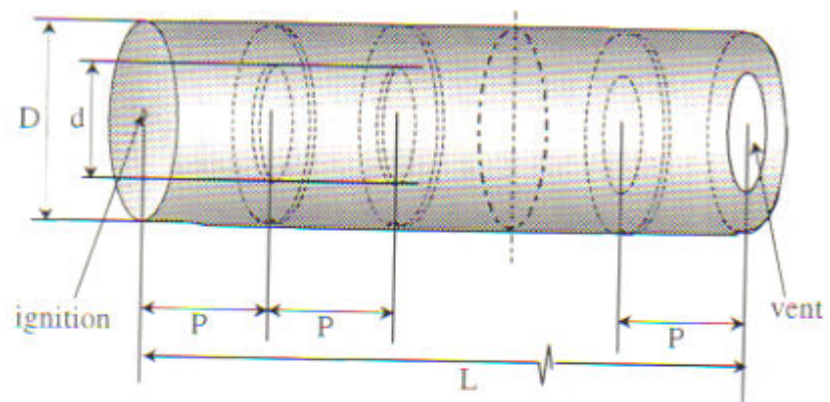
$$\text{BLOCKAGE RATIO } BR = 1 - \left(\frac{d}{D}\right)^2 = 0.43$$

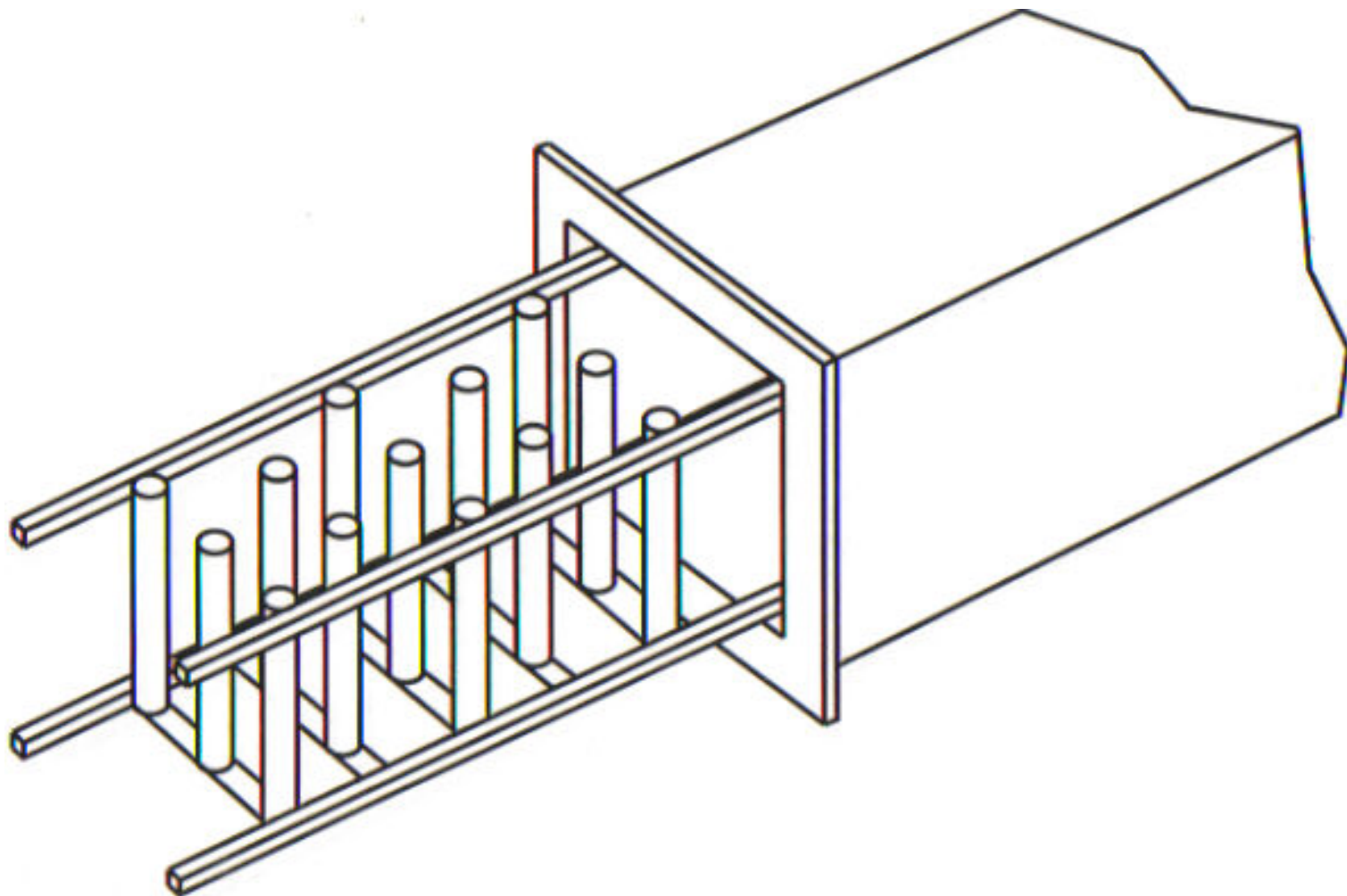
GAS RECIRCULATION SYSTEM



TUBE DIAMETER $D = 15$ cm, OBSTACLE PITCH $P = 15$ cm

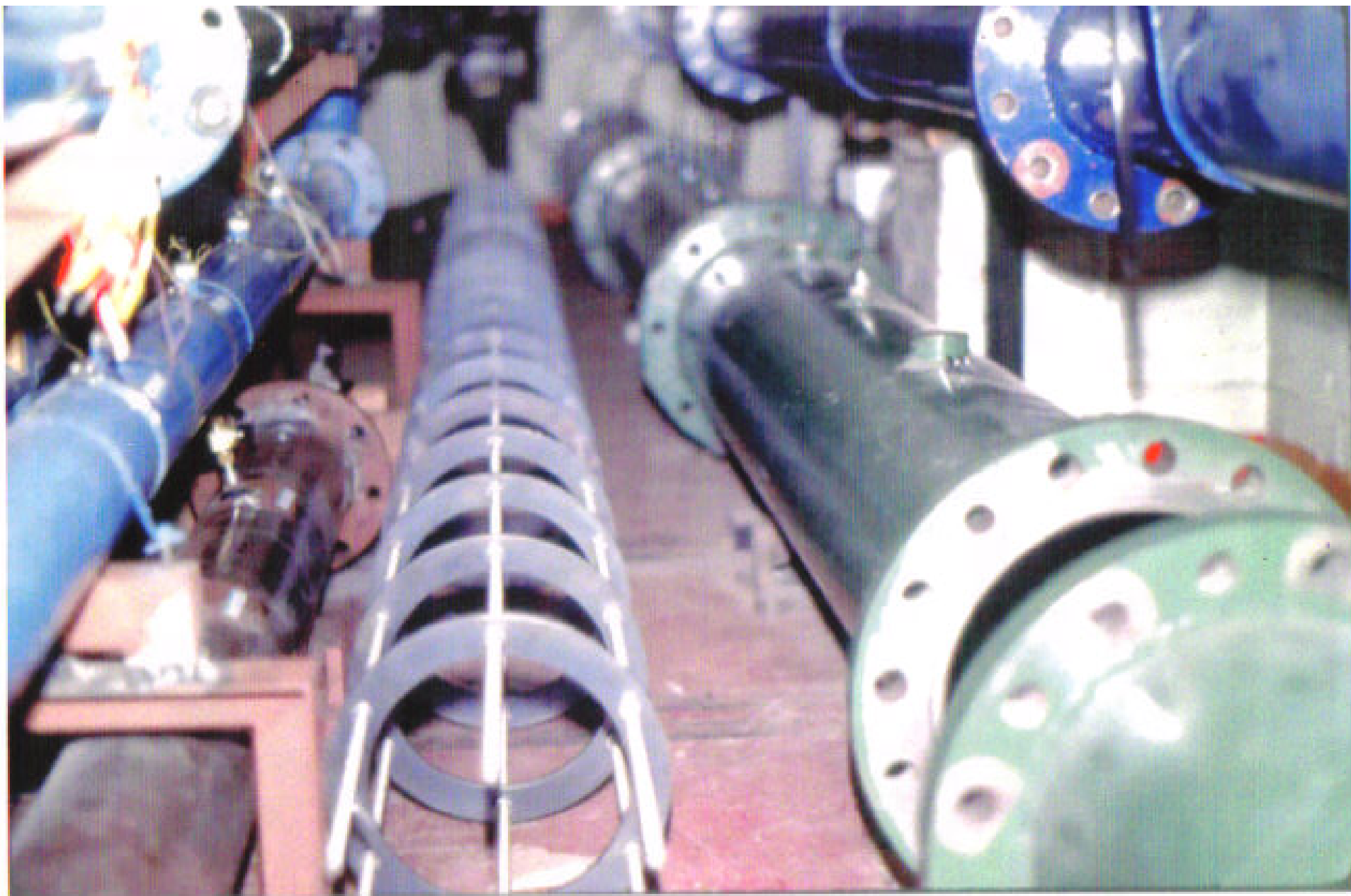
$$\text{BLOCKAGE RATIO } BR = 1 - \left(\frac{d}{D}\right)^2 = 0.39$$

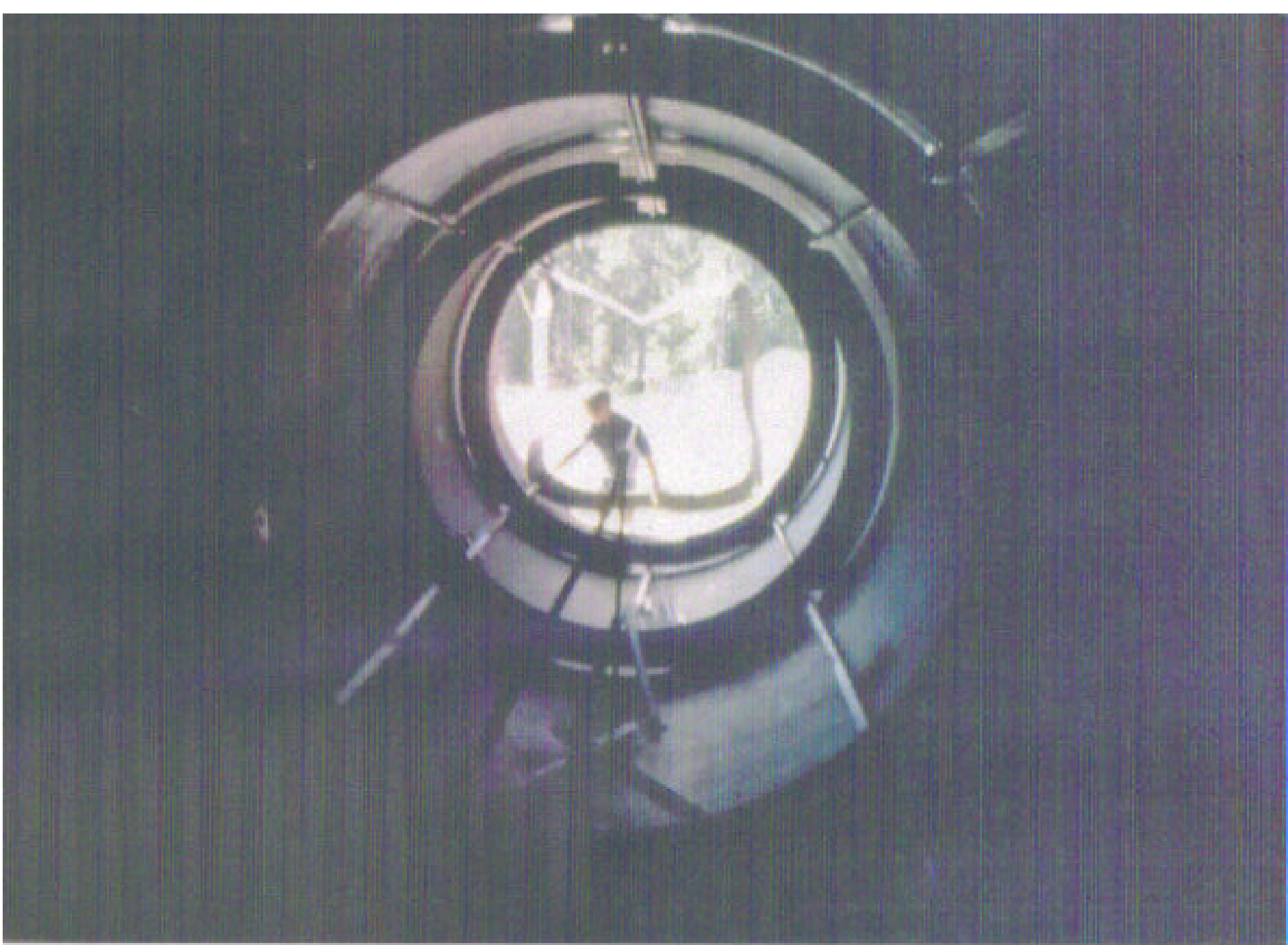


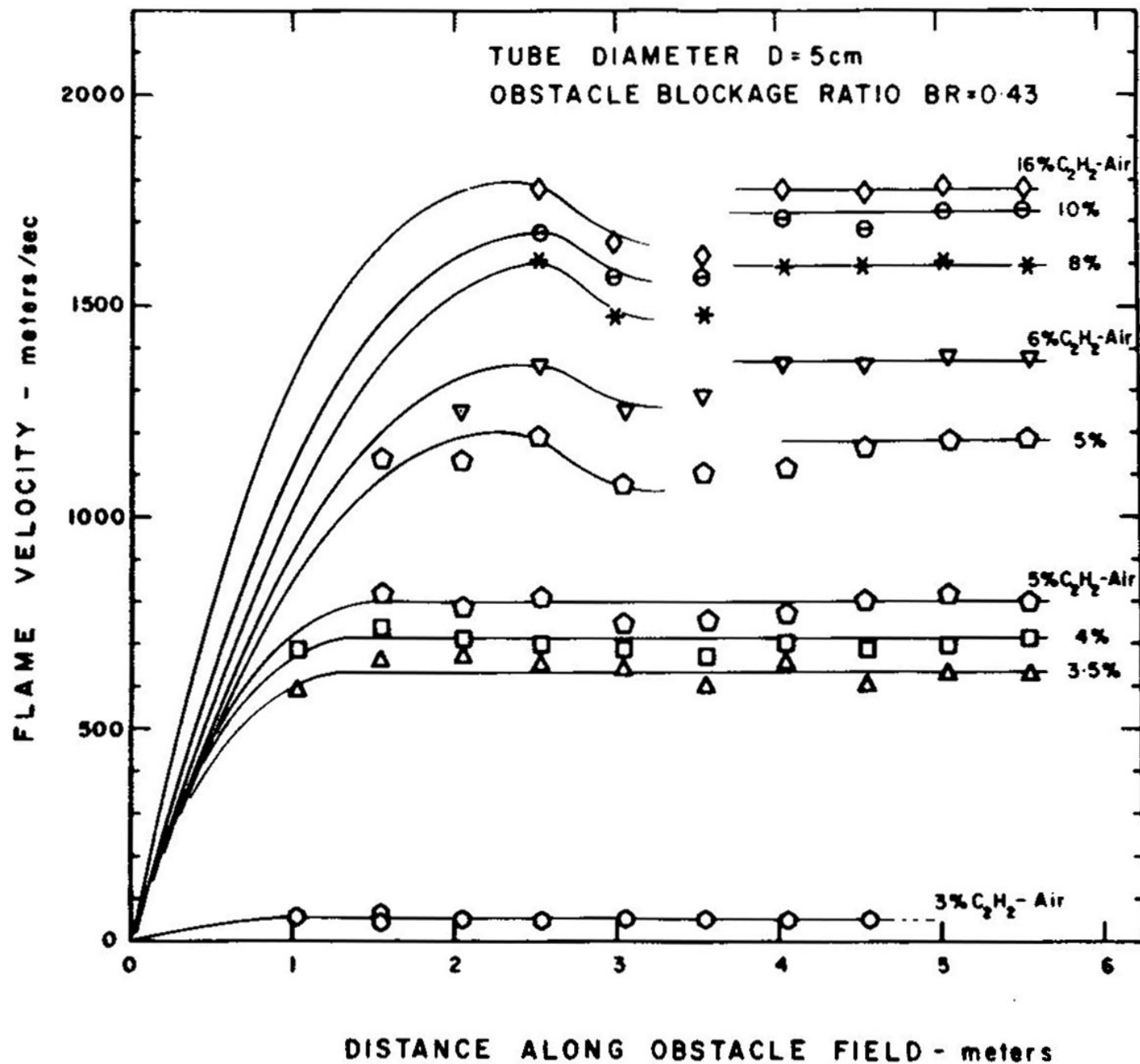


Three-dimensional view of the Tube-Obstacle Assembly



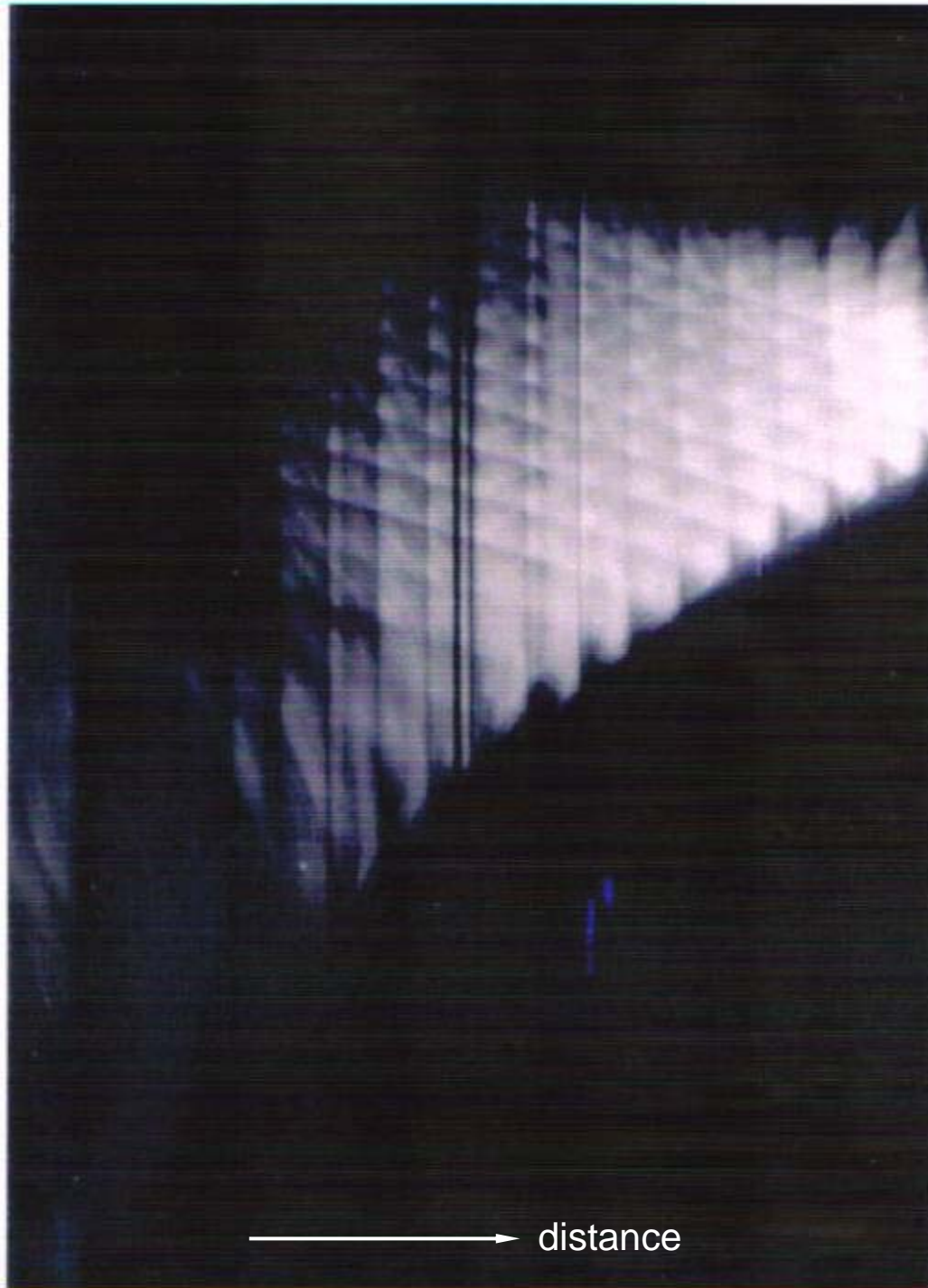






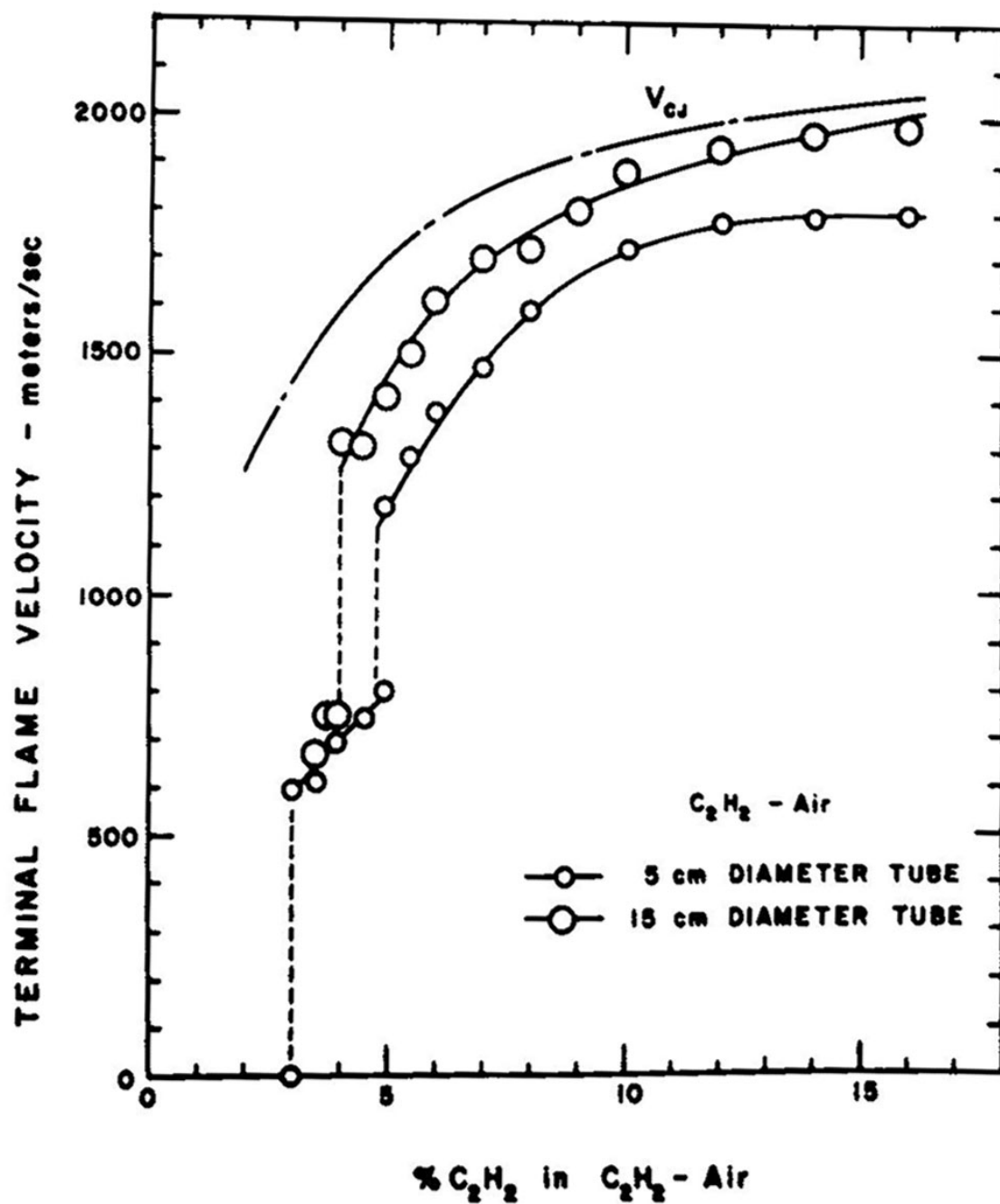
time ↑

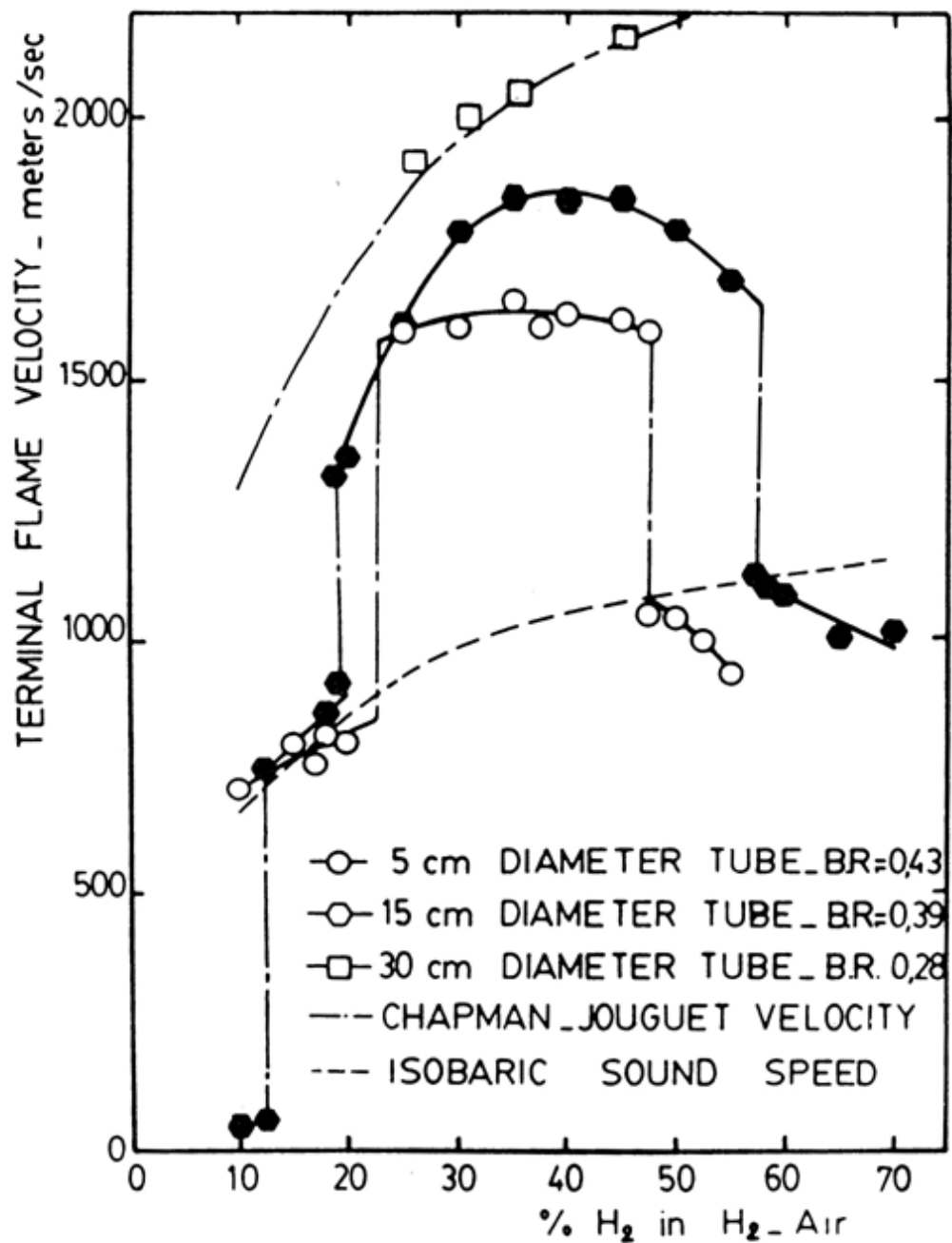
→ distance

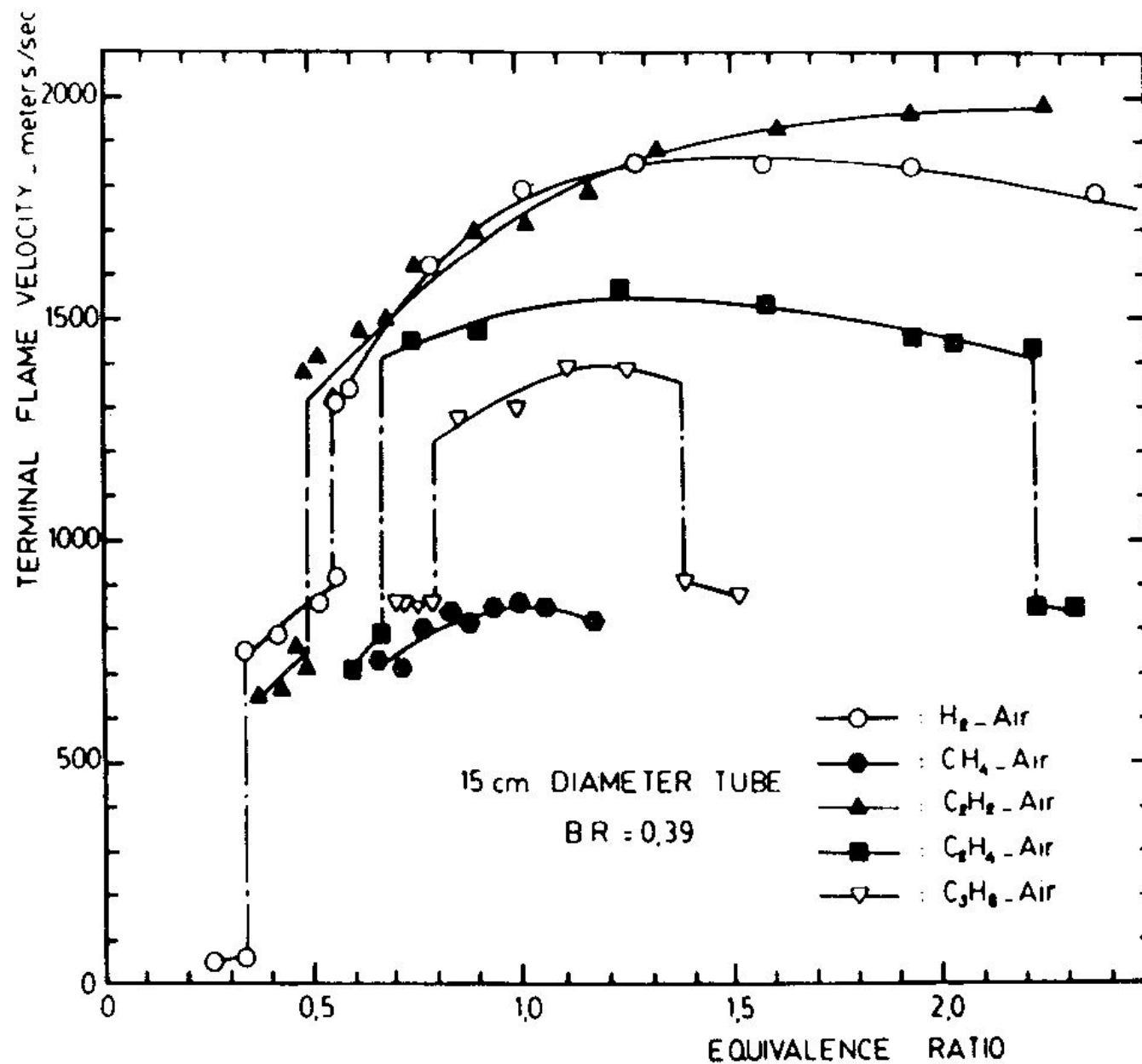


Findings from Rough Tube Experiments

- rapid acceleration to a quasi-steady velocity
- steady velocity is not too sensitive to tube diameter or obstacle configuration
- distinct transition from steady velocity to a higher value when mixture sensitivity varies







Three Distinct Regimes

- turbulent deflagration < 100 m/s
- sonic regime
deflagration speed \sim sound speed of products
 ~ 1000 m/s ($\sim \frac{1}{2} V_{CJ}$)
- quasi-detonation or detonation
 $\sim V_{CJ}$ with large velocity deficit

Three parameters that can characterize the condition for onset of detonation:

1. critical deflagration speed
2. tube diameter
3. sensitivity of mixture

Table 1 Transition within obstacle field

Mixture	D, cm	d, mm	λ , mm	λ / d
4.75% C ₂ H ₂ -air	5	37.4	19.8	0.51
22% H ₂ -air	5	37.4	30.7	0.82
47.5% H ₂ -air	5	37.4	41.2	1.10
6% C ₂ H ₄ -air	5	37.4	37.8	1.01
9% C ₂ H ₄ -air	5	37.4	30.1	0.81
4% C ₂ H ₂ -air	15	114.0	58.3	0.51
3.25% C ₃ H ₈ -air	15	114.0	112.0	0.98
5.5% C ₃ H ₈ -air	15	114.0	116.0	1.02

No Transition

Mixture	D, cm	d, mm	λ min, mm	λ / d
C ₃ H ₈ -air	5	37.4	52.5	1.40
CH ₄ -air	5	37.4	300.0	8.02
CH ₄ -air	15	114.0	300.0	2.63

Table 2 Transition in smooth-walled tube

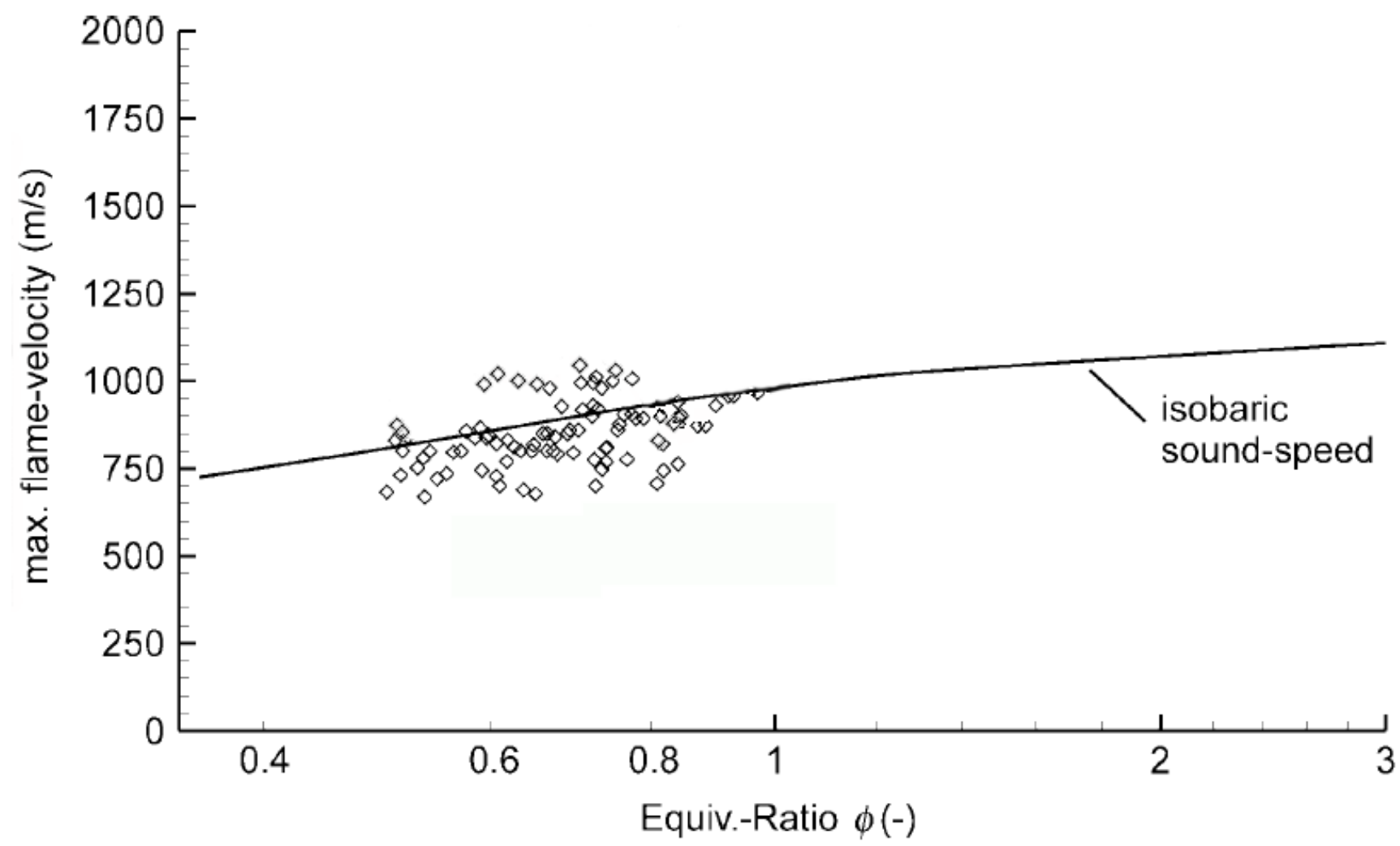
Mixture	D, cm	λ , mm	λ / D
4% C ₂ H ₂ -air	5	58.3	1.18
5% C ₂ H ₄ -air	5	65.1	1.32
10% C ₂ H ₄ -air	5	39.7	0.80
4% C ₃ H ₈ -air	5	52.2	1.06
5% C ₃ H ₈ -air	5	59.0	1.19
20% H ₂ -air	5	55.4	1.12
51% H ₂ -air	5	52.5	1.06

Critical Deflagration Speed for Onset of Detonation

$$\sim \frac{1}{2} V_{CJ}$$

\sim sound speed of products

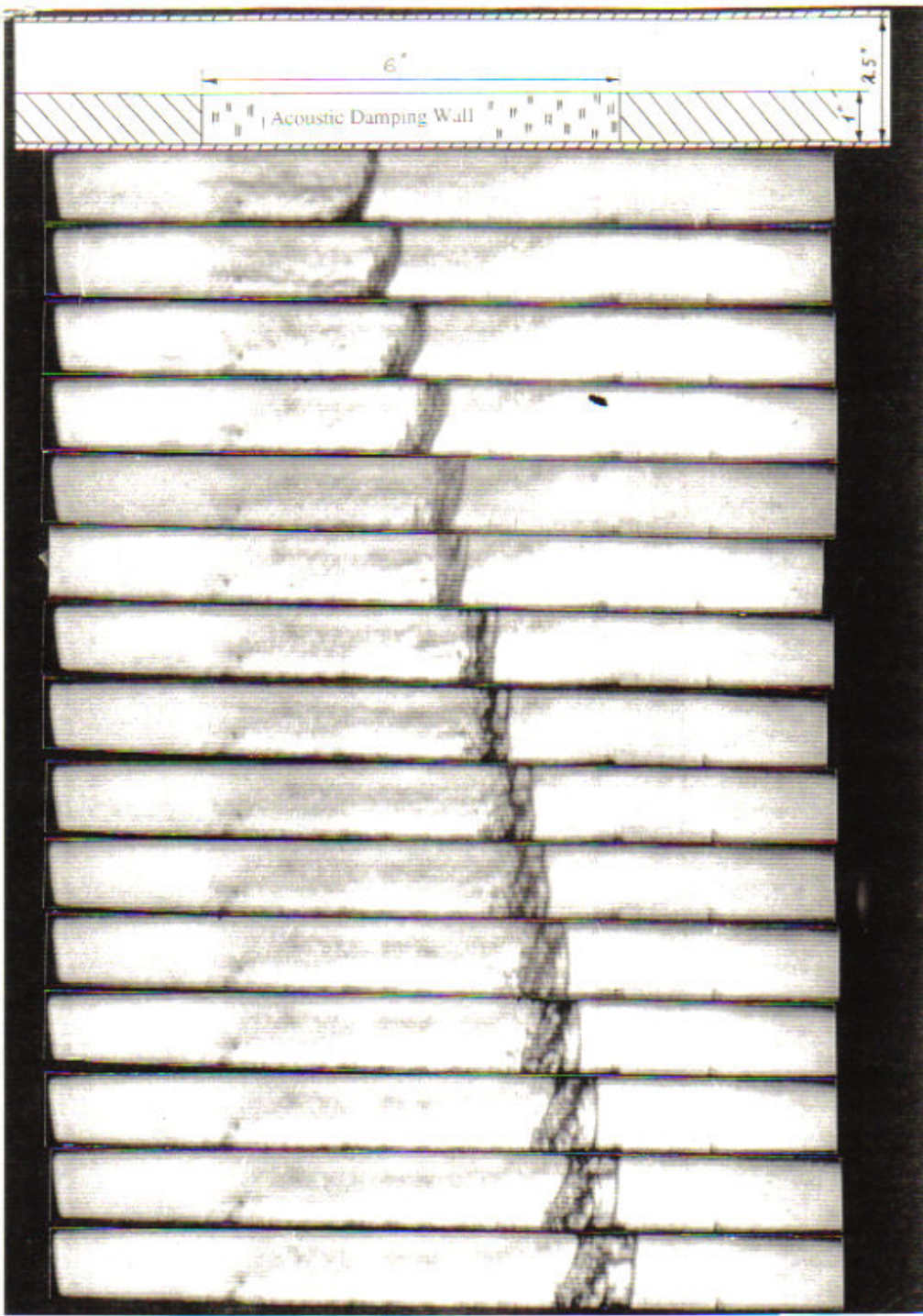
Eder & Brehm (2001)



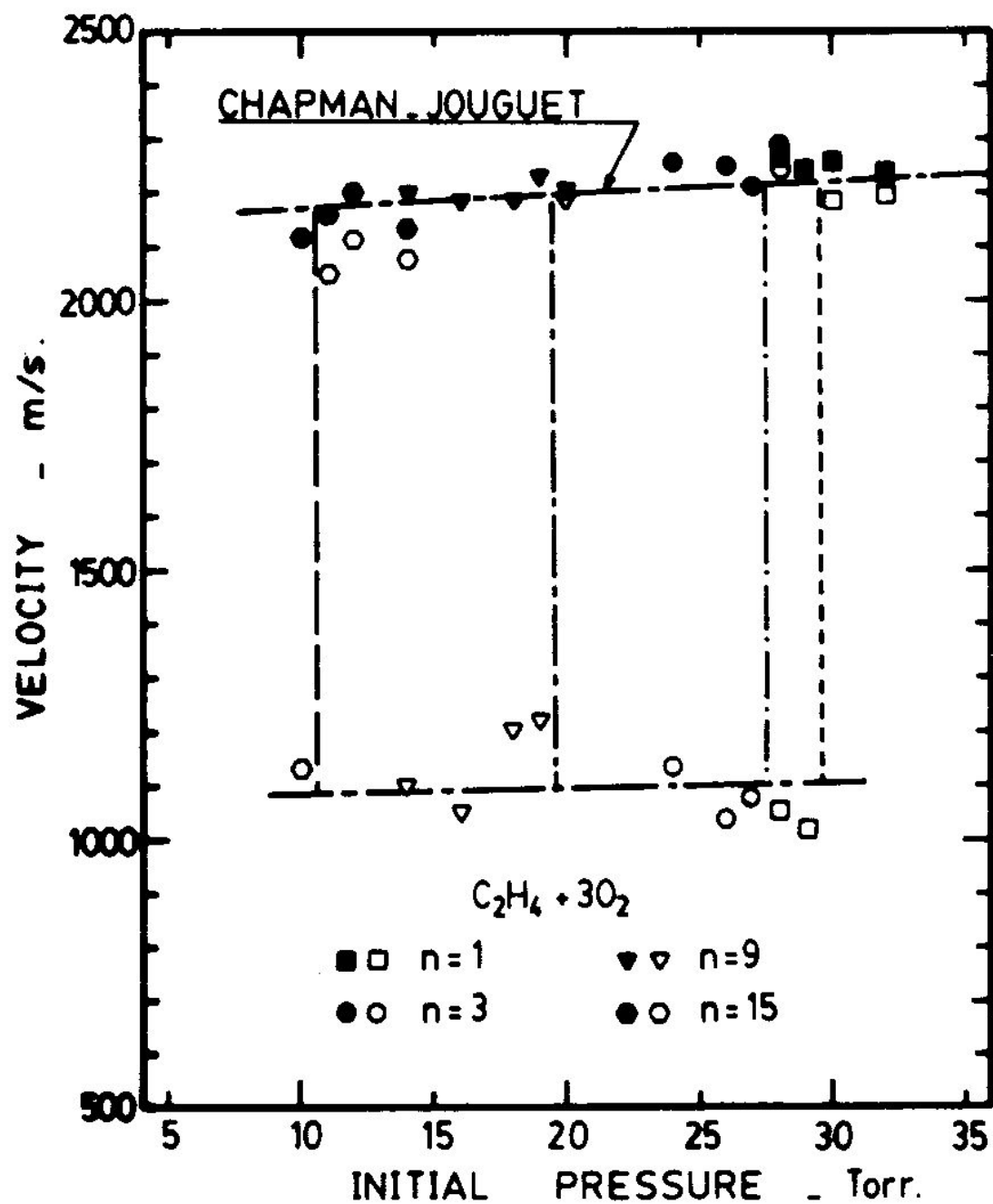
Vasil'ev (2006)

Mixture	c_0 , m/sec	P_{CJ}	σ_{CJ}	P_V	σ_P	P_{def}	σ_{def}	π^*	M_{inc}	M_{ref}	M_0
$C_2H_2 + 2.5O_2$	330	33.83	1.84	17.07	0.07	0.48	0.036	18.2	3.95	2.1	7.34
$C_2H_2 + \text{air (stoichiometric ratio)}$	347	19.11	1.82	9.77	0.12	0.48	0.062	10.6	3.05	1.8	5.38
$C_2H_4 + 3O_2$	328	33.43	1.85	16.87	0.07	0.48	0.036	17.8	—	—	7.24
$C_2H_4 + \text{air (stoichiometric ratio)}$	347	18.35	1.81	9.38	0.12	0.48	0.064	10.1	2.95	1.8	5.26
$2H_2 + O_2$	537	18.79	1.84	9.59	0.12	0.49	0.062	10.4	3	1.8	5.28
$H_2 + \text{air (stoichiometric ratio)}$	409	15.58	1.8	8	0.15	0.48	0.076	9	2.8	1.75	4.82
$CH_4 + 2O_2$	355	29.32	1.85	14.84	0.08	0.49	0.04	15.8	3.65	2.05	6.73
$CH_4 + \text{air (stoichiometric ratio)}$	354	17.17	1.81	8.79	0.13	0.47	0.069	9.6	2.9	1.75	5.09

(confined) $0.33 \leq M_{crit} \leq 0.56 M_{CJ}$ (unconfined)

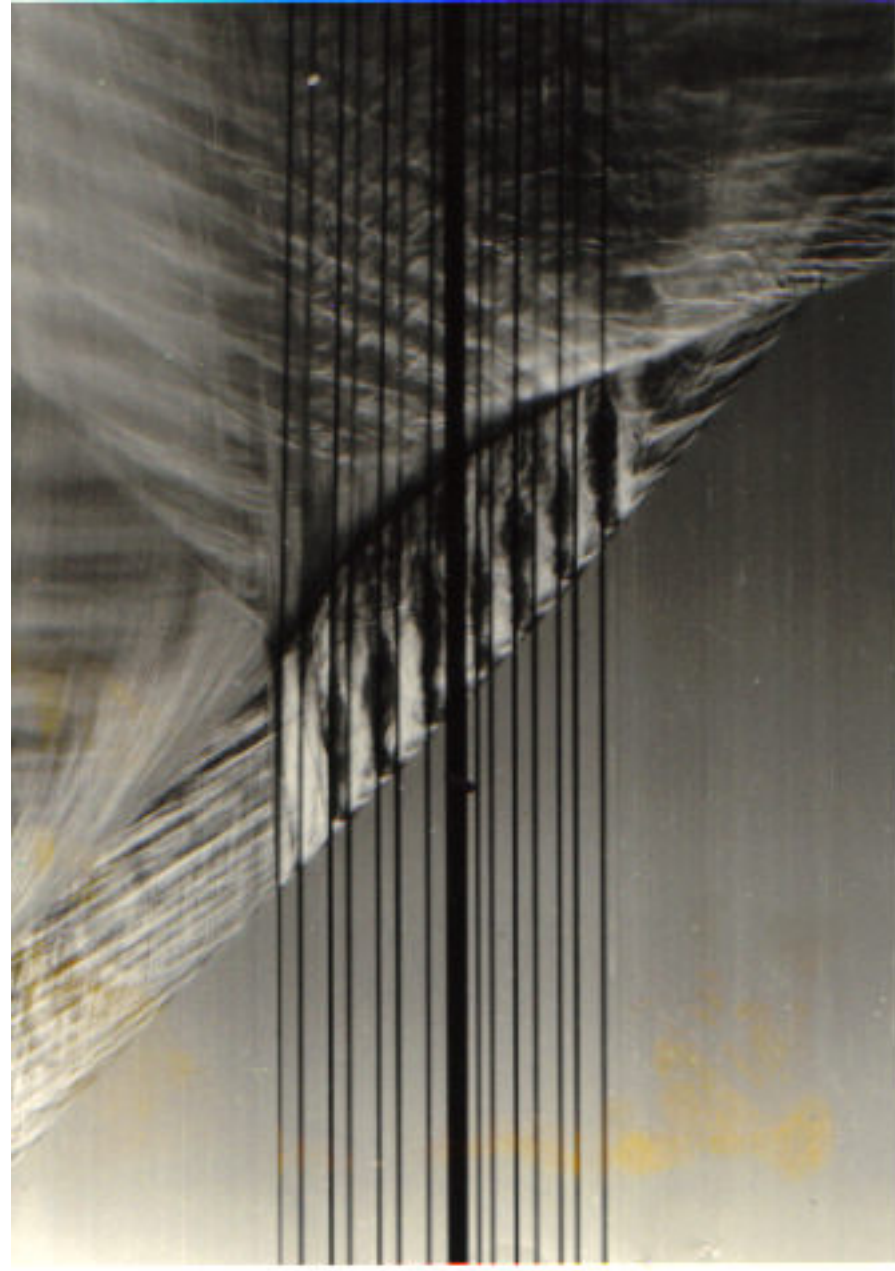






Mechanism of Onset of Detonation in Rough (Obstacle-Filled) Tubes

- turbulence from obstacles
- pressure waves



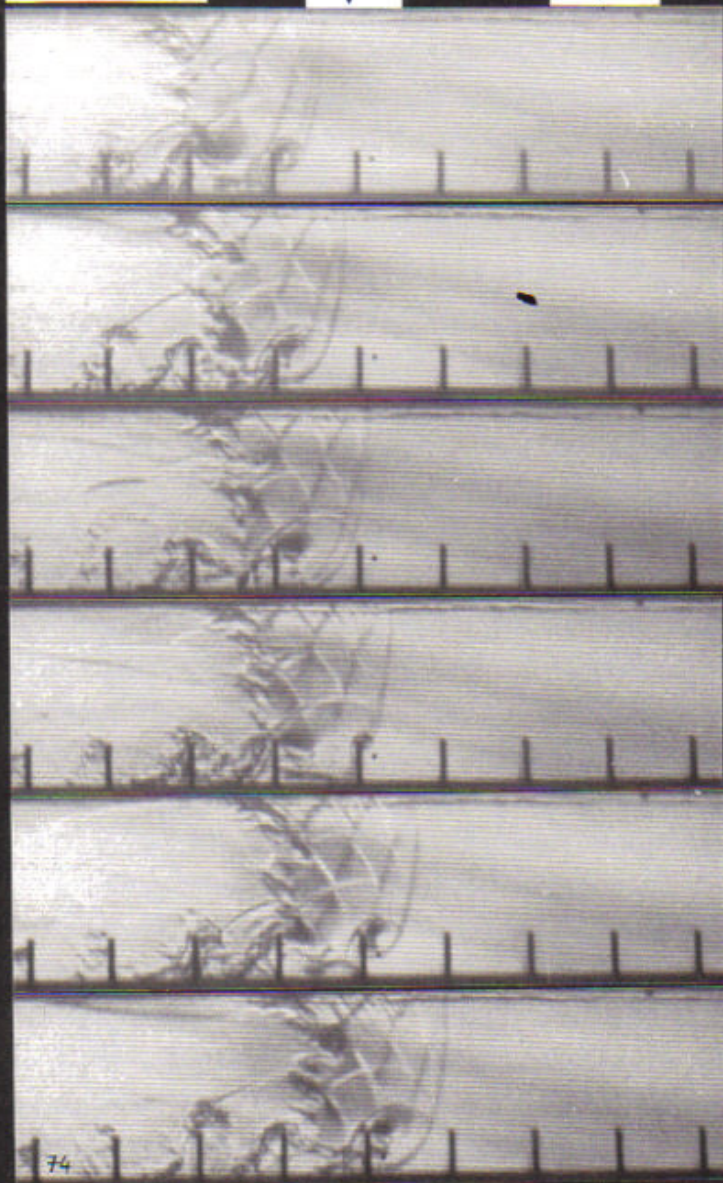
10cm

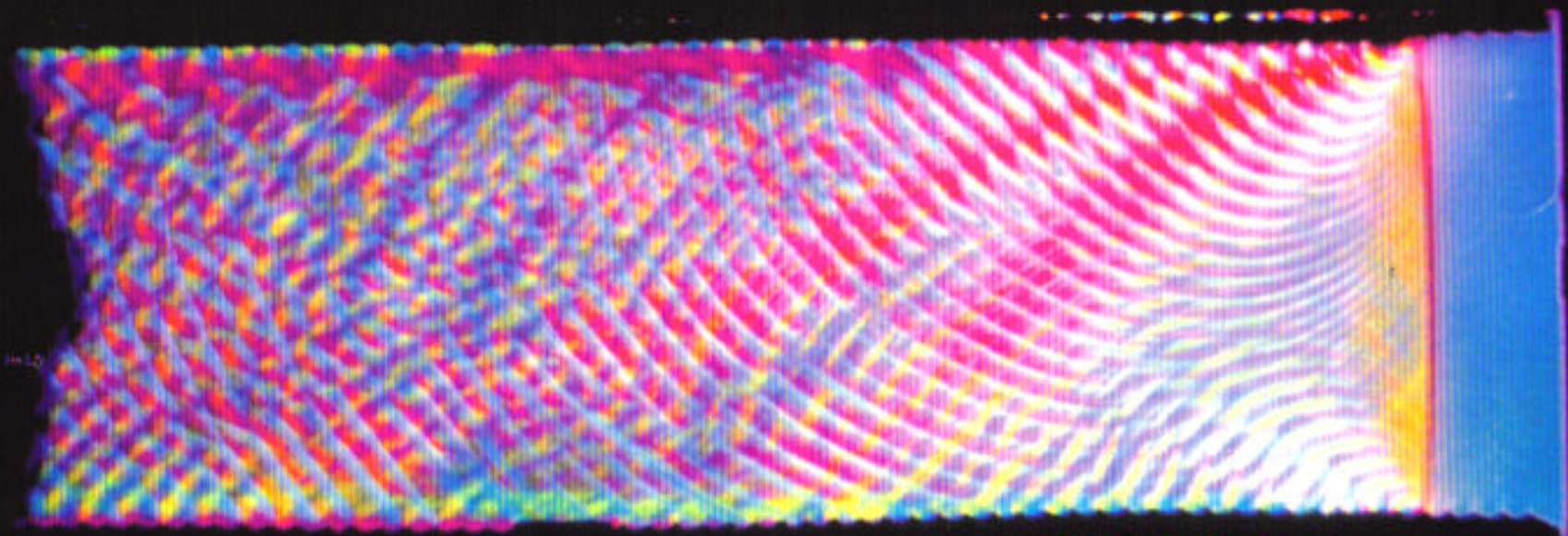
0.1ms

100 torr 5.93 μ s

PT 1
↓

PT 2
↓





0.25m

$2\text{H}_2 + \text{O}_2$ 150 Torr

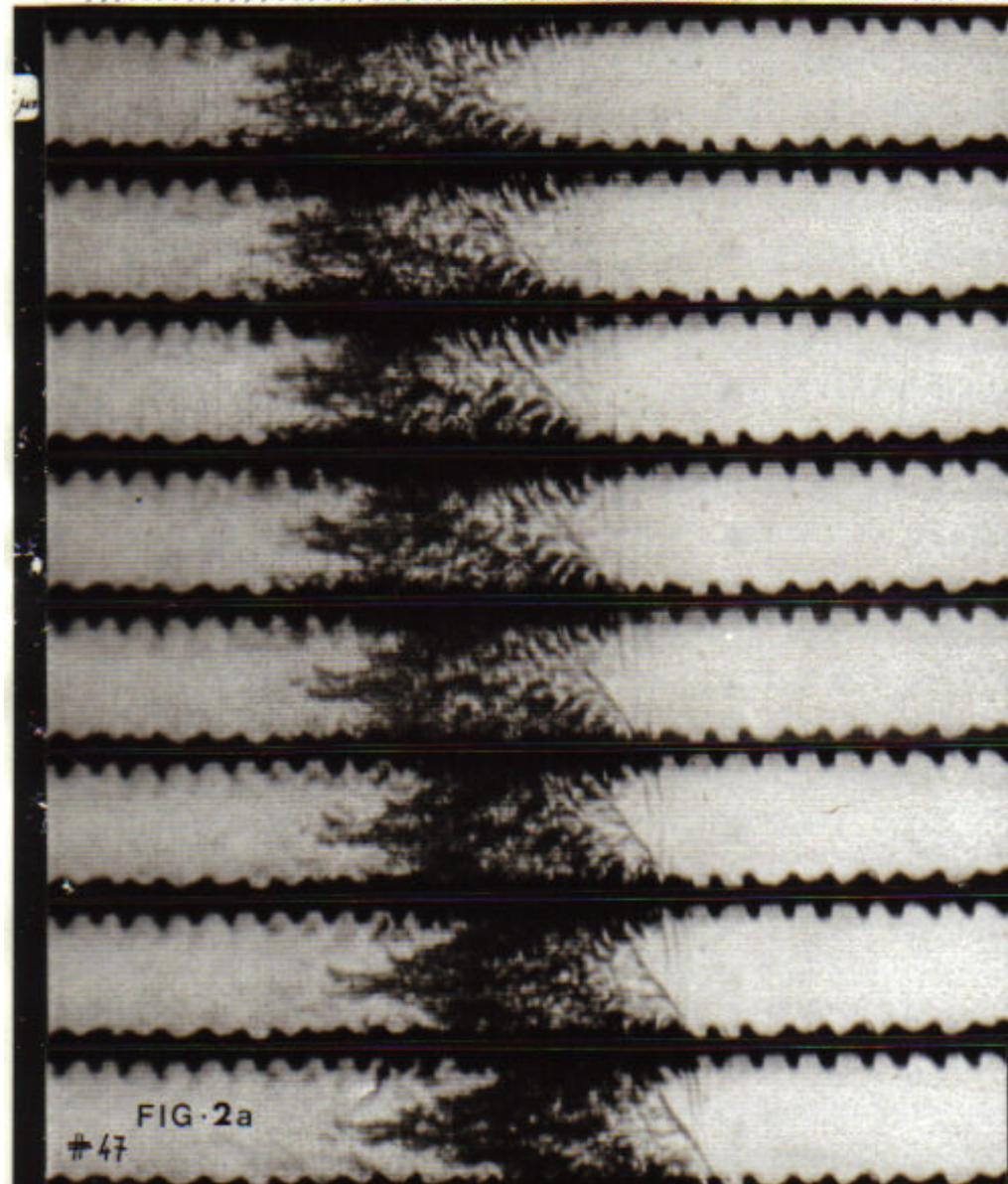
5mm

32mm

37mm

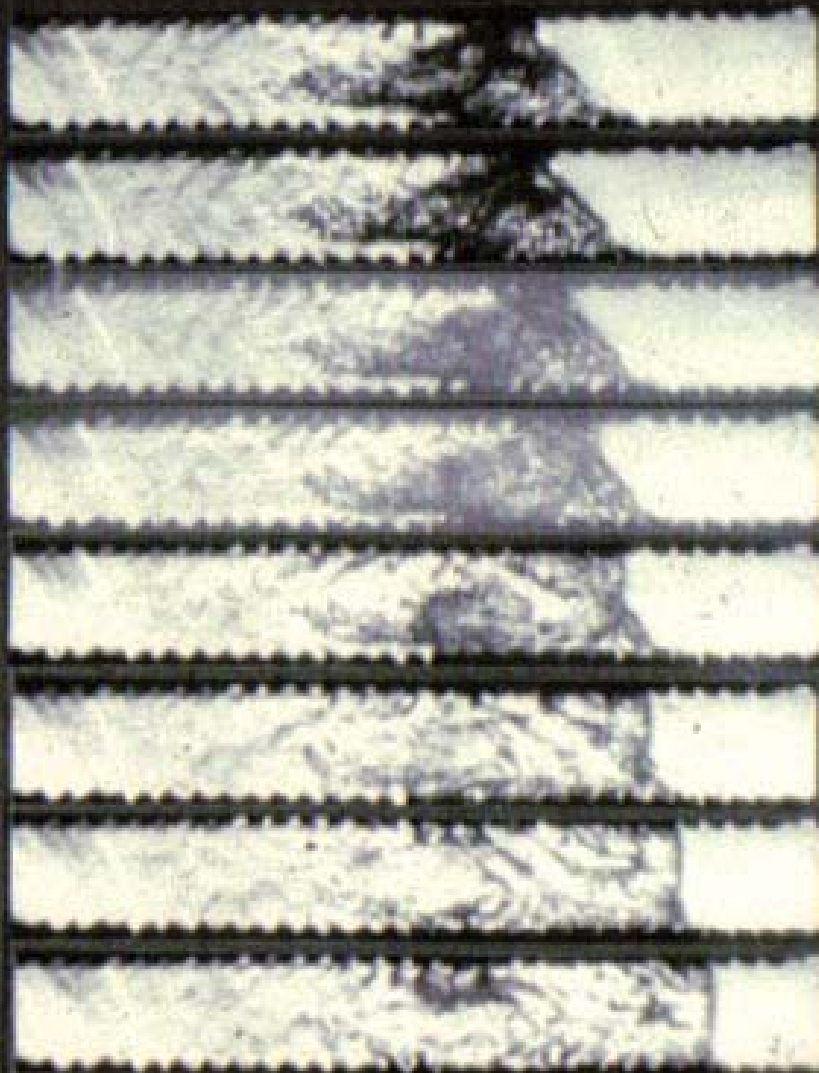
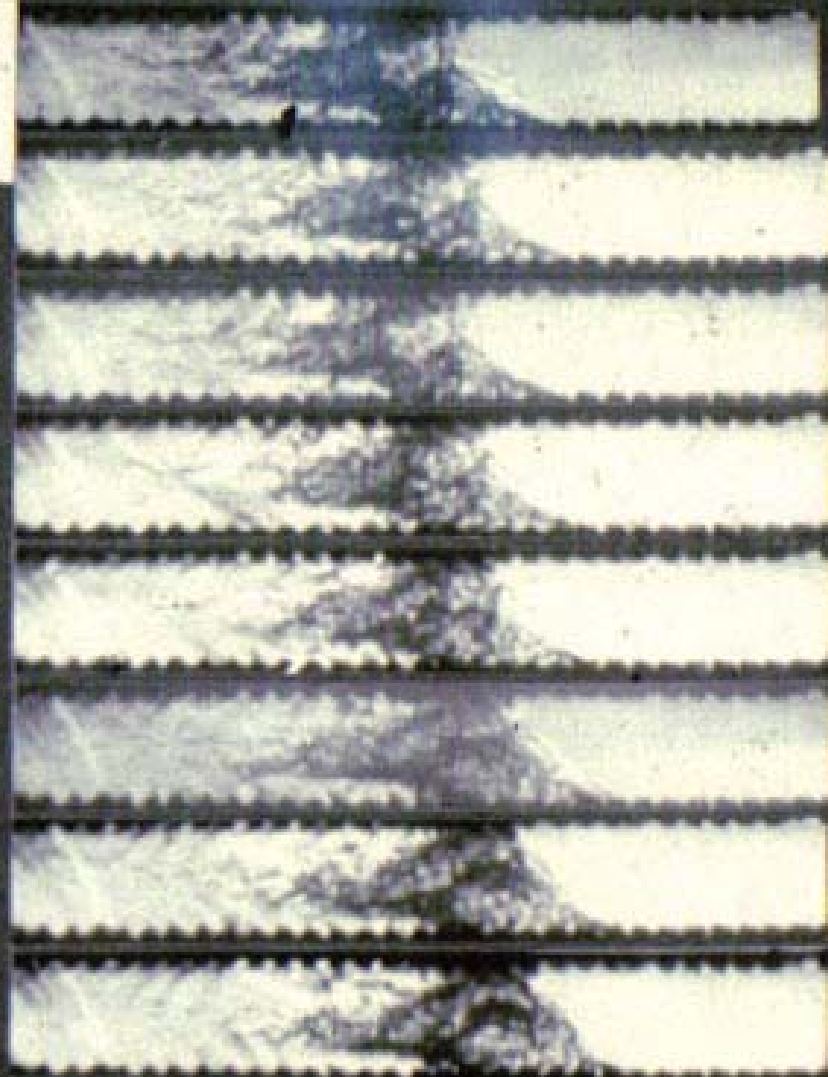
FIG. 2a

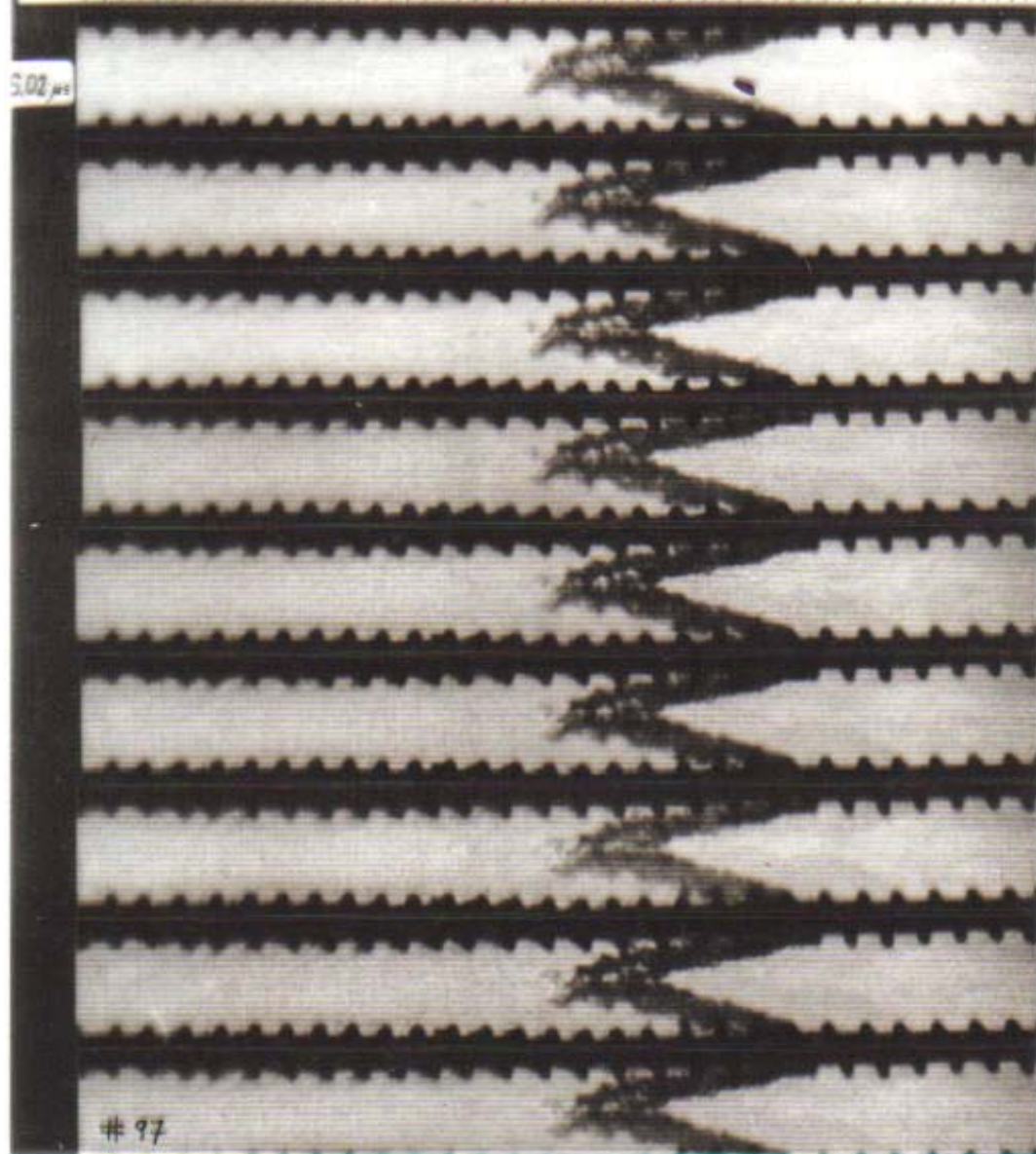
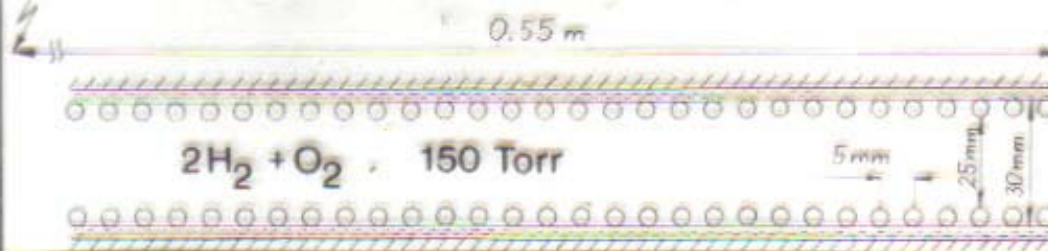
#47





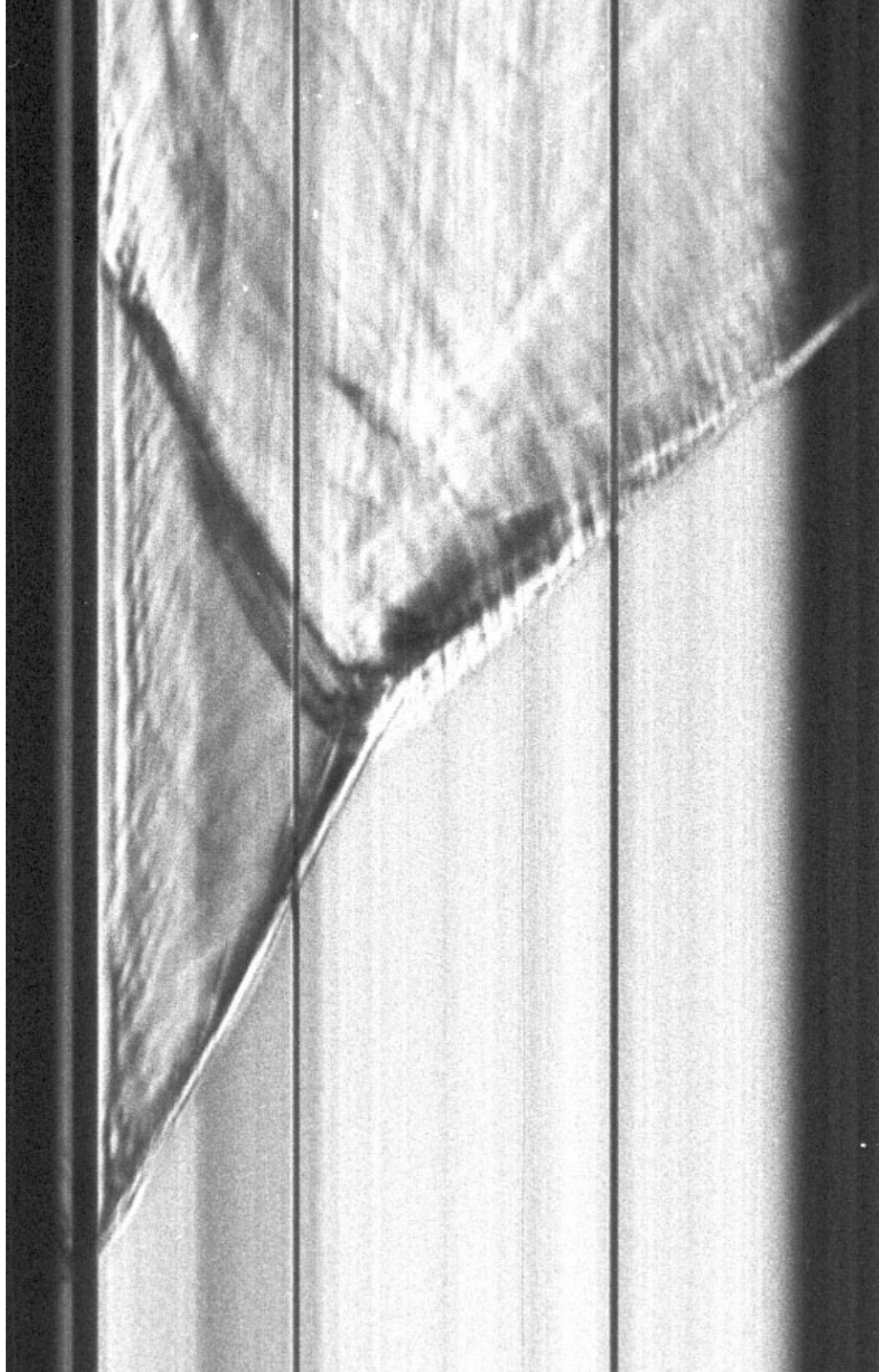
2010
1017

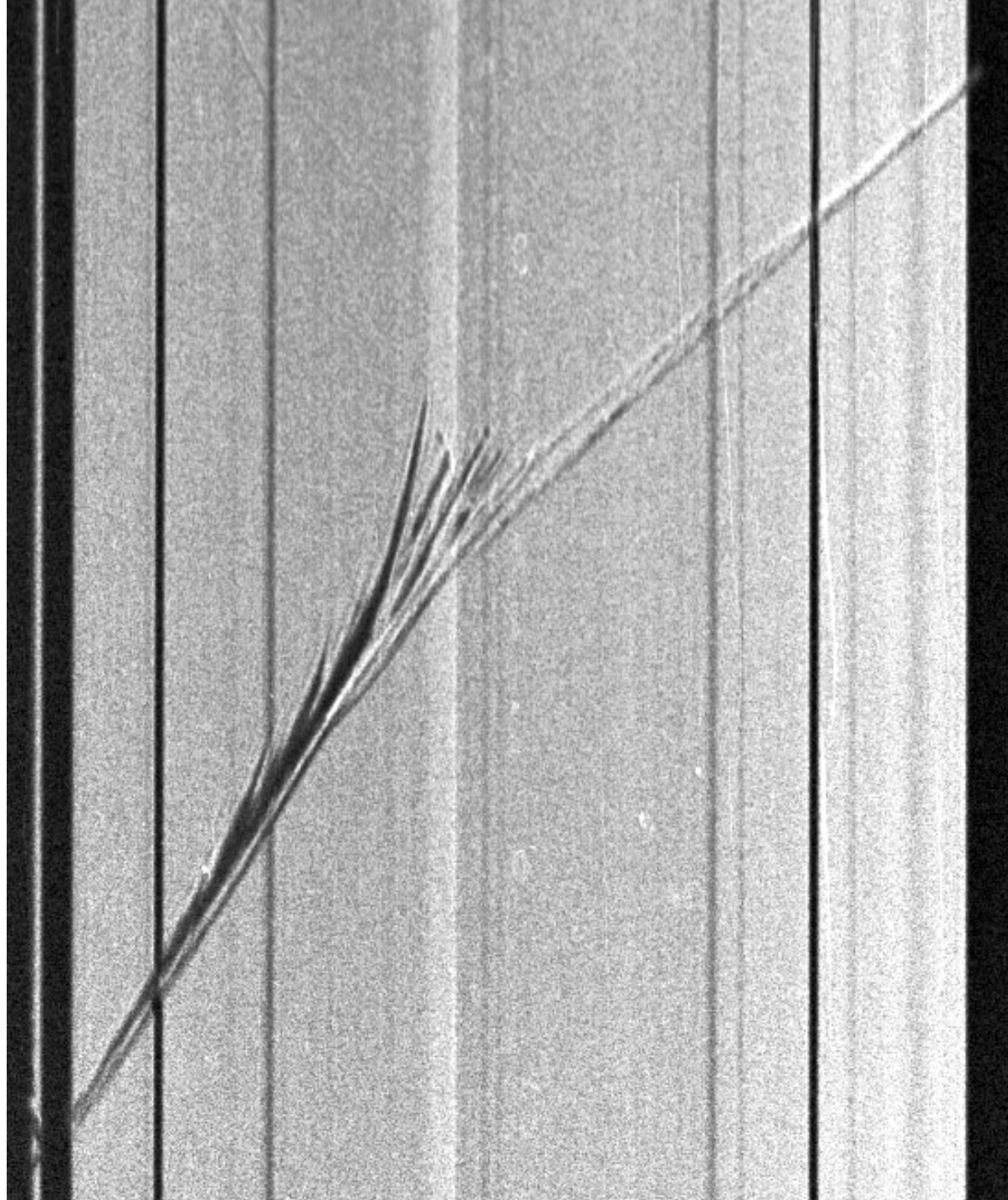


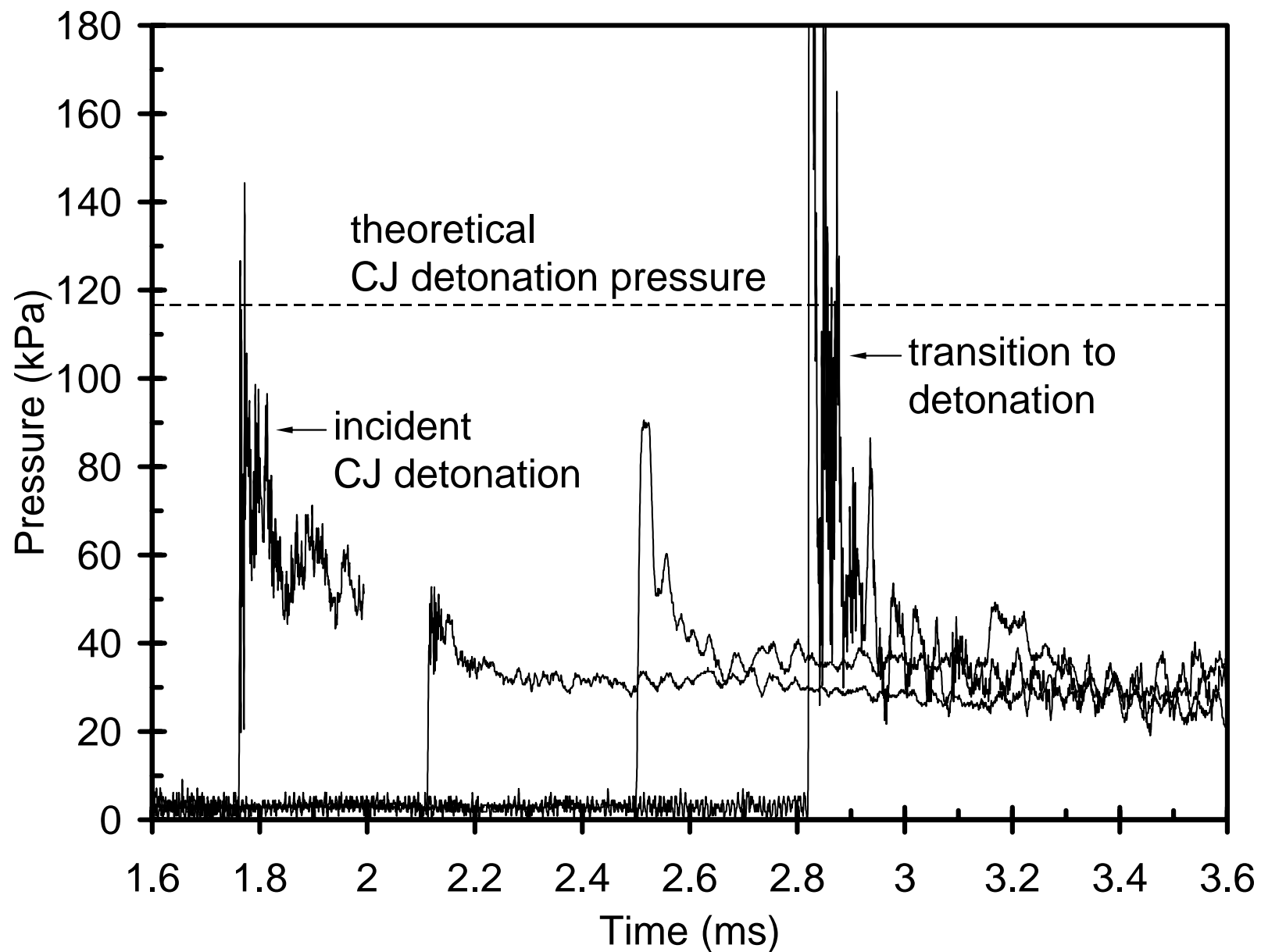


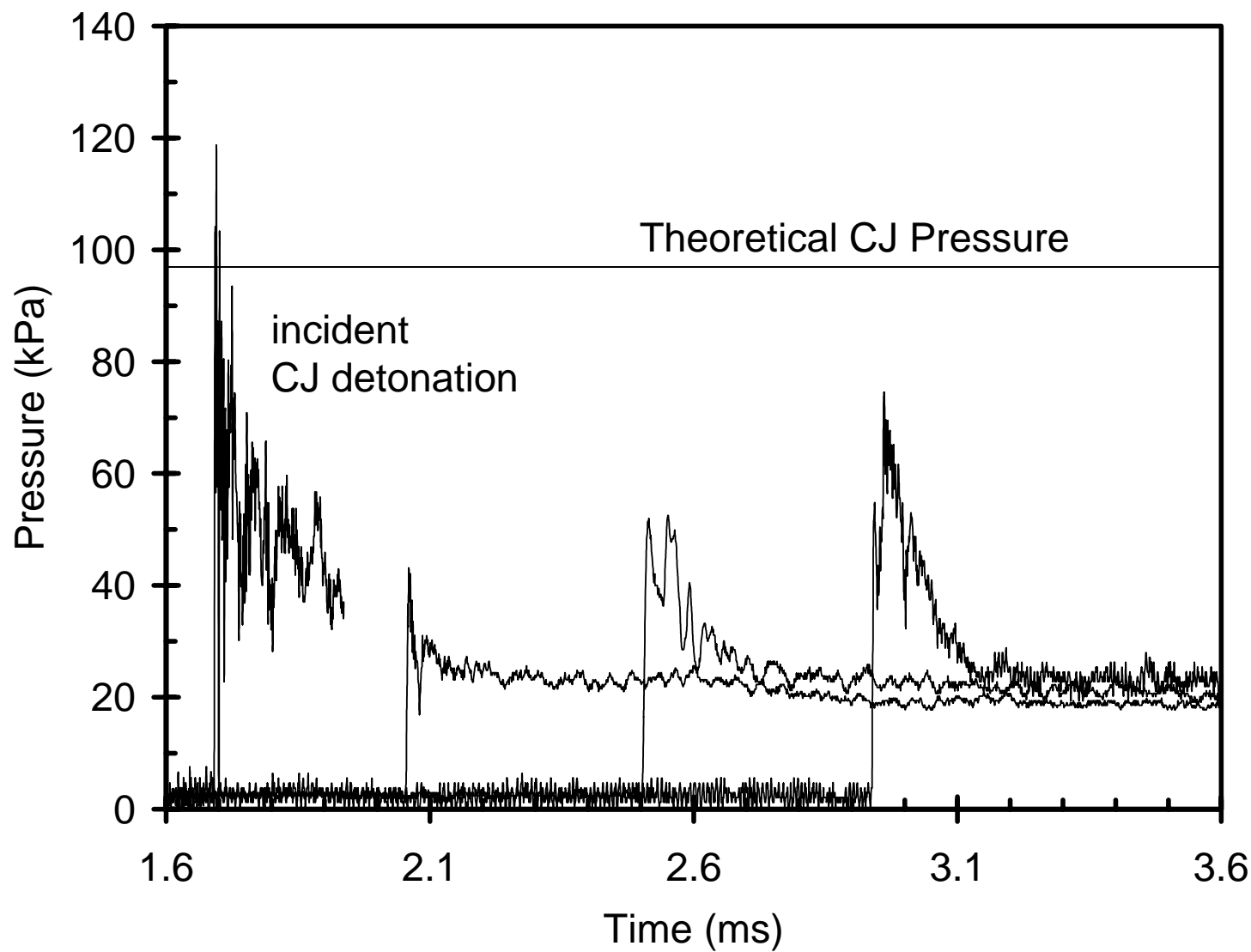
Two Modes of Onset of Detonation

1. unstable mixture: local explosion, SWACER mechanism evidenced by formation of detonation waves
2. progressive wave amplification resonant coupling with turbulent reaction zone

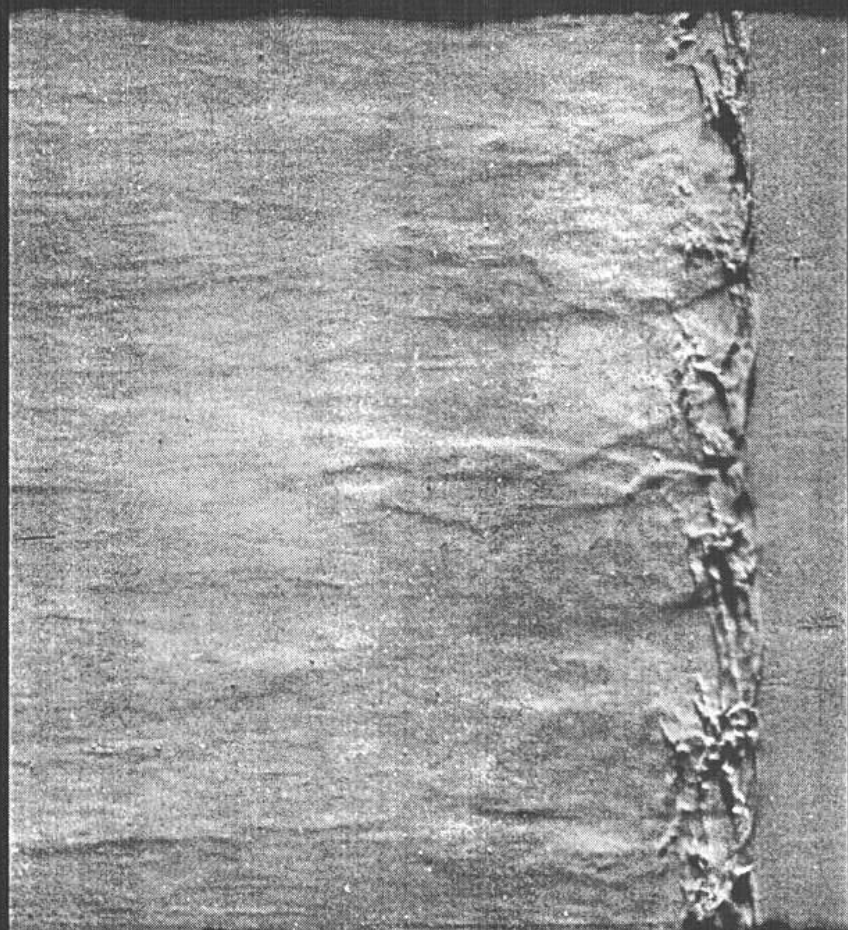




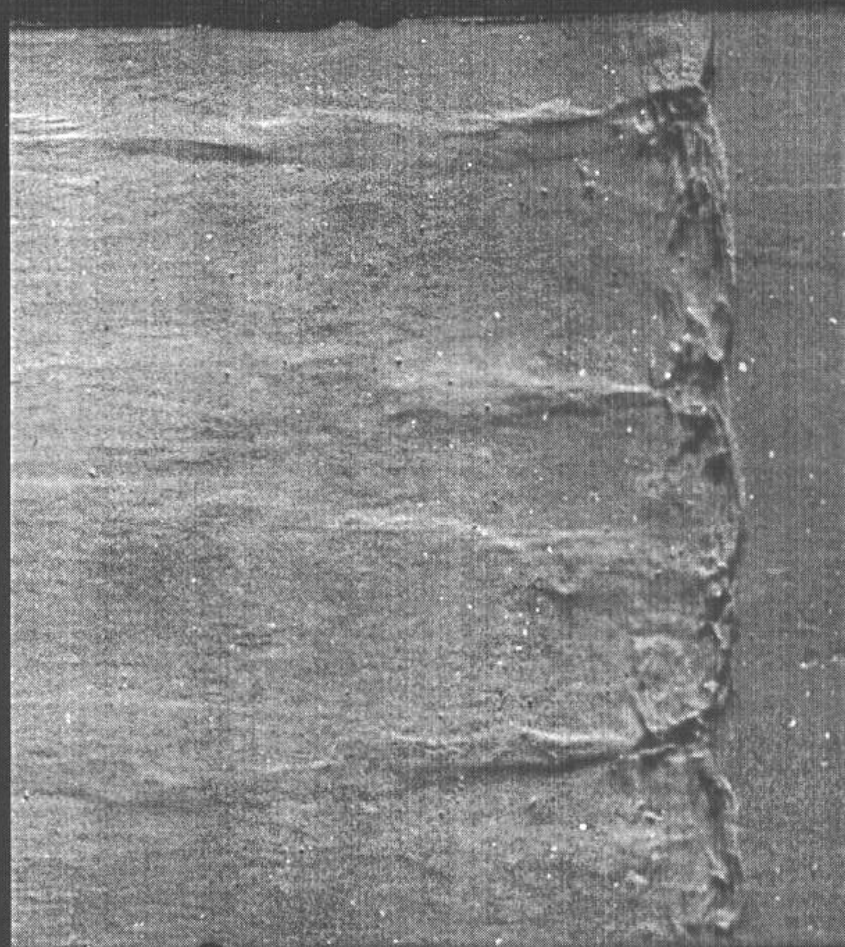




- detonation mechanism is resonant coupling between transverse pressure waves and chemical reactions
- transition means setting up the conditions for the resonant coupling to occur



$\text{C}_2\text{H}_2 + 9.5\text{O}_2$, 5.5kPa



$\text{C}_2\text{H}_2 + 2.5\text{O}_2 + 10.5\text{Ar}$, 5kPa

- turbulent combustion brings the deflagration to maximum speed; Chapman-Jouguet deflagration $\sim \frac{1}{2} V_{CJ}$
- transition to detonation requires the resonant coupling between transverse pressure fluctuations and the chemical reactions

- Chapman-Jouguet deflagration speed is not governed by reaction rate (hence turbulence)
- turbulent combustion rate must be fast enough to pressurize reaction zone
- gasdynamic expansion drives the deflagration like a CJ detonation
- hence, sound speed energetic parameters dominate and not turbulence

Outstanding Problems in DDT

- quantify the pre-detonation state
(thermodynamic, turbulence, chemical kinetics)
- theory for the development of local explosions centers
from hydrodynamic fluctuations
- condition for rapid amplification of pressure waves
(SWACER)