

التبخر Evaporation

Evaporation: is the process in which a liquid changes to the gaseous state (vapor), below the boiling point through the transference of heat energy.

هو العملية التي يتحول فيها السائل إلى الحالة الغازية (بخار) عند سطح الحرارة قبل نقطة الغليان من خلال انتقال الطاقة الحرارية.

The heat absorbed by a unit mass of Liquid required to change it to vapor state, is called "Latent heat of evaporation"

الطاقة الحرارية الممتصة من سائل بكمية تحويله إلى بخار تدعى «الحرارة الكامنة للتبخر» ولتعبير عن هذا بالمعادلة

$$L_h = 597.3 - 0.564 T_c \quad \text{----- (1)}$$

حيث T درجة الحرارة السيليزية. L_h بـ cal/gr
لا حظ أنه بإمكانك تحويله إلى (KJ/gr) :

$$L_h = 2.50 - 0.00236 T_c$$

حيث T كل درجة (cal) تعادل 4.186 جول / أو :

$$\boxed{\text{Cal} = 4.186 \text{ J}}$$

هو مضاعف جول إعتناك (4.186)

The rate of evaporation (معدل التبخر) is dependent on :

- (i) The vapor pressure at the water and air above . معدل التبخر يتغير مع
- (ii) Air & water temperatures . زيادة درجة الحرارة تزيد من معدل التبخر
- (iii) Wind speed . يزداد مع زيادة سرعة الرياح
- (iv) atmospheric pressure . انخفاضه يضاعفه زيادته في الضغط الجوي
- (v) water quality (dissolved salt) . انخفاضه يضاعفه زيادته في الملح
- (vi) Heat Storage in water body . الحرارة المخزنة في الجسم المائي تؤثر على التبخر

Evaporation Estimation

② تقدير التبخر

The estimation of evaporation is of utmost important in many Hydrologic problems associated with planning and operation of reservoirs and irrigation systems.

يعد تقدير التبخر غاية في الأهمية لما له من الدور الهام في تخطيط وتشغيل السدود وأنظمة الري.

The estimation of evaporation can be carried out by :

- (1) measurement by evaporimeter - القياس بأجهزة القياس
- (2) Empirical evaporation equations - المعادلات التجريبية
- (3) Analytical methods - الطرق التحليلية

① Evaporimeters

القياس باستخدام مقاييس التبخر

All evaporimeters are of type of Pan evaporation :

- 1) Class A Pan evaporation ($\phi = 121 \text{ cm}$, $d = 25.5$) A - الدنار صنف A
- 2) Colorado Sunken Pan evaporation - إنار كولورادو الغاطس
- 3) U.S.G Pan (Floating Pan evaporation) - الوعاء الطواف

Evaporation Pans are not exact models for large reservoirs, because :

- (i) They differ in the heat-storing capacity and heat-transfer from the side and bottom.
- (ii) The height of the rim in the pan affects the wind action.
- (iii) The heat transfer is different in pan & reservoir due to Pan material - and thus;

$$\text{Evap}_{\text{Lake}} = \text{Evap}_{\text{Pan}} * C_p \quad \text{--- (2) } C_p = \text{معامل انار التبخر}$$

C_p = coefficient of Pan ranged from 0.6 - 0.8 for class A.

③

The evaporation pans are usually installed in such location according to World Meteorological Organization (WMO) recommendation:

1. Arid Zones - on every 30 000 km² المناطق الجافة
2. Humid temperate climate - one every 50 000 km² المناطق المعتدلة الرطبة
3. Cold regions - One every 100 000 km² المناطق الباردة

② Empirical equations for evaporation معادلات التجريبية للتبخر

A large number of empirical equations are available, but most of these equations are based on Dalton-type equation.

John Dalton 1802 stated that

$$E_L = C (e_s - e_a)$$

E_L = Lake evaporation in (mm/day)

C = Coefficient

e_a = actual vapor pressure of air above water surface at specified height, in (mm Hg).

e_s = saturated vapor pressure at the water surface temperature, in (mm/day).

This basic equation was modified many times and finally takes the following form:

$$E_L = K f(u) (e_s - e_a) \quad \text{--- (3)}$$

$\text{mm/day} \quad \quad \quad \begin{matrix} \uparrow \text{mm Hg} & \downarrow \text{mm Hg} \end{matrix}$

K = coefficient

$f(u)$ = wind speed correction function (Dimensionless)

Meyer Formula (1915)

$$E_L = K_M (e_s - e_a) \left(1 + \frac{u_g}{16}\right) \text{ mm/day} \quad (4)$$

E_L , e_s & e_a as defined previously. $K_M = 0.36$ for deep lake; 0.5 for shallow water.
 u_g = wind velocity (monthly mean) in (km/hr) measured at height 9m above ground surface.

If the wind velocity was known at z_0 , one can estimate a wind velocity at a specified height z , by using,

$$\frac{u}{u_0} = \left(\frac{z}{z_0}\right)^k \quad (5)$$

k : Power Law coefficient varied from 0.1 $\rightarrow$ 0.6.
 Use $k = 1/4$, for natural ground roughness

Rohwer Formula (1931)

$$K = 0.771(1.465 - 7.3 \times 10^{-4} P_a) \approx 0.70 \text{ for } P_a = 760 \text{ mm Hg}$$

$$f(u) = (0.44 + 0.0733 u_0)$$

$$E_L = K f(u) (e_s - e_a) \quad (6)$$

P_a = mean barometric pressure in (mm Hg).

u_0 = mean wind velocity in (km/hr) at ground level which can be taken at 0.6m above ground surface.

These empirical equations are simple to use and permit the use of standard meteorological data. However, the various limitation of these formulae, they can, at best, give an approximate values of evaporation.

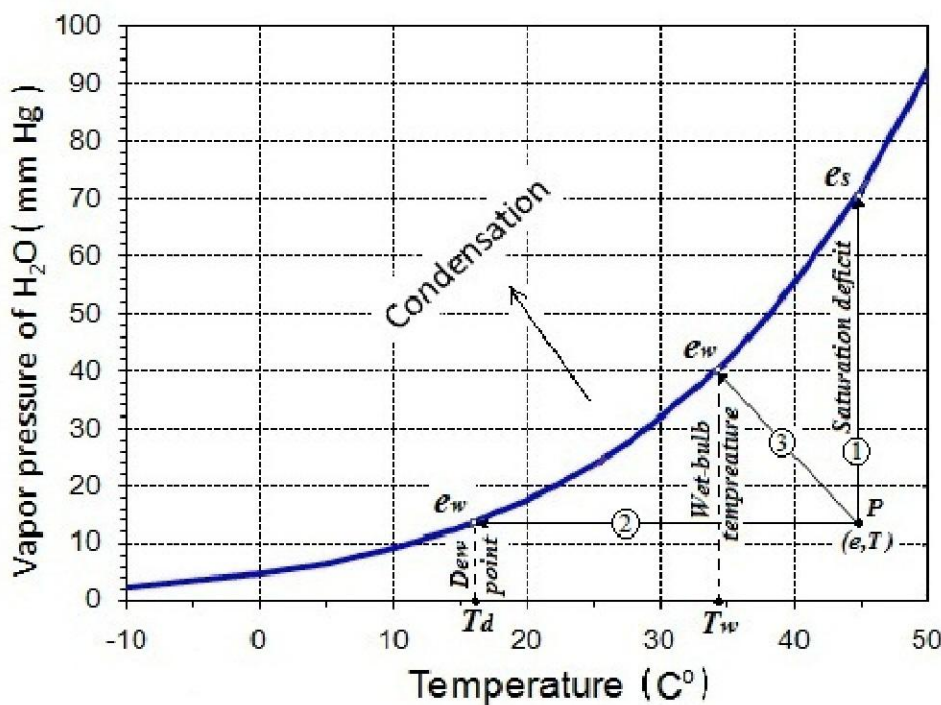
هذه المعادلات التجريبية بسيطة الاستخدام وتسمح باستخدام المقادير
 المناخية الشائعة، لكن المحددات المعقدة لهذه المعادلات تجعل نتائج
 تقريبية فقط في أحسن الأحوال.

لفرض تطبيق الساتورة آتفة الذكر يجب فونة قيم e_s عند
درجة حرارة

In using empirical equations, the saturated vapor pressure at a given temperature should be known. (e_s),

Relation between $e_s - T$ علاقة بين e_s ودرجة الحرارة

Suppose an evaporating surface of water in a closed system and enveloped in air.



evaporating of water into air will take place until a state of equilibrium is reached when $e_a = e_s$ and no more absorption occurs.

حالة التوازن بين بخار الماء في الهواء والسطح
ضغط في راء يمكن ضغط البخار السبع ويتغير بتغير درجة الحرارة كما هو مبين بالشكل

1. Since P lies below saturation Vapor Pressure curve, the air can absorb more water vapor, and if it did so, the position P move vertically (path ①) until e_s curve intersection, with no change in temperature.
2. if no change in a humidity of air, while it was cooled, the P move horizontally along path ② until saturated line was intersected. Now, e_s will takes place at new temperature T_d called dew-point. Cooling beyond this temperature resulted Condensation.
3. if water allowed to evaporate freely, P will move diagonally along path ③ until saturated vapor pressure curve is reached at point defined by (e_w, T_w) . T_w called wet-bulb temp, which is an original air temperature that can be cooled by evaporating water into it. It can be found by bulb thermometer.

Relative humidity (hr)

الرطوبة النسبية

تمثل كمية الرطوبة فيما الهواد الى كمية اللوطة في حالة الاتساع ويعبر عنه :-

$$\% hr = \frac{e_a}{e_s} \times 100 \quad \text{----- (7)}$$

كان قياس e_a باستخدام الحرار في البصلة الرطبة وقياس e_s جواز قياس الرطوبة "طوب" (Psychrometer) حيث يقيس الهواء درجة حرارة البصلة t_w (wet bulb temp.) وفي الدرجة التي يصل اليها الهواء عند تبريده بواسطة

$$\frac{(e_w - e_a)}{(t_w - t_a)} = \gamma \quad \text{----- (8)}$$

$\begin{matrix} \nearrow \text{mmHg} & \nearrow \text{mmHg} \\ e_w & e_a \end{matrix}$
 $\begin{matrix} \uparrow t_c & \uparrow t_c \\ t_w & t_a \end{matrix}$

e_w = saturated vapor pressure corresponding to wet-bulb temperature

t_w = wet-bulb temperature

t_a = dry-bulb temperature

e_a = corresponding air vapor pressure to " t_a ".

γ = Psychrometer constant ثابت الرطبة

$$\gamma = 6.5 \times 10^{-4} \text{ Pa } / ^\circ \text{C} \quad \text{----- (9)}$$

ملاحظة

$$\gamma = 0.149 \text{ mmHg } / ^\circ \text{C}$$

$$\gamma = 0.66 \text{ mb } / ^\circ \text{C}$$

$$\gamma = 0.066 \text{ kPa } / ^\circ \text{C}$$

P_a = atmospheric pressure.

① e_s يجب ان يكون في نفس الوحد التي يكون لافق

② t_w يقيس في الرطبة

③ t حرارة الهواد الكاف في الحرار

④ e_a يجب ان يكون في نفس الوحد التي يكون لافق

Note that:

$$mb = \frac{1}{10} \text{ kPa} \Rightarrow mb = 100 \text{ Pa} \Rightarrow P_a = 10^{-5} \text{ bar}$$

$$mb = \frac{3}{4} \text{ mmHg} \Rightarrow \text{mmHg} = \frac{4}{3} mb$$

$$\text{mmHg} = \frac{2}{15} \text{ kPa} \Rightarrow \text{kPa} = 7.5 \text{ mmHg}$$

How To calculate e_s as a function of T :

After reviewing the variation of e_s with T , an attempt was made to introduce equation that can be calculated e_s and its gradient as a function of T .

$$e_s = 4.58 e^{\left(\frac{17.27 T}{237.3 + T}\right)} \quad \text{----- (10)}$$

$$\Delta = \frac{de_s}{dT} = \frac{4098 e_w}{(237.3 + T)^2} \quad \text{----- (11)}$$

e_s in mm Hg, Δ in mm Hg/ $^{\circ}\text{C}$.

If e_s required to be in Pa instead of mm Hg, then the factor 4.58 should be replaced by 611, whereas Δ remains as it is.

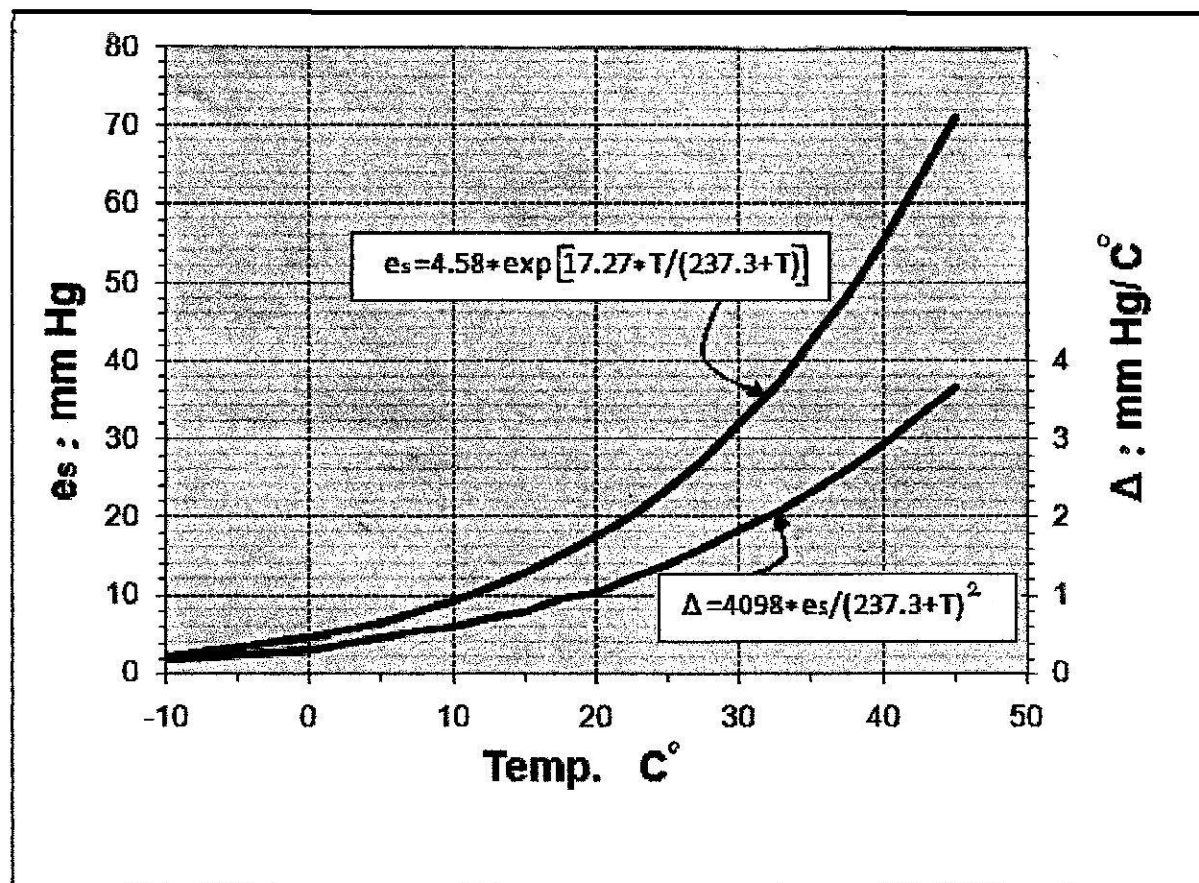


Fig : Saturation Vapor pressure of water in air as a function of temperature

Example

A Lake of 250 hectares surface area, had the following average values of climate parameters during a week of measurements,

- relative humidity = 40%
- wind velocity at 1.0 m above ground surface = 16 km/hr.
- Pan evaporation = 72 mm in the week.
- Pan coefficient = 0.8
- Temperature = 20 °C

- (a) estimate average daily evaporation from Lake using Meyer formula.
- (b) estimate accuracy of this method compared with pan evaporation.
- (c) estimate the volume of water evaporated from lake in that week.

Solution:

$$1 \text{ ha} = 10\,000 \text{ m}^2$$

(a) from eq (10); $e_s = 4.58 e^{\left(\frac{17.27(20)}{237.3+20}\right)} = 17.53 \text{ mm Hg.}$

From eq. (7);

$$e_a = h_r \times e_s = 0.4 (17.53) = 7.01 \text{ mm Hg.}$$

From eq. (5);

$$\frac{u_2}{u_1} = \left(\frac{9}{1}\right)^{1/4} \Rightarrow u_2 = 16 \times 9^{1/4} = 21.9 \text{ km/hr}$$

From eq (4); $K_M = 0.36$ since the Lake is large.

$$E_L = 0.36 (17.53 - 7.01) \left(1 + \frac{21.9}{16}\right) = 8.97 \text{ mm/day}$$

(b) Daily evaporation from Pan evaporimeter = $\frac{72}{7} = 10.28 \text{ mm/day}$

From eq (2)

$$E_{\text{actual}} = 10.28 \times 0.8 = 8.23 \text{ mm/day}$$

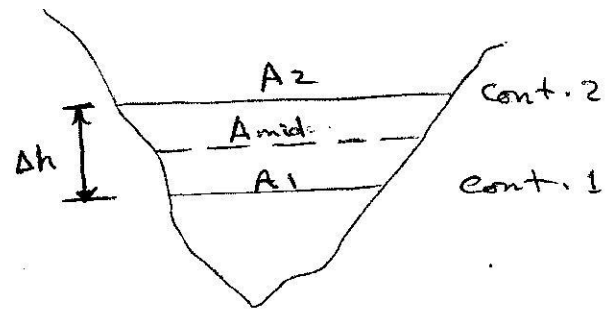
$$\%E = \frac{|\text{actual} - \text{calculated}|}{\text{actual}} \times 100 = \frac{|8.23 - 8.97|}{8.23} \times 100 = 9\%$$

(c) Volume of loss by evaporation = $\frac{8.23}{1000} \times 7 \times 250 \times 10\,000 = 144,025 \text{ m}^3$

$$\text{or} = 0.8 \times \frac{72}{1000} \times 250 \times 10\,000 = 144,000 \text{ m}^3$$

Why the two answers are different?

Note that the surface area of the Lake is assumed to be constant. Is it real??



$$\Delta S = \text{Volume change} = \Delta V$$

$$\Delta V = \Delta h * \bar{A}$$

There are more than rule to calculate $\bar{A} = f(A_1, A_2)$

① for uniform shape:

$$\bar{A} = \frac{A_1 + A_2}{2}$$

$$\textcircled{2} \quad \bar{A} = \frac{A_1 + A_2 + \sqrt{A_1 A_2}}{3} \quad \text{Cone formula}$$

$$\textcircled{3} \quad \bar{A} = \frac{A_1 + A_2 + 4 A_{\text{mid}}}{6} \quad \text{Prismoidal formula}$$

A_m is midway area between two successive contours.

The practical formula for use is the cone formula

$$\Delta S = \frac{h}{3} (A_1 + A_2 + \sqrt{A_1 A_2})$$

③ Analytical methods of estimation ⑨ الطرق التحليلية لتقدير التبخر

The Lake evaporation can be estimated using analytical methods which may be classified into three categories;

- (i) water-budget الموازنة المائية
- (ii) Energy-balance method طريقة التوازن الحراري
- (iii) Mass-transfer method طريقة انتقال الكتلة

(i) Water Budget method

It is the simplest method of the three analytical methods, but it has a least reliable. It involves writing hydrological continuity equation معادلة الاستمرارية الهيدرولوجية for lake and determining evaporation from knowledge of other variables.

$$V_p + V_{qi} - V_{qo} + V_{gi} - V_{go} - V_E - V_T = \Delta S \quad \text{--- (12)}$$

V_p = Volume of daily precipitation over the lake.

V_{qi} = Volume of daily inflow into lake.

V_{qo} = Volume of daily outflow from the lake.

V_{gi} = Volume of daily groundwater inflow.

V_{go} = Volume of daily groundwater outflow (see page).

V_E = Volume of daily Lake evaporation

V_T = Volume of daily transpiration.

ΔS = increase in volume of storage in day.

All units are in m^3 or (m) depth over surface area of lake.

$$\Rightarrow E = P + (Q_i - Q_o) + (G_i - G_o) - T_L - \Delta S \quad \text{--- (13)}$$

All parameters must be had same unit, if not any different unit of such parameter should be converted.

Note that, P, Q_i, Q_o, D_s can be measured while G_i, G_o, T_L were estimated, T_L can be neglected in some lakes.

EL can be well-estimated from eq(13), if the unit of time is kept large, say weeks, months.

عندما يكون الوقت كبيراً، مثلاً أسابيع، أشهر، يمكن تقدير E_L جيداً باستخدام المعادلة (13)، إذا كان وحدة الزمن كبيرة، مثلاً أسابيع، أشهر، يمكن تجاهل T_L في بعض البحيرات.

(ii) Energy Budget method

طريقة موازنة الطاقة

It is application of Law of conservation of energy. The energy available for evaporation is determined by considering incoming energy, outgoing energy and energy stored in the water body over known time interval.

الطريقة هي تطبيق لقانون حفظ الطاقة، فالطاقة المتوافقة للبخار كحد أقصى
الطاقة الواردة والطاقة الخارجة والطاقة المخزنة في الجسم المائي خلال فترة زمنية معروفة.

R_a = incident solar radiation outside atmosphere, it is a function of the latitude and period of the year, see table in next pages.

H_c = incident solar radiation reached the water surface.

H_b = back radiation from water body (long wave)

H_a = sensible heat transfer from water surface to air

H_e = heat energy used up in evaporation.

$$= \rho L_h E_L \Rightarrow \text{cal/cm}^2$$

$\frac{\text{gr}}{\text{cm}^3} \times \frac{\text{cal}}{\text{gr}} \times \text{cm or mm}$

H_g = Heat flux into ground

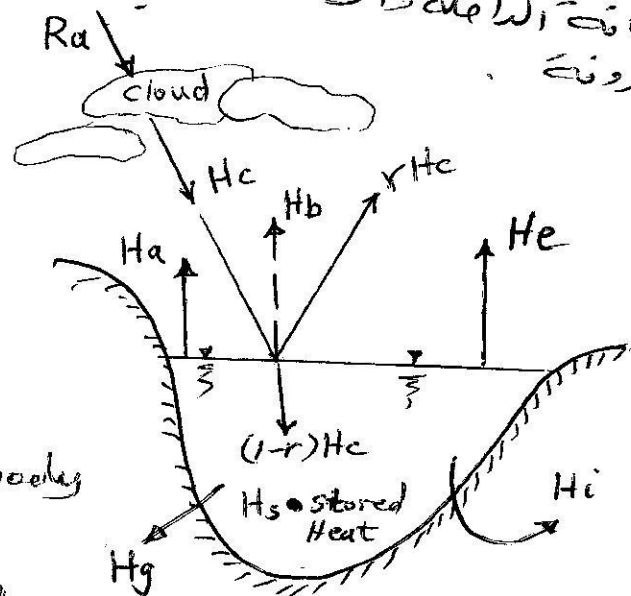


Fig: Energy Balance in Water Body.

H_i = net heat conducted out of the system (advected energy)
 الطاقة المنقولة خارج النظام (الطاقة المنقولة خارج النظام)

H_s = heat stored in the system (water body)
 and thus ;

H_n = Net heat energy received by water surface

$$\left. \begin{aligned} H_n &= H_s + H_i + H_g + H_a + H_e \\ \text{also } H_n &= (1-r) H_c - H_b \end{aligned} \right\} \text{--- (13)}$$

where r = coefficient of reflection (albedo)
 $r = 0.05$ from water surface.

All terms of energy - Balance method in $(\text{cal}/\text{cm}^2, \text{day})$ - If time interval is short, (H_s, H_i) can be neglected, since they are small.
 All terms except H_a can be measured or evaluated.
 جميع المصطلحات باستثناء H_a يمكن قياسها أو تقييمها.

$$\beta = \frac{H_a}{H_e} \quad ; \quad \beta = \text{Bowen's ratio} \text{--- (14)} \quad ; \quad \beta = \frac{H_a}{H_e}$$

$$\text{where } \beta = \gamma \frac{T_o - T_a}{e_s - e_a} \text{--- (15)}$$

T_o = temperature of water surface in $^{\circ}\text{C}$.

T_a = temperature of air in $^{\circ}\text{C}$

e_s = saturated vapor pressure in mm Hg at T_o

e_a = actual vapor pressure in mm Hg at T_a

γ = constant estimated by eq (9) $\approx 0.49 \text{ mm Hg}/^{\circ}\text{C}$

Hence ;

$$\text{by using: } \beta = \frac{H_a}{\rho L_h E_L} \text{ and From eq (13);}$$

$$E_L = \frac{H_n - H_g - H_s - H_i}{\rho L_h (1 + \beta)} \text{--- (16)}$$

it was found that the evaporation in a lake estimated by energy-Budget gives satisfactory result with errors of order 5% if the budget was applied to period less than week.

(iii) Mass transfer method

طريقة انتقال الكتلة

This method is based on the concept of turbulent transfer of water vapor mass from an evaporating surface to the atmosphere. It assumes that the diffusion of vapor pressure is analogue to temperature distribution. The equation similar to Dalton form can be adopted as;

$$E_L = \alpha (e_s - e_a) \quad \text{--- (17)}$$

Where α is vapor transfer coefficient, and

$$\alpha = \frac{0.102 U_a}{\left[\ln \left(\frac{z_a}{z_0} \right) \right]^2} \quad ; \quad \begin{array}{l} z_0 = \text{roughness height of surface} \\ \text{cm} \quad (0.01 - 0.06) \text{ cm} \end{array}$$

z_a = elevation corresponding to
cm velocity U_a (m/s)

Any other details is out of the current study.

Methods of Reduce evaporation: طرق تقليل التبخر

LOSSES

(1) reduce surface area. تقليل المساحة السطحية

(2) use mechanical covers. استخدام غطاء ميكانيكي

(3) use chemical films. الاستخدام الأفلام الكيميائية

hexadecanol } use these (استخدام الكحول سداسي عشر الكربون)

octadecanol }

- strong and flexible قوي ومرن
- If punctured due to impact, the film closes back soon after. يتم إغلاقه إذا تم ثقبه
- colourless, odourless, and nontoxic عديم اللون والرائحة

Evapotranspiration Equations

معادلات التبخر

تعتبر في دراسة مناخية معادلة بيان كمية هبوط النقط
معادلات بلاني - كروك وثورنتويت في صيغتين كذا - الرطوبة

Penman Equation:

معادلة بلاني

Penman's equation is based on combination of energy-balance and mass-transfer approach. This equation is modified many times and still modified by investigators.

$$PET = \frac{\Delta H_n + \gamma E_a}{\Delta + \gamma} \quad \text{----- (18)}$$

PET = daily evapotranspiration (mm/day)

Δ = slope of $e_s - T$ curve (see eq (11) Page 7) $\frac{\text{mmHg}}{^\circ\text{C}}$

γ = Psychrometric constant (see equation (9) Page 6) $= 0.49 \frac{\text{mmHg}}{^\circ\text{C}}$

H_n = net radiation in mm of evaporable water/day. It is the same as used in energy-budget (see eq. (13) Page 11).

E_a = Parameter quoted from mass-transfer, including wind velocity and saturation deficit.
 في الهواء من حيث انتقال الماء وكمية التبخر

$$E_a = 0.35 \left(1 + \frac{u_2}{160} \right) (e_s - e_a) \quad \text{----- (19)}$$

u_2 = mean wind velocity at 2m above ground surface in (km/day)

H_n can be calculated as follow:

R_a = incident solar radiation outside atmosphere. It is a function of period of a year and Latitude. Table of R_a is given for North part of earth whereas it is same for South part of earth. (mm/day)

$$H_c = R_a \left(a + b \frac{n}{N} \right) \quad \text{----- (20)} \quad \frac{\text{mm/day}}{\text{of water}}$$

H_c = as defined previously in Energy-budget (Page 10), incident radiation on water surface.

but; $H_n = (1-r) H_c - H_b$

$$\text{Thus; } H_b = \sigma T_a^4 \left(0.56 - 2.54 \sqrt{\frac{e_a}{p_0}} \right) \left(0.1 + 0.9 \frac{n}{N} \right) \quad \text{----- (21)}$$

(13)

$a = \text{Constant depends on latitude } \phi \Rightarrow a = 0.29 \cos \phi$

$b = \text{constant, average value is of } 0.52$

$n = \text{actual duration of actual sunshine, hrs}$ فترة الساعات الفعلية

$N = \text{max. possible of sunshine as a function of latitude and period of year, hrs (see table below)}$

تقدير أقصى مدة من الإشعاع الشمسي الممكنة في أي بلد صريحيه على أساس
ملا بين القيتين .

Table 1 ; Mean Monthly Hours of Possible Sunshine, N

North latitude	Jan	Feb	Mar	Apr	May	Jun	July	Aug	Sept	Oct	Nov	Dec
0°	12.1	12.1	12.1	12.1	12.1	12.1	12.1	12.1	12.1	12.1	12.1	12.1
10°	11.6	11.8	12.1	12.4	12.6	12.7	12.6	12.4	12.9	11.9	11.7	11.5
20°	11.1	11.5	12.0	12.6	13.1	13.3	13.2	12.8	12.3	11.7	11.2	10.9
30°	10.4	11.1	12.0	12.9	13.7	14.1	13.9	13.2	12.4	11.5	10.6	10.2
40°	9.6	10.7	11.9	13.2	14.4	15.0	14.7	13.8	12.5	11.2	10.0	9.4
50°	8.6	10.1	11.8	13.8	15.4	16.4	16.0	14.5	12.7	10.8	9.1	8.1

Table 2 ; Mean Monthly Solar Radiation (in mm/day) at top of atmosphere, R_a

North latitude	Jan	Feb	Mar	Apr	May	Jun	July	Aug	Sept	Oct	Nov	Dec
0°	14.5	15.0	15.2	14.7	13.9	13.4	13.5	14.2	14.9	15.0	14.6	14.3
10°	12.8	13.9	14.8	15.2	15.0	14.8	14.8	15.0	14.9	14.1	13.1	12.4
20°	10.8	12.3	13.9	15.2	15.7	15.8	15.7	15.3	14.4	12.9	11.2	10.3
30°	8.5	10.5	12.7	14.8	16.0	16.5	16.2	15.3	13.5	11.3	9.1	7.9
40°	6.0	8.3	11.0	13.9	15.9	16.7	16.3	14.8	12.2	9.3	6.7	5.4
50°	3.6	5.9	9.1	12.7	15.4	16.7	16.1	13.9	10.5	7.1	4.3	3.0

$\sigma = \text{Stefan-Boltzman Constant} = 2.0 \times 10^{-8} \text{ mm/day} / \text{K}^4$

one can convert mm of water/day to $\text{cal/cm}^2 \cdot \text{day}$

أي طارة لكل وحدة مساحة في اليوم الواحد حسب التوزيع الآتي

$$58.5 \text{ mm/day of H}_2\text{O} = \frac{\text{cal}}{\text{cm}^2 \cdot \text{day}}$$

and thus; $41.86 \frac{\text{cal}}{\text{cm}^2 \cdot \text{day}} = \frac{\text{KJ}}{\text{day} \cdot \text{m}^2}$

$$28.35 \text{ mm/day} = \frac{\text{Watt}}{\text{m}^2}$$

$$2449.44 \text{ mm/day} = \frac{\text{KJ}}{\text{day} \cdot \text{m}^2}$$

$$\Rightarrow \sigma = 11.703 \times 10^{-8} \frac{\text{cal}}{\text{cm}^2 \cdot \text{day}}$$

$$= 4.89888 \times 10^{-8} \frac{\text{KJ}}{\text{day} \cdot \text{m}^2}$$

$$= 5.67 \times 10^{-8} \frac{\text{J}}{\text{sec} \cdot \text{m}^2} = \frac{\text{Watt}}{\text{m}^2}$$

هذه القيمة المرادفة تعيد إذا أُعطي الكيلو
بني اللوحات المعتمدة (mm/day)

T_a = mean air temperature in degree Kelvin, K° .
 $K^\circ = 273 + C^\circ$.

r = reflection coefficient (albedo). The values of r the surface of incident, and given as;

Table 3; Reflection coefficient (albedo)

Surface	Range of r values
Close ground corps	0.15-0.25
Bare lands	0.05-0.45
Water surface	0.05
Snow	0.45-0.95

لحساب معدل التبخر من سطح الأرض باستخدام المعادلة (13) باستخدام البيانات التالية:

Thus eq (13) can be rewritten as;

$$H_n = R_a \left(a + b \frac{n}{N} \right) (1 - r) - \sigma T_a^4 \left(0.56 - 2.536 \sqrt{\frac{e_a}{P_0}} \right) \left(0.1 + 0.9 \frac{n}{N} \right) \quad (2)$$

Note that P_0 = standard barometric pressure = 760 mmHg

Penman equation (18) can be used for evaporation estimation from water surface by setting $r = 0.05$.

Example: given data as;

- mean monthly temperature = $19^\circ C$ (Nov.), Latitude = $28^\circ 4' N$
- mean relative humidity = 75%, Elevation = 230 m amsl.
- observed sunshine hours = 9 hr.
- wind velocity at 2 m height = 85 km/day
- Nature of surface cover = close-ground green cover.

- Calculate the vapotranspiration by Penman's equation
- Calculate the daily evaporation from water body (Lake).

From eq (10) & (11) $\Rightarrow e_s = 16.47 \text{ mmHg}$

$\Delta = 1.03 \text{ mmHg}/C^\circ$

$e_a = 0.75 \times 16.47 = 12.35 \text{ mmHg}$

$a = 0.29 \cos(28^\circ 4') = 0.29 \cos(28.067) = 0.2559$

$\sigma = 2.0 \times 10^{-8} \text{ mm/day} \cdot K^\circ$ $b = 0.52$

$T_a = 273 + 19 = 292 K^\circ$

$\sigma T_a^4 = 14.54$, From table (3) $r = 0.25$ (used as max.)

$R_a = 9.506 \text{ mm/day}^{11.2}$ by interpolation! $\frac{11.2 - 9.1}{20 - 30} = \frac{11.2 - R_a}{20 - 28.067}$ (from Table)

From Table (1); $N = -\left(\frac{11.2 - 10.6}{10}\right) 8.067 + 10.6 = 10.716 \text{ hrs.}$

From eq. (22);

$$H_n = (1 - 0.25)(9.506) \left(\frac{0.2559 + 0.52 \frac{9.0}{10.716}}{10.716} \right) - (14.54)(0.56 - 2.536 \sqrt{\frac{12.35}{760}}) * \\ H_n = 4.938 - 2.946 \quad (0.1 + 0.9 \frac{9.0}{10.716})$$

$$H_n = 1.992 \text{ mm of water/day}$$

From eq. (19) evaporating parameter E_a is;

$$E_a = 0.35 \left(1 + \frac{85}{160} \right) (16.47 - 12.35) = 2.208 \text{ mm/day}$$

① From eq. (18); $PET = \frac{1.03(1.992) + 0.49(2.208)}{1.03 + 0.49} = \underline{\underline{2.061 \text{ mm/d}}}$

② $H_n = 4.938 \frac{1 - 0.05}{1 - 0.25} - 2.946 = 3.308 \text{ mm/day}$ ← $\text{بقيت المياه في التربة}$

thus; $PET = \frac{1.03(3.308) + 0.49(2.208)}{1.52} = \underline{\underline{2.953 \text{ mm/day}}}$

Q: why PET from Lake evaporation is greater than PET from green-crop area?



استنتاج معادلة بنمان

Derivation of Penman Equation

Penman assumed that the water vapor transferred in a way similar to temperature dispersion (eddy-turbulence), i.e;

$$(e'_s - e_a) \propto (t'_a - t_a)$$

and thus; $(e'_s - e_a) = \text{const.} (t'_a - t_a)$

Recalling Eq.(15);

$$\beta = \gamma \frac{t'_a - t_a}{e'_s - e_a} ; \quad \text{here :} \quad \text{const.} = \frac{\gamma}{\beta}$$

While from Eq. (14): $\beta = \frac{H_a}{H_e}$

Thus; $H_n = H_e + H_a = (1 + \beta)H_e$; Assuming H_i , H_g & H_s are neglected.

$$\rightarrow H_n = \frac{H_n}{(1+\beta)} = \frac{H_n}{1+\gamma \left(\frac{t'_a - t_a}{e'_s - e_a} \right)}$$

Note that;

e'_s is a saturated vapor pressure at t'_a (which is the temperature of air close to water surface. t'_a is difficult to measure, and thus it was replaced by t_a and consequently e'_s by e_s).

According to Dalton; $E_o = C f(u) (e'_s - e_a)$ which can be easily convert to the ;

$E_a = C f(u) (e_s - e_a)$, where E_o is potential evaporation equivalent to H_e , whereas E_a is evaporation from water body.

How to substitute t_a, t'_a, e_s and e'_s

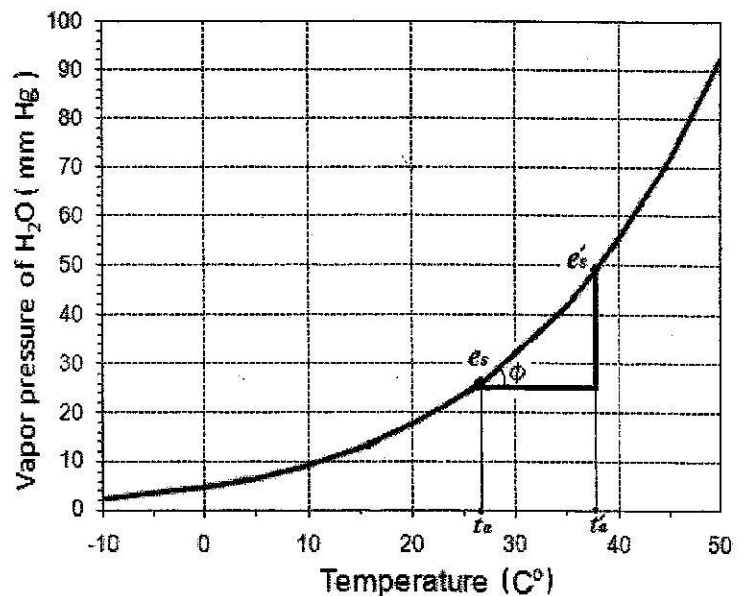
$\Delta = \tan \Phi$, slope of the vapor pressure -temp. curve at t_a ;

$$\Delta = \frac{t'_a - t_a}{e'_s - e_s}$$

And thus;

$$t'_a - t_a = \frac{e'_s - e_s}{\Delta}$$

$$H_e = \frac{H_n}{1 + \frac{\gamma}{\Delta} \left(\frac{e'_s - e_s}{e'_s - e_a} \right)}$$



$$H_e = \frac{H_n}{1 + \frac{\gamma}{\Delta} \left(\frac{(e'_s - e_a) - (e_s - e_a)}{e'_s - e_a} \right)} = \frac{H_n}{1 + \frac{\gamma}{\Delta} \left(1 - \frac{e_s - e_a}{e'_s - e_a} \right)}$$

Since :

$$\frac{E_a}{E_o} = \frac{E_a}{H_e} = \frac{e_s - e_a}{e'_s - e_a}$$

$$H_e = \frac{H_n}{1 + \frac{\gamma}{\Delta} \left(1 - \frac{E_a}{H_e} \right)}$$

$$\Delta H_n + \gamma E_a = (\gamma + \Delta) H_e$$

$$H_e = \frac{\Delta H_n + \gamma E_a}{\gamma + \Delta}$$

①

Universal Law of gases:

From Boyle & Charles laws, one can get the universal Law of gases as:

$$P = \rho R T$$

①

P = Pressure

Pa (N/m²)

ρ = mass density

kg/m³

R = universal gas constant, $R = 286.8 \frac{J}{kg \cdot K^\circ}$ for air.

T = temperature in $K^\circ = C^\circ + 273$

expansion or compression take place according to the law of thermodynamics

① isothermal (constant temperature) يتمدد أو يضغط في درجة حرارة ثابتة
الغاز مع الاحتفاظ بدرجة الحرارة ثابتة
 from eq (1); $T = \text{constant} \Rightarrow \frac{P}{\rho} = \text{constant}$ or;

$$\frac{P}{\rho} = \text{constant}$$

② where is no heat exchange لا يوجد تبادل حراري (عملية أديباتية)
isentropic

$$\frac{P}{\rho^n} = \text{constant} \quad \text{or} \quad \frac{P}{\rho^n} = \text{constant}$$

where n = is adiabatic exponent ≈ 1.4 for air.

Temperature Lapse rate

It is a change of temperature versus elevation, i.e. $\frac{dT}{dz}$?

$$\text{Since } P = \rho R T = \gamma/g R T \quad \text{--- (2)}$$

$$\text{set in general } \frac{P}{\rho^n} = \text{constant} \quad \text{--- (3)}$$

$$\text{i.e.; } \frac{\gamma R T}{g \rho^n} = \text{constant}$$

(2)

$$\text{or } T = \text{const.} \cdot \frac{g \gamma^{n-1}}{R}$$

$$\frac{dT}{d\gamma} = \text{const.} \cdot \frac{g}{R} (n-1) \gamma^{n-2}$$

$$\frac{dT}{d\gamma} = \frac{\gamma R T}{g \gamma^n} \cdot \frac{g}{R} (n-1) \gamma^{n-2}$$

$$\Rightarrow \frac{dT}{d\gamma} = (n-1) \frac{T}{\gamma} \quad \text{--- (4)}$$

$$\text{and; } \frac{T}{\gamma} \frac{d\gamma}{dT} = \frac{1}{n-1}$$

$$\text{From eq (2) } dP = d\left(\frac{\gamma}{g} R T\right)$$

$$\text{but from principle of fluid mechanic ; } \frac{dP}{dz} = -\gamma$$

$$\Rightarrow d\left(\frac{\gamma}{g} R T\right) = -\gamma dz$$

$$\left[R T d\gamma + \gamma R dT = -\gamma g dz \right] \times \frac{1}{\gamma g dT}$$

$$\frac{dz}{dT} = -\frac{R}{g} \left(\frac{T}{\gamma} \frac{d\gamma}{dT} + 1 \right)$$

Subs eq (4) in above expression yields

$$\frac{dz}{dT} = -\frac{R}{g} \left(\frac{n}{n-1} \right)$$

$$\text{or } \frac{dT}{dz} = -\frac{g}{nR} (n-1) \quad \text{--- (5)}$$

For $n > 1$, which normal case for lower portion of atmosphere (Troposphere) $\Rightarrow \frac{dT}{dz} < 0$ (negative)

This occurred from elevation $0_m - 11019 \text{ m}$.

For standard atmosphere ($P = 101.3 \times 10^3 \text{ Pa}$, sea level, $T = 15^\circ \text{C}$) and take $R = 286.8 \frac{\text{J}}{\text{kg} \cdot \text{K}}$ $\Rightarrow n = 1.235$, Thus $\frac{dT}{dz} = -0.0065^\circ \text{C/m}$

For adiabatic case where $n = 1.4$ $\frac{dT}{dz} = -0.0098^\circ \text{C/m}$

(3)

For 11019 m to 20000 m (Stratosphere) — الغلاف الزهري

the temperature has been observed to be essentially constant at $(-56.5^\circ\text{C}) \Rightarrow n=1.0$ and $\frac{dT}{dz} = 0.0$

وقد أُنشأ القرآن الكريم إلى حالة الانحجار في الطبقات العليا وذكر أن جبال من بلع (برد).

سورة النور

أَلَمْ تَرَ أَنَّ اللَّهَ يُزْجِي سَحَابًا ثُمَّ يُؤَلِّفُ بَيْنَهُ ثُمَّ يَجْعَلُهُ رُكَّامًا فَتَرَى الْوَدْقَ يَخْرُجُ مِنْ خِلَالِهِ وَيُنْزِلُ مِنْ السَّمَاءِ مِثْرًا فَجِبَالٌ مِنْهَا مِنْ تَرْدٍ مُصِيبٌ بِهِمْ مِنْ يَسَاءٍ وَيُضْرِبُهُ رَعْنٌ مِّنْ يَسَاءٍ يَكَادُ سَنَا بَرْقُهُ يَذْهَبُ بِالْأَبْصَارِ ﴿١٧﴾

example 1: Calculate the pressure and weight density of air in the u.s. atmosphere at altitude 10 km above sea level.

Solution u.s. standard atmosphere are:

$$P_1 = 101.3 \times 10^3 \text{ Pa}, \gamma_1 = 12.01 \text{ N/m}^3, T_1 = 15 + 273 = 288 \text{ K}$$

Since $\frac{P}{\gamma^n} = \text{constant}$, for first 11 km $n = 1.235$

$$\Rightarrow \frac{P}{\gamma^{1.235}} = C \Rightarrow C = \frac{101.3 \times 10^3}{(12.01)^{1.235}} = 4702.987$$

Note that $dP = -\gamma dz$ (from fluid principle)

$$\Rightarrow \frac{dP}{\gamma} = -dz$$

$$\text{Thus: } \left(\frac{P}{\gamma^{1.235}} \right)^{\frac{1}{1.235}} = (4702.987)^{\frac{1}{1.235}}$$

$$\left[\frac{P^{0.81}}{\gamma} = 941 \right] \times \frac{dP}{P^{0.81}}$$

$$\Rightarrow \frac{dP}{P^{0.81}} = \frac{941}{P^{0.81}} dP = -dz$$

$$\Rightarrow \int_{P_1}^{P_2} \frac{941}{P^{0.81}} dP = - \int_{z_1}^{z_2} dz$$

$$\Rightarrow z_2 - z_1 = 0 = \int \frac{941}{P^{0.81}} dP$$

$$10 \times 10^3 = \frac{941}{0.19} \left[(101.3 \times 10^3)^{\frac{3}{0.19}} - P_2 \right]$$

$$\Rightarrow P_2 = 26305 \text{ KPa}$$

هذا الحد نستبدل

$$z_2 - z_1 = \frac{n}{n-1} C^{\frac{1}{n}} \left[\frac{P_1^{\frac{n-1}{n}}}{P_1^{\frac{1}{n}}} - \frac{P_2^{\frac{n-1}{n}}}{P_2^{\frac{1}{n}}} \right]$$

نفسها: C ثابتة
دالة: $\frac{P}{\gamma^n} = C$

$$P_1 = 101.3 \times 10^3$$

④

$$\frac{\Delta T}{\Delta z} = -\alpha \Rightarrow \Delta T = -0.0065 \Delta z$$

$$T_1 - T_2 = 0.0065 (z_2 - z_1)$$

$$T_1 - T_2 = 10000 (0.0065)$$

$$15 - T_2 = 65$$

$$\Rightarrow T_2 = 15 - 65 = -50^\circ \text{C} \quad (223 \text{ K})$$

$$P = \frac{\gamma}{g} RT \Rightarrow \gamma = \frac{gP}{RT} = \frac{9.81 (26.805 \times 10^3)}{286.8 \times 223} = 4.03 \text{ N/m}^3$$

one can derive a direct relationship between P&T and solve it by applying the given data.

$$\frac{dP}{\gamma} = -dz \quad \text{and } \gamma = \frac{\rho P}{RT} \text{ from eq (2)}$$

$$\Rightarrow \frac{dp}{p} = - \frac{g}{RT} dz$$

but $\frac{dz}{dT} = \frac{1}{-\alpha}$ where $\alpha = 0.0065$ [from eq. (5) after subst.]

$$\frac{dP}{P} = \frac{g}{R\alpha} \left(\frac{dT}{T} \right)$$

by integrating both sides yields:

$$\ln\left(\frac{P_2}{P_1}\right) = \frac{g}{R\alpha} \ln\left(\frac{T_2}{T_1}\right) \quad \text{or}$$

$$\frac{P_2}{P_1} = \left(\frac{T_2}{T_1} \right)^{\frac{\gamma}{\gamma-1}}$$

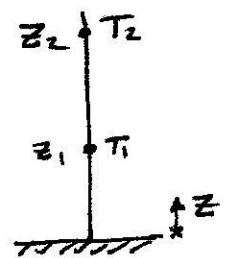
وکن علیہ استقام ت بار ۱۰

Now for example above:

Now for example above,

$$\frac{P_2}{P_1} = \frac{26305}{1013000}, \left(\frac{T_2}{T_1}\right)^{9/\gamma_a} = \left(\frac{T_2}{288}\right)^{5.2623}$$

$$T_2 = 223 \text{ K} \Rightarrow T_2 = -50^\circ \text{C}$$



example 2

(5)

Columns of air are heated by sun, above the standard values of atmosphere. So that the air is buoyant. If the air temperature is 17°C and it is buoyant rise is adiabatic, at what level will the rising air be the temperature and density in equilibrium [stop rising buoyantly]

sol.

$$T_{\text{standard}} = 15^{\circ} - 0.0065 Z \quad (\text{standard atmosphere}).$$

for adiabatic ($n=1.4$)

$$T_{\text{thermal}} = 17^{\circ} - 0.0098 Z \quad (\text{adiabatic condition}).$$

$$T_{\text{standard}} = T_{\text{thermal}} \quad [\text{for equilibrium } T_e \text{ \& } Z_e]$$

$$\Rightarrow 2 = 0.0033 Z_e \Rightarrow Z_e = 607 \text{ m.}$$

Density of vapor:

At same temperature and pressure, the specific gravity of water vapor to dry air is 0.622.

كثافة بخار الماء إلى كثافة الهواء الجاف تساوي (0.622) لنفس درجة الحرارة والضغط

$$\frac{P_v}{P_d} = 0.622 \quad \text{--- (7)}$$

Since P_d from eq. (2) is;

$$P_d = \frac{P_d}{R_d T}, \text{ therefore, for } e = P_d$$

$$P_v = 0.622 \frac{e}{R_d T} \quad \text{--- (8)}$$

According to Dalton's Law the pressure exerted by a gas (vapor pressure in this case) is independent on the pressure of other gases, hence;

$$P_a = P_d + e \quad \text{--- (9)}$$

moist air dry air Vapor

$$\text{or } P_d = P_a - e$$

$$\text{but } P_d = P_d + P_v \quad \left(P_a = \frac{m_e + m_d}{V} \right) \quad \begin{matrix} m_e = \text{vapor mass} \\ m_d = \text{dry air mass} \end{matrix}$$

(6)

$$\Rightarrow P_a = \frac{P_d}{R_d T} + \frac{0.622 e}{R_d T}$$

$$P_a = \frac{P_a - e}{R_d T} + \frac{0.622 e}{R_d T}$$

or: $P_a = \frac{P_a}{R_d T} \left(1 - 0.37 \frac{e}{P_a}\right)$ — (10)

equation (10) shows that moist air is actually lighter than dry air for same values of pressure and temperature.
 $P_a < P_d$ for $P_d = P_a$

The term $\left(1 - 0.37 \frac{e}{P_a}\right)$ is less than one, since $\frac{e}{P_a} < 1$.

Thus: $(P_a < P_d)$ أي أن الهواء الرطب أخف من الهواء الجاف.
 دليلاً على ذلك المعادلة (10) يوجد فيها أن $P_a < P_d$.

The R_d sometimes is slightly different with R_a in general one can assume $R_d = R_a$, if not

$$P_v = 0.622 \frac{e}{R_d T} = \frac{e}{R_v T} \Rightarrow R_v = \frac{R_d}{0.622}$$

Specific Humidity

The specific humidity, g_v is the ratio of density of vapor to a density of moist air. هي نسبة كثافة البخار إلى كثافة الهواء الرطب.

$$g_v = \frac{P_v}{P_a} \quad (11)$$

Since from eq (9); $P_a = P_d + e$

$$\Rightarrow P_a = P_d R_d T + P_v R_v T$$

$$P_a = \left[P_d + \frac{P_v}{0.622}\right] R_d T$$

where $R_v = R_d / 0.622$

From eq (10) and eq (11);

$$g_v = 0.622 \frac{e}{(P_a - 0.378 e)}$$

or $\left\{ g_v \approx 0.622 \frac{e}{P_a} \right\}$ — (13)

since, $P_a = P_a R_a T$ with recalling eq. (12)

$$\Rightarrow \left(P_d + \frac{P_v}{0.622}\right) R_d = P_a R_a$$

(7)

$$\Rightarrow \left(\frac{P_d}{P_a} + \frac{P_v}{0.622 P_a} \right) R_d = R_a$$

$$\text{but } P_a = P_d + P_v \Rightarrow \frac{P_d}{P_a} = 1 - \frac{P_v}{P_a} = 1 - q_v \quad (14)$$

$$\therefore \frac{R_a}{R_d} = \left(1 - q_v + \frac{q_v}{0.622} \right)$$

$$\text{or } R_a = R_d (1 + 0.608 q_v) \quad R_a \geq R_d \quad (15)$$

example:

If the moist air pressure ($P_a = 100 \text{ kPa}$), air temperature 20°C , Vapor pressure is 1871 N/m^2 , Calculate:

* Saturated vapor pressure, relative humidity, specific humidity and air density & dry density.

$$e_s = k e^{\frac{(17.27 T)}{237.3 + T(^\circ\text{C})}}$$

; $k = 611$ for $P = \text{N/m}^2$
 $k = 4.58$ for P in mm Hg

$$+ e_s = 611 e^{\frac{(17.27 \times 20)}{(237.3 + 20)}}$$

$$= 2339 \text{ N/m}^2 (\text{Pa})$$

$$+ \text{hr} = \frac{e}{e_s} = \frac{1871}{2339} = 0.8 = 80\%$$

$$+ q_v = 0.622 \frac{e}{P_a} = 0.622 \frac{1871}{100000} = 0.0116 \text{ Kg of vapor / Kg of moisture}$$

• q_v is the ratio of the mass of water vapor to the mass of dry air.

$P_a = P_a R_a T$; and From eq (15):

$$P_a \approx P_a \cdot R_d (1 + 0.608 q_v) T$$

$$100000 = P_a [286.8 (1 + 0.608 \times 0.0116)] (20 + 273)$$

$$10^5 = P_a (288.8) (293) \Rightarrow P_a \approx 1.180 \text{ Kg/m}^3$$

$$\text{From eq (14)} P_d = P_a (1 - q_v)$$

$$P_d = 1.18 (1 - 0.0116) = 1.168 \text{ Kg/m}^3$$

المطر القابل الهطول (8)

Precipitable water (MP) is amount (mass) of moisture in an atmospheric column. يعرف بأنه كمية الرطوبة (بما الماء) في عمود الهواء الجوي والتي تتحول إلى مطر وتسمى المطر القابل (القابل للتساقط) وهو يرمز له MP وتعبير بوحدة الكتلة فهذا كتلة عمود الهواء في (Kg in the column of atmosphere) إن كمية الماء الحظي في كتلة عمود الهواء الجوي تمثل معدل الرطوبة النوعية $\bar{q}_v$ average specific humid. ومن ثمة الآتي:-

$$\bar{q}_v = \frac{MP}{M_a} \Rightarrow MP = \bar{q}_v M_a \text{ or ;}$$

$$MP = \int_{z_1}^{z_2} \bar{q}_v \rho_a A dz \quad (16)$$

Eq(16) can be reduced to simple form by invoking ($dp = -\gamma dz$)

$$MP = \int_{z_1}^{z_2} \bar{q}_v \rho_a A \left(-\frac{dp}{\gamma_a} \right)$$

$$\text{or ; } MP = -\frac{A}{g} \int_{z_1}^{z_2} \bar{q}_v dp \quad (17)$$

Due to variation of $\bar{q}_v$, T , ρ_a and γ with variation of z as shown in Figure, Eq(16) & Eq(17) was recommended to solve MP incrementally as;

$$MP = \sum_{i=1}^n \Delta MP = \sum_{i=1}^n \bar{q}_{vi} \bar{\rho}_{ai} \Delta z_i, \text{ where } \bar{q}_{vi} \text{ is an average value over } \Delta z_i$$

$$MP = -\frac{A}{g} \sum_{i=1}^n \bar{q}_{vi} \Delta P_i \quad (18)$$

Example: Calculate the precipitable water for saturated atmospheric column up to 10km over 1m² area (ground surface) - The surface pressure is 101.3 kPa, and surface temperature is 30°C, lapse rate is (0.0065)°C/m ($R_d = 286.8 \frac{KJ}{Kg \cdot K^\circ}$).

Solution: Divide the column in 5 increments to get small value of $\bar{q}_v$ over increment

thus; $R_a = R_d$ since $R_a = R_d (1 + 0.608 \bar{q}_v)$ [acceptable assumption].

MP = 76.905 Kg in a column as shown in table below.

1	2	3	4	5	6	7	8	9	10
Z (km)	T (°C)	T (K)	P (pa)	e (pa)	ρ_a (kg/m ³)	q_v	$\bar{q}_v$	$\Delta P/g$	$-A/g \cdot \bar{q}_v (\Delta P)$
0	30	303	101300	4244.5	1.147	0.0261			
2	17	290	80425	1938.4	0.958	0.0150	0.0205	-2127.9	43.679
4	4	277	63179	813.5	0.791	0.0080	0.0115	-1758.0	20.217
6	-9	264	49059	309.3	0.646	0.0039	0.0060	-1439.4	8.586
8	-22	251	37611	104.6	0.522	0.0017	0.0028	-1167.0	3.298
10	-35	238	28430	30.8	0.416	0.0007	0.0012	-935.9	1.125
								$\sum MP =$	76.905

$$\Delta z = \frac{10}{5} = 2 \text{ km ; } T_{i+1} = T_i - \alpha(z_{i+1} - z_i) \text{ ; } T_{10} = T_c + 273 \text{ ; } P_{i+1} = P_i \left(\frac{T_{i+1} K^\circ}{T_i K^\circ} \right)^{\frac{g}{R_d \alpha}}$$

$$e = 611 \exp \left(\frac{17.27 T_c}{237.2 + T_c} \right) \text{ ; } \rho_a = \frac{P_a}{R_d T_{10}} (1 - 0.378 \frac{e}{P_a}) \text{ ; } \bar{q}_v = \frac{e}{P_a} = 0.622 \frac{e}{P_a}$$

Kg, in column of atmosphere

$$\text{Col. 1: } \Delta z = \frac{10 \text{ km}}{5} = 2 \text{ km}$$

$$\text{Col. 2: } T_2 = T_1 - \alpha(z_2)$$

$$\text{Col. 3: } T K^\circ = T C^\circ + 273$$

$$\text{Col. 4: } P_2 = P_1 \left(\frac{T_2 K^\circ}{T_1 K^\circ} \right)^{(3/\alpha \Delta z)}$$

$$\text{Col. 5: } e = 611 \exp \left(\frac{17.27 T C^\circ}{237.3 + T C^\circ} \right)$$

$$\text{Col. 6: } P_a = \frac{P_2}{R_d T_{K^\circ}} \left(1 - 0.378 \frac{e}{P_a} \right)$$

$$\text{Col. 7: } q_u = \frac{A_c}{P_a} = 0.622 e / P_a$$

$$\text{Col. 8: } \overline{q_u} = (q_{u,i} + q_{u,im}) / 2$$

$$\text{Col. 9: } \frac{P_{em} - P_i}{g}$$

$$\text{Col. 10: } (\text{Col. 8} \times \text{Col. 9})$$

which the sum of
all cells of col. 10
is equal to ΣM_p .

Note that $A = 1.0 \text{ m}^2$