Question: "What is the difference between thermodynamics and a stick?

Answer: "A stick has two ends and no beginning. Thermodynamics has two 'beginnings' (the first and second law) and no end."

Leeture 2: - Review of Thermodynamics

Two fundamental assumptions behind thermodynamic treatment of a system.

(1) Continuum

2) Equilibrium (State) 0: So what does this mean for a fluid flow? Define

① Mechanical equilibrium (P)
② Thermal
① Chemical
① (N)

Thermodynamic
② quillibrium
② +② +③
② +③

P=p(+,T) Thermal eq" of state

e= e(+,T) Caloric eq" of state

Any functional form that satisfies Reciprocity relations is valid:

(A) Ideal / Perfect (B) Real gas. VP4 = RT High denisity gases Van-der Waals, P = 8 RT Gas Constent specific to a gas. a: Which model is valid for hypersonic flows? It can be shown experimentally OR from statistical mechanics that for ideal gas,

E = E(T)

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egas E = constant. T, then the is called calorically perfect

1 First law dE = da + dW Intensive de = dq + dw RHS -> Path variables LHS -> State variable So now, the question is, how de 9 calculate Algo,

how do 9 link Q/q x 1/2 to state variables?

-> Answer depends on the kind of process we are working with.

First law applied to a reversible

We know, de= dg -pd+ 9 défine C = dq. Specific heat

Since defining C for any two processes will yield C for all processes, 9 define,  $c_{v} = \frac{dq}{d\tau} |_{v}$ ,  $c_{p} = \frac{dq}{d\tau} |_{p}$ 

For a thermally perfect gas, Cp(T) - Cv(T) = R

In addition to E, we define enthalpy (H) as H = E + PY h = e + pt It can be shown that I'll aw can be enpressed as dh = dq + +dp h = h(T) under what assumptions? - Thermally prespect h= const. T => Glorically Adiabatic => Isentropic reversible  $\frac{S_{2}}{S_{1}} = \left(\frac{T_{2}}{T_{1}}\right)^{(1/r-1)}$  $\frac{\rho_2/\rho_1}{\rho_1} = \left(\frac{\tau_2/\tau_1}{\tau_1}\right)$