This is a formal presentation of the formulæ that describe those properties of water vapour in air that concern conservators. The derivation of these equations is explained concisely and the useful equations are emphasised by bold type. This datasheet is intended for reference, not as a friendly introduction to the topic!

## Water vapour pressure

In a closed container partly filled with water there will be some water vapour in the space above the water. The concentration of water vapour depends only on the temperature. It is not dependent on the amount of water and is only very slightly influenced by the presence of air in the container.

The water vapour exerts a pressure on the walls of the container. The empirical equations given below give a good approximation to the saturation water vapour pressure at temperatures within the limits of the earth's climate.

Saturation vapour pressure, $p_{s}$, in pascals:
$p_{s}=\mathbf{6 1 0 . 7 8} \times \exp (t /(t+238.3) \times \mathbf{1 7 . 2 6 9 4})$
where $\mathbf{t}$ is the temperature in degrees Celsius
Reference: Tetens, O., 1930: Uber einige meteorologische Begriffe. Zeitschrift fur Geophysik, Vol. 6:297. There are more accurate formula given in standard reference works, such as the Smithsonian Tables, but this version is adequate for all but high accuracy laboratory studies.

The svp below freezing can be corrected after using the equation above, thus:
$p_{\mathrm{s}}$ ice $=-4.86+\mathbf{0 . 8 5 5} \mathrm{p}_{\mathrm{s}}+\mathbf{0 . 0 0 0 2 4 4} \mathrm{p}_{\mathrm{s}}{ }^{2}$
The next formula gives a direct result for the saturation vapour pressure over ice:
$p_{s}$ ice $=\exp (-6140.4 /(273+t)+28.916)$
The dew point can be calculated from the actual vapour pressure (vp) by using the inverse of Tetens' equation:
$\mathrm{dp}=(\mathbf{2 4 1 . 8 8} \times \ln (\mathrm{vp} / \mathbf{6 1 0 . 7 8})) /(\mathbf{1 7 . 5 5 8}-\ln (\mathrm{vp} / \mathbf{6 1 0 . 7 8})$
The pascal is the SI unit of pressure $=$ newtons $/ \mathrm{m}^{2}$. Atmospheric pressure is about 100,000 Pa (standard atmospheric pressure is defined as $101,300 \mathrm{~Pa}$ ).

## Water vapour concentration

The relationship between vapour pressure and concentration is defined for any gas by the equation:
$\mathrm{p}=\mathrm{nRT} / \mathrm{V}$
p is the pressure in $\mathrm{Pa}, \mathrm{V}$ is the volume in cubic metres, T is the temperature in degrees Kelvin (degrees Celsius +273.16 ), n is the quantity of gas expressed in molar mass ( 0.018 kg in the case of water ), R is the gas constant: $8.31 \mathrm{Joules} / \mathrm{mol} / \mathrm{m}^{3}$

To convert the water vapour pressure to concentration in $\mathrm{kg} / \mathrm{m}^{3}:(\mathrm{Kg} / 0.018) / \mathrm{V}=\mathrm{p} / \mathrm{RT}$ $\mathbf{k g} / \mathbf{m}^{\mathbf{3}}=\mathbf{0 . 0 0 2 1 6 6} \times \mathbf{p} /(\mathbf{t}+\mathbf{2 7 3 . 1 6})$ where $p$ is the actual vapour pressure

## Relative Humidity

The Relative Humidity ( RH ) is the ratio of the actual water vapour pressure to the saturation water vapour pressure at the prevailing temperature.
$\mathbf{R H}=\mathbf{p} / \mathbf{p}_{\text {s }}$
RH is usually expressed as a percentage rather than as a fraction. In the biological literature, however, the RH is often expressed as a fraction and is then called the water activity.

The RH is a ratio. It does not define the water content of the air unless the temperature is given. The reason RH is so much used in conservation is that most organic materials have an equilibrium water content that is mainly determined by the RH and is only slightly influenced by temperature.

Notice that air is not involved in the definition of RH. Airless space can have a RH. Air is the transporter of water vapour in the atmosphere and in air conditioning systems, so the phrase "RH of the air" is commonly used, and only occasionally misleading. The independence of RH from atmospheric pressure is not important on the ground, but it does have some relevance to calculations concerning air transport of works of art and conservation by freeze drying.

## The Dew Point

The water vapour content of air is often quoted as dew point. This is the temperature to which the air must be cooled before dew condenses from it. At this temperature the actual water vapour content of the air is equal to the saturation water vapour pressure. The dew point is usually calculated from the RH. First one calculates $\mathbf{p}_{s}$, the saturation vapour pressure at the ambient temperature. The actual water vapour pressure, $\mathbf{p}_{\mathrm{a}}$, is:
$\mathbf{p}_{\mathrm{a}}=\mathbf{p}_{\mathrm{s}} \times \mathbf{R H} \% / 100$
The next step is to calculate the temperature at which $\mathbf{p}_{\mathrm{a}}$ would be the saturation vapour pressure. This means running backwards the equation given above for deriving saturation vapour pressure from temperature:

Let $\mathrm{w}=\ln \left(\mathrm{p}_{\mathrm{a}} / 610.78\right)$
Dew point $=w \times 238.3 /(17.294-w)$
This calculation is often used to judge the probability of condensation on windows and within walls and roofs of humidified buildings.

The dew point can also be measured directly by cooling a mirror until it fogs. The RH is then given by the ratio

RH $=100 \times \mathbf{p}_{\text {s }}$ dewpoint $/ \mathbf{p}_{\mathrm{s}}$ ambient

## Concentration of water vapour in air

It is sometimes convenient to quote water vapour concentration as $\mathrm{kg} / \mathrm{kg}$ of dry air. This is used in air conditioning calculations and is quoted on psychrometric charts. The following calculations for water vapour concentration in air apply at ground level.
Dry air has a molar mass of 0.029 kg . It is denser than water vapour, which has a molar mass of 0.018 kg . Therefore, humid air is lighter than dry air. If the total atmospheric pressure is P and the water vapour pressure is p , the partial pressure of the dry air component is $\mathrm{P}-\mathrm{p}$. The weight ratio of the two components, water vapour and dry air is:
kg water vapour $/ \mathrm{kg}$ dry air $=0.018 \times \mathrm{p} /(0.029 \times(\mathrm{P}-\mathrm{p}))$

$$
=0.62 \times p /(P-p)
$$

At room temperature P - p is nearly equal to P , which at ground level is close to $100,000 \mathrm{~Pa}$, so, approximately:
kg water vapour $/ \mathrm{kg}$ dry air $=0.62 \times 10^{-5} \times \mathrm{p}$

## Thermal properties of damp air

The heat content, usually called the enthalpy, of air rises with increasing water content. This hidden heat, called latent heat by air conditioning engineers, has to be supplied or removed in order to change the relative humidity of air, even at a constant temperature. This is relevant to conservators. The transfer of heat from an air stream to a wet surface, which releases water vapour to the air stream at the same time as it cools it, is the basis for psychrometry and many other microclimatic phenomena. Control of heat transfer can be used to control the drying and wetting of materials during conservation treatment.
The enthalpy of dry air is not known. Air at zero degrees celsius is defined to have zero enthalpy. The enthalpy, in $\mathrm{kJ} / \mathrm{kg}$, at any temperature, t , between 0 and 60 C is approximately:
$\mathrm{h}=1.007 \mathrm{t}-0.026$ below zero: $h=1.005 t$
The enthalpy of liquid water is also defined to be zero at zero degrees celsius. To turn liquid water to vapour at the same temperature requires a very considerable amount of heat energy: $2501 \mathrm{~kJ} / \mathrm{kg}$ at 0 C

At temperature $t$ the heat content of water vapour is:
$\mathrm{h}_{\mathrm{w}}=2501+1.84 \mathrm{t}$
Notice that water vapour, once generated, also requires more heat than dry air to raise its temperature further: $1.84 \mathrm{~kJ} / \mathrm{kg}$.C against about $1 \mathrm{~kJ} / \mathrm{kg}$.C for dry air.

The enthalpy of moist air, in $\mathrm{kJ} / \mathrm{kg}$, is therefore:
$\mathbf{h}=(\mathbf{1 . 0 0 7} \times \mathbf{t}-\mathbf{0 . 0 2 6})+\mathbf{g} \times(\mathbf{2 5 0 1}+\mathbf{1 . 8 4 \times t})$
$g$ is the water content in $\mathrm{kg} / \mathrm{kg}$ of dry air

## The Psychrometer

The final formula in this collection is the psychrometric equation. The psychrometer is the nearest to an absolute method of measuring RH that the conservator ever needs. It is more reliable than electronic devices, because it depends on the calibration of thermometers or temperature sensors, which are much more reliable than electrical RH sensors.

The psychrometer, or wet and dry bulb thermometer, responds to the RH of the air in this way: Unsaturated air evaporates water from the wet wick. The heat required to evaporate the water into the air stream is taken from the air stream, which cools in contact with the wet surface, thus cooling the thermometer beneath it. An equilibrium wet surface temperature is reached which is very roughly half way between ambient temperature and dew point temperature. The air's potential to absorb water is proportional to the difference between the mole fraction, $\mathrm{m}_{\mathrm{a}}$, of water vapour in the ambient air and the mole fraction, $\mathrm{m}_{\mathrm{w}}$, of water vapour in the saturated air at the wet surface. It is this capacity to carry away water vapour which drives the temperature down to $\mathrm{t}_{\mathrm{w}}$, the wet thermometer temperature, from the ambient temperature $\mathrm{t}_{\mathrm{a}}$ :
$\left(\mathrm{m}_{\mathrm{w}}-\mathrm{m}_{\mathrm{a}}\right)=\mathrm{B}\left(\mathrm{t}_{\mathrm{a}}-\mathrm{t}_{\mathrm{w}}\right)$
$B$ is a constant, whose numerical value can be derived theoretically by some rather complicated physics (see the reference below).

The water vapour concentration is expressed here as mole fraction in air, rather than as vapour pressure. Air is involved in the psychrometric equation, because it brings the heat required to evaporate water from the wet surface. The constant B is therefore dependent on total air pressure, $P$. However the mole fraction, $m$, is simply the ratio of vapour pressure $p$ to total pressure $P: p / P$. The air pressure is the same for both ambient air and air in contact with the wet surface, so the constant $B$ can be modified to a new value, $A$, which incorporates the pressure, allowing the molar fractions to be replaced by the corresponding vapour pressures: $\mathrm{p}_{\mathrm{w}}-\mathrm{p}_{\mathrm{a}}=\mathrm{Ax}\left(\mathrm{t}_{\mathrm{a}}-\mathrm{t}_{\mathrm{w}}\right)$

The relative humidity (as already defined) is the ratio of $\mathrm{p}_{\mathrm{a}}$, the actual water vapour pressure of the air, to $\mathrm{p}_{\mathrm{s}}$, the saturation water vapour pressure at ambient temperature.
$\mathbf{R H} \%=100 \times p_{a} / p_{s}=100 \times\left(p_{w}-\left(\mathbf{t}_{\mathbf{a}}-\mathbf{t}_{w}\right) \times \mathbf{6 3}\right) / \mathbf{p}_{\mathrm{s}}$
When the wet thermometer is frozen the constant changes to 56
The psychrometric constant is taken from: R.G.Wylie \& T.Lalas, "Accurate psychrometer coefficients for wet and ice covered cylinders in laminar transverse air streams", in Moisture and Humidity 1985, published by the Instrument Society of America, pp 37-56. These values are slightly lower than those in general use.

There are tables and slide rules for calculating RH from the psychrometer but a programmable calculator is very handy for this job. Psychrometric charts have graphical versions of all these formulæ and don't need electricity.

To check your program, take air at 20C and 15.7C wet bulb temperature. The RH is $65 \%$. The water vapour pressure is 1500 Pa. The water vapour concentration in $\mathrm{kg} / \mathrm{m} 3$ is 0.011 , in $\mathrm{kg} / \mathrm{kg}$ it is 0.009 . The dew point is $13 C$.

