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Energy performance and climate dependency of technologies for fresh water production from atmospheric water vapour

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Extraction of water vapour from atmospheric air and condensing it to liquid water for human usage is an imaginative solution to the water scarcity problem. Atmospheric water vapour is a large and readily accessible fresh water source able to fulfil human water needs. Many systems that draw water vapour from the air with water collecting surfaces, desiccant materials such as zeolites, silica gels, MOFs, polymers and salts and aids such as membranes have been proposed. Much progress has been made in increasing water harvesting efficiency, reducing cost and improving applicability especially in the extreme atmospheric conditions of arid regions. But all these systems are energy intensive and this energy demand for water production is an important element of the water-energy nexus. In this paper the intrinsic energy requirements of water vapour capturing processes in different atmospheric conditions are quantified as the specific water yield ($\text{L kW}^{-1} \text{h}^{-1}$). Distinction is made between passive systems that use natural phenomena like solar energy directly, and active systems with human transformation of the energy vector. The generation of thermoelectric energy involves water use and may even lead to overall water consumption instead of production. Technologies involving air cooling to provoke condensation of the water vapour reach specific water yields of $1\text{--}4 \text{ L kW}^{-1} \text{h}^{-1}$ but their application is strongly dependent on atmospheric conditions. A specific water yield of $0.1\text{--}1 \text{ L kW}^{-1} \text{h}^{-1}$ is commonly achieved for an ad/absorption-desorption cycle with a desiccant material. Depending on climate conditions, either passive systems with desiccants or active cooling of condensation surfaces is energy wise the optimum choice. The intrinsic energy requirements of atmospheric water harvesting are more than hundred times larger than seawater desalination. Fundamentally new concepts are needed to make atmospheric water an affordable fresh water source.

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Water impact

Water-from-air technologies have the potential to overcome water scarcity at any geographic location. Existing technologies relying on absorption, adsorption or condensation of water vapour contained in the atmosphere are very energy demanding and complicate the water-energy nexus. New emerging concepts in materials science pave the way to new less energy demanding sustainable water production technologies.

Introduction

Human society critically depends on the availability of fresh water. Fresh water unfortunately represents only a small fraction of the water available on earth. Most of the water is saline. According to the global water distribution on Earth (Fig. 1), saline water in seas and oceans accounts for *ca.* 97.5% of the 1.4×10^{18} ton of water on our planet. Only 2.5% of all water is fresh water. Close to 70% of it is frozen in glaciers, ice caps, and permafrost and is not readily available for humans. Another 30% is located in underground lakes. The most useful part of the fresh water on earth representing only 0.4% of the total is di-

vided over lakes (67.4%), soils (12.2%), swamps (8.5%), rivers (1.6%) and atmospheric water vapour (9.6%). A small amount of *ca.* 0.8% is contained in living organisms.^{1–3} These figures show that although our planet has plenty of water, fresh water resources and especially the readily available ones such as lakes, rivers and accessible ground water are only a tiny part of the global water resources.

Water on Earth is subjected to a hydrological cycle in which water continuously changes geographical location by evaporation or sublimation, atmospheric transport, and deposition elsewhere by condensation or freezing. This hydrological cycle interconverts saline and fresh water. The hydrological transportation cycle moves a quantity of 45 500 billion metric ton (bmt) of water yearly.³ The residence time of water in the stages of this cycle varies from a short

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average stay of 9 days in the atmosphere to thousands of years in oceans, seas and underground lakes.^{3,4} The global annual water withdrawal by human activity amounts to *ca.* 4000 bmt per year, much smaller than the mass of water engaged in the hydrological cycle. Water withdrawn from surface and underground fresh water sources is either consumed or returned to the source short after usage. The consumed part is evaporated or transported to elsewhere in the world in goods and food in which it is contained.^{2,3} Human interventions for water production accelerate the hydrological cycle as it reduces the water residence time in fresh water states such as ground water. Human water use, even with increasing demand in the future by a growing human population will not disturb the global natural hydrological cycle given its magnitude. Ultimately all water withdrawn by human activity is rendered to the hydrological cycle. The water supply problem arises from the uneven spreading of the hydrological cycle on earth. In many places on a local scale the natural water supply does not match the fresh water demand at all times.^{2,3,5}

Surface water accounts for about two third of the fresh water withdrawn by humans, and water from underground reservoirs for the other third.² The volume of water withdrawal according to different types of human activities in 2014 and a prediction for 2040 are presented in Fig. 2. The agriculture sector currently is by far the main fresh water user, with a volume of *ca.* 2800 bmt, representing a share of *ca.* 70%.^{2,6} The water intensity in this sector is evident considering *e.g.* the water demand of the production of a kilogram of beef cattle which requires about 15 000 L water.^{7,8} In 2014, the water volume for municipal usage, energy generation and industry were about equal (Fig. 2). The energy sector requires water for primary energy production and especially for cooling in thermally driven electricity production. In the chemical industry, water serves as a solvent, and it is a source of H and O atoms for chemicals and materials.²

The share of withdrawn water returned to the source is dependent on the application (Fig. 2). Irrigation in agriculture is an application where an important share of the water is not returned locally. In the municipal, industrial and power



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Dr. Jan Rongé holds a PhD in Bioscience Engineering since 2016. He is now a postdoctoral fellow of the Research Foundation – Flanders (FWO) at COK-KAT, KU Leuven. His main research interests are photo-electrochemistry, vapour phase electrocatalysis and water capture from air. In 2014, he demonstrated the first device producing hydrogen from sunlight and water vapour captured from ambient air. The team further devel-

oped this concept towards a full-scale panel and they presented a first prototype in 2018.



Johan Martens

Johan Martens is an expert in water and solar fuels. The atmosphere is a source of the essential chemical elements hydrogen, oxygen, nitrogen and carbon. Capturing water from atmospheric air is important for fresh water production, but also for sustainable synthesis of chemicals, like hydrogen and oxygen gas and ammonia using solar energy. This paper is one of a series of contributions on radical innovation in sustainable basic

chemicals and fuels synthesis from atmospheric air compounds. Johan Martens is holder of an ERC Advanced Grant on this topic.



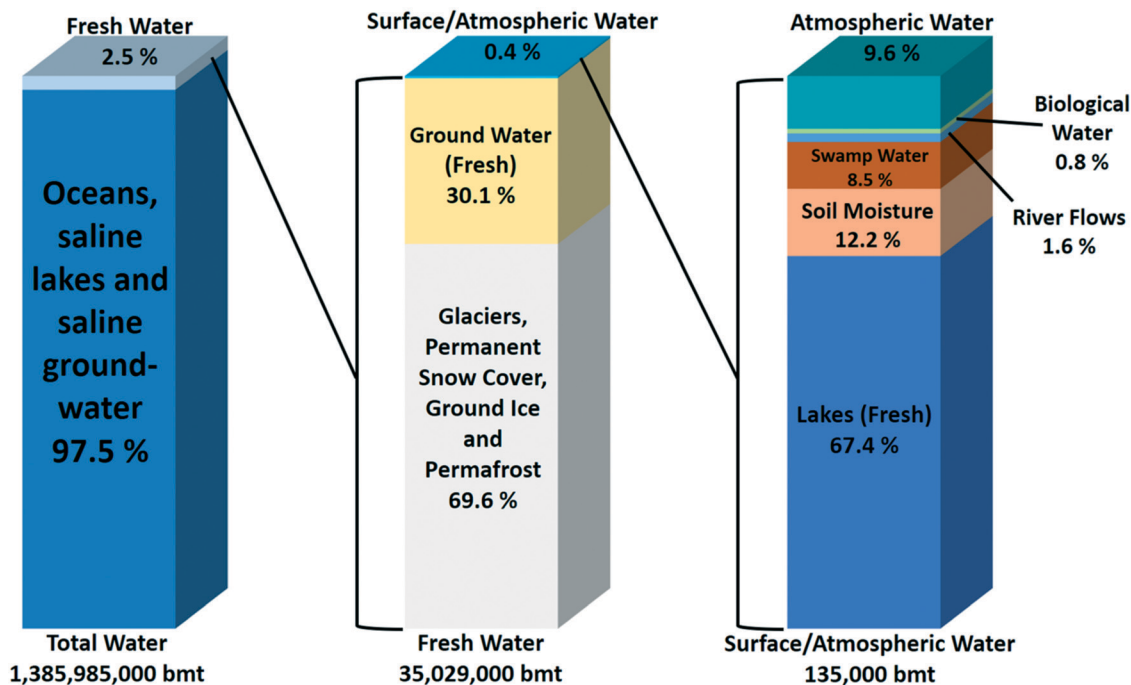


Fig. 1 Estimated global water distribution.¹

and energy applications, the share of returned water is larger. Nevertheless, overall the consumed amounts are substantial, and this volume is expected to increase by the year 2040 (Fig. 2). An increase by 10% of water withdrawal and a 20% rise in water consumption is expected to occur in the period from 2014 to 2040.² Municipal water use is primarily dependent on the living standard. Developing countries are facing a strong rise in water needs for the municipal sector.^{2,6,9}

With a growing human population water scarcity is a growing global concern and it is an obstacle for sustainable development. Water scarcity occurs when water resources are below 1000 m³ *per capita* per year.^{2,5} Currently, water shortage affects about one billion people worldwide and more than three billion people are expected to face this problem by

2025.² A growing world population, more intensive water usage *per capita*, desertification and an increasing salt content of fresh water aggravate the water scarcity problem.^{10,11} The most severe and extreme conditions occur at the local scale, and the ongoing climate change deepens the severity of these local problems.^{6,12}

Water management is complex because all users compete for the same water resources.⁶ Water is an atypical commodity since scarcity of this good is not always reflected in its price. Artificial low pricing leads to unsustainable and inefficient water usage.^{2,11} Because of government subsidies water prices are often artificially low, like for energy. Countries with the lowest water and energy prices rank the highest in terms of water consumption and tend to have the least sustainable water production. The strong correlation between water and energy demand complicates the problem even more. The water to be returned to the natural hydrological cycle is in most cases polluted and needs wastewater treatment. Water purification is also energy demanding. Increasing volumes entail enhanced energy consumption and consequently, additional water usage in cooling processes.² Mastering of this water-energy nexus is essential to achieve sustainable resource management.

There are three ways to resolve local fresh water shortage. First, when the supply from local fresh water sources is insufficient, water transportation over long distances from areas with water abundance is an obvious solution. Second, seawater is an inexhaustible water source (Fig. 1). Desalination of seawater and transportation with pipelines inland is an example of this solution.^{9,13,14}

Finally, an important observation is that atmospheric water is largely sufficient to cover the human fresh water needs (Fig. 1). Liquefaction of water vapour captured locally from

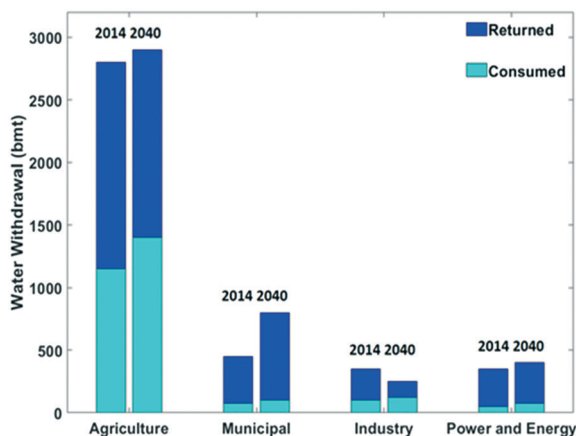


Fig. 2 Water withdrawal in different sectors split into a consumed and returned fraction in billion metric ton (bmt) in 2014 and a forecast for 2040. Data extracted from the IEA (2016).²



atmospheric air is a way to avoid the need of long distance transportation, because it is ubiquitous. Water vapour in the atmosphere is rather clean, and after condensation an attractive source of fresh water, similar to natural precipitation. The atmosphere is an inexhaustible source of water because of the hydrological cycle which replenishes the atmosphere swiftly with water vapour.¹⁴ The idea of a decentralised water supply system relying on water extraction from the air has been appealing for decades. Some authors forecasted a bright future for this technology.^{15–17} The main obstacle for widespread implementation of water-from-air technology is the high energy demand, although at locations with abundant solar energy this is seen as a minor concern.^{16,18}

In this review a thermodynamic analysis of the energy requirement of water production from air is provided. The boundary conditions depending on geographical location can vary substantially and the variability of the water content of atmospheric air and climatological conditions are part of the analysis. Implementing water-from-air technology would have a large impact on the water-energy nexus.

Parameters of water vapour in atmospheric air

Relative humidity (Φ), absolute humidity (ω) and dew-point temperature (T_d) are the essential properties of atmospheric air to be used as source of water. Air is a gas mixture, composed of nitrogen, oxygen and argon gas as major components (in this order), and water vapour in variable contents. The relative humidity (Φ) is the ratio of the partial pressure of water vapour (P_w) over the saturation pressure (P_s) (eqn (1)).^{19,20}

$$\Phi = \frac{P_w}{P_s(T)} \quad (1)$$

The dew-point (T_d) is the temperature at the onset of condensation and at which the relative humidity reaches 100%. Relative

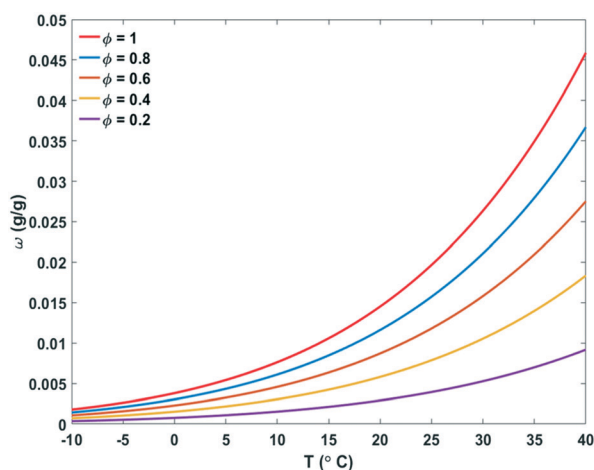


Fig. 3 Absolute humidity (ω) and relative humidity (Φ) of air depending on temperature (T). Unsaturated air becomes saturated by lowering the temperature or by increasing the absolute humidity.²⁰

humidity and temperature determine the absolute amount of water contained in the air, denoted absolute humidity (ω). The absolute humidity represents the maximum extractable amount of water. The absolute humidity at different relative humidity levels is plotted against temperature in Fig. 3. The upper curve at 100% relative humidity is the saturation curve where condensation occurs.^{19,20} Unsaturated air can reach saturation by cooling or by humidification.

The mathematical relation between relative humidity (Φ), absolute humidity (ω), total air pressure (P) and temperature (T) is given in eqn (2). The dew-point T_d can be obtained from this equation by solving for T and setting $\Phi = 1$ for a given absolute humidity and air pressure.²⁰

$$\Phi = \frac{\omega P}{(0.622 + \omega)P_s(T)} \quad (2)$$

The water vapour saturation pressure P_s at temperature T ($^{\circ}\text{C}$) can be described mathematically with the empirical AERK Magnus formulation, such as *e.g.* in the range of $T = [-40\text{ }^{\circ}\text{C}, 50\text{ }^{\circ}\text{C}]$ by eqn (3).²¹

$$P_s(T) = 610.94 \exp\left(\frac{17.625T}{243.04 + T}\right) \quad (3)$$

The total air pressure P is simply the sum of the partial pressure of dry air P_a and P_w , the vapour pressure of water in the air (eqn (4)).²⁰ For simplicity, P_a is taken as 100 000 Pa, *i.e.* at 1 bar.

$$P = P_a + P_w \quad (4)$$

The total moist air enthalpy consists of a dry air term H_a and a water vapour term H_{wv} (eqn (5)). The heat capacity of air $C_{p,a}$ is taken as a constant of $1.005\text{ kJ kg}^{-1}\text{ }^{\circ}\text{C}$, which is a reasonable assumption in the temperature range $[-10\text{ }^{\circ}\text{C}, 50\text{ }^{\circ}\text{C}]$. The average heat capacity of water vapour $C_{p,wv}$ in the interval $[-10\text{ }^{\circ}\text{C}, 50\text{ }^{\circ}\text{C}]$ equals $1.82\text{ kJ kg}^{-1}\text{ }^{\circ}\text{C}$ and the water vapour enthalpy at $0\text{ }^{\circ}\text{C}$ is 2500.9 kJ kg^{-1} .²⁰

$$H = H_a + \omega H_{wv} \quad (5)$$

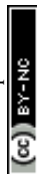
$$H_a = C_{p,a}T \quad (6)$$

$$H_{wv} = H_{wv}(0\text{ }^{\circ}\text{C}) + C_{p,wv}T \quad (7)$$

The heat of condensation of water (ΔH^{vap}) at atmospheric pressure and normal boiling temperature ($100\text{ }^{\circ}\text{C}$) equals 2257 kJ kg^{-1} , which is a large amount of energy.²² This large amount of energy has to be removed to enable the phase change of water from vapour to liquid.

Principles of water extraction from air

Atmospheric water generators can be divided into categories (Fig. 4). Water vapour can be extracted from the air either by



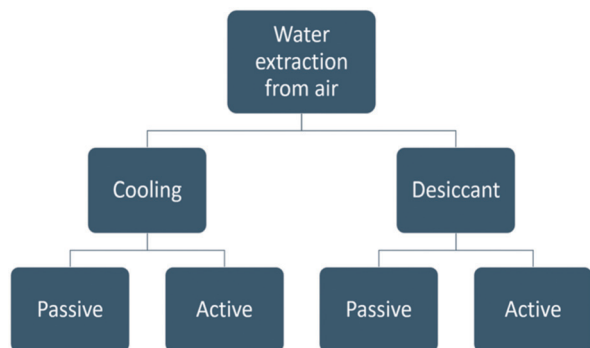


Fig. 4 Subdivision of water-from-air extraction systems.

cooling and condensing moisture out of the air, or else, by using a desiccant material. Each category is subdivided into a passive and active way to apply the water extraction method. The term passive indicates water is captured with the help of natural phenomena and without an energy delivery system. In the active case, an external energy vector is implemented to assist the water extraction process. The two most important principles are active cooling and passive water extraction by water vapour ad- or absorption on a desiccant.

In the next sections the energy requirements of the different concepts are discussed. A theoretical performance evaluation will be based on the specific water yield SY_{water} ($\text{L kW}^{-1} \text{h}^{-1}$) which represents the volume of liquid water which can be produced with an energy input of 1 kW h.

Technologies for water extraction from the air

Passive cooling technologies

Dew and fog harvesting are examples of passive cooling. In this section, the advantages and disadvantages of water collection through artificial dew and fog formation and the way to generate the required spontaneous cooling are discussed.

Principle of dew formation. Passive radiative cooling of a condensation surface to a surface temperature below the

dew-point leads to deposition of liquid water originating from water vapour in the air (Fig. 5A). Insects adapted this principle to harvest water from atmospheric air. The Namib beetle is a popular example of an insect using passive dew harvesting to provide water for itself and to survive in extreme arid conditions.^{23,24}

Any surface outdoor emits energy by radiation of electromagnetic waves in the infrared wavelength region to the sky.²⁵ The available cooling power lies between 25 and 100 W m^{-2} on cloudless nights. A theoretical maximum water production of 0.8 L m^{-2} per day under most optimal conditions has been estimated.^{24,26} Consequently, a large surface area exposed to the sky is needed to collect artificial dew in significant amounts.²⁶

The dew harvesting process faces the problem of inconsistent production caused by the critical condition of sufficient radiative cooling to reach a surface temperature below the dew-point. In addition, wind may interfere with the condensation process. Air flow provides a larger flow of water vapour toward the cold surface, but depending on the temperature of the wind, it also can counteract the radiative cooling. Wind can lift dew drops from the surface and reduce this way the harvesting.^{26,27} The inconsistency of water production is the main reason why the technique is applied only in specific regions with suitable climate for dew formation.²⁸ The energy efficiency is less relevant for this system since no artificial energy input is needed.

Principle of fog collection. Fog consists of tiny water droplets suspended in the atmosphere. These droplets nucleate homogeneously in saturated air. Droplet nucleation is favoured by the presence of fine suspended particles acting as nucleation centre.²⁹

Fog collection is the capturing of the already existing liquid water droplets of air aerosol on a mesh material (Fig. 5B). The droplets grow by coalescence on the mesh and are finally detached from the mesh and collected by gravity.²⁹ The amount of fog that can be harvested depends on relative humidity, absolute humidity, wind velocity and the frequency of fog occurrence.²⁹ Fog harvesting, like dew collection, is a

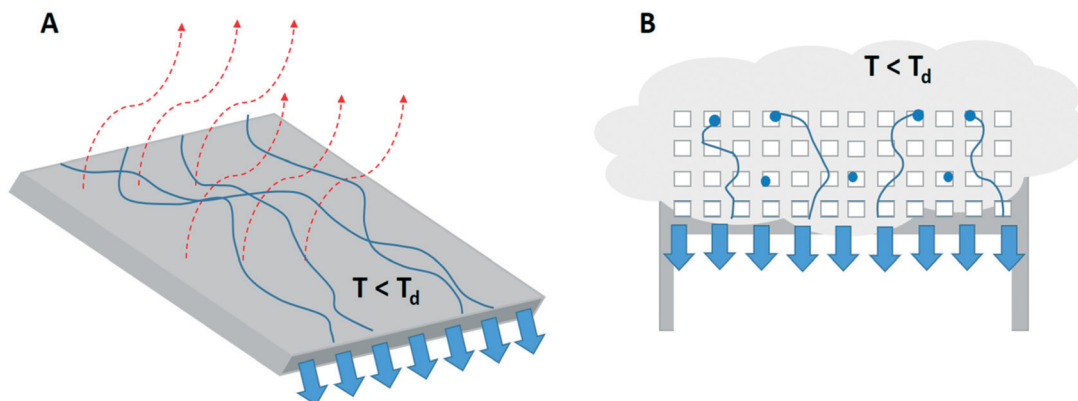


Fig. 5 A visual presentation of the process of A) dew harvesting by droplet formation and dripping off a surface cooled below the dew-point by passive radiative cooling (red arrows) and B) fog harvesting where fog droplets (in blue) hit a mesh structure, coalesce and detach from the mesh by gravity.^{23,24,29}



relatively inexpensive, easy and definitely helpful approach in combatting water scarcity, but it is critically depending on climatologic conditions.³⁰ The water collection capacity is depending on external factors rendering the water yield unpredictable.

Dew and fog collection devices and applications. In literature several reports on the performance of dew and fog collection devices are available. A typical water yield of 0.3–0.6 L m⁻² per day from passive dew harvesting has been achieved in arid environments.³¹ Berkowicz *et al.* were able to produce peak quantities of water amounting to 0.5 L m⁻² per day in Jerusalem. At that location the probability of dew formation on daily basis is around 50%.²⁸ The effectiveness of dew harvesting on islands with humid climate is also documented. In an investigation on French Polynesia Islands, on yearly basis the water volume from dew harvesting represented *ca.* 3% of the total rainfall.²⁶ In the dry seasons on islands of Croatia, the contribution of dew water compared to rainwater was in the range of 38–120%.²⁴ A dew plant built in Kothara (India) is able to collect 20 L of dew per square meter per year. Besides the collection of dew, the surface also serves a rain capturing function. A yearly average of 300 L m⁻² of rain can be collected.³² This reveals the limited added value of dew collection even in areas where rain falls only during a few days in a year.

Special surfaces have been designed for dew harvesting. Thickett *et al.* illustrated condensation rate improvements during dew harvesting by a micro patterned coating. Water condenses on the hydrophilic coating part and flows consecutively over the hydrophobic area when the droplet detaches from the surface.³³ Patterned polymers applied as a condenser coating were shown to promote water vapour condensation rates by 57%.³⁴ The principle is bio-inspired as the same principle is used by the *Stenocara* beetle in the Namib Desert.³³ Beysens *et al.* reached a 20% increase in dew yield by application of a condenser foil of TiO₂ and BaSO₄ microspheres in polyethylene.²⁷

Fog serves as a water source for plants and animals such as *e.g.* in the Atacama Desert in Chile. In these arid areas there is almost no rainfall. On coastal locations at altitudes of 600–1200 m in Chile fog stays in the air for a long time. In those areas, fog extraction can yield water at rates of 1 to 15 L m⁻² per day. A maximum of 6.3 L m⁻² per day was captured by a polypropylene coated mesh by Cáceres *et al.*²⁹ An average fog collection of 4.6 L m⁻² per day was reported in South Africa.³⁵ Commonly used mesh types are made of nylon or Teflon.³⁶ Park *et al.* optimised the mesh design to improve fog collection. Based on an extrapolation of reported experimental data at a site in Chile, a production of 12 L m⁻² per day is estimated.³⁰

Capturing fog requires less space compared to dew. Fog capture nets can be placed vertically which reduces the overall lateral space requirements.

When fog or dew is common at a specific location it can be a convenient way of providing water to small communities, but for agriculture purposes, the collected quantities are too low.³⁷ Temporal water storage is needed to equilibrate

supply and demand when using a dew, fog or even rainwater capturing approach.³²

Conductive heat loss to the underground is an alternative way of cooling a condensation surface. In a device operating according to this principle, air is forced to enter an underground metal coil connected to a reservoir for collecting the condensate. The metal coil is in thermal contact to the underground.^{38,39} The minimum required temperature difference between the underground and atmospheric air derived from the temperature dependence of the relative humidity (Fig. 3) is presented in Table 1.

Air at $T = 15\text{--}25$ °C needs to be cooled by *ca.* 14 °C at $\Phi = 40\%$, 8 °C at $\Phi = 60\%$ and 3.5 °C at $\Phi = 80\%$ to initiate condensation. This seems to be a small temperature difference, but it is difficult to maintain. Underground heat conduction is slow, and inefficient to dissipate the condensation heat. The temperature of the metal coil will be increased by the water condensation process. The temperature difference between air and underground condensation surface vanishes rapidly.

Consequently, the underground temperature needs to be below that of the condensation coil to keep the condensation coil below the critical temperature needed for water vapour condensation. Literature data on temperature differences between atmospheric air at ground level and underground at 2 m deepness are available for a location in Lecce, Italy.⁴⁰ Locations with different thermal conductivity of the underground are compared. The temperature difference shows seasonal fluctuations, as expected. The temperature difference between air and underground reaches a maximum of about 8 °C in July. Air at relative humidity below 60% will never reach saturation under these conditions. At 80% relative humidity, temperature differences of 3.5 °C will only be reached in certain months.⁴⁰ Active cooling of the condensation coils appears to be needed to reach the desired temperature differences.

Active cooling technologies

In this section, active air cooling technologies for water extraction from air are presented. The thermodynamics of these processes and the impact of the climate on the specific water yield and energy efficiency are analysed. Theoretical limitations of the specific water yield depending on ambient temperature and relative and absolute humidity are presented. Energy efficiencies of practical devices based on the reverse Carnot cycle, thermoelectric coolers, self-filling water bottles

Table 1 Required temperature difference to initiate condensation for different relative humidity and temperature conditions of ambient air

Φ_{amb} [%]	T_{amb} [°C]	$T_{\text{amb}} - T_{\text{d}}$ [°C]
80	25	3.7
80	15	3.4
60	25	8.3
60	15	7.7
40	25	14.5
40	15	13.5



Critical review

and membrane assisted water condensation technologies are estimated.

Process description and thermodynamic analysis. In most cases, active air cooling is based on an evaporation–condensation cycle of a coolant to extract heat from a cold reservoir (evaporator of the coolant) and transportation of the extracted heat to a hot reservoir (condenser of the coolant), illustrated in Fig. 6. The evaporation and condensation of the coolant are driven by decompression and compression, respectively. The air enters the refrigerant evaporator section, where it is cooled below the dew-point. In this way water vapour contained in the inlet air condenses and liquid water is harvested.²⁰

This water extraction concept is based on a reverse Carnot cycle of a coolant. The coefficient of performance of the cooling unit ($COP_{cooling}$) is defined in eqn (8), with T_c the temperature of the cold reservoir and T_h the temperature of the hot reservoir. The hot reservoir is typically at ambient temperature, T_{amb} .²⁰

$$COP_{cooling} = \frac{T_c}{T_h - T_c} \quad (8)$$

Typical refrigerants are 1,1,1,2-tetrafluoroethane, isobutane and ammonia.^{41,42} Advanced concepts for applying this principle maximizing $COP_{cooling}$ have been proposed.⁴³

Thermoelectric coolers (TEC) are an alternative to the evaporation–condensation cycle of a coolant. This electricity based technology forces heat transport by application of a voltage difference between a semiconductor pn-junction. The electric current causes cooling on one side of the semiconductor assembly, and heating on the other side.⁴⁴ TEC based cooling processes are generally less energy efficient, but the absence of a recirculating fluid is a practical advantage.⁴⁵ In-depth analysis of cooling technologies is beyond the scope of this review.^{43,46–48}

The general process scheme of an active cooling device to condense water vapour out of air is presented in Fig. 7. The theoretical maximum amount of liquid water that can be recovered from air for a given power supply in a 100% efficient process without heat recovery is given by SY_{water} , a convenient physical quantity to estimate the influence of the climate on the operation of an air cooling system:

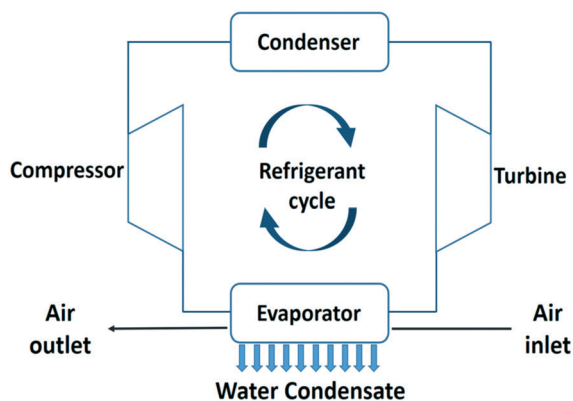


Fig. 6 Process flow diagram of an air cooler and water condensation unit using a refrigerant evaporation–condensation cycle.²⁰

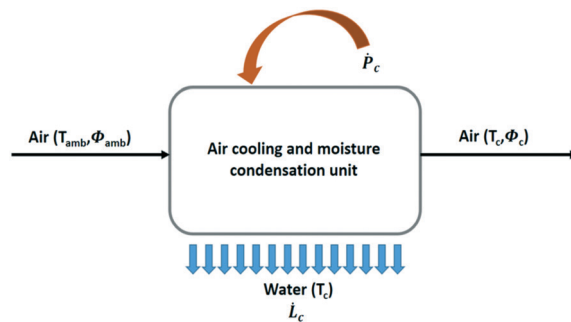


Fig. 7 Process diagram of a water from air unit with air cooling and water condensation.

$$SY_{water} = \frac{\dot{L}_c}{\dot{P}_c} \quad (9)$$

In which \dot{L}_c is the rate of water production ($L h^{-1}$) and \dot{P}_c , the total power supply (kW). \dot{L}_c and \dot{P}_c values are obtained from eqn (10) and (11), respectively.

$$\dot{L}_c = \rho_a \dot{V}_a (\omega_{amb} - \omega_c) \quad (10)$$

In eqn (10) ρ_a represents the air density ($kg m^{-3}$) and \dot{V}_a , the volumetric air flow rate ($m^3 h^{-1}$). The power supply \dot{P}_c has two contributions, *viz.* the energy needed for cooling and water condensation, and the ventilator energy, \dot{P}_v :

$$\dot{P}_c = \frac{Q_c}{COP_{cooling}} + \dot{P}_v \quad (11)$$

The theoretical energy needed for cooling and condensation Q_c (kW), expressed in eqn (12), is obtained from the heat balance over the air cooling and moisture condensation device (Fig. 7). $C_{p,w}$ ($kJ kg^{-1} K^{-1}$) stands for the heat capacity of liquid water and \dot{V}_a the volumetric air flow expressed in ($m^3 s^{-1}$).

$$Q_c = \rho_a \dot{V}_a (H_{amb} - H_c - (\omega_{amb} - \omega_c) C_{p,w} T_c) \quad (12)$$

Dividing Q_c by the performance of the Carnot refrigeration cycle, $COP_{cooling}$ (eqn (8)) gives a value for the energy need for cooling and condensation (eqn (11)).

The condensation temperature T_c has a strong impact on the specific water yield. This is illustrated by comparing two cases in Fig. 8. The first case deals with a humid hot climate characterized by high relative humidity of 80% at a temperature of 25 °C. The second case refers to a desert climate with low relative humidity of 30% at a temperature of 30 °C. Temperatures below 0 °C in the air cooling and moisture condensation unit were not considered because these would involve the complication of ice formation. Practically, it would complicate the water production. Ventilator power values \dot{P}_v and air flow rates \dot{V}_a were taken from axial ventilator specifications.⁴⁹

In the hot humid climate case the dew-point T_d is *ca.* 21 °C (Fig. 8A). The optimal condensation temperature for



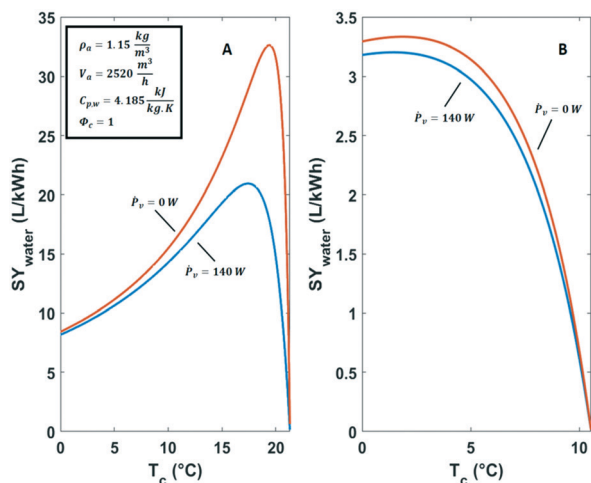


Fig. 8 Specific water yield of active cooling technologies against the condensation temperature in A) a humid climate ($\phi = 80\%$, $T = 25\text{ }^{\circ}\text{C}$) and B) a dry climate ($\phi = 30\%$, $T = 30\text{ }^{\circ}\text{C}$).

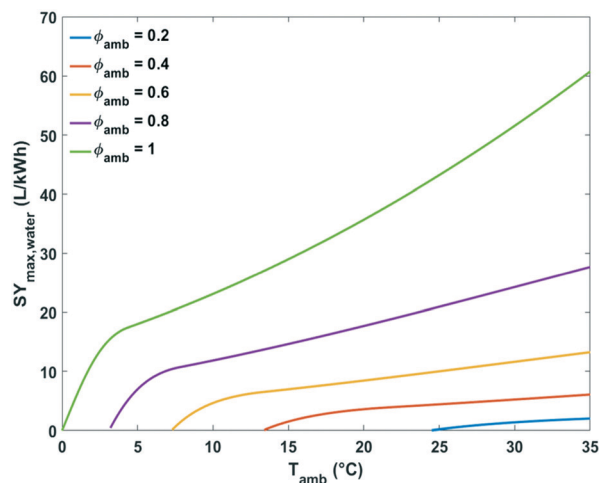


Fig. 9 Plot of maximum specific water yield as a function of ambient temperature at different relative humidity levels for water extraction from the air by active cooling.

energy efficient water capturing from air is lower than the dew-point. The specific water yield under these conditions shows a clear maximum at $17\text{ }^{\circ}\text{C}$ (Fig. 8A). In the situation of Fig. 8A the maximum specific water yield ($\text{SY}_{\text{max,water}}$), amounts to $21\text{ L kW}^{-1}\text{ h}^{-1}$. When the ventilator power \dot{P}_v (eqn (11)) is omitted the optimum specific water yield is $32.6\text{ L kW}^{-1}\text{ h}^{-1}$. The situation in a dry climate is different (Fig. 8B). There is a shallow optimum energy efficiency around $3\text{ }^{\circ}\text{C}$. The yield of liquid water per kW h is small and does not surpass $3.3\text{ L kW}^{-1}\text{ h}^{-1}$ (Fig. 8B). This is mainly a result of the high sensible heat removal requirements.⁵⁰ The contribution of \dot{P}_v is negligible in this case.

In both climates (Fig. 8), water production \dot{L}_c can be raised by reducing the condensation temperature T_c at the cost of increasing the power supply and operating at lower energy efficiency.

The impact of ambient temperature, absolute humidity and relative humidity on the energy efficiency of water production from air by cooling and condensation was analysed in more detail. The maximum specific water yield for a given climate condition was estimated at the corresponding optimal condensation temperature $T_{c,\text{opt}}$ (eqn (13)).

$$\text{SY}_{\text{max,water}} = \text{SY}_{\text{water}}(T_{c,\text{opt}}) \quad (13)$$

For relative humidity values of 20, 40, 60, 80 and 100%, the specific water yield at optimum condensation temperature is plotted against ambient temperature in Fig. 9. The higher the relative humidity, the higher the maximum specific water yield, and the more water can be condensed from air at a given ambient temperature. Evidently, at 100% relative humidity, it takes little energy to produce dew on the collector surface.

At the lowest possible ambient temperatures for each case the maximum specific water yields are very low. One of the reasons is that the absolute water content of cold air is low. In addition, the most optimal condensation temperature in

this instance lies below $0\text{ }^{\circ}\text{C}$ which is impractical. The minimal ambient air temperature necessary for avoiding water freezing in the cooling section of the device at optimal energetic performance was estimated at 3, 7, 13 and $24\text{ }^{\circ}\text{C}$ for relative humidity of 80, 60, 40 and 20%, respectively (Fig. 9).

Above these critical temperatures, the specific water yield increases significantly with increasing ambient temperature. The higher the relative humidity, the more significant the increase (Fig. 9).

Comprehensive plots showing the influence of absolute humidity at different relative humidity values and relative humidity at different ambient temperatures on maximum specific water yield are presented in Fig. 10 and 11, respectively. The specific water yield is little influenced by absolute humidity (Fig. 10). Relative humidity governs the energy intensity to extract water from the air with active cooling

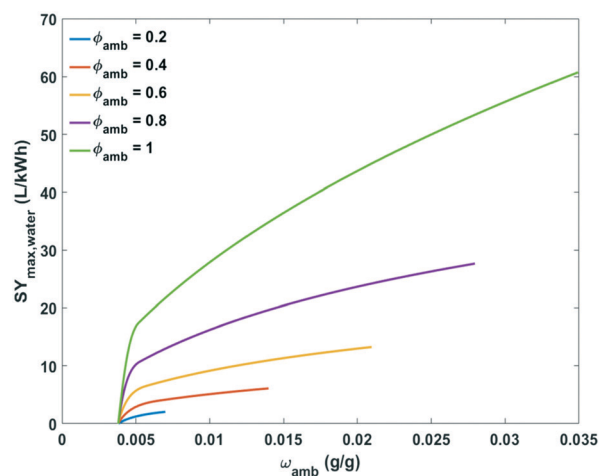


Fig. 10 Plot of maximum specific water yield against absolute humidity at different relative humidity levels for water extraction from the air by active cooling.



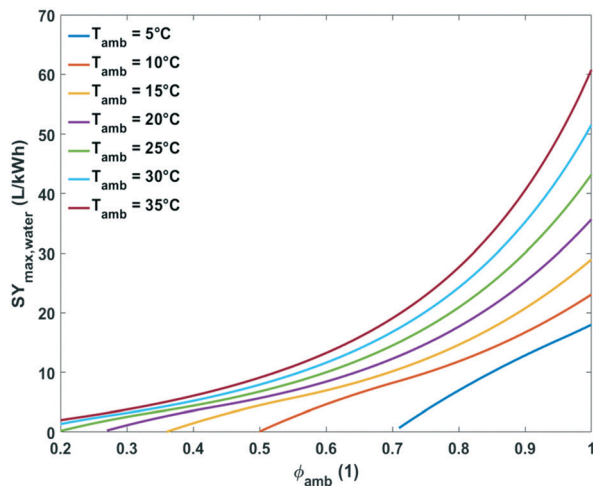


Fig. 11 Plot of maximum specific water yield against ambient relative humidity at different ambient temperature levels for water extraction from the air by active cooling.

technology. The maximum specific water yield increases with relative humidity in an exponential way (Fig. 11).

Active cooling devices. Water extraction from air is very common in air conditioning. Dehumidifiers based on the process of active air cooling to force condensation of water vapour contained in the air are established technology. Air dehumidifiers have however been designed for energy efficient air conditioning, and not for maximizing the water yield. Adaptation according to the purpose of the device is needed. Dehumidifiers are rated according to the energy needed for water removal.⁵¹ The specific water yields of devices designed for air conditioning are in the range of 1.7–4.2 L kW⁻¹ h⁻¹.^{52,53} Devices designed for air dehumidification when applied for water production likely are operated at suboptimal conditions and fixed T_c values, and produce water at the expense of increasing the energy consumption.

Some performances of devices are given in Table 2. Provided the relevant data were available, the efficiency (eqn (14)) of reported devices was estimated based on reported specific water yield and an estimated theoretical maximum, defined in eqn (13):

$$\eta_c = \frac{SY_{\text{water}}}{SY_{\text{max,water}}} \quad (14)$$

The efficiency of commercial dehumidifiers is *ca.* 40% (Table 2). This relatively high efficiency can be explained by the maturity of the technology. For devices dedicated to water

Table 2 Specifications of commercial dehumidifiers (D) and active condensation technologies (C) to extract water from the air

Type/ref.	\dot{P}_c (W)	\dot{L}_c (L d ⁻¹)	SY_{water} (L kW ⁻¹ h ⁻¹)	η_c (%)	Measuring conditions (ϕ , T)
(D) ⁵³	530	50	4.2	40.2	60%/27 °C
(C) ⁵⁹	450	32	2.96	—	—
(C) ⁶⁰	—	800	4	—	—
(C) ⁵⁵	61	1.58	1.1	4.3	80%/33 °C

production insufficient data are available for making the estimation.

Portable self-filling water bottles are miniaturised versions of the air cooling and moisture condensation technique. Air driven by a fan is sent inside the bottle where the moisture condenses onto thermoelectric cooling surfaces. The device is powered by a small photovoltaic (PV) panel to minimize size and weight which is essential for a portable bottle.⁵⁴ A power input of 313 W is required to produce 0.5 L h⁻¹ when only condensation heat needs to be removed and when taking a maximum COP_{cooling} value of 1 for a TEC device.^{22,45,55} At full irradiation, a PV panel typically delivers 180 W m⁻², therefore around 1.7 m² of solar panel is required to supply enough power.

The specific water yield of TEC based air cooling is estimated to be in the range of 0.14–1.1 L kW⁻¹ h⁻¹.^{55–58}

Membrane assisted water condensation. Membranes are capable of selectively separating water molecules from the air. The membrane separation concept in combination with active cooling is shown in Fig. 12. Air moisture permeates from the feed side to the permeate side by a pressure difference generated by a vacuum pump. The performance critically depends on the water permeability of the membrane, which needs to favour water permeation over the other gas components of air. The permeate is transferred to the condenser to produce liquid water. This setup reduces the total cooling requirement by increasing the relative humidity of the inlet air of the condenser.¹⁰

The addition of a membrane module (Fig. 12) has been reported to double the specific water yield. The specific water yield of the modelled membrane dehumidifier setup reached 6.18 L kW⁻¹ h⁻¹ with a 5000 m² membrane surface area.¹⁰ A liquid membrane with hygroscopic salts on a carbon black nanoparticle support has been reported effective in this application.⁶¹ Polymer electrolyte membranes are used for electrochemical based air dehumidification, an alternative to the pressure driven membrane dehumidification process. In this process a net water flux over the membrane is created by splitting water electrochemically into oxygen gas and protons at the anode side of the membrane and combining these protons again with the oxygen gas at the cathode side to make water. In addition, water moves through the membrane by the effect of electro-osmosis. Back diffusion lowers the performance of this set-up.⁶²

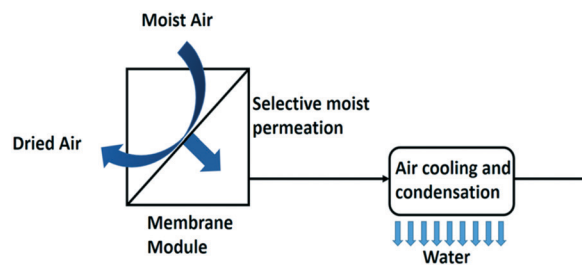


Fig. 12 Membrane assisted water condensation setup for water extraction from air.¹⁰



Conclusions. Active air cooling technologies to cause water vapour contained in air to condense is an established technology in air conditioning and it can be adapted to serve water production. The optimum settings for water production and air conditioning are different, and dedicated devices according to the application are needed. Specific water yields are in the range $2\text{--}4\text{ L kW}^{-1}\text{ h}^{-1}$. Miniaturised portable systems exist, but are less energy efficient.

Hot humid climates like in tropical regions offer the best use cases for this way of producing water. Water scarcity rarely occurs in such climates and the need of water-from-air technology is limited. Close to 100% relative humidity, active cooling may not be needed as dew and fog formation frequently occur under these conditions, and these are easier to collect than water vapour. At the other extreme, in dry climates the dew-point of humid air is below $0\text{ }^{\circ}\text{C}$ which makes the cooling concept impractical because of the need of defrosting the condenser for collecting the harvested water. The condensation temperature could be enhanced above freezing temperature by implementation of a membrane module to increase the relative humidity prior to condensation. The membrane/condenser combination significantly improves the specific water yield. The reduction of energy consumption in the cooling unit can balance the additional investment in a membrane module.

Passive desiccant technologies

Trapping water in solid and liquid desiccants is an alternative to condensation by cooling. Liquid desiccants take up water by absorption. Solid desiccants adsorb water molecules in their porous matrix. Especially in dry and arid regions where the water condensation temperature is too low for practical application, the use of desiccants is an attractive alternative. In this section, the thermodynamics of water-from-air technologies with desiccants are reviewed. The use of the day–night cycle for reversing the water sorption process is discussed.

Process description and thermodynamic analysis. The temperature difference between day and night can be used to conceive an adsorption–desorption cycle, illustrated in Fig. 13.⁶³ The desiccant adsorbs water vapour from the air at night. During the day, the temperature of the reservoir with saturated adsorbent rises by solar heating causing desorption of water. The desorbing water vapour is collected by condensation on a colder surface in a separate cold reservoir.⁶³ In passive systems, the condenser temperature is critical.^{64,65} Shielding the water condensing surface from sunlight is needed to produce water.⁶⁶ Alayli *et al.* in 1987 were among the first to describe such passive water extraction system from air using a desiccant.⁶³

The determination of the water extraction capacity from adsorption and desorption branches of the adsorption isotherm are explained in Fig. 14. During the night the desiccant is saturated with water vapour according to the isotherm pertaining to the cold temperature, T_{ads} . During the day, the volume holding the saturated desiccant is heated by sunlight to T_{des} , which also lowers the relative humidity. This causes desorption according to the adsorption isotherm at the desorption temperature. The difference between both states (q_{ads} at T_{ads}) and (q_{des} at T_{des}) defines the extractable amount of water in a night and day cycle.⁶⁶ The adsorption isotherm of a desiccant at T_{ads} and T_{des} defines the suitability of an adsorbent to serve atmospheric water harvesting.

The shifting of the adsorption isotherm on the relative humidity axis with temperature is dependent on the isosteric heat of adsorption.⁶⁷ A large isosteric heat of adsorption results in a large shift and a large amount of water that can be produced in the cycle.

The theoretical maximum specific water yield of passive desiccant based systems was estimated for the process and its parameters depicted in Fig. 13. If included, a ventilator with the same specifications as for the previously discussed active air cooling systems was considered (see legend of Fig. 8). The fan delivers air to the desiccant during the night and transports the hot humid air to the

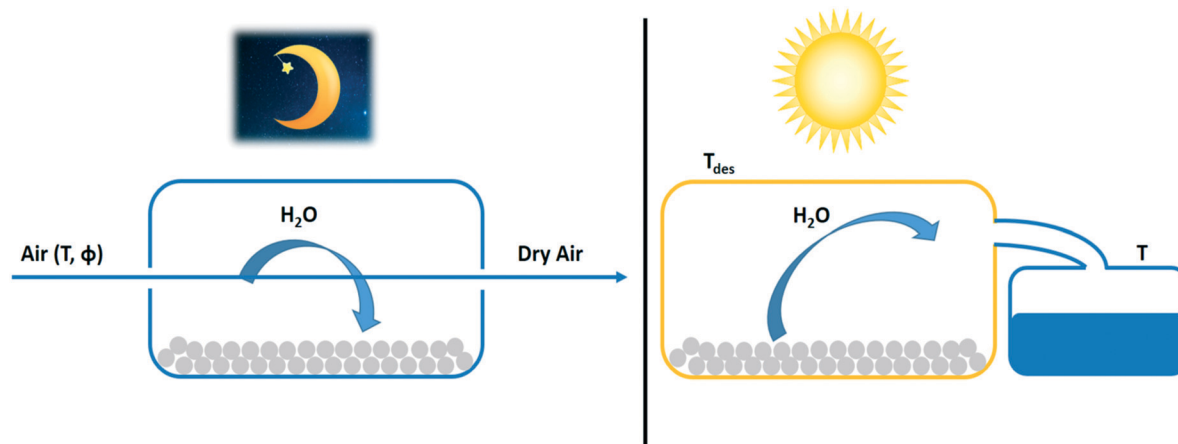


Fig. 13 Night and day cycle of passive water extraction from air with a desiccant (grey spheres). Water adsorbs at night from cold air on desiccant material and desorbs by solar heating during the day. Water vapour desorbing from the adsorbent is condensed in a separate container.⁶³



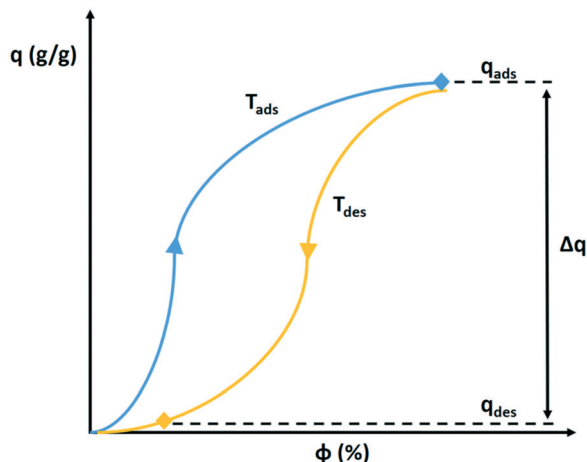


Fig. 14 Hypothetical adsorption and desorption isotherm of water vapour on a solid desiccant. The extractable amount of water Δq corresponds to the difference in adsorbed amount at the relative humidity during night at cold temperature T_{ads} and during the day at high temperature T_{des} .⁶⁶

condenser during the day. All water of the inlet air is considered to be adsorbed. The adsorption heat is assumed to be dissipated and no energy input is considered to be needed for this step. During the day, the temperature is raised by solar heating. The condenser at the end of the process (Fig. 13) is operated at constant ambient temperature and its energy need is neglected in the thermodynamic analysis.

The heat of desorption is dependent on the desiccant material, temperature and uptake at which the process is operated, reflected in the adsorption isotherm. A water desorption enthalpy of 3.3 kJ g^{-1} is assumed here, which corresponds to an average value for zeolites and MOF materials at high relative humidity. The desorption heat can be significantly higher at extremely low relative humidities.^{68,69} The day and night temperatures are considered to be constant for simplicity. The desorption temperature is selected in the range of $40\text{--}80 \text{ }^\circ\text{C}$.⁷⁰⁻⁷²

The specific water yield of a desiccant based process (eqn (15)) is defined as

$$\text{SY}_{\text{max,water}} = \frac{\dot{L}_d}{\dot{P}_d} \quad (15)$$

The water production \dot{L}_d (L h^{-1}), equals the water volume captured during the night.

$$\dot{L}_d = \rho_a \dot{V}_a \omega_{\text{amb}} \quad (16)$$

In the estimation of the total power supply \dot{P}_d (kW), the contribution of the sensible heat losses of heating the air itself and the desiccant mass are not included. This is a reasonable simplification since the sensible heat requirements are much lower than the water desorption enthalpy.

$$\dot{P}_d = (\Delta H^{\text{des}} + C_{p,w}(T_{\text{des}} - T_{\text{atm}}))\dot{L}_d + \dot{P}_v \quad (17)$$

The estimated maximum specific water yield is plotted in Fig. 15 against ambient temperature at desorption temperatures of 40 , 60 and $80 \text{ }^\circ\text{C}$, and neglecting ventilator energy input. The desorption temperature shows a limited effect on performance.

The heat of water desorption from the adsorbent has the strongest impact on the specific water yield (Fig. 16). Using adsorbents with a lower heat of desorption enhances the specific water yield. Ventilator power requirements are negligible relative to the heat of desorption.

Passive atmospheric water capturing devices with desiccants. Reports on water yields with passive devices using desiccants at different locations and climates are summarised in Table 3. Literature typically reports water yields per surface area, or per weight of desiccant. Specific water yields on energy basis are seldom reported. Estimates of specific water yields and efficiencies (eqn (18)) were derived from the available literature data, and assuming the following parameter values for $\text{SY}_{\text{max,water}}$: $\Delta H^{\text{des}} = 3300 \text{ kJ L}^{-1}$, $\dot{P}_v = 0 \text{ W}$, and no sensible heating.

$$\eta_d = \frac{\text{SY}_{\text{water}}}{\text{SY}_{\text{max,water}}} \quad (18)$$

The maximum specific water yield ($\text{SY}_{\text{max,water}}$) amounts to $1.09 \text{ L kW}^{-1} \text{ h}^{-1}$. Note that this maximum and the actual energy efficiency will deviate because the desorption energy of the materials may be different from the assumed value.

Calcium chloride solutions either unsupported or supported on solids like the ordered mesoporous silica material MCM-41 and sawdust have been investigated.^{14,25,73,74} MOF-801 is a porous crystalline metal organic framework consisting of zirconium fumarate.⁶⁴ A tubular solar still filled with CaCl_2 was reported to produce water at a yield of 0.47 L m^{-2} in extreme climate conditions at $25\text{--}35\%$ relative humidity.⁷⁵ Metal-organic framework MIL-101(Cr) encapsulating LiCl has been successfully demonstrated in a water harvesting device.⁷⁶ A large specific water yield was reached with LiCl encapsulated in hollow carbon spheres. The carbon

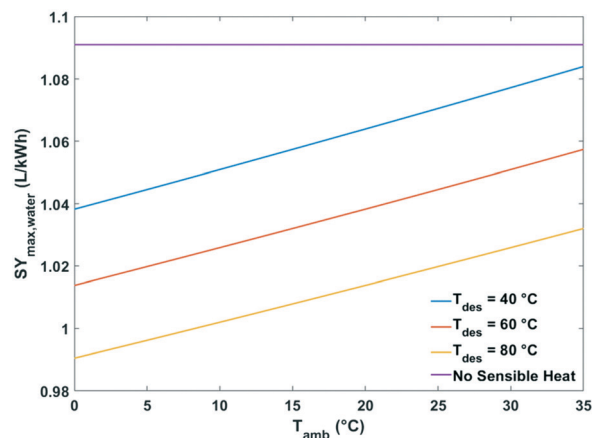


Fig. 15 Maximum specific water yield of a passive solid desiccant based water-from-air extraction device against ambient temperature. The curves pertain to different desorption temperatures.



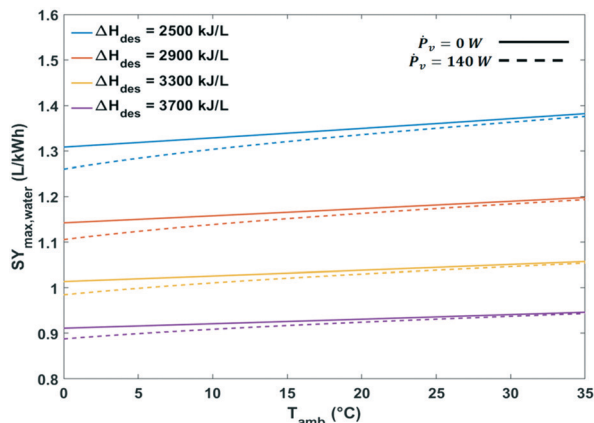


Fig. 16 Specific water yield of a passive solid desiccant based water from air extraction device against ambient temperature ($\phi = 40\%$) at different heat of desorption values at $T_{des} = 60\text{ }^{\circ}\text{C}$.

collects solar heat efficiently and conducts it directly to the desiccant.⁷⁷ Carbon nanotubes (CNT) behaved similarly in polyacrylamide-CNT- CaCl_2 composite material.⁷⁸ The addition of gravel to a desiccant material to help store thermal energy has also shown to improve the specific water yield.⁷⁹

The water harvesting with these systems is in the range of $0.1\text{--}0.6\text{ L kg}^{-1}$, and $0.3\text{--}2\text{ L m}^{-2}$ (Table 3). The thermodynamic efficiencies of these passive solar heating devices are in the range of 10–70%. Direct comparison of the performances of the different desiccants is not possible since the experimental conditions were different.^{14,25,64,73,74} Heating of the condenser surface by sunlight, heat losses to the surroundings and an excess of available solar energy relative to adsorbed water quantity can lead to drops in energy efficiency.^{75,77,80}

Thermo-responsive polymers. Thermo-responsive polymers are a particular category of desiccants. They undergo a drastic volume change due to a conformational reorganisation of the polymer chains in response to a temperature change. This conformational change makes the material more hydrophobic and this alteration causes water expulsion. In this instance liquid water is collected, which is

energetically a big advantage over desiccants from which the collected water can only be recovered as a vapour.^{81,82} Poly(*N*-isopropylacrylamide) (poly-NIPAM) is such a thermo-responsive polymer. The water capacity can be enhanced by adding a second hydrophilic compound. Matsumoto *et al.* reported an interpenetrating polymer network comprising poly-NIPAM and sodium alginate with water adsorption capacity of 0.6 g g^{-1} at $27\text{ }^{\circ}\text{C}$ and 80% relative humidity. When heated to $50\text{ }^{\circ}\text{C}$, which surpasses the hydrophilicity switching temperature of poly-NIPAM, approximately 20% of the absorbed moisture was ejected by an oozing process.⁸¹ A hydrogel based on poly-NIPAM, calcium alginate and agarose helices shows a similar oozing process at $40\text{ }^{\circ}\text{C}$.⁸³

Strong synergy between poly-NIPAM thermo-responsive polymer and chloride-doped polypyrrole (PPy-Cl) was recently achieved in a super moisture-absorbent gel (SMAG).⁸²

Water vapour is adsorbed on the hygroscopic PPy-Cl phase and transferred to the super-hydrophilic poly-NIPAM, which can store large amounts of liquid water. The water capacity amounts to 6.7, 3.4, and 0.7 g g^{-1} at a relative humidity of 90, 60, and 30%, respectively. A temperature rise above hydrophilic/hydrophobic switching temperature of the SMAG ($40\text{ }^{\circ}\text{C}$) leads to shrinkage and expulsion of liquid water. In cyclic operation of SMAG up to 50% of the water content of the saturated SMAG can be collected. Specific water yields of 0.21, 3.71 and $9.28\text{ L kW}^{-1}\text{ h}^{-1}$ were achieved at relative humidities of 30, 60 and 90%, respectively. At low relative humidity, the specific water yield of SMAG ($0.21\text{ L kW}^{-1}\text{ h}^{-1}$) is in the range of CaCl_2 systems and MOFs (Table 3). The performance of SMAG in passive conditions at high relative humidity is spectacular and potentially surpasses the performance of active cooling (discussed higher), but these excellent SMAG performances in real conditions outdoor still needs further investigation.⁸²

Kinetics of water uptake by desiccants. Water uptake kinetics matter to the efficiency of the water collecting process. The adsorption step is found to be slower than desorption.⁷⁷ In addition, concentrating the solar heat can enhance desorption kinetics.⁸⁴ This makes the adsorption step the bottleneck of this kind of system. The saturation process of typical

Table 3 Harvesting potential of different hygroscopic materials

Ref.	Geographic location	Desiccant material	Water harvested per cycle (L kg^{-1}) and (L m^{-2})		SY_{water} ($\text{L kW}^{-1}\text{ h}^{-1}$)	η_d^a (%)
14	Egypt	CaCl_2 (liquid)	0.52	1.12	0.24^b	22
73	Shanghai	CaCl_2 in MCM-41 matrix (solid)	0.54	1.34	0.27^c	25
74	India	CaCl_2 in sawdust (solid)	0.18	0.5	0.16^d	15
25	Egypt	CaCl_2 (liquid)	—	0.63	0.19^e	17
64	Tempe, Arizona	MOF-801 (solid)	0.28	0.34	0.15^f	14
76	Shanghai	LiCl in MIL-101(Cr) (solid)	0.45	0.4	0.37^g	34
77	Thuwal	LiCl in hollow carbon spheres (solid)	0.61	0.50	0.71^h	65
78	Thuwal	PAM-CNT- CaCl_2 hydrogel	0.57	2	0.75^i	69
75	Hail City	CaCl_2 (anhydrous)	0.13	0.47	0.12^j	11

^a $\text{SY}_{\text{max,water}} = 1.09\text{ L kW}^{-1}\text{ h}^{-1}$ ($\Delta H_{des} = 3300\text{ kJ L}^{-1}$, $\dot{P}_v = 0\text{ W}$, no sensible heat). ^b $\Phi_{amb} = 36\text{--}54\%$, $T_{amb} = 25\text{--}35\text{ }^{\circ}\text{C}$, solar flux: $0.2\text{--}0.8\text{ kW m}^{-2}$. ^c Daily total radiation: 17.8 MJ m^{-2} . ^d $\Phi_{amb} = 40\text{--}80\%$, $T_{amb} = 25\text{--}30\text{ }^{\circ}\text{C}$, solar flux: $0.3\text{--}0.88\text{ kW m}^{-2}$. ^e $T_{amb} = 20\text{--}33\text{ }^{\circ}\text{C}$. ^f $\Phi_{amb} = 26\text{--}34\%$, $T_{amb} = 20\text{--}32\text{ }^{\circ}\text{C}$, solar flux: 1.1 kW m^{-2} . ^g $\Phi_{amb} = 30\%$, $T_{amb} = 30\text{ }^{\circ}\text{C}$, solar flux: $0.6\text{--}0.8\text{ kW m}^{-2}$. ^h Data of first cycle batch-process, solar flux: 0.525 kW m^{-2} . ⁱ $\Phi_{amb} = 60\text{--}70\%$, $T_{amb} = 26\text{ }^{\circ}\text{C}$, average solar flux: 1.06 kW m^{-2} . ^j $\Phi_{amb} = 25\text{--}35\%$, $T_{amb} = 30\text{ }^{\circ}\text{C}$, average solar flux: 0.603 kW m^{-2} .



adsorbent materials is illustrated at low relative humidity ($\Phi = 30\%$) in Fig. 17, and at high relative humidity ($\Phi = 90\%$) in Fig. 18. It should be remarked that kinetic data from literature is difficult to compare because there are many experimental parameters involved, like ad- or absorbent bed geometry, particle size, temperature and pre-treatment conditions.^{71,82,85–87} Fig. 17 and 18 are meant to show general trends. At low relative humidity ($\Phi = 30\%$) LiCl salt doubles its weight by water absorption after 2800 minutes. SMAG reaches 80% of its capacity already after only 180 minutes, and is fully hydrated (0.7 g g^{-1}) after 2000 minutes.⁸² MOF-303 and zeolite 13X show similar water adsorption kinetics.^{71,87} They are slower than SMAG, and the capacity is more limited due to the smaller pore volume in a much less expandable framework.

At 90% relative humidity (Fig. 18) the SMAG has by far the highest capacity (6.7 g g^{-1}), reached in about 2000 minutes. After 100 minutes the water uptake reaches already 5.4 g g^{-1} .⁸² LiCl salt keeps on absorbing water vapour and reaches a water content of *ca.* 2.5 g g^{-1} after 3000 minutes. A nanoporous super hygroscopic hydrogel based on an amorphous oxygen and zinc matrix is able to absorb 3.6 g g^{-1} after 720 min.⁸⁵ The water adsorption capacity of zeolite 13X is limited. Zeolites are known for their high affinity for water, but the capacity is limited to the pore volume, which for zeolites does not exceed 0.4 mL g^{-1} .⁸⁷ Silica gel is a well-known desiccant. It has high capacity, but the adsorption kinetics are extremely slow. In the example of a mesoporous silica gel in Fig. 18, after 10 000 minutes there is hardly 0.7 g g^{-1} adsorbed.⁸⁶

Water recovery from desiccants. Ad- and absorbent materials have largely different regeneration requirements (Table 4). Water is strongly adsorbed in zeolites, which need to be heated above $100 \text{ }^\circ\text{C}$ for desorbing water vapour, which makes them less practicable for capturing water vapour from the air.^{71,87,88} Hydrated salts also need substantial heating for water evaporation.⁶⁸ MOFs and Amorphous Zn–O hydrogel are less demanding with respect to desorption temperature.^{65,71,85,89} Thermo-responsive polymers have the attractive feature of a hydrophilic/hydrophobic phase transition at temperatures below $50 \text{ }^\circ\text{C}$. This conformational change is fast. Using SMAG, a water production cycle can proceed within 1

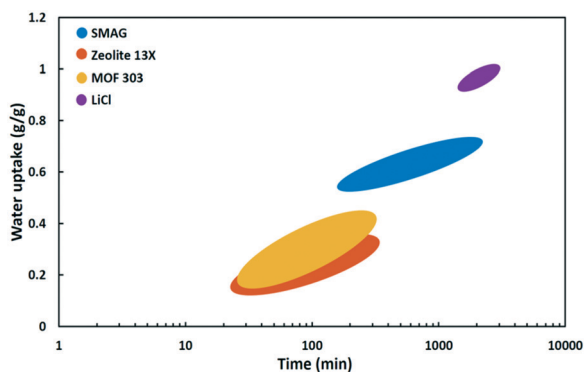


Fig. 17 Water uptake as a function of time at $\Phi = 30\%$ of SMAG (polymer),⁸² zeolite 13X,⁸⁷ MOF-303 (ref. 71) and LiCl (salt).

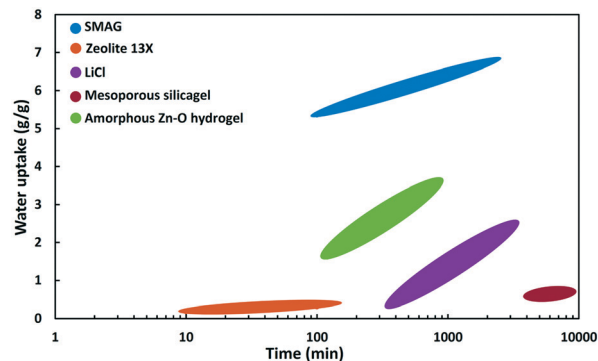


Fig. 18 Water uptake as a function of time at $\Phi = 90\%$ of SMAG (polymer),⁸² zeolite 13X,⁸⁷ hydrogel,⁸⁵ mesoporous silica gel⁸⁶ and LiCl (salt).

hour, with water capturing during 50 minutes, and water expulsion during 10 minutes. Evacuating all water from a SMAG necessitates heating like for other materials.⁸²

Estimation of the maximum specific water yield when using liquid desiccants is more complicated than for a solid adsorbent. Concentrated solutions of inorganic salt are typical examples of liquid absorbents.⁹⁰ Such solutions keep on absorbing water such that in a practical timeframe of a night and day cycle thermodynamic absorption equilibrium is never reached. It may explain some contradicting statements in literature. Some authors state that liquid desiccants are able to capture more water on weight basis than solid adsorbents and are more productive.⁷⁰ Aqueous desiccants would have advantages over solid desiccants when solar heating is used for regeneration because the regeneration temperature of liquid desiccants is generally lower than for solid adsorbents.^{61,70,91} Other authors claim solid adsorbents to be the best choice.^{64,73} The regeneration requirements of liquid and solid desiccants are relatively similar.

Conclusions. Passive desiccant based water harvesting is appealing because of its technical simplicity. It may offer a decentralised solution for small communities in dry climates, where the condensation by cooling principle is not applicable. The strong water affinity of desiccants is advantageous for trapping water vapour, but the high desorption enthalpy of water on the desiccant causes high energy needs, and it limits the theoretical maximum specific water yield. Traditional adsorbents and salts have been used in the past with limited success. In recent years there has been innovation in desiccant materials, and especially MOFs and thermo-responsive polymers bear great promise. The advent of better performing desiccants gives new stimuli to the development of superior water capturing processes with reduced energy consumption.

Active desiccant technologies

Passive desiccant systems rely on the day-night cycle and produce water discontinuously.⁷¹ Continuous water production is made possible by implementing an artificial energy supply.



Table 4 Regeneration temperatures and recovered water fraction for different desiccant materials

Ref.	Desiccant material	Regeneration temperature (°C)	Fraction desorbed (%)
72	Silica type 3A	90	95
71, 88	Zeolite 13X	120; 600	83; 100
71	MOF-303	85	100
65, 89	MOF-801	85	100
82	SMAG	40; 160	40; 100
68	MgSO ₄	80	31
73	CaCl ₂ in MCM-41 matrix	80	90
92	CaCl ₂ in alginate matrix	100; 150	90; 100
68	CaCl ₂	80	31
68	FeCl ₃	80	57
93	LiCl impregnated in cloth	85	100
85	Amorphous Zn-O hydrogel	55	99

This section provides a description of such active water capturing devices with desiccants. The principles of the use of desiccants are similar to the passive systems. The difference resides in the regeneration which is activated by an auxiliary energy source. The maximum specific water yield of the active desiccant technology will therefore be comparable with the energy requirements of the passive desiccant case.

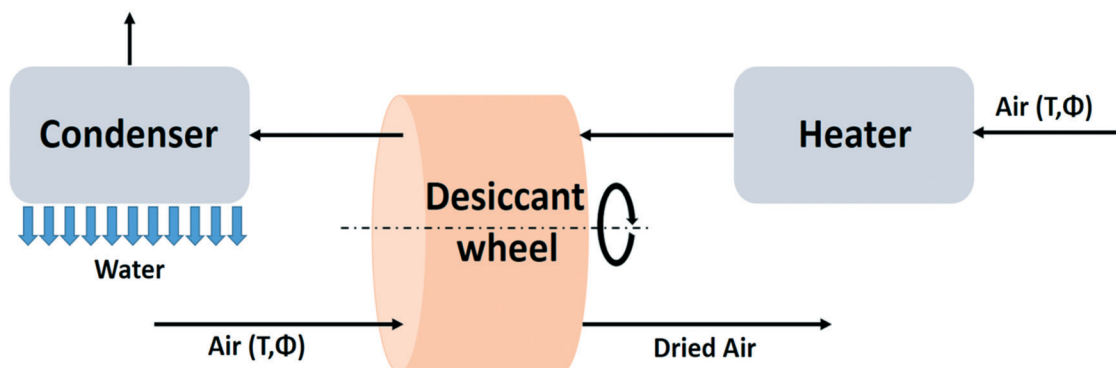
Process description of active desiccant technology. Continuous regeneration of the desiccant can be achieved by fixing the solid desiccant in a wheel (Fig. 19). The turning of the wheel causes the desiccant to be alternately positioned in the adsorption and in the regeneration section. The air with its water vapour enters the desiccant wheel which extracts the moisture. The rotating wheel moves the saturated solid desiccant to the regeneration section where the adsorbed water is evacuated by hot air. The desorbing moisture is collected by condensation in a downstream condenser.⁹⁴

Some modifications of this principle have been proposed. Instead of a wheel a two-bed system can be used with the beds alternating between adsorption and regeneration mode.⁹⁰ A discontinuous process is another alternative.⁹⁵ The desiccant wheel is not suited for handling liquid desiccants. In that case pumping equipment is installed to transport the desiccant material from absorption to regeneration section of the unit.⁹⁴ Complex systems with a liquid desiccant have been analysed with mathematical models.^{96,97}

Moving from a passive to an active desiccant based system has its trade-offs. The desiccant is used more efficiently in the active system, which reduces the desiccant material costs for a given water production rate. Active systems are technically more complicated, and require an external energy supply. The footprint of active desiccant based systems is smaller than for passive systems, but for providing the auxiliary energy, solar panels and solar collectors occupy additional space. The principles of active water extraction using a desiccant have already been put to practice long time ago in the field of air conditioning. A review of the early days of desiccant dehumidification technology is provided in a review of desiccant dehumidification technology by A. Pesaran.⁹⁴

Active atmospheric water capturing devices with desiccants. One example of an active desiccant based device is a panel which produces drinking water from air, by using solar thermal energy and photovoltaic electricity. The so-called hydropanel has two arrays of panels which each produce 2–5 L of water per day depending on relative humidity, temperature and solar irradiation conditions. One panel has a surface area of approximately 2.9 m².⁹⁸

The technology is based on an adsorption/desorption cycle by using a desiccant wheel.⁹⁹ The middle section produces electricity by photovoltaic panels. The regeneration heat is produced in the heat absorbing panels positioned left and right of it. The panel produces water only during daytime.¹⁰⁰ The major

**Fig. 19** Process flow diagram of the active desiccant technology based on the desiccant wheel design.⁹⁴

advantage of this standalone device is its applicability in isolated and remote areas where grid power is unavailable.

From the data provided in different environmental conditions, the specific water yield is estimated at 0.14–0.32 L kW⁻¹ h⁻¹.¹⁰⁰ These values are in the range of passive desiccant based devices with classic adsorbents (Table 3).

Similar specific water yields of 0.3–1 L kW⁻¹ h⁻¹ depending on environmental parameters have been reported for related devices. Internal heat recovery increases the dehumidifier performance.^{101,102} Wang *et al.* reached an SY_{water} of 0.84 L kW⁻¹ h⁻¹ by using LiCl enclosed in an active carbon felt matrix as adsorbent and an electric heater for the discontinuous desorption stage.⁹⁵ The high efficiency can be explained by the humid conditions under which this excellent performance was achieved in combination with an efficient heater. An active desiccant based device reported by Hanikel *et al.* with MOF-303 adsorbent has been demonstrated in the Mojave Desert. Even under those challenging climate conditions a specific water yield of 0.19 L kW⁻¹ h⁻¹ could be reached.⁷¹

Conclusions. The use of ab- and adsorbent materials can be maximised by involving auxiliary energy. That energy can be used to move the adsorbent physically from adsorption to desorption sections of the device, and to provide heat for regeneration. The active system has some practical advantages, like the disconnection of the water production from the day-night cycle. Energetically, it does provide some benefits over the passive system mainly by the ability to use heat regeneration. The specific water yield amounts to 1 L kW⁻¹ h⁻¹ at the maximum. The energy performance is inferior to systems involving water condensation by active cooling. The reason is the large heat of desorption of water from adsorbents, being larger than the heat of condensation.

Water from air and the water-energy nexus

The energy demand of water-from-air technologies is significant, and the potential of water harvesting from the atmosphere is part of the water-energy nexus. The climate dictates the technology to be advised. Dew and fog harvesting show potential in areas with periods with 100% relative humidity. When this condition is not met, other water-from-air technologies have to be considered. From the analysis of the different concepts in this review, passive desiccant technologies and active cooling emerged as most promising technologies. The climatological conditions favouring active cooling *versus* passive desiccant technology depending on relative humidity and ambient temperature are indicated in Fig. 20. Passive desiccant based systems with desorption temperature of 80 °C (Fig. 15) are considered; and active cooling according to the description in Fig. 9. Active cooling is the preferred choice in hot and wet climates, a generally recommended guideline in dehumidification technology.¹⁰³ This conclusion was also reached previously by Gido *et al.*³⁷ In dry and hot regions especially vulnerable to water scarcity, desiccant tech-

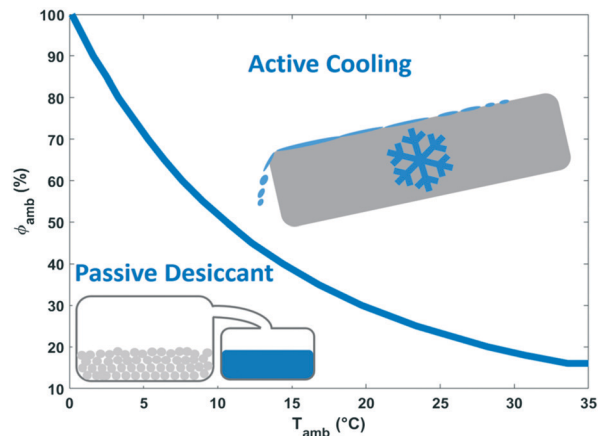


Fig. 20 Optimal regions for active air cooling and passive desiccant technologies as a function of ambient temperature and relative humidity values, based on the theoretical maximum specific water yield.

nologies are the best and only choice when considering the atmosphere as a water source.

Fig. 21 summarises the specific water yield reached for different types of water-from-air technologies relative to the maximum specific water yield. Relatively high specific water yields can only be reached with active cooling. Opting for desiccant technology in arid regions will lead to low specific water yields. A specific water yield of 0.1–4 L kW⁻¹ h⁻¹ is characteristic for water-from-air technology. This is very low compared to conventional water production techniques like reverse osmosis desalination, which reaches specific water yields of 250 L kW⁻¹ h⁻¹.¹⁰⁴

The limited productivity according to the state-of-the-art (~1 L m⁻², Table 3), limits the application potential of passive desiccant water extraction to small communities.⁷⁰ Applications with high water demand like agriculture are problematic.

The active desiccant case could overcome the scaling problem for water production when use is made of grid electricity. In that case electricity generation should not involve water consumption. Thermoelectric power plants consume on average 1.25 L kW⁻¹ h⁻¹ (ref. 105) which exceeds even the specific water yield to be expected (Table 3). Innovation in adsorbent

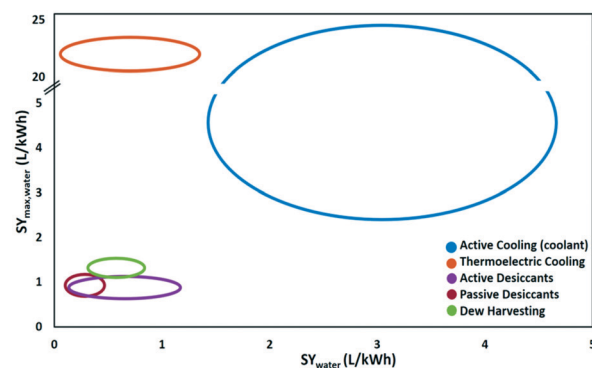


Fig. 21 The maximum specific water yield and the achieved specific water yield of five different water-from-air technologies.



materials and concepts for water-from-air technologies is needed. The advent of thermo-responsive polymers and their enhanced specific water yields reached at laboratory scale is promising.

Conclusion

The maximum specific water yield ($\text{L kW}^{-1} \text{h}^{-1}$) is a consistent parameter to evaluate water-from-air technologies and indicate the preferred climate conditions for their operation.

The use of dew and fog harvesting is advantageous when the air carries airborne liquid water droplets. But in humid climate with frequent dew and fog, water scarcity is seldom a problem. Unpredictable and low water output per surface area are major drawbacks of this inexpensive technology. A continuous water output is achieved with the active cooling approach by using forced air convection over a condenser. Active cooling is the most energy efficient way to extract water from the air in mid to high relative humidity regions and warm climates. In this approach the maximum specific water yield decreases exponentially with decreasing relative humidity. The cooling principle cannot be applied in low relative humidity climates because of dew-points below $0\text{ }^{\circ}\text{C}$ causing freezing of the collected water. An active cooling system can be favourably assisted by a membrane module to enhance the relative humidity artificially. A more common approach in dry climates is the application of desiccant materials. Research in this area has mostly been concentrated on new material development. This led to the introduction of several new desiccant materials like MOFs which show lower desorption temperatures, improved adsorption kinetics and better applicability in extremely arid regions with a relative humidity around 30%. Salts like CaCl_2 and LiCl have been shown to be efficient in these harsh conditions as well. The specific water yields achieved with desiccants is relatively low and the theoretical limits are within reach. Despite the recent advances in desiccant materials and process engineering the intrinsic energy intensity remains a major obstacle. The introduction of fundamentally new concepts is needed to cause a step change in energy performance. Some thermo-responsive polymers display hydrophilic to hydrophobic phase transitions at temperatures when water is still liquid. A promising specific water yield of *ca.* $9\text{ L kW}^{-1} \text{h}^{-1}$ at 90% relative humidity has been reported, surpassing current cooling and desiccant based technologies. The performance under more challenging conditions of low relative humidity remains to be demonstrated. There is plenty of water in the atmosphere, but harvesting it in an energy efficient way remains one of the grand scientific challenges. Introducing fundamentally new concepts may be key in making this technology a viable solution to combat water scarcity.

Conflicts of interest

There are no conflicts to declare.

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