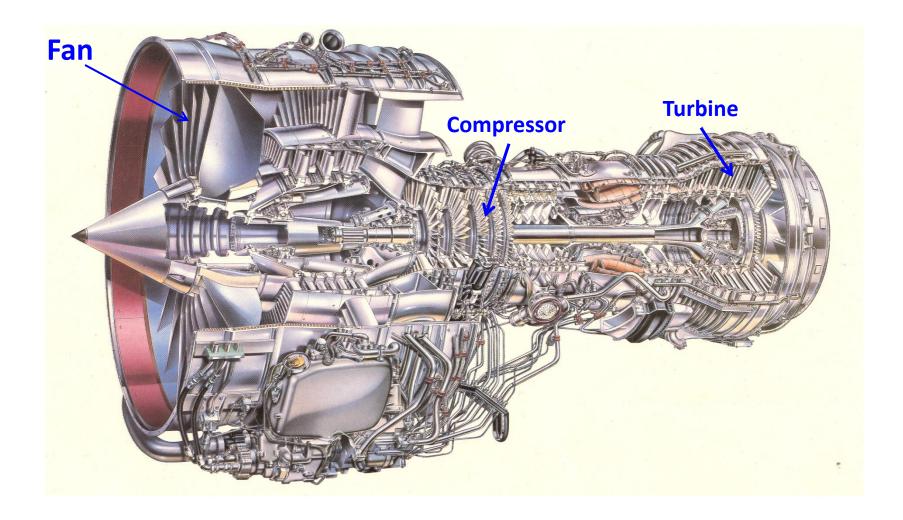
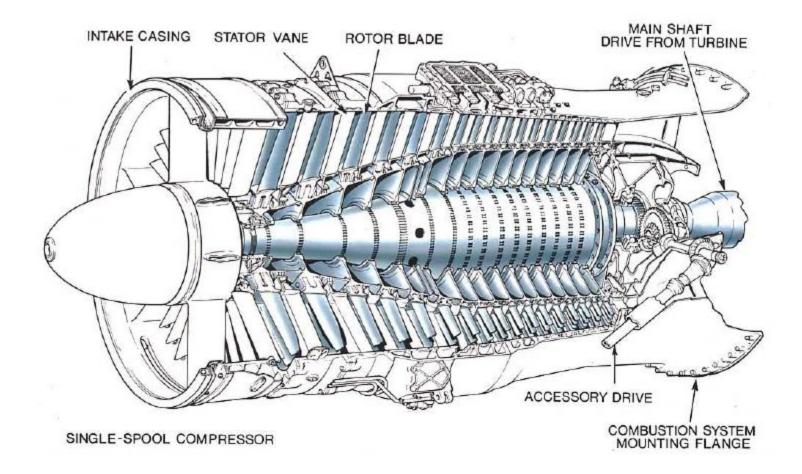
- Recap: Lecture 1: 21<sup>st</sup> July 2015, 1530-hrs.
  - Introduction
  - Aims of the course
  - Course contents
  - Text books/references
  - Evaluation scheme and course schedule

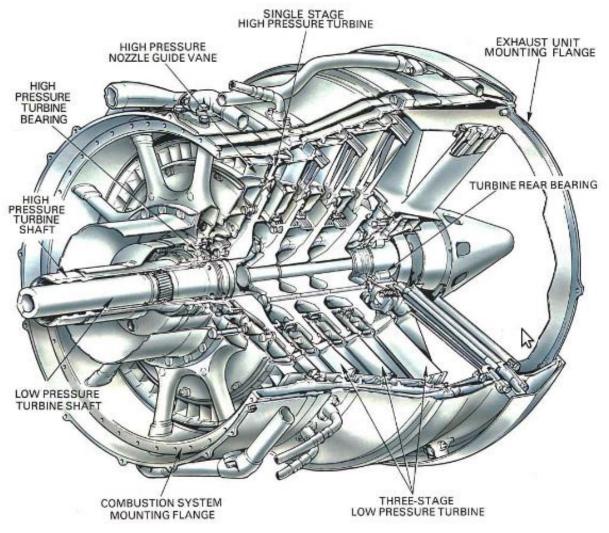


A modern high bypass turbofan engine



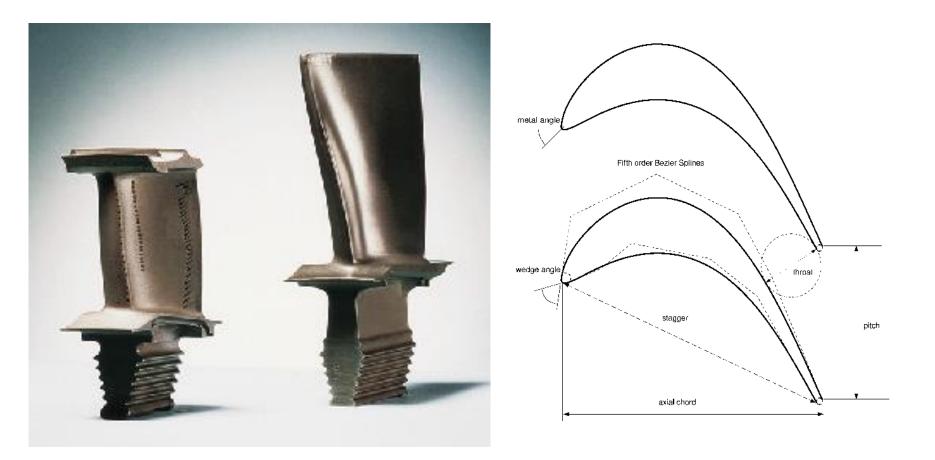
#### Typical multi-stage axial flow compressor

Source: The Jet Engine, Roll Royce Plc, 1996



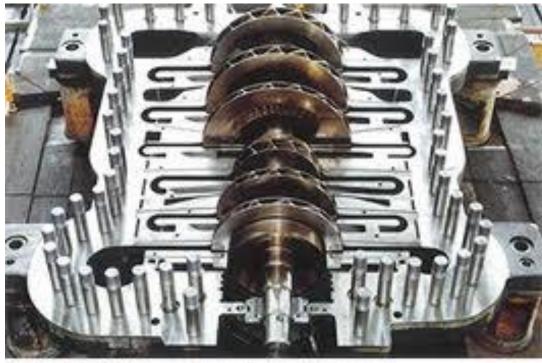
#### A twin turbine and shaft system

Source: The Jet Engine, Rolls Royce, 1996



#### Turbine nozzle and rotor blade geometries



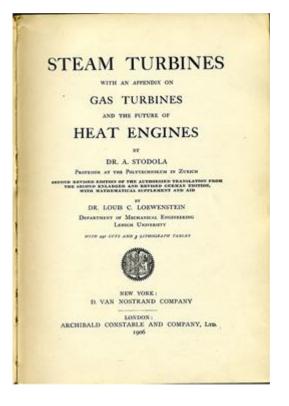


Source: Mitsubishi Heavy Industries, Ltd.

#### Centrifugal compressor

	787 Engines: GEnx-1B Trent 1000	767 Engines: GE CF6-80C2 RR RB211-524G/H
Bypass Ratio	~10	~5
<b>Overall Pressure Ratio</b>	~50	~33
Thrust Class	53,000–74,000 lbf	53,000–63,000 lbf
Fan Diameter	111–112 in	86–93 in
Specific Fuel Consumption	15% lower	Base
Noise	ICAO Chapter 4	ICAO Chapter 3
Emissions	CAEP/8 (2014)	CAEP/2

Source: http://www.boeing.com/commercial/aeromagazine/articles/2012\_q3/2/

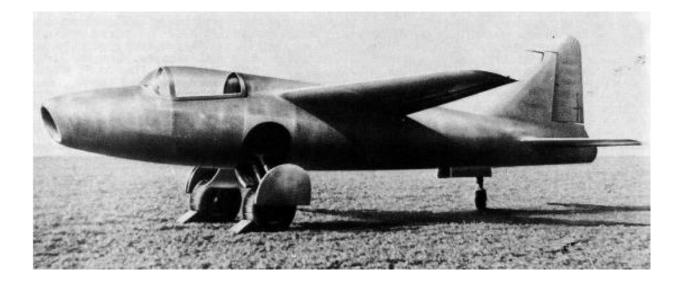




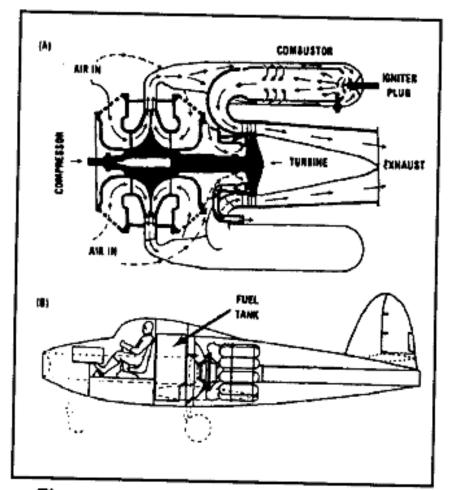
*Commissioning of world's first industrial gas turbine, Neuchatel, 1939 (Stodola at age 80)* 



Frank Whittle and Hans von Ohain



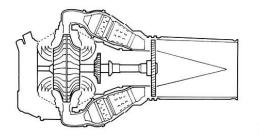
#### First turbojet-powered aircraft – Ohain's engine on He 178 (1939)



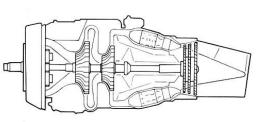
PHOTOVALITY CHOTOVALITY

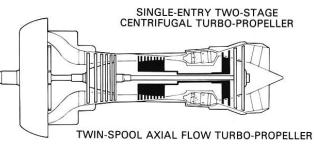
Gloster Whittle E-28/39

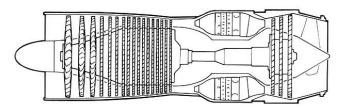
Figure 2-5. A – Whittle's Reverse-Flow Combustion Chamber. B – Fuselage Arrangement of the E28/39 Experimental Powered the Gloster E28/39 Britain on 15 May 1941.



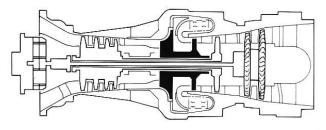
DOUBLE-ENTRY SINGLE-STAGE CENTRIFUGAL TURBO-JET







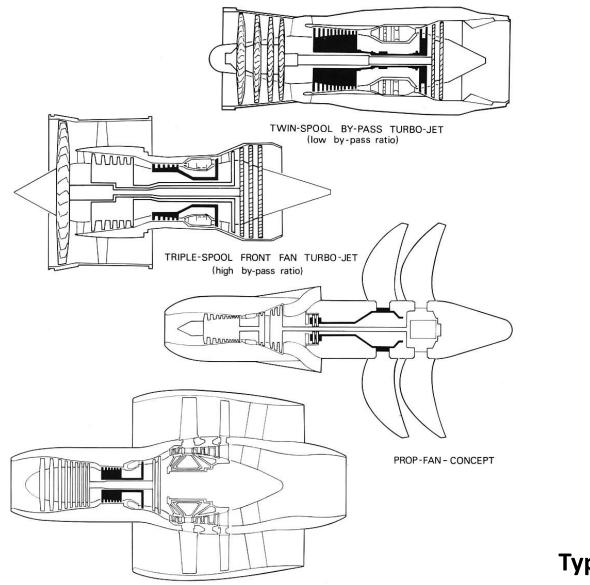
SINGLE-SPOOL AXIAL FLOW TURBO-JET



TWIN-SPOOL TURBO-SHAFT (with free-power turbine)

#### **Types of Gas Turbine Engines**

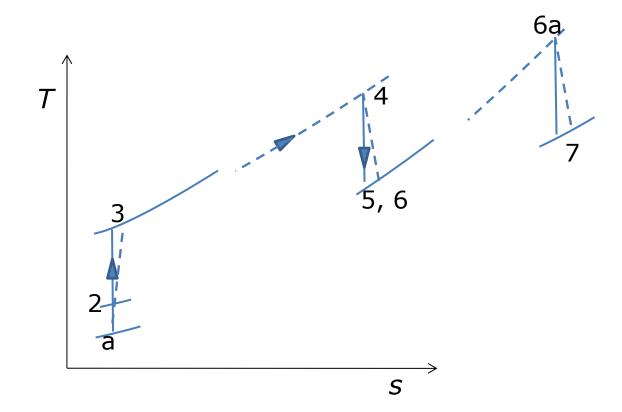
Courtesy: The Jet Engine: Rolls Royce, 1996



#### **Types of Gas Turbine Engines**

CONTRA-ROTATING FAN - CONCEPT (high by-pass ratio)

#### **Real cycle for turbojet engines**



Real turbojet cycle (with afterburning) on a T-s diagram

#### **Review of Thermodynamic Concepts**

#### Total energy of a system

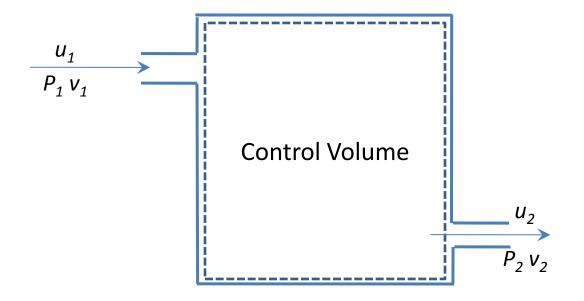
 In the absence of magnetic, electric, and surface tension effects, the total energy of a system consists of the kinetic, potential, and internal energies

$$E = U + KE + PE = U + \frac{mV^2}{2} + mgz \quad (kJ)$$

or, on a unit mass basis

$$e = u + ke + pe = u + \frac{V^2}{2} + gz$$
 (kJ/kg)

## Enthalpy



The combination u+Pv is frequently encountered in the analysis of control volumes

# Enthalpy

- The combination of internal energy u and Pv is often encountered in the analysis of control volumes
- Enthalpy is a combination property Enthalpy, h = u + Pv (kJ/kg)

 $H = U + PV \quad (kJ)$ 

- Enthalpy is also often referred to as heat content
- Process in which enthalpy is constant: isenthalpic process

## Entropy

- Entropy is an extensive property of a system and sometimes is referred to as total entropy. Entropy per unit mass, designated s, is an intensive property and has the unit kJ/kg.K
- The entropy change of a system during a process can be determined by

$$\Delta S = S_2 - S_1 = \int_1^2 \left(\frac{\delta Q}{T}\right)_{\text{int. rev.}} \quad (\text{kJ/kg})$$

## Entropy

- Entropy is a property, and like all other properties, it has fixed values at fixed states.
- Therefore, the entropy change *dS* between two specified states is the same no matter what path, reversible or irreversible.

#### **Temperature-entropy plot**

$$dS = \frac{dQ_{rev}}{T}$$

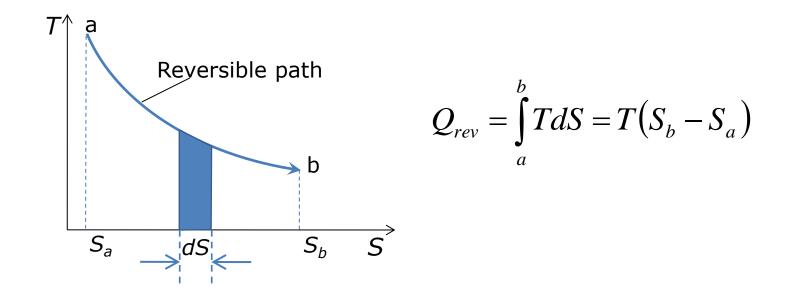
If the process is reversible and adiabatic,  $dQ_{rev} = 0$ 

 $\therefore dS = 0 \text{ or } S = \text{constant}$ 

• A reversible adiabatic process is, therefore, and isentropic process.

$$dQ_{rev} = TdS$$
  
or,  $Q_{rev} = \int TdS$ 

#### **Temperature-entropy plot**



• The area under the reversible path on the T-S plot represents heat transfer during that process.

#### **Isentropic processes**

- A process where,  $\Delta s=0$
- An isentropic process can serve as an appropriate model for actual processes.
- Isentropic processes enable us to define efficiencies for processes to compare the actual performance of these devices to the performance under idealized conditions.
- A reversible adiabatic process is necessarily isentropic, but an isentropic process is not necessarily a reversible adiabatic process.

#### **TdS equations**

 From the first law for an internally reversible process, we know that

$$dQ_{\rm int\,rev} - dW_{\rm int\,rev,out} = dU$$

Since,  $dQ_{int rev} = TdS$  and  $dW_{int rev,out} = PdV$ TdS = dU + PdV or, Tds = du + Pdv

• This is known as the first TdS equation.

#### **TdS equations**

• From the definition of enthalpy, we know that, h = u + Pv

or, dh = du + Pdv + vdPsince, Tds = du + Pdv,

Tds = dh - vdP

This is known as the second TdS equation.

#### Entropy change of ideal gases

For ideal gases we know that,

 $du = c_v dT$ , P = RT / vFrom the *TdS* relations, dT = dv

$$ds = c_v \frac{dT}{T} + R \frac{dv}{v}$$

• The entropy change for a process,

$$s_2 - s_1 = \int_{1}^{2} c_v(T) \frac{dT}{T} + R \ln \frac{v_2}{v_1}$$

#### Entropy change of ideal gases

If we use these relations,  $dh = c_p dT$ , v = RT / PThen, from the *TdS* relations,

$$s_2 - s_1 = \int_{1}^{2} c_p(T) \frac{dT}{T} - R \ln \frac{P_2}{P_1}$$

• Usually, we assume average values of  $c_p$ and  $c_v$  in the above equations and thus can replace  $c_p(T)$  with  $c_{p,av}$  and  $c_v(T)$  with  $c_{v,av}$ .

# Energy analysis of steady flow systems

For single entry and exit devices,

$$\dot{Q} - \dot{W} = \dot{m} \left[ h_2 - h_1 + \frac{V_2^2 - V_1^2}{2} + g(z_2 - z_1) \right]$$

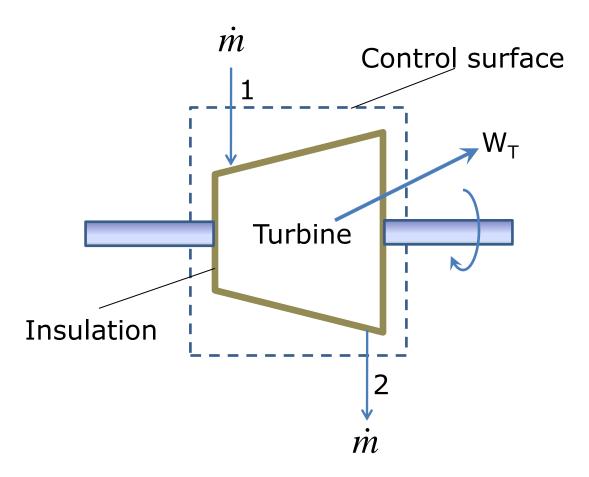
or per unit mass,

$$\dot{q} - \dot{w} = h_2 - h_1 + \frac{V_2^2 - V_1^2}{2} + g(z_2 - z_1)$$

#### **Turbines and compressors**

- Pumps, compressors and fans: used to increase the pressure of a fluid and require work input.
- Turbines generate work.
- Q, KE and PE may or may not be zero.
- Usually *PE* is negligibly small.

#### **Turbines and compressors**



#### **Turbines and compressors**

 For a turbine for eg., the energy equation would be:

$$\dot{m}(h_{1} + \frac{V_{1}^{2}}{2} + gz_{1}) = \dot{W}_{out} + \dot{m}(h_{2} + \frac{V_{2}^{2}}{2} + gz_{2})$$
  
If KE and PE are negligible,  
 $\dot{W}_{out} = \dot{m}(h_{1} - h_{2})$ 

- Enthalpy represents the total energy of a fluid in the absence of potential and kinetic energies.
- For high speed flows, though potential energy may be negligible, but not kinetic energy.
- Combination of enthalpy and KE is called stagnation enthalpy (or total enthalpy)

$$h_0 = h + V^2/2$$
 (kJ/kg)

Stagnation enthalpy Static enthalpy Kinetic energy

- Consider a steady flow through a duct (no shaft work, heat transfer etc.).
- The steady flow energy equation for this is:  $h_1 + V_1^2/2 = h_2 + V_2^2/2$ or,  $h_{01} = h_{02}$
- That is in the absence of any heat and work interactions, the stagnation enthalpy remains a constant during a steady flow process.

• If the fluid were brought to rest at state 2,

 $h_1 + V_1^2/2 = h_2 = h_{02}$ 

- The stagnation enthalpy represents the enthalpy of a fluid when it is brought to rest adiabatically.
- During a stagnation process, the kinetic energy of a fluid is converted to enthalpy (internal energy + flow energy), which results in an increase in the fluid temperature and pressure.

• When the fluid is approximated as an ideal gas with constant specific heats,

$$c_p T_0 = c_p T + V^2/2$$
  
or,  $T_0 = T + V^2/2c_p$ 

- *T<sub>0</sub>* is called the stagnation temperature and represents the temperature an ideal gas attains when it is brought to rest adiabatically.
- The term  $V^2/2c_p$  corresponds to the temperature rise during such a process and is called the dynamic temperature.

- The pressure a fluid attains when brought to rest isentropically is called the stagnation pressure,  $P_0$ .
- For ideal gases, from isentropic relations,

$$\frac{P_0}{P} = \left(\frac{T_0}{T}\right)^{\gamma/(\gamma-1)}$$

Similarly, for density wehave,

$$\frac{\rho_0}{\rho} = \left(\frac{T_0}{T}\right)^{1/(\gamma-1)}$$