

BULK MODULI OF SOLIDS IMPACTED BY SHOCKWAVES

TABLE II. Sound velocities and compressibilities.

Specimen material	ρ_0 (g/cm ³)	C_0 (m/sec)	Intercept U_s (m/sec)	Longitudinal Sound velocity (m/sec)	Incident $U_s(H_2O)$ (m/sec)	Transmitted $U_s(H_2O)$ (m/sec)	K (cm ² /dyn)
Brass (Cu Zn)							
36 at.% Zn	8.40	3815	3860	4700 ¹²	1493	1479	0.08×10^{-11}
Aluminum	2.70	5454	5350	6420 ¹²	1480	1476	0.13×10^{-11}
Boron Nitride	2.20	2628	2831		1501	1496	0.66×10^{-11}
Vinylidene Chloride	1.60	2690			1473	1466	0.86×10^{-11}
TNT	1.61	2574	2390	2586 ± 80^{13}	1495	1463	0.94×10^{-11}
Plexiglas	1.18	2760	2710	2770^8 2650^{10}	1497	1492	1.11×10^{-11} $1.3 \times 10^{-11 9}$ $1.7 \times 10^{-11 10}$ $2.5-3.0 \times 10^{-11 11}$
Nylon	1.15	2616	2300	2650^8	1474	1480	1.27×10^{-11}
Epoxy-2	1.20	2535			1476	1458	1.30×10^{-11}
Epoxy-3	1.12	2402			1498	1463	1.55×10^{-11}
Epoxy-1	1.18	2254	2240		1500	1475	1.67×10^{-11}
Epoxy-4	1.16	2098			1490	1468	1.96×10^{-11}
Teflon	2.03	1453	1991	1340 ± 47^8	1494	1467	2.33×10^{-11}

this method, intense light is obtained from the explosion of a 5.1-cm-long 0.012-cm-diam tungsten wire threaded within a 1-mm-bore capillary tube. This light is collimated by a field lens positioned behind the aquarium. The shock motion is delineated by the disturbance it produces in the light that reaches the film through the smear-camera slit. The tungsten wire is exploded by the discharge of a 4- μ F capacitor charged to 4000 V. The backlighting is usually delayed to occur $\sim 150 \mu$ sec after the initiation of the incident shock from the detonator. A camera writing speed of ~ 1.3 mm/ μ sec is best for recording wave propagation in most liquids, using the above experimental geometry. Figure 2 is a typical record resulting from the experimental arrangement. With proper interpretation, it allows measuring the velocity of the incident wave striking the specimen, the transit time of the wave⁶ in the specimen, and the velocity of the emerging wave in the water below.

II. RESULTS AND DISCUSSION

Table II gives the results obtained for several materials and lists the density, ρ_0 ; the sound velocity in the specimen, C_0 ; the velocity of the incident wave in the water, Incident $U_s(H_2O)$; and the velocity of the wave transmitted from the specimen to the water below, Transmitted $U_s(H_2O)$. Table II also contains values of the reciprocal of the bulk modulus, i.e., the initial compressibility, K , determined for $V=V_0$ in Eq. 16, and experimental values of the intercept of the linear U_s-u_p curve.

The combined errors produced by camera speed and magnification ratio measurements result in an error in the measured velocities not exceeding 0.3%. The velocity data listed are the average of three experiments run with each specimen material, except TNT and

⁶ The wave propagation in transparent material, e.g., Plexiglas, is made visible by this technique.

polymethyl methacrylate (Plexiglas), for which five experiments each were conducted. The maximum mean velocity deviation was $\pm 0.6\%$, and this is considered the maximum error to be expected in the measurement of velocity.

A temperature variation of $\pm 2^\circ\text{C}$ was possible during these determinations, principally because the experiments were conducted in bomb-proof facilities at times when temperature control was difficult to maintain. However, large errors due to temperature variations were avoided by adjusting the water temperature in the aquariums to 20°C at the time of firing.

The average values of the incident water-wave velocity and transmitted water-wave velocity in Table I are ~ 1490 and ~ 1470 m/sec, respectively. The velocity of sound⁷ in distilled water at 20°C is 1481.63 m/sec. Therefore, considering the errors in measuring the weak

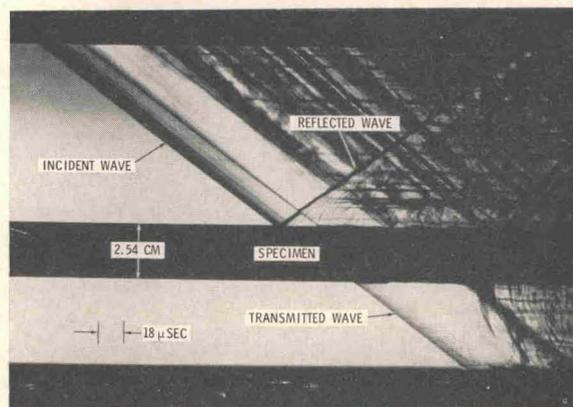


FIG. 2. Smear-camera shadowgraph of weak shock waves in water, obtained using the arrangement in Fig. 1. The upper and lower traces, respectively, are from the incident and transmitted waves in the water. The central dark bend denotes the specimen material.

⁷ W. D. Wilson, J. Acoust. Soc. Amer. 31, 1067-1072 (1959).

shock propagation, the waves entering and leaving the specimen material may be treated as sonic. This view is supported by pressure measurements using piezoelectric gauges, which indicate that the incident pressure at the water-specimen interface does not exceed 50 bars.

For Plexiglas, the weak shock-velocity data compare favorably with ultrasonic measurements of longitudinal waves. For example, Auberges and Reinhart⁸ give 2770 m/sec as compared to the weak shock measurement of 2760 ± 15 m/sec. From this measurement, we find $k = 1.11 \times 10^{-11}$ cm²/dyn. Kolsky's⁹ results from pressure bar experiments give $k = 1.3 \times 10^{-11}$. Wada's¹⁰ measurements of the velocities of longitudinal waves (2650 m/sec) and transverse waves (1240 m/sec) at 1.08×10^6 Hz give $k = 1.7 \times 10^{-11}$ cm²/dyn. Pullen *et al.*¹¹ find much greater compressibility for molded Plexiglas specimens, with K ranging from 2.5×10^{-11} cm²/dyn to 3.0×10^{-11} cm²/dyn. These discrepancies in the compressibility and the close agreement of the weak shock measurement to the ultrasonic value for longitudinal waves in nylon, TNT, and Teflon in (Table II), indicate the method does actually give values associated with longitudinal waves. The C_0 data of Table II then would be 10%–30% greater than the bulk sound velocities calculated from the elastic constants (see Eq. 14). This would, of course, result in erroneous values of the initial dynamic compressibility in Table II, compared to static data.

However, the conditions of shock propagation are not isentropic. Dynamic compressibilities may be different from the corresponding static values obtained under equal pressures. Also, dynamic and static yield strengths of a material do not necessarily have the same value. For solids with significant shear strength, which can contribute to the material stiffness, the values of C_0 obtained by the shock-attenuation technique represent elastic-wave compressions and should agree with the velocities of low-pressure longitudinal waves. At the critical stress (the Hugoniot elastic limit), the specimen

material passes from elastic compression behavior to plastic- or fluidlike flow. In reaching this point, the elastic-wave data in the shock-velocity-particle-velocity plane follow a relation with its limiting wave velocity the zero-pressure longitudinal sound speed. For the low-amplitude limit of the plastic wave, the limiting velocity is the bulk sound speed. For aluminum, however, the measured C_0 value in Table I is 5454 m/sec. This value is 15% less than the longitudinal sound velocity, (6420 m/sec), but it exceeds the ultrasonic bulk sound velocity (5375 m/sec) by only 1.5% and probably indicates the shock in aluminum attenuated to a value close to the yield stress. Similar results are obtained for brass (Cu Zn alloy containing 36 at.% Zn) as compared to the ultrasonic data available only for a Cu Zn alloy¹² containing 30 at.% Zn.

The C_0 value for Teflon, however, differs considerably ($\sim 30\%$) in Table II from the intercept U_s , which was obtained by a linear extrapolation of shock data in the 25-kbar to 290-kbar range. This result indicates the U_s-u_p relation for Teflon may not be linear and the possibility of a phase change exists. Several investigators including Bridgman¹³ have reported that Teflon undergoes a transition at 6.5 kbar. For the other pressed materials in Table II, e.g., TNT,¹⁴ BN,¹⁵ and vinylidene chloride, where the incident weak shock stress approaches the yield strength, the velocities measured apparently are close to the bulk sound velocities and the compressibilities in Table II are probably accurate values.

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¹² *American Institute of Physics Handbook* (McGraw-Hill Book Co., New York, 1957).

¹³ J. B. Ramsay and A. Popolato, 4th Symp. on Detonation, U. S. Naval Ordnance Lab., Silver Spring, Md., 4th, (Oct. 1965).

¹⁴ P. W. Bridgman, Proc. Amer. Acad. Arts Sci. **76**, 71–80 (1948).

¹⁵ Boron nitride transforms under shock compression at 122 kbar. Its U_s-u_p data are related linearly below the transition pressure. See N. L. Coleburn and J. W. Forbes, J. Chem. Phys. **48**, 555–559 (1968).

⁸ M. Auberges and J. S. Reinhart, J. Appl. Phys. **32**, 219–222 (1961).

⁹ H. Kolsky, Proc. Soc. **B62**, 676–700 (1949).

¹⁰ Y. Wada, J. Phys. Soc. Japan, **13**, 1390–1398 (1958).

¹¹ W. J. Pullen, J. Roberts, and T. E. Whall, Polymer **5**, 471–478 (1964).