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Feed gas humidity: a vital parameter affecting a cold atmospheric-pressure plasma jet and plasma-treated human skin cells

J Winter^{1,2,3}, K Wende^{1,2,3}, K Masur^{1,2}, S Iseni^{1,2}, M Dünnbier^{1,2},
M U Hammer^{1,2}, H Tresp^{1,2}, K-D Weltmann² and S Reuter^{1,2}

¹ Centre for Innovation Competence plasmatis, Felix-Hausdorff-Str. 2, 17489 Greifswald, Germany

² Leibniz Institute for Plasma Science and Technology INP Greifswald e.V., Felix-Hausdorff-Str. 2, 17489 Greifswald, Germany

E-mail: winter@inp-greifswald.de and kristian.wende@inp-greifswald.de

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Abstract

In this study, the effect of feed gas humidity on the reactive component generation of an atmospheric-pressure argon plasma jet and its effect on human skin cells are investigated. Feed gas humidity is identified as one key parameter that strongly influences stability and reproducibility of plasma medical studies. The plasma jet is investigated by absorption spectroscopy in the ultraviolet and infrared spectral region for its ozone production depending on the humidity concentration in the feed gas. By optical emission spectroscopy the dependence of present excited plasma species such as hydroxyl radicals, molecular nitrogen, argon and atomic oxygen on the feed gas humidity is investigated. As an interface layer between the plasma jet effluent and the biological cell, a buffer solution is treated and the hydrogen peroxide (H_2O_2) production is studied with two independent colorimetric assays as a function of humidity admixture to the feed gas. Ultimately, the effect of varying feed gas humidity on the cell viability of indirect plasma treated adherent HaCAT cells is investigated. The highest viability is found for the driest feed gas condition. Furthermore, this work shows answers for the relevance of unwanted—or intended—feed gas humidity in plasma medical experiments and their comparatively large relevance with respect to ambient humidity. The findings will lead to more reproducible experiments in the field of plasma medicine.

1. Introduction

The past decade has seen the rise of atmospheric-pressure low-temperature plasmas. These plasmas operated under ambient conditions find a wide range of applications, e.g. in surface treatment of polymers [1–3], treatment of woven fabrics [4] and surface functionalization [5], and also as actuators for the control of air flow in aeronautics [6]. Atmospheric-pressure plasmas are increasingly used for deposition of thin films [7] or generation of carbon nanotubes [8]. Advantages are that large and expensive vacuum equipment can be omitted and high reactive species density can be achieved due to the high density

under ambient conditions. High focus is set on application of atmospheric-pressure plasmas in the medical sector. In this sector, a large scientific community has evolved investigating applications of wound treatment, biocompatible surfaces, or disinfection, inactivation of pathogens, sterilization of medical equipment as well as the inactivation of bacteria in biofilms, see e.g. [9–12]. The high reactivity at low gas temperatures due to the strong non-equilibrium properties of these plasmas can be utilized to generate a reactive chemistry, usually available only at high temperatures, on temperature-sensitive surfaces such as human skin.

In particular in the medical sector, plasma sources are operated in ambient air, introducing humidity into the active plasma zone. Humidity plays an important role

³ These authors contributed equally to this work.

in plasma chemical processes since dissociation of water (H_2O) opens a wide variety of subsequent plasma chemical reactions [13]. This is especially true for atmospheric-pressure plasmas operated in humid ambient air but also for noble gas driven atmospheric-pressure plasma jets [14–16]. In the case of plasma jets, humidity is either introduced directly into the working gas as impurity or diffuses into the effluent when the jet is operated in a humid environment. In biomedical applications, changing humidity conditions not only has an influence on the plasma but also on the plasma-treated biological targets, such as cells or cell structures [17]. The products originating from the water reaction pathways, for example hydroxyl (OH), atomic oxygen (O) and hydrogen peroxide (H_2O_2) are potent to generate oxidative stress in biological samples. This is already known from decontamination experiments, where bacteria were plasma-treated under different humidity settings [18]. For eukaryotic cells relevant for plasma medicine (e.g. human skin cells) no such data exist. Hence, the focus of this work is not only the investigation of feed gas humidity influence on the reactive component generation of an atmospheric-pressure argon plasma jet by optical emission and absorption spectroscopy but also the observation of *in vitro* human keratinocyte behaviour to indirect plasma treatment. It is shown in this work that increasing feed gas humidity leads to a significant viability decrease in the investigated cell line. In order to get a first glimpse into the governing mechanisms, the H_2O_2 concentration in the liquid growth medium is also monitored. The increase in H_2O_2 concentration with feed gas humidity suggests a correlation between H_2O_2 and cell viability.

In addition to the fundamental research aspects, it is shown that gas supply tubes made of polyurethane (PUR) are a major source of feed gas humidity and must be considered from the practical point of view. Moreover, the large relevance of feed gas humidity with respect to ambient humidity is emphasized in this paper.

2. Experimental setup

2.1. Plasma jet

The investigated plasma source is a cold atmospheric-pressure plasma jet (kinpen, neoplas GmbH, Greifswald, Germany). It consists of a centred rod electrode inside a ceramic capillary and a grounded ring electrode [19, 20]. When a sinusoidal voltage signal of 2–6 kV_{pp} with a frequency of about 1 MHz is applied to the centred electrode, plasma is ignited within the core region of the jet. Also an up to 10 mm long effluent becomes visible after ignition. The working gas is argon (Ar purity 99.999%) with a gas flow rate of 3 standard litres per minute (slm). Furthermore, the humidity concentration of the feed gas is changed with the setup described in detail in section 2.3.

2.2. Humidity measurements in gas supply tubes

Feed gas tubes made of PUR can absorb a significant amount of water when they are exposed to a humid environment. In

Figure 1. Experimental setup for measuring the water concentration that originates from a previously humidified tube.

operation, dry gas from the gas bottle becomes humidified while streaming through such previously humidified feed gas tubes. Hence, a significant number of water molecules are transported by the feed gas into the core plasma region of the investigated plasma jet. In order to determine the number of water molecules that originate from feed gas tubes the experimental setup in figure 1 was used. A 3 m long PUR tube (Norgren, Germany) with inner and outer diameters of 4 mm and 6 mm, respectively, was enclosed by a plastic box. The air atmosphere inside the box was set to a relative humidity level of 50% (corresponds to an absolute H_2O concentration of 1.4%) by means of a humidified constant air flow rate of 1 slm. The measurement procedure had three phases. In the first phase dry argon gas at a flow rate of 1 slm was led through the 3 m tube for 12 h. Therefore, valves B were opened and valve A was closed. After this drying phase the second phase began in which valves B were closed and valve A was opened. Now, the dried argon filled tube was exposed to the humidified atmosphere in the box for another 12 h. Meanwhile the argon gas kept flowing through valve A. This provides unchanged dry conditions for the supply tubes. In the third phase, valves B were opened and valve A was closed again, so that dry argon gas flew through the humidified tube. The gas humidity was measured over time with a chilled mirror dew point hygrometer (DewMaster, EdgeTech, USA) with an uncertainty of about 2%. The humidity concentration detection limit of this hygrometer was 20 parts per million (ppm). For comparison, a 3 m long stainless steel pipe with the same dimensions as the PUR tube was analysed with the same procedure.

2.3. Feed gas humidification and optical emission spectroscopy

To investigate the effect feed gas humidity has on emission spectra and on biological samples the humidity of the feed gas was controlled. Due to the difficulty in adjusting humidification by gas tubes, a controlled feed gas humidification was performed with a bubbler setup. Dry argon gas was split into two channels, a main gas channel and a channel which was guided through a water reservoir.

Figure 2. Scheme of the optical emission spectroscopy setup and of gas humidification.

Behind the bubbler, both argon gas streams were mixed and the resulting humidity was analysed by the chilled mirror dew point hygrometer already described in section 2.2. Afterwards the humidified feed gas was led into the plasma jet. By changing the argon flow rate through the bubbler in the range 0–0.5 slm the feed gas humidity was precisely adjusted between 20 and 1300 ppm. To provide a constant total argon flow rate of 3 slm the dry gas channel flow rate was appropriately reduced.

A sketch of the humidification setup in combination with the optical emission spectroscopy (OES) setup is displayed in figure 2. The OES was performed using a UV and a NIR sensitive spectrometer (AvaSpec-2048-USB2, Avantes) with a spectral resolution of 0.7 nm, respectively. The axially resolved emission spectra with a spatial resolution of 0.5 mm were obtained for different humidity concentrations. Particularly four spectral regions were evaluated in this work corresponding to the emission of argon atoms (Ar 4p–4s, 695–698 nm), hydroxyl radicals ($\text{OH } A^2\Sigma^+ \rightarrow X^2\Pi$, 305–313 nm), nitrogen molecules ($\text{N}_2 C^3\Pi_u \rightarrow B^3\Pi_g$, 330–338 nm) and atomic oxygen (O 3p–3s, 775–778 nm). Integrating the respective emission intensities for different axial positions (0–10 mm, step size: 0.5 mm) and different feed gas humidity concentrations over the given spectral ranges and normalizing to the maximal value yields an emission intensity map for each species.

Furthermore, the plasma jet was equipped with a shielding device that enables the shielding of the effluent plasma from ambient species [21]. In order to investigate whether feed gas humidity differentially influences the plasma composition than ambient humidity two experiments were performed using this shielded jet. Firstly, dry air as shielding gas and humidified argon with a humidity concentration of 490 ppm as feed gas were used. In this situation, the humidity concentration in the effluent is only determined by the feed gas humidity. Secondly, the opposite situation, namely humidified shielding gas with a humidity concentration of 20 500 ppm (corresponds to a relative humidity level of 72%) and dry feed gas, was investigated. As a measure for the impact of both situations the excited OH emission was analysed with the setup explained above.

2.4. Ozone net production rate

The ozone net production rate of the investigated plasma jet operated with 3 slm argon plus 1% oxygen was determined for different feed gas humidity concentrations using UV and IR absorption spectroscopy techniques identical to the setups reported in [20, 22]. For a comparison of both techniques, the measurement setup-independent net production rate was calculated, considering the velocity profile of the gas for UV absorption and the total flow rate for IR absorption. Details of the production rate calculation are also given in the two papers already mentioned above. The humidification of the feed gas was performed with the bubbler setup described in section 2.3.

2.5. Plasma treatment and hydrogen peroxide determination in RPMI solution

The plasma treatment was performed in a polystyrene Petri dish (53 mm diameter TPP, Trasdingen, CH) filled with 5 ml complete cell growth medium (Roswell Park Memorial Institute 1640-RPMI + 8% foetal calf serum +1% penicillin/streptomycin solution). In order to treat the liquid at different positions the plasma jet was mounted on an xy-table, and the plasma jet effluent was moved in a meandering pattern over the liquid for a constant treatment time of 40 s. The distance between the nozzle and the liquid surface was set to 9 mm. The H_2O_2 concentration of the treated liquid was analysed using commercial H_2O_2 test stripes (Merckoquant 110011, Merck) and via an Amplex Red homogeneous assay. Aliquots of 50 μl (Amplex Red assay) and 30 μl (test stripes) were taken immediately after 40 s plasma treatment time. The experiment was repeated three times for each humidity setting.

The H_2O_2 test stripes were read out by a digital microscope camera (Conrad, Germany) in order to analyse their colour value. A calibration procedure was conducted prior to the experiments. Therefore, 13 different H_2O_2 concentrations from 0 to 6 mg l^{-1} were produced by diluting 10M H_2O_2 stock solution (340 g l^{-1} , Merck KGAA) in RPMI and H_2O . The sum of the colour values of the red and green camera channel for different H_2O_2 concentrations is plotted in figure 3. Regardless of whether the calibration was performed in RPMI or H_2O almost the same linear calibration characteristic was observed. The linear approximation of the data allows the determination of an unknown H_2O_2 concentration by measuring the colour change of a test stripe.

Another method to detect H_2O_2 in liquid is the usage of a homogeneous assay (Amplex Red Hydrogen Peroxide/Peroxidase Assay Kit, Invitrogen, Ltd). The kit uses the Amplex Red reagent (10-acetyl-3,7-dihydroxyphenoxazine). After 40 s plasma treatment time 50 μl of the treated solution were transferred into a well of a 96-well plate. The well already contains a volume of 50 μl reaction buffer solution according to the protocol provided by the company. Furthermore, a standard curve with known H_2O_2 concentrations was also prepared in the 96-well plate. The absorption and fluorescence signals at wavelengths of 560 nm and 590 nm, respectively, were obtained by means of a multiplate reader (Tecan infinite M200, Tecan Group Ltd, Switzerland). By comparing the results of the standard curve with the samples taken at different

Figure 3. H_2O_2 test stripe calibration curve.

feed gas humidity levels the plasma treatment induced H_2O_2 concentrations after 40 s treatment time were determined. By comparison with the test stripe method the Amplex Red assay is more sophisticated but also more sensitive. The H_2O_2 test stripe lower detection limit is about 0.5 mg l^{-1} , whereas a significant lower detection limit of 0.2 ng l^{-1} is reported for the Amplex Red assay [23].

2.6. Cell viability assay

The viability of adherent human keratinocytes (HaCaT cells) was determined using the conversion of fluorescence dye resazurin by metabolic active cells into the highly fluorescent resorufin. Details of this assay are given in [24]. The cells were cultivated in the same medium that was used for the H_2O_2 measurements.

After 40 s plasma treatment of 5 ml RPMI solution under the equivalent conditions as described for the H_2O_2 measurements, the plasma-treated medium was immediately transferred into one row of wells of the cell culture plate and serially diluted with complete medium immediately (six parallel wells per treatment time) to obtain different treatment time equivalents. Analogous to the procedure described in [24] the cells were cultivated for a further 71 h and fluorescence intensity after resazurin addition ($100 \mu\text{M}$ resazurin in complete RPMI w/o phenol red) was analysed by a Tecan M200 multiplate reader after 1 h of incubation. As a measure of viability the 50% inhibition value was determined for every investigated feed gas humidity level (example for 20 ppm see figure 4).

3. Results and discussion

3.1. Humidity originating from feed gas tubes

The humidity concentration in dependence on the gas tube flushing time is displayed in figure 5 for the two investigated materials, namely PUR and stainless steel. For the PUR tube an initial humidity concentration of more than 1000 ppm depicts a high porosity and storage capacity of this material for water molecules. With increasing flushing time a drying of the inner

Figure 4. Cell viability in dependence on the treatment time equivalent for a feed gas humidity level of below 20 ppm. The 50% value was determined via sigmoidal regression (here 41.7 s treatment time equivalent).

tube surface occurs. However, it takes more than 3 h of flushing time to decrease the feed gas humidity concentration of a 3 m long PUR tube under a level of 70 ppm, which from a practical point of view can be considered as dry. In contrast, the initial feed gas humidity concentration for the investigated metal tube of the same length is by one order of magnitude lower than for the polymeric tube. Already after a couple of minutes the humidity concentration is significantly lower than 70 ppm. A consequence of these findings is that not only dry but also stable humidity conditions in the feed gas can be easily achieved with metal tubes. However, metal tubes are expensive and by far not as practical as polymeric tubes. Thus, in the majority of cases, the usage of polymeric tubes can possibly not be avoided totally. In experiments that require stable humidity concentrations it is therefore important to consider polymeric feed gas tubes as a significant source of humidity. From a practical point of view the length of these tubes should be limited as far as possible in order to reduce the drying phase before the experiments.

3.2. Optical emission spectroscopy

The emission spectrum of the jet effluent at an axial position of 4 mm and for a feed gas humidity concentration of 350 ppm shows several emission bands and atomic lines in the ultraviolet (figure 6) and near infrared spectral range (figure 7). In the UV spectral range, emission bands of OH ($A^2\Sigma^+ - X^2\Pi$) and $\text{N}_2(C^3\Pi_u - B^3\Pi_g)$ originate from the dissociation of feed gas water molecules and the diffusion of ambient nitrogen into the effluent [21, 24].

In the NIR spectral range, a set of atomic argon lines $\text{Ar}(4p-4s)$ and the triplet of atomic oxygen lines $\text{O}(3p^5P-3s^5S)$ around 777 nm were measured. The atomic oxygen can either originate from water dissociation alongside with the formation of hydrogen molecules or be generated by dissociation of ambient molecular oxygen that diffuses into the effluent. Focusing on the effluent no lines of atomic hydrogen

Figure 5. Humidity concentration in argon feed gas (flow rate: 1 slm) in dependence on the flushing time through a 3 m long tube made of either PUR or stainless steel.

Figure 7. NIR emission spectrum at an axial position of 4 mm and for a feed gas humidity concentration of 350 ppm.

Figure 6. UV emission spectrum at an axial position of 4 mm and for a feed gas humidity concentration of 350 ppm.

were found. However, end-on OES measurements (not shown here) reveal the presence of atomic hydrogen (H_α and H_β lines) in the active plasma zone and thus confirm water dissociation at least in that region.

Since humidity concentration variation changes not only the plasma chemistry but also the excitation conditions, a strong influence on the emission intensity of the investigated species is expected. Furthermore, the density of molecular oxygen and nitrogen varies with axial distance whereby a spatial dependence on the emission intensity also occurs. Hence, the integral emission intensity was evaluated for both parameters. The result is presented in figure 8 for the four discussed species Ar, OH, N_2 and O. Every emission intensity map is normalized to the maximum value, respectively. The dots in the contour plot give the parameter settings at which the measurements were actually conducted. The intermediate values were calculated by 2D linear interpolation.

It has to be mentioned that careful interpretation is necessary when evaluating OES data. Since emission intensities in figure 8 are integrated over band structures

(OH, N_2) and over atomic lines (Ar, O) their resulting intensities cannot be directly compared. However, by evaluating the respective dependences on the axial position and on the humidity concentration valuable information is obtained despite the lack of quantitative comparability.

With increasing humidity concentration in the feed gas the emission of argon and oxygen lines and the band emission of nitrogen decrease. Two different effects are responsible for this finding. Firstly, with increasing humidity concentration in the feed gas quenching of excited states by water molecules becomes more likely [25]. This is illustrated by the quenching rate coefficients and decay rates given in table 1. For a humidity concentration of 1000 ppm (corresponding to a H_2O density of $2.47 \times 10^{16} \text{ cm}^{-3}$ at atmospheric pressure), the decay rate of $N_2(C^3\Pi_u, \nu = 0)$ due to quenching by water molecules is in the same order of magnitude as the radiative decay rate and therefore plays an important role for depopulating the excited nitrogen state. Unfortunately, no data on the quenching rate coefficients by water vapour were found in the literature for the $O(3p^5P)$ and $Ar(2p_2)$ states. However, assuming that the unknown coefficients are roughly comparable to the known quenching rate coefficients of $O(3p^3P)$ and $Ar(1s_5)$ water molecules also play a major role in the depopulation of these excited atomic states.

Secondly, it is known that the admixture of molecular gases leads to a reduction in electron temperature [36] and therefore affects the formation of photons and energetic species such as argon metastables $Ar(1s_5)$ in the active plasma zone. These photons and also the metastable argon atoms with energies of 11.55 eV and a lifetime on the order of ten microseconds transport energy from the active plasma zone into the effluent and induce dissociation and excitation processes of other species [37, 38]. If, for instance, the formation of Ar metastables is impeded by water molecules, Ar metastable induced excitation processes in the effluent are also impeded and the emission intensity decreases. Hence, the emission intensity of excited Ar, O and N_2 decreases with water vapour admixture.

The emission of OH ($A^2\Sigma^+, \nu = 0$) behaves differently from that of the other species. For small water admixtures

Figure 8. Integrated emission signals originating from Ar, OH, N₂ and O in dependence on the axial distance and feed gas humidity concentration. The dots in the contour plot give the position and humidity concentration where the measurements were conducted. The intermediate values were obtained by linear interpolation. Due to different integration intervals (wavelength integration ranges are given in parentheses below the species label), intensities of different species cannot be directly compared.

an increase in OH emission is observed up to a humidity concentration level of about 400 ppm. This is similar to the results of Nikiforov *et al* on a DBD argon jet, who found an emission maximum of OH ($A^2\Sigma^+, v = 0$) at a water vapour concentration of 350 ppm. Presumably, the OH density increases with rising H₂O admixture due to direct dissociation of water molecules by electrons. This effect would lead to a stronger emission signal provided that excitation mechanisms do not change significantly by small H₂O admixtures. However, when increasing the humidity concentration beyond 400 ppm quenching of excited OH (see table 1) and the reduced electron temperature become relevant. Both counter the emission gain by increased OH density and a maximum evolves. Of course, no information on OH densities is given in this paper and therefore conclusions concerning OH densities are speculative. However, a rising OH density is a reasonable explanation for the formation of the observed maximum. Otherwise, it would mean that for increasing feed gas humidity up to 400 ppm more OH molecules become

excited than for the dry condition despite de-excitation losses by increased quenching and reduced electron temperature.

Interesting results were also found by analysing the axial dependence of the emission signals. Except for nitrogen, the highest emission intensity was always found close to the nozzle. From there on, the intensity decreases towards longer nozzle distances. Nitrogen behaves differently because no significant nitrogen density is present close to the nozzle. Hence, no nitrogen excitation can occur in this region. With a longer nozzle distance, the diffusion of ambient nitrogen into the effluent drastically increases [21, 24] and a stronger emission intensity is obtained. However, since the excitation of nitrogen becomes less likely with increasing distance from the nozzle, an emission maximum is reached at an axial position of about 5.5 mm. For larger distances the emission signal decreases monotonically despite a still increasing nitrogen density.

Analogous to ambient nitrogen ambient oxygen also diffuses into the effluent and is potentially dissociated by

Table 1. Quenching rate coefficients for different excited species with state energies E , when H_2O is the quenching molecule. Furthermore, the decay rates due to radiative decay and due to quenching (for a humidity concentration of 1000 ppm) are given.

Excited state	E (eV)	Quenching rate coefficient for H_2O ($10^{-10} \text{ cm}^3 \text{ s}^{-1}$)	Decay rate (10^7 s^{-1})	
			Radiative	Due to H_2O quenching ^a
N_2 ($C^3\Pi_u, \nu = 0$)	11.18 [26]	7.1 ± 0.7 [27]	2.39 ± 0.08 [27] 2.63 [28]	1.75 ± 0.17
OH ($A^2\Sigma^+, \nu = 0$)	4.25 [29]	5.66 ± 0.1 [30]	0.139 [30]	1.40 ± 0.02
$\text{O}(3p^3P)$	10.99 [31]	49 ± 3 [32] 9.4 ± 1.5 [33]	2.69 ± 0.24 [32]	12.1 ± 0.74 2.32 ± 0.37
$\text{O}(3p^5P)$	10.74 [31]	n/a	2.56 [34]	—
$\text{Ar}(1s_5)$	11.55 [31]	4.8 [35]	Metastable	1.19
$\text{Ar}(2p_2)$	13.33 [31]	n/a	4.35 [31]	—

^a Calculated for a H_2O concentration of 1000 ppm.

radiation or high energetic particles. However, the highest emission intensity is found in front of the jet nozzle. This indicates that, in addition to the atomic oxygen production in the jet effluent, atomic oxygen is also produced inside the active plasma zone. Analogous to the discussion of feed gas humidity originating from the gas supply tubes (see section 3.1), it is reasonable to assume that even for the dry condition, water molecules are still present in the feed gas in a concentration between 0 ppm and 20 ppm (the detection limit of hygrometer). Hence, atomic oxygen is generated from water dissociation in the active plasma zone and becomes excited by electron collisions. This explains the observed emission signal of atomic oxygen in front of the jet nozzle.

OES was also used to indicate whether feed gas humidity has a larger effect on the OH emission than ambient air humidity. Figure 9 shows the integrated emission intensity of OH for the two investigated conditions. When dry feed gas and humidified shielding gas are used the axial intensity increases with distance to the nozzle. Similar to the discussion for nitrogen, the OH density continuously increases with distance. Since excitation is limited in the axial direction the emission profile forms a maximum. Identical to nitrogen emission, the highest emission signal was found at a distance of 5.5 mm. In contrast, when humidified feed gas and dry shielding gas are used the emission signal decreases steadily in the axial direction and no maximum evolves. Particularly interesting is the fact that although the feed gas humidity concentration in this case is by a factor of 42 lower compared with the humidity concentration in the wet shielding gas, a higher OH emission signal is detected (for $z < 5$ mm).

In conclusion, ambient air humidity influences the chemical and physical processes in the plasma effluent and must be considered especially for applications that require stable plasma conditions. In fact, much more important than controlling the ambient humidity is to consider the feed gas humidity since it affects the plasma substantially.

3.3. Ozone net production rate depends on feed gas humidity

In addition to other reactive species such as atomic oxygen or hydroxyl radicals, ozone (O_3) also plays a key role in plasma chemical processes. Under a condition where molecular oxygen density is significantly higher than the atomic oxygen density, which is usually the case for the effluent region of the

Figure 9. Comparison of the axial OH emission profiles for two conditions: firstly, dry shielding gas and humidified feed gas; secondly, humidified shielding gas and dry feed gas.

plasma jet, ozone can be interpreted as an indicator for atomic oxygen since it is mainly produced by the following reaction:

where M is an arbitrary collision partner, e.g. argon. Since the emission of excited atomic oxygen is linked to the humidity concentration in the feed gas it is reasonable to assume that the production of atomic oxygen and therefore the production of O_3 are also affected by feed gas humidity changes. The O_3 measurement result of the two applied absorption spectroscopic techniques (UV and IR laser absorption spectroscopy) is presented in figure 10. In order to compare both methods the ozone net production rate of the jet was calculated and plotted in this figure. Despite the different techniques remarkable agreement between both methods is ascertained. Generally, the highest O_3 net production rate of $(3 \dots 4) \times 10^{17} \text{ s}^{-1}$ is found for the driest condition. With increasing humidity concentration up to 200 ppm a steep decrease in ozone net production rate is obtained. For humidity concentrations larger than 200 ppm a decrease in ozone net production is still observed but is not as strong as for lower humidity concentrations. For more than 800 ppm the O_3 net production rate drops below $1 \times 10^{17} \text{ s}^{-1}$. A further

Figure 10. Ozone net production rate in dependence on the feed gas humidity concentration. The measurements were made with UV and IR absorption techniques.

humidity increase does not change the O_3 net production rate significantly.

The decrease in ozone net production with increasing feed gas humidity means that a higher humidity level either inhibits the production and/or favours the destruction of ozone. Without a plasma chemical model or an O atom density measurement it is difficult to disentangle both mechanisms. However, since ozone is mainly formed by one single reaction (equation (1)), atomic oxygen acts as a bottleneck for the ozone production. If the atomic oxygen density is affected by the feed gas humidity, an increase in feed gas humidity will consequently influence the O_3 formation as well. A possible mechanism for the reduction of excited atomic oxygen density by water molecules is given by the following reactions [39]:

Both reduce the amount of atomic oxygen available for reaction (1). This results in a lower ozone production with increasing water molecule admixture. However, not only a lower atomic oxygen concentration but also different humidity-depending destruction mechanisms reduce the ozone concentration and lead to the observed ozone behaviour of figure 10. One candidate is the reaction of ozone with OH in a cyclic process [40]:

where a single OH molecule is able to destroy more than one ozone molecule. According to the discussion in section 3.2, the OH density most likely increases with humidity in order to explain the observed maximum in the OH emission signal. Hence, the ozone destruction cycle of equations (4) and (5) becomes prominent when the water concentration in the feed gas rises. Furthermore, reactions with hydrogen also lead to ozone decomposition [40]:

Figure 11. H_2O_2 concentration in a plasma-treated liquid increases with feed gas humidity. Merckoquant stripes and the Amplex Red assay were utilized for H_2O_2 detection.

Since hydrogen can only originate from water dissociation an increase in feed gas humidity will favour the generation of atomic hydrogen, which in turn will contribute to the ozone decomposition.

3.4. H_2O_2 concentration in RPMI increases with feed gas humidity

The increase in feed gas humidity not only changes the optical emission spectrum and the ozone production of the jet but also affects the plasma-treated liquid. In figure 11, the H_2O_2 concentration is displayed for different feed gas humidity concentrations. The results of both utilized methods, namely the Merckoquant stripes and the Amplex Red assay, show remarkable agreement and indicate an increase in H_2O_2 concentration in the liquid with increasing feed gas humidity. For the highest investigated humidity setting a H_2O_2 concentration of almost 4 mg l^{-1} is measured after 40 s plasma exposure. Under dry gas conditions an H_2O_2 concentration of around 0.5 mg l^{-1} is still measured with the Amplex Red assay. Owing to the limited sensitivity no reliable data are obtained from the test stripes for this condition. For both humidity settings (driest and most humid), the H_2O_2 production rate is calculated assuming a linear increase in the H_2O_2 concentration with treatment time. Production rates of $1.8 \times 10^{-3} \text{ g h}^{-1}$ for the highest humidity setting and $2.0 \times 10^{-4} \text{ g h}^{-1}$ for the driest feed gas condition are obtained. A similar value of $3 \times 10^{-4} \text{ g h}^{-1}$ was obtained by Ikawa *et al* using a plasma jet driven in dry helium [41].

Different mechanisms are known on how hydrogen peroxide is generated in liquids [42, 43]. A basic pathway is the recombination of hydroxyl radicals in the liquid. These OH molecules are either generated directly inside the liquid by radiation, excited species or electrons from the plasma or are primarily produced in the plasma phase and recombine afterwards on the liquid surface to H_2O_2 . Another possibility is that H_2O_2 is produced in the gas phase and simply dissolved in the liquid. In that case the H_2O_2 concentration in the gas phase must increase with increasing humidity level. Support

Figure 12. Cell viability determined by resazurin assay in dependence on the feed gas humidity concentration. The cells were treated indirectly.

for this theory is found in the model results of Liu *et al* where an increase in H_2O_2 with humidity is predicted [16]. However, this model is computed for helium, whereas in this work an argon plasma jet was used. Measurements of H_2O_2 in the gas phase are necessary and planned for future work in order to identify a correlation between H_2O_2 in the gas and liquid phases.

3.5. Cell viability after indirect plasma treatment is impeded with higher humidity

In figure 12, the influence of different feed gas humidity settings on the viability of human skin cells is presented. Cell viability detection utilized in this study relies on a cellular supply of reduction equivalents to reduce a tracer compound (resazurin, containing an N-oxide) enzymatically yielding a highly fluorescent reporter dye (resofurin). Hence, cells need to be both living and metabolically active to be detected, which allows the conclusion on their actual viability rather than their number only. Cell viability and cell division rates strongly depend on the general condition of the cells, which will be influenced by many factors, such as nutrition, availability of oxygen, and also on harming influences such as oxidative stress. Under this condition, cells face imbalance between reactive oxygen species (ROS) concentration and their ability to detoxify the reactive intermediates or to repair the resulting damage. Some ROS or their degradation products are known to be highly reactive with biomolecules such as proteins, lipids and DNA. In addition to a direct random modification of these molecules by covalent implementation of functional groups or altering their redox status, indirect effects can be transmitted via cellular redox sensors such as the different peroxiredoxins [44]. In turn, these molecules activate cellular antioxidant defence mechanisms, which allow the cells to withstand a certain ROS excess. From figure 12 it is evident that the plasma generated by the feed gas with a content of humidity affects the cell viability even more than the plasma created from the dry feed gas. Considering the findings of the H_2O_2 concentration

measurements of section 3.4 a correlation between cell viability and H_2O_2 concentration becomes obvious. This means when the H_2O_2 concentration is low, the cell viability is high. With increasing H_2O_2 concentration a decrease in cell viability is also observed. In conclusion, by applying the plasma source in our eukaryotic cell model the cells were exposed to ROS and, hence, to oxidative stress, which exceed the cells' capacity for detoxification and repair. Although H_2O_2 plays an important role in the oxidative stress that affects the cell viability, other reactive species such as OH or O_2^- might also be involved in the observed effects. Cell viability could be influenced by mechanical stress or electric fields derived from the plasma source [45, 46]. Since indirect plasma treatment was applied, these and other stress factors such as UV radiation were excluded for our experimental conditions.

4. Conclusion

In this paper, the significant influence of feed gas humidity introduced into an atmospheric-pressure argon plasma jet is demonstrated on the axially resolved optical emission spectrum of the jet effluent, the generation of hydrogen peroxide in a plasma-treated cell growth medium and on human skin cells charged with this treated medium. Optical emission spectroscopy reveals a strong dependence of present excited plasma species such as hydroxyl radicals, molecular nitrogen, argon and atomic oxygen on the feed gas humidity. Furthermore, it was demonstrated by this spectroscopic method that small feed gas humidity concentrations (below 1000 ppm) have a much larger effect on OH emission than high ambient humidity concentrations. From this finding it is concluded that whenever humid feed gas is applied, ambient humidity can be neglected for processes inside the effluent region. However, when the feed gas is dry, ambient humidity becomes important and must be considered in such a situation. The question whether used feed gas is dry or humid is complicated by the fact that polymeric supply tubes have a major contribution to the feed gas humidity. It is shown that a vast amount of water vapour diffuses through PUR tube walls and hence increases the feed gas humidity. Compared with metal tubes a ten times higher concentration was found for the polymeric tubes. Depending on the length of the feed gas tubes, a constantly changing feed gas humidity prevents stable plasma conditions for up to several hours.

Furthermore, plasma treatment of the liquid cell growth medium with increasing feed gas humidity leads to an almost linear increase in H_2O_2 concentration in the liquid. When human skin cells get challenged with this medium a strong correlation between cell viability and H_2O_2 concentration is revealed, indicative of an oxidative stress response.

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