

# Humid air and condensation

*The first documented study of humid air seems to be the description made by Charles Leroy (1751), a physician from Montpellier (France). He explained in 1751 to the Royal Academy of Sciences that water can be dissolved in air as a function of air temperature, with the higher temperature corresponding to greater dissolution. To support his claim, he described several experiments. The most demonstrative one involves a bottle of air closed at daytime temperature. Once cooled at night, the air becomes unable to retain all the water dissolved at the higher temperature of the day: the excess water leads to the formation of condensed droplets that are clearly visible inside the bottle.*

*Indeed, air is never completely dry; it always contains, in an invisible way, water vapor whose concentration varies according to its temperature. In addition to steam, humid air can also contain water in a visible condensed state: liquid (fog droplets), solid (frosted fog). In these last cases, where vapor and condensed phases coexist, the humid air is said to be supersaturated.*

*We can therefore consider that humid air is formed of (1) dry air unlikely to condense under the temperature and pressure conditions considered here, and (2) water vapor likely to condense into liquid or ice. Dry air is composed primarily of nitrogen ( $\approx 78\%$ ) and oxygen ( $\approx 21\%$ ). For usual conditions of temperature and pressure encountered at the earth's surface, both gases are far from their critical point coordinates and both fluids can be accepted as perfect gases. Air will therefore be considered as a single perfect gas. Water is also far from its critical point coordinates and can also be considered as a perfect gas.*

## 1. Partial pressures: Dalton's law

The partial pressure of a gas is the pressure that the gas would have if it were alone in a volume  $V$ . Dry air and water being both ideal gases, the total pressure is equal to the sum of the partial pressures: It is Dalton's law. With  $p_a$  (resp  $p_v$ ) the partial pressure of air (resp. water) and  $p_m$  the total pressure, we obtain  $p_m = p_a + p_v$ . This rule of additivity corresponds to neglect the intermolecular forces between the gas molecules. The pressure being due to the impacts of the gas molecules in movement, the total pressure is simply the addition of the impacts of each type of molecules.

## 2. Equation of state

The equation of state for dry air and water vapor is the perfect gas equation  $p_i V = n_i R T$  where the subscript  $i$  represents air ( $i=a$ ) or water vapor ( $i=v$ );  $n_i = m_i / M_i$  is the number of moles ( $i$ ) of molar mass  $M_i$  and mass  $m_i$  in a volume  $V$ ,  $R = 8.314 \text{ J.mole}^{-1}.\text{K}^{-1}$  is the molar constant of gases. The properties of moist air at a given partial pressure of water vapor can be deduced from this simple equation of state.

## 3. Saturation ; condensation

Consider the process of cooling at constant pressure  $p_m$  of a mass of moist air that contains a given mass of water. The conservation of mass requires that the total mass and the mass of water vapor remain constant during the process. The same applies to the number of moles of water vapor and the corresponding mole fraction  $n_v / n = p_v / p_m$  (from the perfect gas equation above and  $n = n_a + n_v$ ). As a result, the pressure of the water vapor remains constant during the cooling process. In the atmosphere, the cooling of moist air therefore occurs **at constant water vapor pressure**.

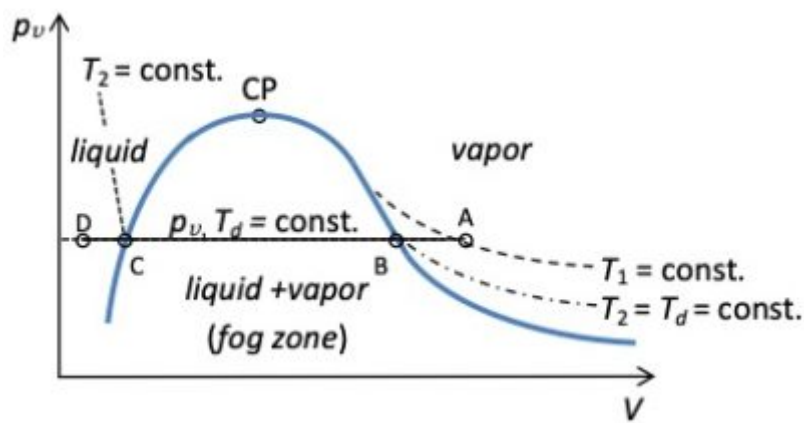


Figure 1. cooling at constant pressure  $p_v$  in the Clapeyron phase diagram. CP: Critical point; B: Dew point;  $T_d$ : Dew point temperature where  $p_v(T_1) = p_s(T_d)$ , the saturated vapor pressure at temperature  $T_d$ . [Source: after Beysens, D. (2018). Dew water. Gistrup: Rivers Publisher]

During cooling, condensation to liquid can occur (see Clapeyron phase diagram in Figure 1). Consider a mass of moist air initially at point A on the  $T_1$  isotherm. When the temperature decreases at constant pressure  $p_v(T_1)$ , its volume also decreases. The liquid-vapor coexistence curve (the saturation curve) is reached (point B) at a certain temperature  $T_2$  and drops of liquid may appear. Point B is called the **dew point** and  $T_2$  is the **dew point temperature  $T_d$** . The vapor pressure is the **saturation pressure** at temperature  $T_d$ ,  $p_s(T_d) = p_v(T_1)$  where condensation of water vapor can begin. When the air is cooled further, condensation continues at constant pressure  $p_v$  and temperature  $T_d$ . The cooling energy only compensates for the release of the latent heat of condensation (Read: [Pressure, Temperature and Heat](#)). In C, all the water contained in the humid air has condensed. The BC zone is the fog zone where liquid droplets coexist with the vapor. Further cooling (to point D) involves only the liquid.

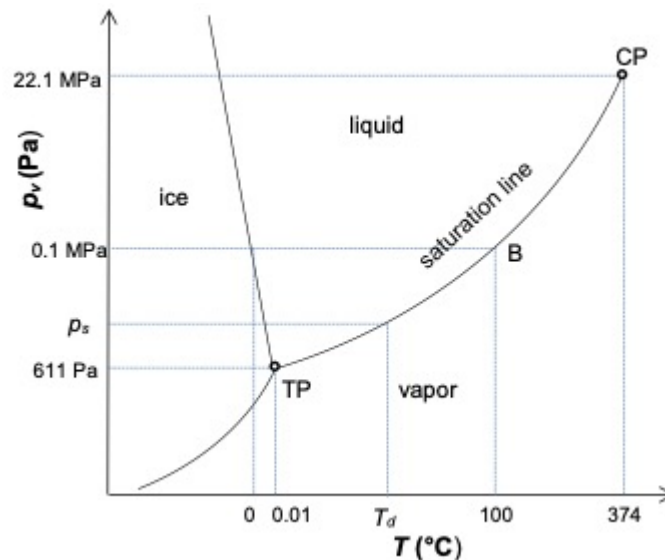


Figure 2. Phase diagram of water highlighting the saturation line  $p_s(T_d)$ . CP: Critical point; TP: Triple point. B: Dew point. [Source: Based on Beysens, D. (2018). dew water. Gistrup: Rivers Publisher]

The liquid-vapor **saturation curve** represents in the  $p_v - T$  plane the liquid-vapor equilibrium (Figure 2). At a given temperature, the maximum pressure above which water vapor turns into liquid water is the saturated vapor pressure  $p_s$ . Therefore, in a given mass of moist air, the vapor pressure can be such that (i)  $p_v < p_s$ : the water in the moist air is in the vapor state; (ii)  $p_v = p_s$ : the water in the moist air is both in the vapor and liquid state because the phase change occurs at constant pressure  $p_s = p_v(T_d) = p_v(T)$ .

Moist air can be saturated (point B in Figure 1) or supersaturated, when liquid droplets (mist) are present (line BC in Figure 1).

Saturated vapor pressure can then be achieved in a given moist air in two ways. (i) By cooling a given mass of moist air: the vapor pressure remains constant at  $p_v$  but  $p_s$  decreases until equality  $p_v = p_s(T_d)$  is satisfied. (ii) Adding a mass of water to a given

volume of humid air at constant temperature: the vapour pressure increases until it reaches  $p_v=p_s$  at the same temperature. If we add more water at constant temperature, we obtain the coexistence of the saturated vapor pressure and the liquid. The humid air is then **supersaturated**.

From the above, we can define the **relative humidity RH** as the ratio, at a given temperature, between the vapor pressure and the saturated vapor pressure,  $RH = p_v(T)/p_s(T)$ . When a given mass of moist air is cooled, RH increases to 100% at the saturation line. When water is added at constant temperature, RH also increases to 100% and reaches the saturation line. Relative humidity is a common index for determining how close a given humid air is to saturation. The higher the RH, the less cooling or mass addition required to achieve condensation.

## Notes

**Cover image.** [Source: royalty free]

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