

For numerical output of this problem, refer to Vol. II of this report filed in the Document Library of Ballistics Research Laboratories.

D. List of Labels

DECIDE

$DX(S)$ = Eulerian space interval in region S at $t = 0$

$$= L(S)/Z\emptyset N(S)$$

$H(S)$ = no. of cells from left boundary through region S

$$= \sum_{L=2}^S Z\emptyset N(L)$$

B5INIT(S)

$RH\emptyset(S)$ = density at zero pressure in region S

$A1, A2, A3$ = coefficients in Eq. (4.5)

$DV(S)$ = $v_2(p, T) - v_1(p, T)$

PM = pressure at which the Hugoniot in phase I intercepts
the phase boundary

$CV1$ = C_{v1}

$CVMIX$ = $C_{v,m}$

$GAMM1(S)$ = Γ

$E\emptyset$ = internal energy at the foot of the Hugoniot

$T\emptyset$ = T_0

$DPDTMX$ = $(\partial p / \partial T)_{v,m}$

$TAU\emptyset$ = $1/\tau$, Eq. (5.11)

VP = specific volume in phase I at $p = PM = v_1(pM, T)$

$V2$ = $v_2(pM, T)$

$CSPS$ = starting value for sound speed

J = index for space grid

$V(J)$ = v_j

$U(J)$ = U_j

$Q(J)$ = q_j

$P(J) = p_j$
 $TLIMA(J) =$ value of Δt_j for next time step
 $CSP(J) =$ sound speed in cell J
 $E(J) = E_j$
 $ENT(J) = s_j$
 $TMP(J) = T_j$
 $NSA(J) =$ switching index
 = 1, phase I
 = 2, mixed phase
 = 3, phase II

MAIN

$X(J) = x_j$ (Fig. 5.2)
 $MASS(J) =$ mass of cell J
 $JSTAR =$ cell label just ahead of shock front at which
 computation stops for each time cycle
 $TIMES = t$
 $CYCLE =$ number of times t has been incremented
 $JCRIT =$ value of J for which $TLIMA$ is minimum
 $LAST =$ switching index for halting program after
 writing last output.
 $PPEAK =$ maximum computed pressure in each cycle
 $TLIMB = TLIMA(JCRIT)$
 $PLEFT =$ pressure applied to left boundary
 $DFNU =$ mass in cell J+1
 $XA = x(t + \Delta t)$
 $VN = v(t + \Delta t)$
 $QA = Q(t + \Delta t)$
 $JPMAX =$ value of J at which p is maximum

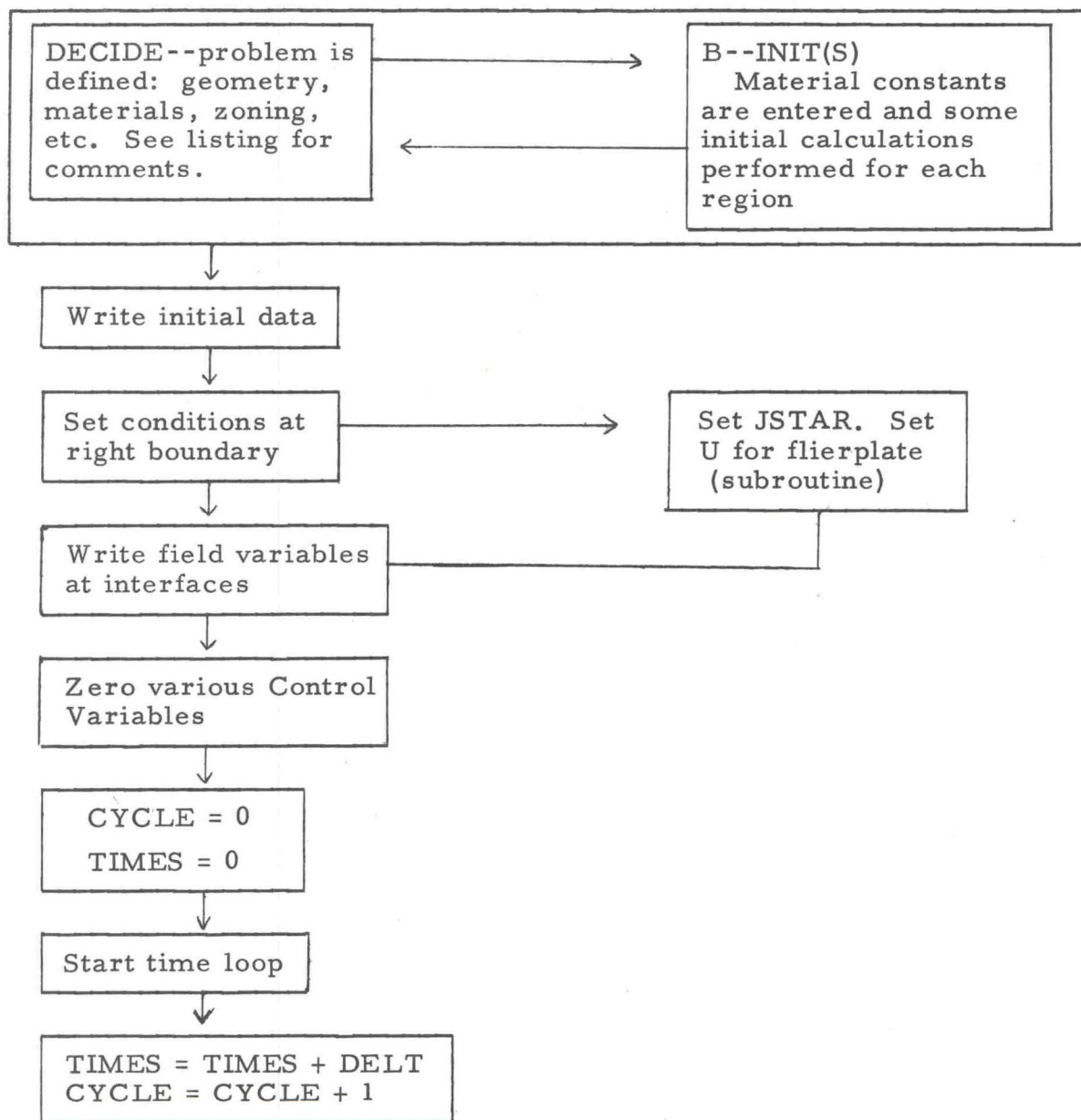


Fig. 1.--FLOW CHART FOR BURN

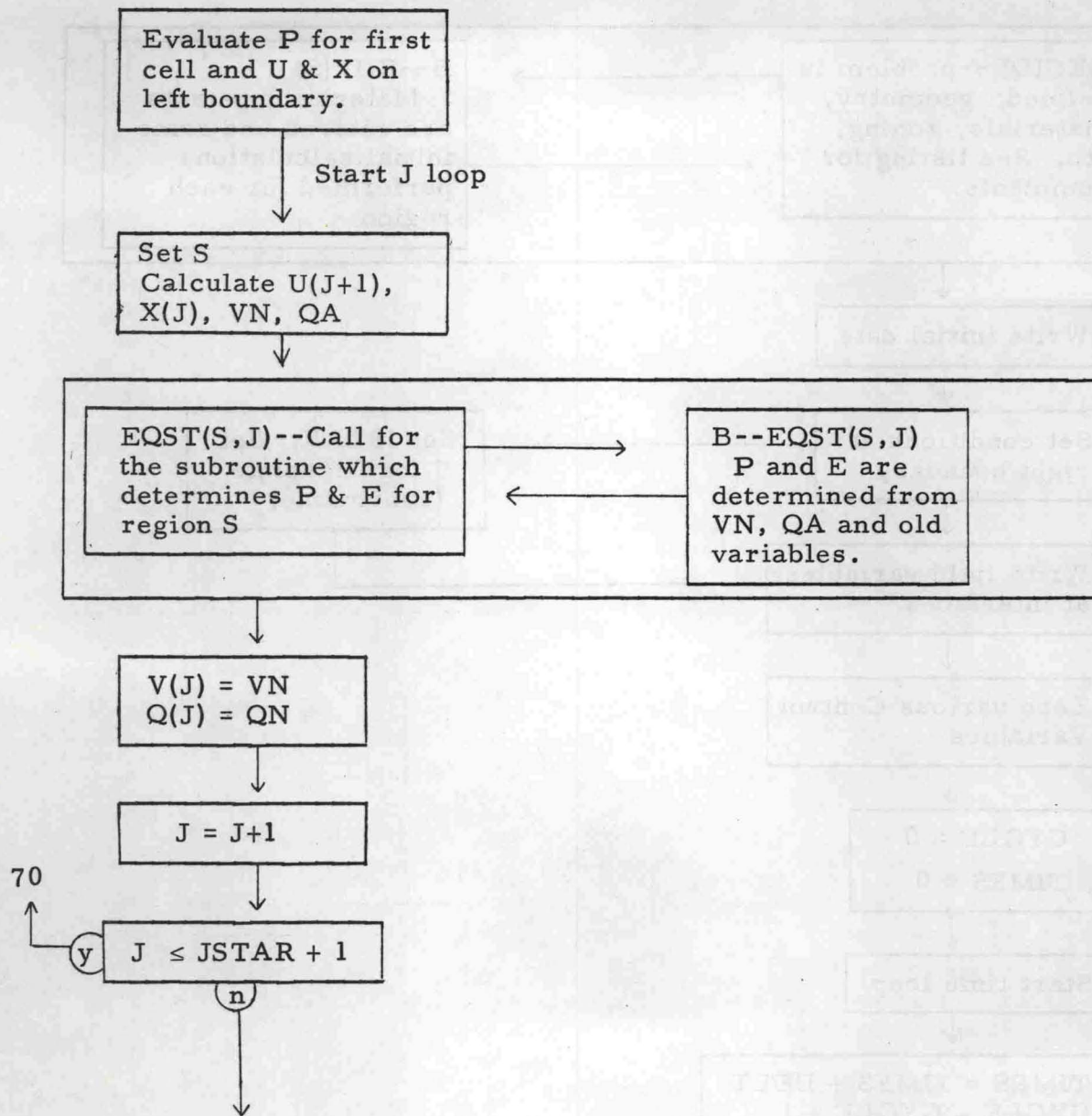


Fig. 1. (b)

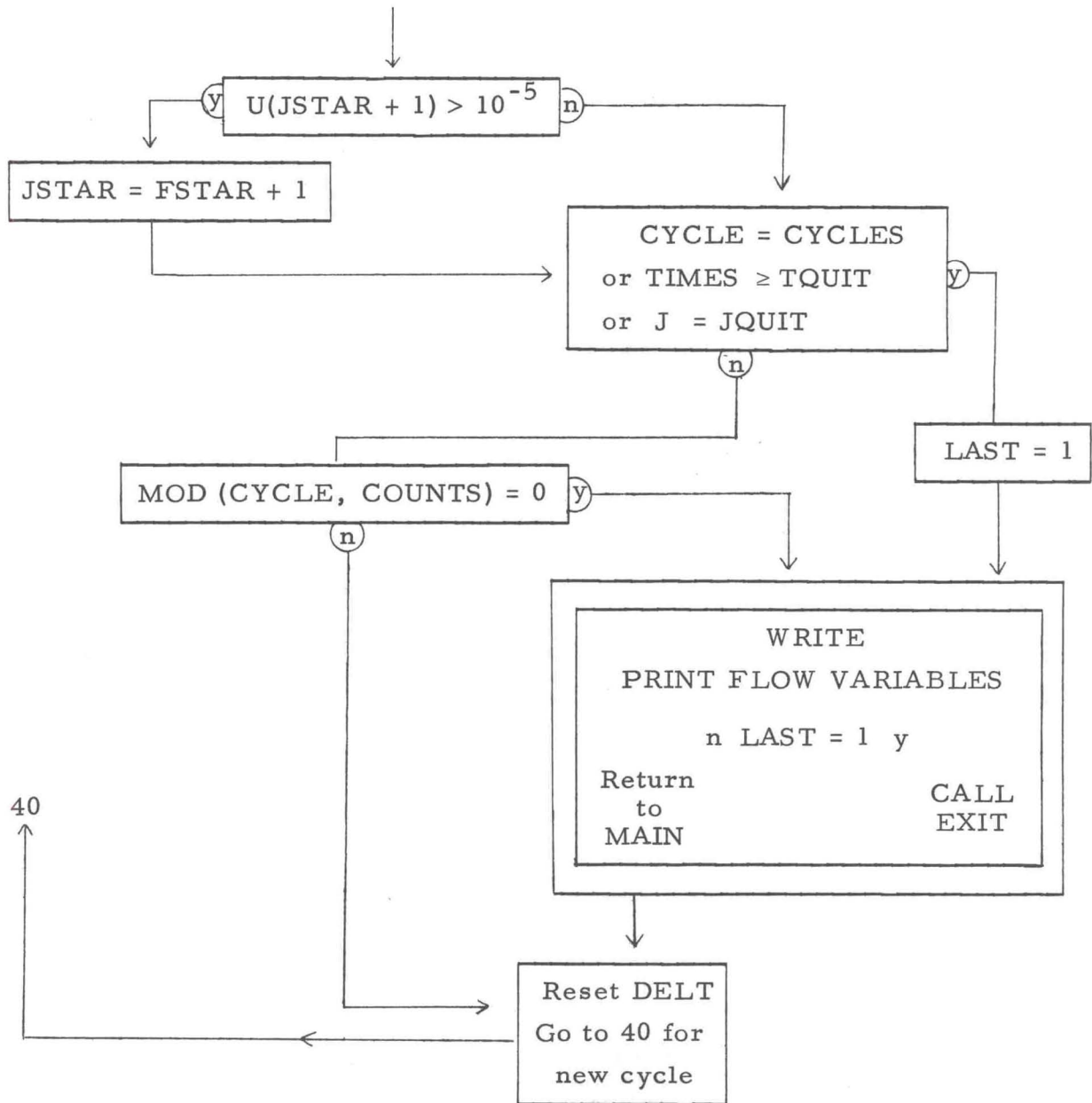


Fig. 1. (c)

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13. ABSTRACT A procedure is described for developing simple approximate equations of state of liquids from Hugoniot P-V relations determined in shock wave measurements. This is applied to a number of liquids and a table of coefficients is given. The formalism of irreversible thermodynamics is applied to time-dependent phase transitions in iron and an approximate set of constitutive relations is obtained in a form suitable for numerical integration with the equations of continuum dynamics. These are applied in an approximate form to study the development of the two-wave structure in iron caused by the α - ϵ phase transition. Finite strain theory is applied to the analysis of shock wave data for quartz, and the results supply enough information to estimate some of the fourth-order elastic constants.		

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