Infrastructure Commission (INFCOM)

Standing Committee on Measurements, Instrumentation and Traceability (SC-MINT)

Expert Team on Quality, Traceability and Calibration (ET-QTC)

Calibration of humidity instruments Part-1: Introduction

1.1 Water vapour concepts and variables to measure



ORGANIZATION

Calibration of humidity instruments

Overview:

- 1. Introduction
 - 1.1 Water vapour concepts and variables to measure
 - 1.2 Water vapour theory
 - 1.3 Humidity scales and conversions
- 2. Description of technology
- 3. Calibration procedures
- 4. Humidity uncertainties



What do we mean by humidity or moisture?

Water vapour mixed in a gas. Water impregnated in a body.

Water content in a solid, liquid, or gas



And for ambient humidity:



Existence of gaseous state water (vapour) in other carrier gas (air)

We quantify the amount of water in the total mixture:

Dew / Frost point temperature, relative humidity

Mixing ratio, mass per unit volume

Mole (amount) fraction
$$X_{\text{vapor}} = \frac{n_v}{n_v + n_a}$$



Dew-point (DP) temperature: is the temperature to which a humid air must be cooled for water vapour to condense into liquid water.

Temperature at which the air is saturated in equilibrium over a flat water surface.





Frost-point (FP) temperature: is the temperature to which a humid air must be cooled for water vapour form frost.

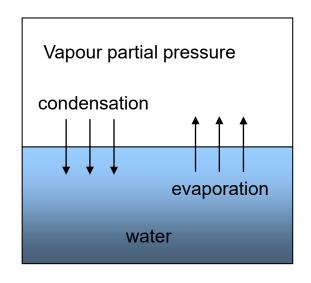
Temperature at which the air is saturated in equilibrium over a flat ice surface.

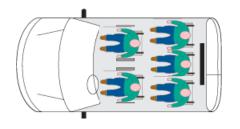
WARNING: In the range **just below 0 °C** where either frost or dew (supercooled water) can form, the dew- and the frost-point temperatures differ.



What is saturation?

It is the result of a dynamic equilibrium and determines a maximum amount of water vapour that space can contain at a certain pressure and temperature.





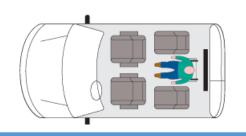
total possible seats to occupy are occupied



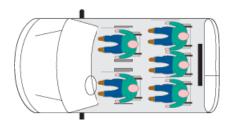
Relative humidity (U): Ratio between the "current" partial vapour pressure $e'(\text{mole fraction } X_v)$ of the mixture (air with vapour) and the saturation vapour pressure $e'_{w/i}$ (mole fraction saturation X_{vs}) over a flat surface of liquid water or ice (w or i) at the same temperature and pressure of the mixture expressed as a percentage.

$$U = \left(\frac{e'}{e'_{w/i}}\right)_{P,T} \times 100 \text{ or } U = \left(\frac{X_v}{X_{vs}}\right)_{P,T} \times 100$$

relative occupied seats in a car (*U*)



current occupied seats (e and X_{v})



total possible seats to occupy $(e_s \text{ and } X_{vs})$



Concise version of main humidity definitions as given in WMO-No.8 (2024) Guide to Instruments and Methods of Observation, Volume 1, Chapter 4:

- Mixing ratio r: ratio between the mass of water vapour and the mass of dry air;
- Specific humidity q: ratio between the mass of water vapour and the mass of moist air;
- Dew-point temperature or dew point t_d: The temperature at which moist air is saturated with water vapour.
- Relative humidity U: ratio in % of the observed vapour pressure to the saturation vapour pressure with respect to water at the same temperature and pressure. The term "relative humidity" is often abbreviated to RH;
- Vapour pressure e': The partial pressure of water vapour in air;
- Saturation vapour pressures e'_w and e'_i: Vapour pressures in air in equilibrium with the surface of water and ice, respectively.



Commonly used units and scales relevant to humidity are given WMO-No.8 (2024) as:

- Mixing ratio r and specific humidity q (dimensionless quotient of masses, in kilogrammes per kilogrammes, kg·kg⁻¹);
- Vapour pressure in air e', e'_w, e'_i and pressure p (in units of pressure, such as hectopascals, hPa. 1 hPa = 1mbar);
- Temperature t, wet-bulb temperature t_w, dewpoint temperature t_d, and frost-point temperature t_f (in degrees Celsius, °C);
- Temperature T, wet-bulb temperature T_w, dewpoint temperature T_d, and frost-point temperature T_f (in kelvins, K, as used for certain humidity calculations, and for expressing differences, rather than for general expression of humidity values);
- Relative humidity U (in percent, %. The alternative symbol, %rh*, is also often used to avoid confusion with other percentages.)

* Or %RH



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1.2 Water vapour theory



Written by Drago Groselj (ARSO)

Adapted by Javier Garcia Skabar (INTI) and Stephanie Bell (NPL)

Presented by Stephanie Bell

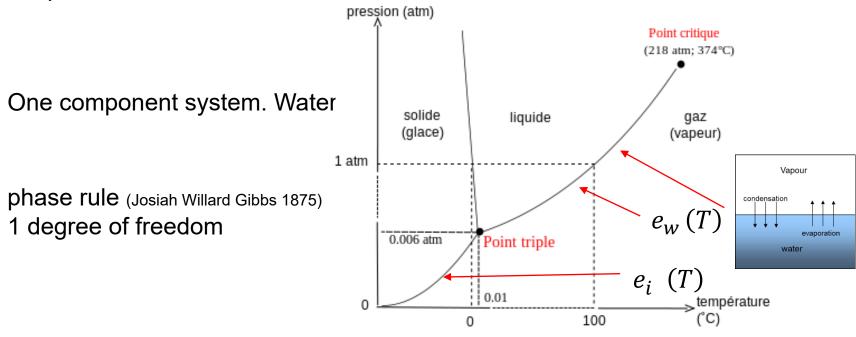
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Saturation pressure vapour, relative humidity and dew- / frost-point temperature,



In this system, the saturation vapour pressure depends only on the temperature

Important: We have an empirical equation for $e_{w/i}(T)$



$$e_{w/i}(T) = exp\left(\sum_{i=0}^6 a_i \cdot T^{i-2} + a_7 \cdot \ln(T)
ight)$$
 (pressure in Pa, temperature in K)

ITS-90 formulations by Hardy (1998) for vapor pressure, frost-point temperature, dew-point temperature, and enhancement factors in the range - 100 to 100 °C

Hardy R., in *Papers and Abstracts from the Third ISHM 1998;* NPL, London, 1998, 1, pp.214 222.

Or similar equation by Sonntag (1990)

Sonntag, D.; Important New Values of the Physical Constants of 1986, Vapor Pressure Formulations based on the ITS-90 and Psychrometer Formulae; Z. Meteorol., 70 (5), 1990, pp. 340-344.

The above are correct in terms of the current temperature scale (ITS-90)

(Older equations by Wexler/Greenspan are not in terms of ITS-90 but can be used approximately)

Wexler, A., Greenspan, L., in *Journal of research of NBS* 80A: 775-785 (1976). Wexler, A., in *J.Res.Natl.Bur.Stand.A, Phys.Chem.*(USA) 81A: 5-20 (1977).



WMO No.8 GUIDE TO INSTRUMENTS AND METHODS OF OBSERVATION - VOLUME I (2023 ed), ANNEX 4.B. FORMULAE FOR THE COMPUTATION OF MEASURES OF HUMIDITY

The following formulae (due to Magnus) are convenient for approximate calculation if the uncertainty is considered

$$e_w(t) = 6.112 \cdot exp(17.62 \cdot t / (243.12 + t))$$

Water, pure phase, -45 °C to +60 °C, uncertainty up to 0.6 %, (k = 2)

$$e_i(t) = 6.112 \cdot exp(22.46 \cdot t / (272.62 + t))$$

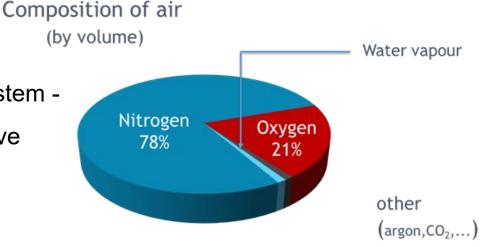
Ice, pure phase, -65 °C to 0 °C, uncertainty up to 1 %, (k = 2)

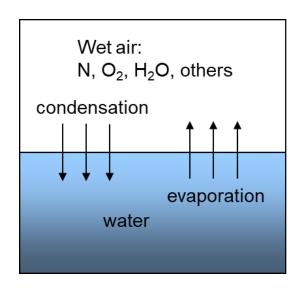
(Pressure in Pa, temperature in °C)

Warning: not for two pressure humidity generators



Wet air is not a one-component system two phases and various non-reactive
components





These change the vapor pressure and the freedom degrees of the system.

The effective saturation vapor pressure $e'_{w/i}$ also depends on the total pressure of the system.

The empirical equation for $e_{w/i}(T)$ must be adjusted, as water vapour does not act as an "ideal gas".



A good approximation at atmospheric pressure and near room temperature is: $e'_{w/i}(P,T) \approx e_{w/i}(T)$

This underestimates effective vapour pressure by about 0.4 %, in room air.

For accurate calculation, an increase factor known as the *water vapour enhancement factor, f* , must be considered due to the interaction in the air-water vapor mixture. Then:

$$e'_{w/i}(P,T) = f(P,T) \cdot e_{w/i}(T)$$

Accurate enhancement factor equations by Hardy (1998):

ITS-90 formulations for vapor pressure, frost-point temperature, dew-point temperature, and enhancement factors in the range –100 °C to 100 °C

Hardy R. in Papers and Abstracts from the Third ISHM 1998: NPL London, 1998, 1, pp. 214-22

Hardy R., in *Papers and Abstracts from the Third ISHM 1998;* NPL, London, 1998, 1, pp.214 222.



For enhancement factor, approximate calculations using older equations (pre-1990) also can be used with acceptable uncertainty.

Water vapour enhancement factor general form of equation:

$$f(P,T) = exp\left(\alpha \cdot \left(1 - \frac{e_{w/i}(T)}{P}\right) + \beta \cdot \left(\frac{P}{e_{w/i}(T)} - 1\right)\right)$$

$$lpha = \sum_{i=0}^3 A_i t^i$$
 $eta = e^{\sum\limits_{i=0}^3 B_i t^i}$ Water 0 °C to 100 °C Water -50 °C to 0 °C lce -100 °C to 0 °C

Hyland R. W., A correlation for the second virial coefficients and enhancement factors for CO₂-Free moist air from –50 °C to 90 °C, in *Journal of research of NBS*, 79A: 551-560 (1975)

Greenspan, L., Functional equations for the enhancement factors for CO₂-Free moist air, in *Journal of Research of the NBS*, 80A, 41-44 (1976).

Goff J.A. standardization of thermodynamic properties of moist air, Heating, piping and air conditioning 21, 118 (1949).



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1.3 Humidity scales and conversions



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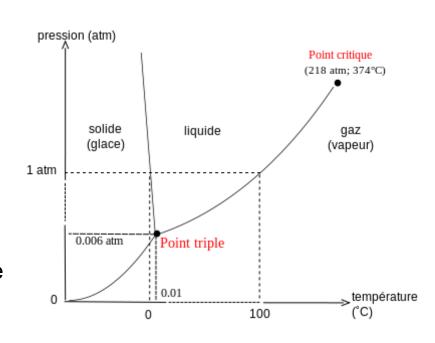


To understand humidity scales and conversions, we combine the following concepts and definitions:

saturation vapour pressure,

the empirical equations or formulae

$$e'_{w/i}(P,T) = f(P,T) \cdot e_{w/i}(T),$$



Relative humidity
$$U = \left(\frac{e'}{e'_{w/i}}\right)_{P,T} \times 100$$
,

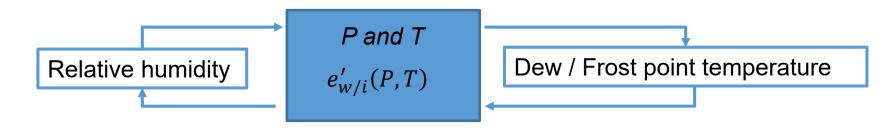
and dew / frost point temperature definitions.



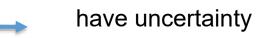
Measuring U or $t_{dew/frost}$ allows indirect evaluation of other humidity variables such as vapour pressure or mixing ratio.

Conversions typically involve a vapour pressure equation.

To convert from U to $t_{dew/frost}$, or the inverse:



The empirical equation f(P,T) and $e_{w/i}(T)$ are approximations and formulas,



"Fundamental" fraction or ratio in terms of mass or moles (amount) of vapour can be directly measured

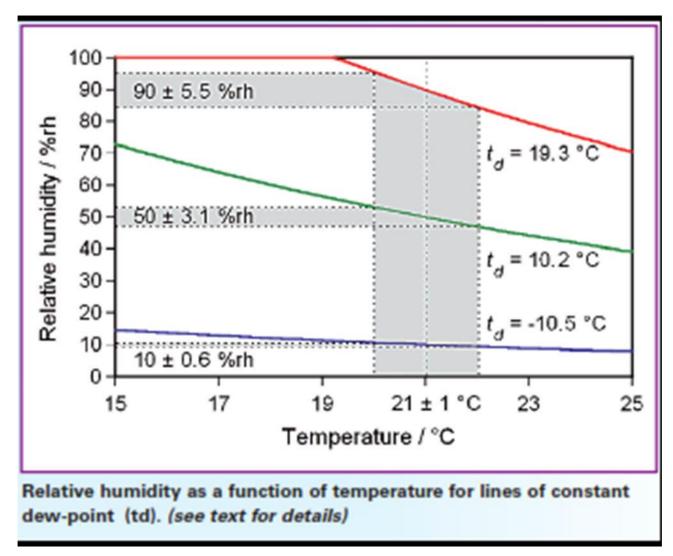


otherwise, can be calculated via f(P,T) and $e_{w/i}(T)$



WORLD

METEOROLOGICAL ORGANIZATION



Ref:: Humidity hassles by Jeremy Lovell-Smith, Measurement Standards Laboratory

Why the scale in dew point temperature?

It is based on a fundamental thermodynamic equilibrium that can be reproduced, with direct link to SI unit of temperature

- It is more direct than measuring relative humidity
- Does not depend on ambient temperature
- The technology for dew point meters is more "reliable" they are more repeatable, less long-term drift.



Thank you.



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