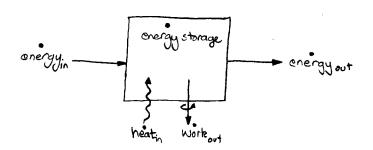
Energy balance is based on the conservation of evergy (first law of thermodynamics) and the net growth of entropy (second law of thermodynamies)

Energy balance is used to track energy through a system. It is a little more complicated than the mass balance because the "transfer" term involves heat and work



The energy balance is usually written as an energy flow rate equation Conservation of energy for a system says:

$$\frac{dE}{dt}\Big|_{system} = \frac{dQ}{dt} - \frac{dW}{dt}$$

$$heat_n \quad work_{out}$$

In words: rate of energy accumulated - rate entering + rate exiting = rate of heat entering - rate of work (done by system) out.

Using c as a symbol for energy, the balance can be arranged as

An open system will ksually have both mass of energy Hows.

A closed system will have only heat (and possibly) work flows.

Energy is the ability to do work. There are different "kinds" of energy based on the kind of work performed

kinetil energy - momentum

potential energy - positivis in gravitational, electrical or magnetic

field

Internal energy - microscopic level associated with KE & PE and Hormal overgy of a molecule. Often expressed as pressure in gases of liquids - temporatu in solids

chemical energy - energy associated with chemical bonds

Total energy is sum of all relevant types of energy for a fixed quantity of mass. E = KE+PE+U. Thus the energy in the balance expression includes all types of energy.

Adding energy to a system often raises its temperature (think of a teapot). The amount of heat added (thermal energy) to raise the temperature of a unit mass of a substance by I degree is called the specific heat of the substance.

Btu - energy to heat 116 water, 1°F & 59°F. keal - energy to heat 1kg water, 1°C & 15°C cal - " 1g water, 1°C @ 15°C The preferred unit is kJ/kg. For water He specific heat is 4.184 kJ/kg. For substances that remain as liquids or solicis during the entire heating event (no phase change), the specific hoals are assentially constant.

basses expand during heating unless their volume is fixed. Specific heals are determined either at constant volume, constant pressure, or both Cv - constant volume sp. heat; Cp - constant pressure sp. heat

Enthalpy of a substance is the sum of its internal energy and its "state" (pt). H = U+ pt

internal "compression" energy energy

When heat is added to a substance, He change in contralpy is given by $\frac{dH}{dT} = \frac{dU}{dT} + \frac{dD}{dT} + \frac{dD}{dT} + \frac{dD}{dT} + \frac{dD}{dT}$

For solids & liquids the pt term is relatively constent (dpt ~ small) so that dH = dU dT.

For a gas either p = const. (balloon) or t = const. (bas cylinder) H = U + Pt = U + mRT

dH = dU + +dP = m CV (Although pressure charges, it cannot do any wark; work = farce o distance)

 $\frac{dH}{dT}\Big|_{p=const} = \frac{dV}{dT} + p\frac{dV}{dT} = mC_p$

Work-some thermal overgy is lost

In many systems of interest, the system is a fluid or solid - or behavior is isovolumetric so that the change in stored energy is mostly internal and can be represented as \(\Delta e_{istored} = m CAT \) Constant pressure hearing

A
$$(b,U)$$
 $H_1 = U_1 + p_1 AL$

Now add heat

 $A \downarrow b_1 U_1'$
 $C \downarrow C \downarrow C \downarrow C$
 $C \downarrow C$
 C

$$AH = H_2 - H_1 = U_2 - U_1 + p_2 A (L + \Delta L) - p_1 AL \quad but \quad p_1 = p_2 \quad so$$

$$= U_2 - U_1 + p_1 A \Delta L$$

$$force * distance = Work$$

DH = DU + Work to expand

Constant Volume hearing

$$\Delta H = H_2 - H_1 = U_2 - U_1 + (p_2 - p_1)AL$$

$$\Delta H = \Delta U + \Delta p_1 AL$$
No work because $V = const.$

Heat transfer

Stored energy from heat transfer is mCAT; the expression assumes C~ constant and that there is no phase change.

Phases: gas, liquid, solid.

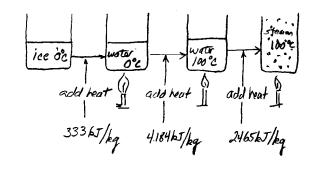
When a phase change occurs, energy is absorbed or released without a change in temperature (water freezes at 0°C; ice notes at 0°C)

(water boils at 100°C; steam condenses at 100°C)

constant pressure sp. treat required to change from solid to liquid (or reverse), is called the enthalpy of twion (latent heat of fusion).

Constant pressure sp. energy required to change from liquid to gas for reverse)

Is called the enthalpy of vaporization,



Total energy to convert

Iky 10e at 0°C to Iky steame 100°C

S33kJ/kg (ice to water)

418kJ/kg (water 0°C to water 100°C)

2465kJ/kg (water 100°C to steam 100°C)

3216 kJ/kg

To reverse process remove chergy in equal amounts

Phose change overgues can be represented as Dephase change = m L

Many problems involve the flow of both mass and energy across the system boundary. In such cases we might write estated = meat mishmass flow rate.

e has units of power (kJ/sec; Btu/hr; work force. distance; force. velocity)

A typical, overyy balance might be expressed as

(Uin + KEIN+PEIN) - (Vour + KEOUT + PEOUT) + HEATIN = AESYSTEM

Example: Coal fired 1000MW power plant. 33% officient. Remaining power last as host; 15% to atmosphere; 85% to cooling water. Cooling water from river Q = 100m3/s, T = 20°C. How much flow is directed from river if DT cooling water is 10°C? What is temp. of river after remixing

Permitting

heat Coal T=20 T=20heat

A electricity T=30 T=20 T=2

Overall energy balance

3000MW +
$$p(1-x)eT_{in}$$
 - 1000 MW - $p(1-x)eT_{out} = 300$ mW
2000MW - $p(1-x)eAT = 300$ mW

$$p(1-x)eAT = 1700$$
MW

$$(1-x) = \frac{1700 \cdot 10^6}{4184 \text{ T/sec}} = \frac{40.63}{1000} \text{ m}^3/\text{sec}$$

Downstream temperature

$$T=20\% \chi \qquad 100 \text{ m/s} \qquad \dot{e}_{10}-\dot{e}_{00}+\dot{e}_{10}+$$

$$6n = e_{out}$$

$$6x = 20^{\circ}C + 4(1-x) = 30^{\circ}C = 4100 = 59.47(20) + 40.63(30) = 24.1^{\circ}C$$

$$100$$

Second law

The second law of thermodynamics states that the rate of enough of entropy in the system plus net rate of entropy flow out of the system is at least equal to overall heat transfer across the boundary and the internal heat production.

Entropy is roughly analogous to the recriprocal of "useful toel".

As ontropy decreuses, a system can do more useful work. As entropy increases, a system does loss useful work and generates more waster heat.

Carnot cycle heat engine
$$\frac{T_{Heather}}{\sqrt{\frac{g_h}{Work}}}$$
 work $M = \frac{Work out}{Work} = 1 - \frac{T_c}{T_h}$

Maximum possible $\eta=1$ when $T_c\to0$ or $T_4\to\infty$. All real systems have $\eta<1$

· Component officiencies can be very high, but overall of is what counts
· waste heat is wasted \$1, but it a 1% increase in of costs more Moon

He & saved, He efficiency will not be improved (economics)
· waste is unavoidable - finding use for waste is beneficial and increases

overall efficiently.

Hart transfer concepts

hot (~~) ioje (

Two objects at different T. Heat will kanster from hot to cold object.

conduction - transfer by direct physical contact convection - transfer by host fluid that carries host

radiation - transfer by heat acting as e-m wave.