

# Ch.9 Compressible flow

high speed  $\rightarrow$  density change  $\rightarrow$  compressible

$\left\{ \begin{array}{l} \text{liquid} \rightarrow \text{almost incompressible} \Leftrightarrow Ma = \frac{V}{a} \ll 1 \\ \text{gas} \rightarrow \text{compressible} \end{array} \right.$

$\hookrightarrow$  gas dynamics

$T$   
speed of sound

## 9.1 Introduction

Incompressible flow :  $\frac{V}{(3)}, P_{(1)}$  (constant pressure)  $\hookrightarrow$  mechanical press.  $\nabla p$   
 $\hookrightarrow$  cont. eq (1) N-S eqs. (3)  $\leftrightarrow$  4 eqs.

compressible flow :  $\frac{V}{(3)}, P_{(1)}, \rho, T$   $\leftarrow$  temperature  
 $\hookrightarrow$  6 unknowns

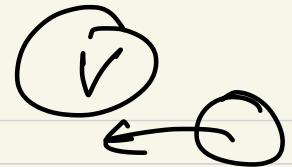
cont. eq (1)  
N-S eqs. (3)

thermodynamic  
press. Change in  $P \rightarrow$  change in  $T$

6 eqs.

$\downarrow$   
energy eq. (1)  
change in  $P \Leftrightarrow$  state eq.  $P = \rho R T$  (1)

- $\boxed{Ma = V/a}$  Mach number



$Ma < \underline{0.3}$

: incomp. flow ( $\rho \doteq \text{const}$ )

$0.3 < Ma < \underline{0.8}$

: subsonic flow (no shock)

$0.8 < Ma < 1.2$

: transonic " (shock)

$1.2 < Ma < 3.0$  : supersonic " (no subsonic)

$Ma > 3.0$  : hypersonic "

- specific-heat ratio,  $k = c_p/c_v$   $k=1.4$  for air

perfect gas,  $P = \rho R T$ ,  $R = c_p - c_v = \text{const}$

$$c_p = \frac{k}{k-1} R$$

For real gas,  $c_p$ ,  $c_v$ ,  $k$  moderately vary with  $T$

$$du = c_v dT$$

$$TdS = du + P d\left(\frac{1}{P}\right)$$

$$dh = c_p dT$$

$$= dh - \frac{1}{P} dp = c_p dT - \frac{1}{P} dp$$

$$\Rightarrow ds = \frac{c_p}{T} dT - \frac{1}{P} dp = \frac{c_p}{T} dT - \frac{R}{P} dp$$

$$s_2 - s_1 = c_p \ln \frac{T_2}{T_1} - R \ln \frac{P_2}{P_1} = c_v \ln \frac{T_2}{T_1} - R \ln \frac{P_2}{P_1}$$

\* Isentropic process :  $ds = 0$

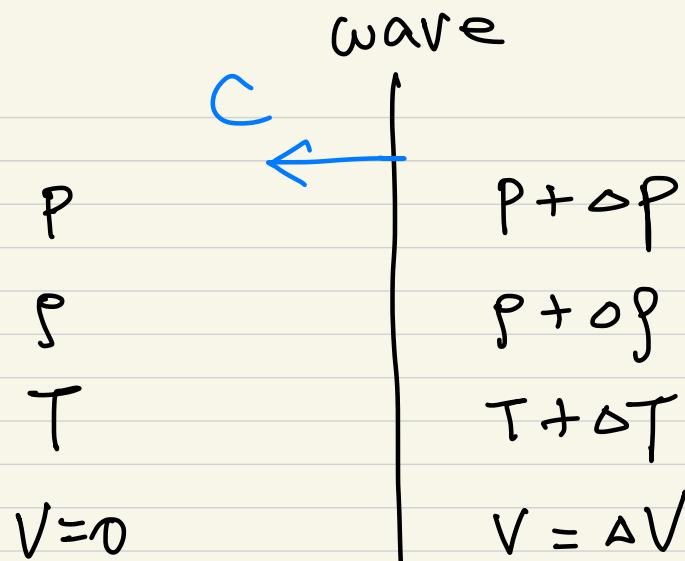
$$\Delta S = 0$$

reversible  
adiabatic

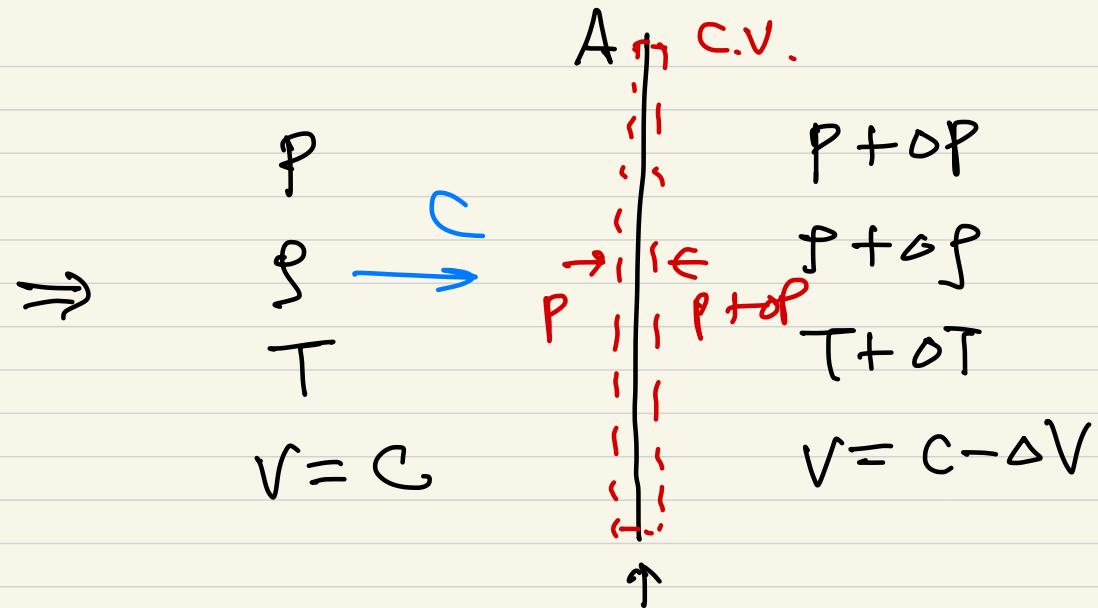
$$\boxed{\frac{P_2}{P_1} = \left(\frac{T_2}{T_1}\right)^{\frac{k-1}{k}} = \left(\frac{P_2}{P_1}\right)^k}$$

## 9.2 Speed of sound

→ a pressure pulse of infinitesimal strength



moving wave  
of frontal area  $A$



fixed wave

$$\text{cont: } \rho c A = (\rho + \Delta \rho)(c - \Delta V) A \Rightarrow \Delta V = c \frac{\Delta \rho}{\rho + \Delta \rho} \quad \text{--- ①}$$

$$\text{mtm: } \sum F = PA - (P + \Delta P)A$$

if  $\Delta \rho = O(\epsilon)$ ,  
 $\Delta V = O(\epsilon)$ .

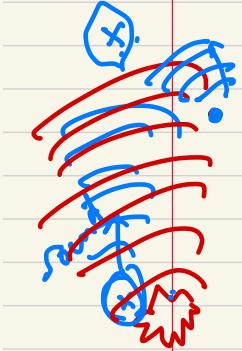
$$= -\rho c^2 A + (P + \Delta P)(c - \Delta V)^2 A$$

$$\Rightarrow \Delta P = \rho c \Delta V \quad \text{--- ②}$$

if  $\Delta V = O(\epsilon)$ ,  
 $\Delta P = O(\epsilon)$

$$\textcircled{1} \rightarrow \textcircled{2}, \quad c^2 = \frac{\partial P}{\partial \rho} \left( 1 + \frac{\partial P}{P} \right)$$

$$P = P(\rho, T)$$

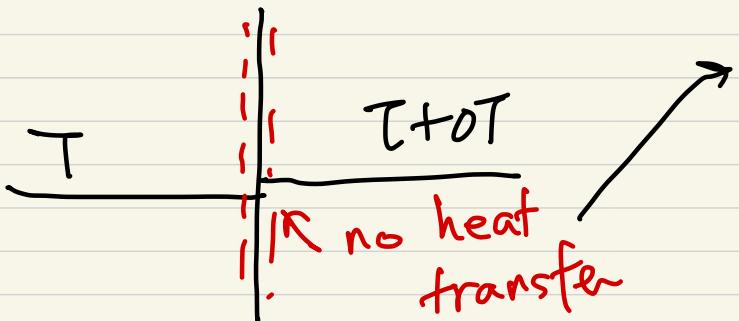


$$\frac{\partial P}{\partial \rho} \rightarrow 0 \quad \Rightarrow \quad a^2 = \frac{\partial P}{\partial \rho} \Big|_S \quad ?$$

: speed of sound

powerful explosion ( $\Delta P = O(1)$ ) waves move  
much faster than sound wave ( $\Delta P = O(\epsilon)$ )

Newton (1686) obtained 'a' using the assumption of  
isothermal process.  $\rightarrow 20\%$  error  $\therefore$  wrong!



adiabatic process

$\Delta T = O(\epsilon) \rightarrow$  reversible process

$\Rightarrow$  isentropic process

$$a^2 = \frac{\partial P}{\partial \rho} \Big|_S$$

$$\Delta S = 0 : \frac{P}{\rho^k} = \text{const} \rightarrow \ln P - k \ln \rho = \text{const.}$$

$$\rightarrow \frac{dP}{P} - k \frac{d\rho}{\rho} = 0$$

$$\rightarrow \frac{\partial P}{\partial \rho} \Big|_S = k \frac{P}{\rho} = kRT = a^2$$

$$\Rightarrow a = \sqrt{kRT}$$

@ 1 atm, 15°C, air  $a = 340 \text{ m/s}$

water      1490 m/s      bulk  
steel      5060 m/s      modulus

### 9.3 Adiabatic and isentropic steady flow

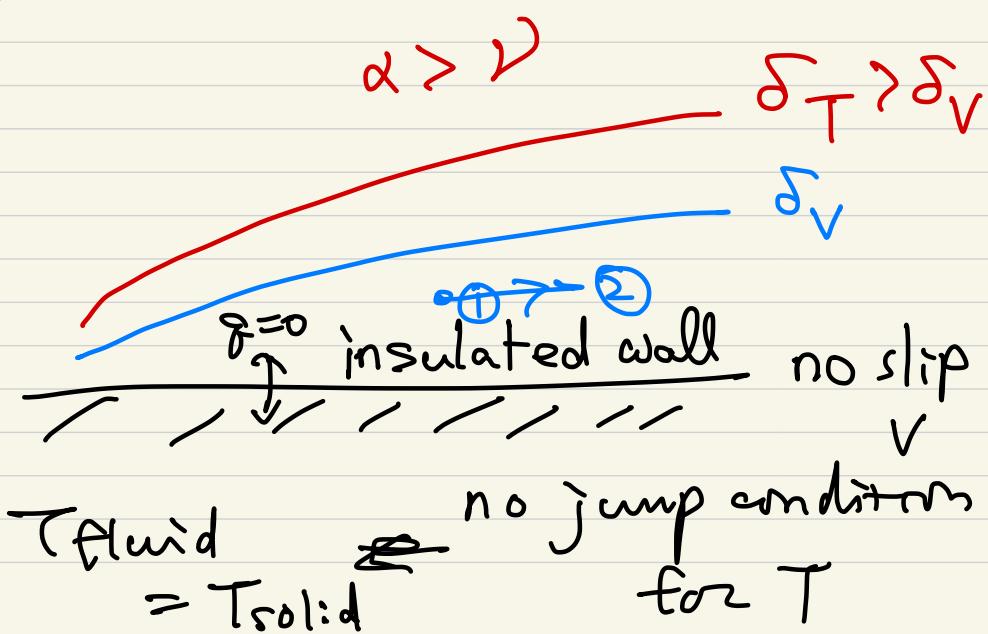
$$Pr = \frac{V}{\alpha} < 1 \quad \text{air } Pr = 0.7 \text{ or } 0.71$$

$$\int \frac{\partial V}{\partial e} = \dots + \mu T^2 V$$

$$\int C_P \frac{\partial T}{\partial e} = \dots + \alpha \nabla T$$

$$V < \alpha$$

$$U_\infty \rightarrow T_\infty$$



energy conservation  $B = E$ ,  $\beta = \frac{dE}{dm} = e$

$$\left( e = e_{\text{internal}} + e_{\text{kinetic}} + e_{\text{potential}} + \dots \right)$$

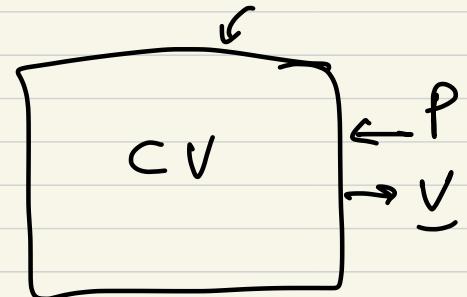
$$= \dot{u} + \frac{1}{2}v^2 + gz + \dots$$

$$\frac{dQ}{dt} - \frac{dW}{dt} = \frac{dE}{dt} = \frac{d}{dt} \int_{CV} e \rho dV + \int_{CS} e \rho (\underline{V} \cdot \underline{n}) dA$$

$$\dot{w} = \frac{dW}{dt} = \dot{w}_{\text{shape}} + \dot{w}_{\text{press}} + \dot{w}_{\text{viscous stress}}$$

$$= \dot{w}_s + \dot{w}_p + \dot{w}_v$$

$$\dot{w}_p = \int_{CS} P (\underline{V} \cdot \underline{n}) dA$$



$$\dot{w}_v = - \int_{CS} \underline{\tau} \cdot \underline{V} dA \quad \underline{\tau} : \text{stress vector on } dA$$

$$\rightarrow \dot{Q} - \dot{w}_s - \dot{w}_v = \frac{\partial}{\partial t} \int_{CV} e \rho dV + \int_{CS} \underline{c} \underline{\tau} + \frac{\underline{P}}{\rho} \cdot \underline{\tau} (\underline{V} \cdot \underline{n}) dA$$

$$= (\dot{u} + \frac{1}{2}v^2 + gz) + \frac{P}{\rho} = h + \frac{1}{2}v^2 + gz$$

enthalpy

1D & steady : ①  $\rightarrow$  ②

$$\dot{Q} - \dot{W}_s - \dot{W}_J = -\dot{m}(h_1 + \frac{1}{2}v_1^2 + qz_1) + \dot{m}(h_2 + \frac{1}{2}v_2^2 + qz_2)$$

$$\rightarrow h_1 + \frac{1}{2}v_1^2 + qz_1 = h_2 + \frac{1}{2}v_2^2 + qz_2 - g + \dot{W}_s(\dot{m} + \dot{W}_J/\dot{m})$$

$\stackrel{\text{1. } \dot{Q}/\dot{m}}{\curvearrowleft}$

inside the bdry layer :  $g \neq 0, \dot{W}_J \neq 0$

outside " " " :  $g = \dot{W}_J = 0$

$$\hookrightarrow h_1 + \frac{1}{2}v_1^2 + qz_1 = h_2 + \frac{1}{2}v_2^2 + qz_2$$

neglect  $g(z_2 - z_1) \approx 0$

$$\rightarrow h_1 + \frac{1}{2}v_1^2 = h_2 + \frac{1}{2}v_2^2 = \text{const} = h_0 = h + \frac{1}{2}v^2$$

stagnation enthalpy  
 $h_0$  varies inside the thermal boundary layer,  
but its average value is the same because of energy  
conservation.

For perfect gas,  $h = c_p T \rightarrow c_p T_0 = c_p T + \frac{1}{2} V^2$

↑ stagnation temperature

$$V_{\max} = (2 c_p T_0)^{\frac{1}{2}} = (2 \frac{kR}{k-1} T_0)^{\frac{1}{2}}$$

$$c_p T_0 = c_p T + \frac{1}{2} U^2 \quad c_p T = \frac{kR}{k-1} T = \frac{a^2}{k-1} \quad (a = \sqrt{kR T})$$

$$\rightarrow \frac{T_0}{T} = 1 + \frac{U^2}{2 c_p T} = 1 + \frac{(k-1) V^2}{2 a^2}$$

$$Ma = \frac{V}{a}$$

$$\rightarrow \boxed{\frac{T_0}{T} = 1 + \frac{k-1}{2} Ma^2}$$

adiabatic

Given  $T$  &  $Ma \rightarrow T_0$

$$\boxed{\frac{a_0}{a} = \left(\frac{T_0}{T}\right)^{\frac{1}{2}} = \left(1 + \frac{k-1}{2} Ma^2\right)^{\frac{1}{2}}}$$

adiabatic

If isentropic process,

$P_0$ : stag. pres.

$\rho_0$ : " density

$$\frac{P_0}{P} = \left(\frac{T_0}{T}\right)^{\frac{k}{k-1}} = \left(1 + \frac{k-1}{2} M_a^2\right)^{\frac{k}{k-1}}$$

$$\frac{P_0}{\rho} = \left(\frac{T_0}{T}\right)^{\frac{1}{k-1}} = \left(1 + \frac{k-1}{2} M_a^2\right)^{\frac{1}{k-1}}$$

1.  $\rightarrow$  2. irrev.  
adibatic

$$T_{01} = T_{02} \rightarrow h_{01} = h_{02}$$

$$a_{01} = a_{02}$$

$$\overbrace{P_1 + \frac{1}{2} \rho_1 V_1^2}^{P_{01}} = P_{02} \neq P_2 + \frac{1}{2} \rho_2 V_2^2$$

$$P_{01} \neq P_{02}$$

isentropic  $\Rightarrow T_{01} = T_{02}, a_{01} = a_{02}, P_{01} = P_{02}, \rho_1 = \rho_2$ .

•  $h + \frac{1}{2} V^2 = \text{const (adiabatic)}$

$$\rightarrow dh + VdV = 0$$

$$TdS = dh - \frac{1}{P} dP \stackrel{\text{isentropic}}{=} 0 \rightarrow dh = \frac{1}{P} dP$$

$$\frac{dp}{\rho} + VdV = 0$$

Bernoulli eq

isentropic process

