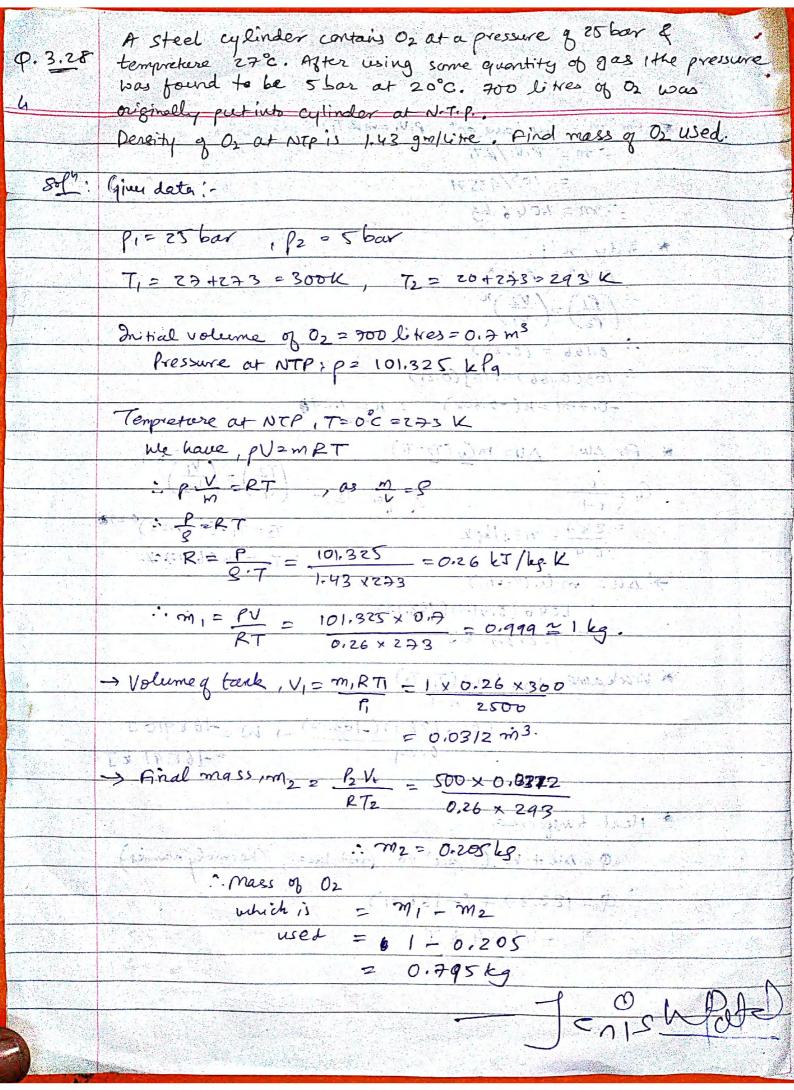
Properties q Gases + Numericals + 3 kg of ethane gas is compressed according to the law 3.14 'pv 1.3 = constant! from 1.013 box 27°c to 8 bar pressure. Defermine is heat transferred iii) work done (iii) Change in Internal energy. Molecular voeight of ethone is 30 belleg mod of (p=1.75 b) bgik for ethere. Assume ethane is a perfect gas [Ro = 8314.4 J/kg.mol K] sofn: given data: m=3/cp polytropic index n=1.3 P1 = 1.013 × 105 N/m2, P2 = 8×105 N/m2 Ti = 27° -> 273 +27 = 300 K M (moveular marght) = 30 kg/bgmod, Cp = 1.75 kJ/kg.K Ro= 8314.4 J/kg.molk. 4 0=3, W=?, DU=? 86 H. 1 V Sa. 077 - W.d * Work done for polytropic process:a w=mf(T1-T2) -) Fer t2, 1 T1 = (P2) 1 T. V AD. 5 N.E $\frac{72}{300} = \left(\frac{8 \times 10^{5}}{1.013 \times 10^{5}}\right)^{\frac{1.3}{1-3}}$ T2 = 300 (7.897) 0.231 T2 = 300 x1.6118 T2= 483.54 K - for Characteristic gas Constant; Ro= Ro = 8314.4 277.14 J/kg.K. = 0.277 LJ/kg.K

a	
	LIWE MRCTI-TE)
	· W = 3 x 0.277 (300 - 483,54)
	1.3-1
	50.(102 51)
guarge dang dankan d	0-3
Martin Service of the Service of Service of the Ser	= -508.4965
	· Also, Change in Internal energy;
	au= mCvaT o mCv(T2-T1)
Manufalia (graphing graphing region - world filth - should diswell him - work world in miles and an analysis of the contract o	= 3 (v (483.54-300)
	- 550.62 Gr
	- Prra :- we have go-are
	: Cu2 Q-R
	= 1.75-0,279 = 1.473 kJ/48.K
	-: DU = 550.62 x 1.4 73
	= 811.06 kJ
	+ Heat hosperred: Q = BUtW
	= \$11.06 - 508.40
	2 302.66 kJ
Q3.15	One cubic metre quir at pressure 1 bour \$ 600 is compressed
	final pressure 6 box & volume 0.25 m³. Determine in mass of (ii) indep 'n' for compression, (iii) Change in intermed energy (civ) heat transferred. CTake T=1.4 PR==877/bg
	(i) indep 'n' for compression, (iii) Change in internal energy
	civ) heat transferred. CTake T=1.4 & R= =875/43
sopu.	Wis 120 W 5 0 35 W3
	P1= 1×105 N/m2, P2= 6×105 N/m2, T1= 60+873= \$335
	4 m=7, n=7, DV=7, Q=7

-3.41	how ept. PIV, = mRTI.	
TI VALL	Porm'i we have ept. PIVIEMRTI.	4
	$= \frac{105}{95574}$	
	= () () / (
	:- m = 1.046 kg	
	er 122	
	$\frac{1}{2}\left(\frac{P_1}{Q_1}\right) = \left(\frac{V_2}{V_1}\right)^{N_1}$	
	$\frac{-\left(\frac{P_1}{P_2}\right) - \left(\frac{V_2}{V_1}\right)^{h}}{0.166} = \left(0.25\right)^{h}$	
	$0.166 = (0.25)^{n}$ $0.166 = 10.25)^{n}$ $0.166 = 10.25)^{n}$ $0.166 = 1.298$	
	$-0.799 = \mu(-0.602) \text{i. } \mu = 1.296$	
	-0.789 = x(-0.602) :. K=1.29	
	$G = \frac{R}{r+1} \qquad A = \frac{R}{r+1$	
	0 L To - (T) 2 (V2)	
	= 287 = 717. Thek : T2 = T1. (1/0.25)0-29	•
	= 287 2127/60	
	0.4 3 1/2 - 1/2 - 1/2 - 1/2 - 1/2 - 1/4 6 217 K	
	7. Bu = 41 00 ((2111)	
	122.37 EJ	
	* Workdone; -Wasma (TI-TZ) Man and January portales	
	n-1	
	0.29 =-168.91 k	7
	0.29 =-168.91 k	
	* fleat transferred;	
	0 = AU + W (Ac + to Gratians a Thermody namics)	
	· Q = 122.37 + (-164.91)	
	4-10-15 (-16 (191) - 1 dilling	
	2 -46.54 kJ	
	1	
1/1		
76		
		and the second second
mention of Carlotte State Section		CHERTY THE PART OF THE



, L		THERMODYNA	MIC PROCESSES	—Je	nie Water
Types of process	Governing	Heat interaction	Work done during state change from 1 to 2. W= Spd.V	Change in Internal energy	Change in Enthalpy AH
Isobaric 1) process P=constant	$\frac{T_2}{T_1} = \frac{V_2}{V_1}$ index $n = 0$	m Cp(T2-Ti)	$W = p(V_2 - V_1)$	Δ4= mQ (T2-Ti)	ΔH=m(p(T2-T1)
Isochoric process 2) Cloustant volume process)	$\frac{T_1}{T_2} = \frac{P_1}{P_2}$ $\text{index } m = \infty$	mGVCT2-TI)	W=0	DU=mCvCT2-TI)	ΔH=m(p(T2-T1)
Jso hermal process (Constant tempreture process)	$P_1V_1=P_2V_2$ index $n=1$	= $\rho_1 V_1 \log_e(\frac{V_2}{V_1})$ = $\rho_1 V_1 \log_e(\frac{\rho_1}{\rho_2})$	$W = P_1 V_1 \log_2 \left(\frac{V_2}{V_1} \right)$ $= P_1 V_1 \log_2 \left(\frac{P_1}{P_2} \right)$	∆u =0	ΔH2O
				1	

L						
	tabes of	Governing equations	Heat interaction	work done during state change from 1 to 2 W= \int^2 pdV	Change in Internal energy DU	change in Enthalpy DH
4)	ADIABATIC PROCESS	$P_1V_1^{\Upsilon} = P_2V_2^{\Upsilon}$ $\left(\frac{T_1}{T_2}\right) = \left(\frac{V_2}{V_1}\right)^{\frac{\gamma}{1}}$ $\frac{T_2}{T_1} = \left(\frac{P_2}{P_1}\right)^{\frac{\gamma}{1}}$ $\text{index } n = \Upsilon$	Ö	$W = P_1V_1 - P_2V_2$ $T - 1$ $also,$ $W = \frac{mR(T_1 - T_2)}{\Upsilon - 1}$	ΔU=m(v(T2-T1)	ΔH=m(p(T2-T1)
5	Polytro pic Process	$\rho_1 V_1 \stackrel{\mathcal{H}}{=} \rho_2 V_2 \stackrel{\mathcal{H}}{=} \left(\frac{T_1}{T_2} \right) = \left(\frac{V_2}{V_1} \right)^{m-1} \stackrel{\mathcal{H}}{=} \left(\frac{T_2}{T_1} \right) = \left(\frac{\rho_2}{\rho_1} \right)^{m-1} \stackrel{\mathcal{H}}{=} \left(\frac{T_2}{\rho_1} \right) = \left(\frac{\rho_2}{\rho_1} \right)^{m-1} \stackrel{\mathcal{H}}{=} \left(\frac{\rho_2}{\rho_1} \right)^{m-$	3	W= P1V1-P2V2	DU=mW(T2-T1)	ΔH=m(p(T2-T1)