

HEATING VENTILATING AIR CONDITIONING GUIDE 1944

TABLE 2. SPECIFIC VOLUME OF DRY AIR^a at 29.921 in. Hg

TEMP F <i>t</i>	CU FT PER LB <i>v_a</i>	TEMP F <i>t</i>	CU FT PER LB <i>v_a</i>	TEMP F <i>t</i>	CU FT PER LB <i>v_a</i>
-96	9.1488	32	12.3888	160	15.6229
-64	9.9597	64	13.1977	192	16.4310
-32	10.7699	96	14.0063	224	17.2389
0	11.5796	128	14.8147	256	19.0467

^aPrepared by John A. Goff.

energy of the molecules. In ordinary calculations not requiring too great accuracy, the effect of intermolecular forces may be ignored and Equation 2 simplified to

$$pv = BT \quad (1)$$

Example 2. Calculate an approximate value for the specific volume of dry air at 64 F, 29.921 in. Hg.

$$\text{Solution. } v = \frac{53.351 \times 523.70}{29.921 \times 0.49115 \times 144} = 13.200 \text{ cu ft per pound.}$$

Note: This answer may be compared with the value in Table 2. The difference is due to intermolecular forces. It should not be concluded, however, that because the effect of intermolecular forces on the volume is so small these forces can be ignored entirely.

The relationship of Equation 1 expresses certain familiar laws approximately true for gases at not too high pressures. Thus, *with temperature constant, the volume of a given weight of gas is inversely proportional to its absolute pressure* is a statement of Boyle's Law. If v_1 denotes the specific volume at absolute pressure p_1 , then *at the same temperature*, the specific volume v_2 at absolute pressure p_2 is approximately,

$$v_2 = v_1 \left(\frac{p_1}{p_2} \right)$$

Also, *with pressure constant the volume of a given weight of gas is directly proportional to its absolute temperature* is a statement of Charles' Law. If v_1 denotes the specific volume at absolute temperature T_1 , then *at the same pressure*, the specific volume v_2 at absolute temperature T_2 is approximately,

$$v_2 = v_1 \left(\frac{T_2}{T_1} \right)$$

Absolute Temperature

For the range 0 to 660 C, the standard temperature scale is the International Centigrade Scale, namely, the readings of a platinum resistance thermometer standardized at the ice-point (0 C), the steam-point (100 C) and the sulphur-point (444.60 C). The corresponding Fahrenheit scale t used in scientific work is derived from the International Centigrade Scale by means of the relation,

$$t = 1.8 (\text{Int. Cent. Temp.}) + 32 \quad (3)$$

Temperatures on the absolute Fahrenheit scale are then obtained by adding 459.70 according to the equation

$$T = t + 459.70 \quad (4)$$

CHAPTER 1. THERMODYNAMICS OF AIR AND WATER MIXTURE

Absolute temperatures computed from Equations 3 and 4 are *practically* identical with the fundamental thermodynamic temperatures to which the zero-pressure values of the product pv for gases are proportional in accordance with Equation 2.

Specific Enthalpy

Most air conditioning processes are of the steady-flow type. In steady flow the energy convected with the fluid crossing a given section is the sum of (a) kinetic energy due to velocity, (b) gravitational energy due to elevation, (c) enthalpy due to the condition of temperature, pressure and composition at a given section. It is clear, therefore, that in order to apply the Law of Conservation of Energy to steady-flow processes, information regarding the enthalpy is needed.

Recent developments in quantum mechanics have made it possible to calculate the zero-pressure specific enthalpy of a gas from spectroscopic measurements, and with a degree of accuracy exceeding that with which this property can be inferred from direct calorimetric measurements. Available data for each gas listed in Table 1 have been assembled and critically examined; and from them have been calculated best values for the specific enthalpy of dry air at zero pressure. These are listed in Table 3. The unit of energy is the Btu which is related to the foot-pound as follows:

$$1 \text{ Btu} = 778.18 \text{ ft-lb}^1 \quad (5)$$

In Table 3 are also listed values of *mean zero-pressure specific heat* for the range 0 to t F. This is simply the increase of specific enthalpy from 0 to t F, divided by the increase of temperature or, with 0 F as the reference point, by the temperature itself. The numerical values indicate that a rounded figure of 0.24 Btu per pound can be used in ordinary calculations.

Applying well known identical relations of thermodynamics to Equation 2, the following expression for specific enthalpy, valid at not too high pressures, is obtained

$$h = h^0 + \left[T^2 \frac{d(A/T)}{dT} \right] p \quad (6)$$

where h^0 denotes specific enthalpy at zero pressure. This equation emphasizes that the effect of pressure on specific enthalpy is not so much due to the second virial coefficient A itself as to its variation with tempera-

TABLE 3. SPECIFIC ENTHALPY OF DRY AIR AT ZERO PRESSURE^a

TEMP F <i>t</i>	SPECIFIC ENTHALPY BTU PER LB <i>h_a⁰</i>	MEAN SPECIFIC HEAT [<i>C_p⁰</i>] _{<i>t</i>}	TEMP F <i>t</i>	SPECIFIC ENTHALPY BTU PER LB <i>h_a⁰</i>	MEAN SPECIFIC HEAT [<i>C_p⁰</i>] _{<i>t</i>}	TEMP F <i>t</i>	SPECIFIC ENTHALPY BTU PER LB <i>h_a⁰</i>	MEAN SPECIFIC HEAT [<i>C_p⁰</i>] _{<i>t</i>}
-96	-22.839	0.2393	32	7.796	0.2395	160	38.529	0.2400
-64	-15.186	0.2393	64	15.466	0.2396	192	46.238	0.2401
-32	-7.529	0.2394	96	23.145	0.2397	224	53.962	0.2403
0	+0.131	0.2394	128	30.831	0.2398	256	61.702	0.2405

^aPrepared by John A. Goff from published data computed from spectroscopic measurements.

¹This conversion factor is not exact by definition, but involves an experimental determination of the relation between the absolute and the standard electrical units of energy. The value 1 int. joule = 1.00019 abs. joule, recommended by Osborne, Stimson and Ginnings [8] was used.