Interactive comment on “Investigation of coastal sea-fog formation using the WIBS (Wideband Integrated Bioaerosol Sensor) technique” by Shane M. Daly et al.

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We thank the reviewers for carefully reading our manuscript and for providing the critical review to improve the manuscript. In the following, we provide responses to both of the reviewers comments and concerns.

Reviewer 1 comments:
Comment: While the timeline link between molecular iodine release at low tide, iodine oxide formation (in daytime) and particle formation is well established (references cited in the paper itself), the study links these two with the formation of coastal mist
for the first time. I would then recommend a slightly more nuanced statement about the novelty of this 3-step time-line in the abstract and conclusions. What is certainly new and exciting about the study is the field observation of I$_2$ adsorbed onto water droplets. This appears to be a previously unknown stabilizing transport mechanism for the dispersal of I$_2$ in the marine environment. This finding is supported by targeted laboratory experiments demonstrating that the non-biological WIBS signals observed in the field are in fact consistent with the adsorption of I$_2$ on the surface of nebulized pure water droplets. **Manuscript changes:** Abstract and conclusion has been modified to reflect the reviewer comments.

**Page 2, Line 17:** While the process of molecular iodine release, particle formation and sea-fog formation have been studied in detail in previous studies, this study provides a potential link of the three phenomena.

**Page 28, Line 4:** The dual field and laboratory study presented here provides a possible real-time profile as observed on several occasions in July 2011 at Haulbowline Island, Co. Cork.

**Comment:** The absence of fluorescence for nebulized seawater in the presence of iodine vapour may well be explained by the formation of trihalide ions as the authors suggest. It would be interesting to know, though, whether the saline solutions prepared for the lab experiments have similar iodide and chloride concentrations to those expected in sea spray, and whether more diluted concentrations would have resulted in increased signal in the FL1 channel. Also, whether changing the residence time of the nebulized sea salt solution in the chamber could have shown some evidence of the kinetics of the removal of the adsorbed I$_2$ molecules at the surface. In any case, the characterisation of the sea salt solution should be included in section 2.3. I would like to see also some information about the pressure, temperature, flow rate, and residence time conditions in the aerosol dispersion chamber. **Response:** This study can indeed be expanded to flow tube experiments where there is a better control of the residence time. In terms of the experiment conditions, there was no pressure
gauge present in the system however the system can be considered to operate under atmospheric pressure. The system was allowed run for 30 minutes before and after each experimental run each experiment using the WIBS4 pump at 2.5 L/min. For all experiments (the iodine vapour with water droplets, sea salt droplets and mixed iodine/water droplets), the air flow in the system was 5.6 L/min for generating the aerosolised water droplets. For the solely iodine vapour experiments, only sublimed iodine was released into the system (No carrier gas), again after the system was pumped down for 30 minutes after each run. With regards to the residence time, there was no adjustable injector to vary reaction distance in the chamber. The only possibility was changing the direction of the injection (pointing the water aerosoliser upwards allowed for more adequate mixing time). **Manuscript changes:** 

Page 8, Line 1: 0.25g of $\geq 99\%$ iodine crystals were measured for each sublimation test, with 0.05 g of refined rock salt from the Wieliczka Salt Mine used for the saltwater mimic tests. A smaller quantity of salt was used to avoid overloading the detector. 

Page 8, Line 24: Before each experiment, the system was pumped down for 30 minutes to remove any residual material using the WIBS-4A total pumping capacity of 2.5 L/min. During the experiment, a flow of 5.6 L/min of compressed air was supplied for the aerosolization into the system. No pressure transducer was present in the system at the time so estimated conditions were of the order of 1 atm at 298 K. Relative humidity was $>70\%$ based from observation of the chamber rather than direct measurement. 

Page 9, Line 7: This study could be later applied at a flow tube experiment where there is a greater control of experiment conditions such as residence time. 

Page 9 Line 11: The WIBS fluorescence data obtained in the experiments were filtered using thresholds most commonly utilized in the literature (ie the mean of forced trigger mode values + $3\sigma$ method) (Hernandez et al., 2016):

**Comment:** The absence of fluorescence when only iodine vapor is admitted in the laboratory chamber is very interesting, but I don’t find very
convincing the argument given by the authors to explain this point. The WIBS-4 instrument is optimized for particle detection, but in its configuration (http://www.dropletmeasurement.com/widebandintegrated-bioaerosol-sensor-wibs-neo) I don’t find any obstacle for the detection of gas-phase fluorescence (perhaps the authors could comment more on that to make it clearer). I$_2$ molecular iodine presents a strong absorption feature in the 170-210 nm spectral range (the Cordes bands, D-X system), with peak absorption cross section of 2E-17 cm$^2$ molecule$^{-1}$ at 188 nm (Myer and Samson, 1970; Roxlo and Mandl, 1980). After absorption of VUV and middle UV photons, fluorescence from the D ion-pair state back to the ground state exhibits an ordinary bound to bound spectrum together with a bound to free diffuse quantum interference spectrum (the McLennan bands) (Tellinghuisen, 1974; Exton and Balla, 2004). Concurrently, a significant fraction of the initial D state population is collisionally transferred to the D’ state at increasing buffer gas pressure, resulting in fluorescence in the D’-A’ band at 340 nm. Since the transition probability of the D-X system at 280 nm is small (absorption cross section of 6E-19 cm$^2$ molecule$^{-1}$ (Saiz-Lopez et al., 2004)), this may explain why I$_2$ is not observed in the gas phase. On the other hand, complexation with water may red-shift the absorption spectrum. E.g. the peak of the I$_2$ VUV band shifts to 203 nm in aqueous solution (Kireev and Shnyrev 2015). This would be a plausible explanation as to why 280nm-pumped fluorescence in the 310 - 400 nm range can be observed when I$_2$ is complexed with water and not in the absence of water droplets. In feel that some spectroscopic discussion in this sense is pertinent. 

**Response**: Work indicated by Alizadeh et al., 2012 showed an I$_2$ peak at 450 nm but also a trace absorption form 250-290 nm (See figure 1 of that paper). O'Driscoll et al., 2008, discusses I$_3^-$ at 288 nm and 352 nm (both broad enough to meet the 280 nm and 370 nm flash lamp requirements of the WIBS). During the UV absorption analysis, the 288 nm and 352 nm peaks were observed, indicative of I$_3^-$. However, the 450 nm I$_2$ peak was also present in the spectra, indicative of some equilibrium between the two. The aerosolized mixture of iodine and water should have given a fluorescent signal if the I$_3^-$ ion was fluorescent as it was present in UV-absorption analysis but this
wasn’t the case. The likely explanation for the observed FL1 fluorescence is the I$_2$ molecule adsorbing to the water droplet surface. Work by (Liu et al., 2004) show that if I$_2$ is complexed with different organic solvents such as toluene, a strong absorption band is present at 280 nm due to the I$_2$ molecule red-shifting with the organic complex. A similar process is possible, with the I$_2$ not residing in gaseous or liquid phase but rather binded to the surface of the water droplet. The reason for no iodine vapour fluorescence is simply due to the vapour not existing in the particle phase for detection by the laser. If the laser doesn’t detect a particle, then the flash lamps are not triggered, hence no fluorescence because no excitation. It’s not stating that iodine vapour doesn’t fluoresce, but that whatever the WIBS saw at Haulbowline wasn’t attributed to that. One possibility is a non-fluorescent particle triggering the PMT with the laser, followed by triggering of the flash lamps, picking up I$_2$ fluorescence from surrounding vapour phase. However, after pumping the chamber down for 30 minutes, no particles are detectable, ensuring sufficient vacuum. **Manuscript changes:** Iodine vapour will be removed from the table as it may cause confusion in relation to the particle measurements.

*Page 21, line 19:* Iodine vapour fluorescence was not measurable because it does not provide particles which are necessary for detection by the internal diode laser of the WIBS. If the laser does not detect a particle of the specified size, the flash lamps are not triggered and therefore fluorescence cannot be detected because no excitation occurs. One possibility is that a non-fluorescent particle could be detected, which activates the flash lamps and a fluorescent signal from the surrounding iodine vapour is detected. However, with the chamber being pumped down for 30 minutes before each experiment, no particles of that size can interfere.

*Page 23, line 21:* The I$_3^-$ peak at 352 nm was not present at all in cold saltwater mimic samples and only in trace amounts (<5% of iodine and water mix) in heated samples while the peak at 288 nm largely remained. Work indicated by Alizadeh et al (2012) showed an I$_2$ peak at 450 nm but also a trace absorption at 250-290 nm. Therefore, it shows the possibility of I$_2$ absorbance at 280 nm even though the expected cross
section of $I_2$ at 288 nm is $<1 \times 10^{18}$ cm$^2$ (Roughly 18 times smaller than the peak absorbance at 203 nm (Saiz-Lopez et al., 2004). The absorbance spectra for each sample is available in the supplementary material.

Page 24, line 19: It is known for $I_2$ in vapour form that the transition probability of the D-X system at 280 nm is $<1 \times 10^{18}$ cm$^2$. This value is about eighteen times smaller than the peak absorbance at 203 nm (Saiz-Lopez et al., 2004). However work by (Liu et al., 2004) show that if $I_2$ is complexed with different organic solvents such as toluene, a strong absorption band is present at 280 nm due to the molecular $I_2$ UV spectrum red-shifting. A similar interaction can be envisioned for iodine bound to the surface of water droplets rather than being present in the vapour phase or simply dissolved inside a droplet.

Comment: Section 2.3 mentions that fluorescence spectra of the solutions were investigated using a Shimadzu RF-6000, but the results of these investigations are not reported. I also find that the solution absorption data is presented in a rather schematic way and that it would be more informative to show absorbance spectra (perhaps in the supplementary material) to demonstrate how efficient is the absorption of the two WIBS wavelengths. Response: The main implications of the spectra: Iodine and Milli-Q water – Gives $I_2$ at 288 nm and 450 nm as well the tri-iodide $I_3^-$ at 288 nm and 352 nm. Iodine and saltwater mimic – Initially only iodine chlorides present. Later waiting periods show retention of the 288 nm and 450 nm peak with the 352 nm reduced. Figures: Figure 1: UV Absorption spectrum of iodine in milliQ water with a range from 250 – 600 nm, Figure 2: UV Absorption spectrum of iodine in saltwater mimic, Figure 3: UV Absorption spectrum of iodine in a saltwater mimic after 4 days. Manuscript changes: The spectra has been added to the supplementary data and reference to the table in the paper.
Comment: The suggested link between I$_2$·(H$_2$O)$_x$ and HIO$_3$ is a problematic one. Sipila et al. 2016 did observe HIO$_3$ and molecular cluster formation in their laboratory experiments, but in the presence of water vapor (no nebulized water droplets). Daly et al. do not report WIBS measurements of a mixture of water and iodine vapor, but it is known that I$_2$ and H$_2$O form a weakly bound complex (Galvez et al. 2013), and under atmospheric conditions only a residual amount of I$_2$ would be complexed with H$_2$O. The presence of I$_2$·(H$_2$O)$_x$ in the laboratory experiments of Sipila et al. is therefore unlikely. Furthermore, HIO$_3$ increase was observed by Sipila et al. in the field at daytime, well after noon, while Daly et al show that I$_2$·(H$_2$O)$_x$ is a night time reservoir which disappear quickly after sunrise. Response: The work by Galvez et al., 2013 suggests a theoretical weakly bound complex of one I$_2$ molecule with one H$_2$O molecule (14 kJ mol$^{-1}$). Further studies should address apply the iodine/water vapour method to check if this is a possibility. However, a different mechanism is suggested here. Several I$_2$ molecules would bind to one H$_2$O droplet as a surface ionic mechanism. The interaction between iodine and oxygen from the water droplet may result in enough partial positivity in the iodine atom to make it susceptible to radical attack, thus starting the reaction process to HIO$_3$. In the paper, the WIBS-4 peak for fluorescent counts at 6 am was during the middle of July (approaching the longest day of the year where sunrise would be before 6 am) whereas the observations by Sipila et al., 2016 were made from August to October. At this point, morning daylight would have started later sometime between 6:30-8:30 am. There is 3 hours of delay between the WIBS signal decrease and the SMPS count increase. If the HIO$_3$ and IO data (Figure 1.) was recorded towards the October period, the later sunrise could correlate the observed HIO$_3$/IO traces to the study here. Manuscript changes: Page 26, line 22: Work by Galvez et al (2013) suggests a theoretical weakly bound complex of one I$_2$ molecule with one H$_2$O molecule (14 kJ mol$^{-1}$). However, the current study likely addresses cases when several I$_2$ molecules bind to one or more water droplets in a surface adsorption mechanism.

Page 27, line 7: In fact, a recent report, which outlines evidence for some coastal
aerosol particle formation being due, in part, to the sequential addition of HIO$_3$ indicates that at Mace Head, Ireland, the production of the oxo-acid has been shown to begin at sunrise reaching a maximum at noon (Sipila et al., 2016). It should be noted that the measurements were made during the August to October period. During that time, the iodic acid is reported to appear at 13.00 with the IO radicals peaking at 15.00 pm due to later sunrise.

**Minor Comments:**

Comment: Page 6, line 20: 0.5-20 m. do you mean micron?
Response: Yes this was meant to be micron.
Paper edit: This change has been included in the paper.

Comment: Page 11, lines 2 and 3: it looks like all these size ranges should be microns rather than meters.
Response: Yes they should be.
Paper edit: This change has been included in the paper.

Comment: All figures. In general, the legends and axis labels are too small and difficult to read, especially in multi-panel figures.
Paper Edit: Each graph has been re-edited to include larger legends and axis labels.

Comment: Figure 9: some of the tidal height data is missing: the 6:00AM and 18:00 tidal values are not shown (as opposed to figure 1 in the supplementary information). Paper edit: This change has been included in the paper.

**Reviewer 2 comments:**
Comment: I think Section 2 needs a more detail on A) the laboratory experiments that took place using the WIBS-4A instrument and B) the preparation for the data for both the field analysis and complementary laboratory studies. Response: The manuscript has been modified to reflect these comments. Manuscript changes: As already indicated in response to reviewer 1

Comment: From my understanding the author uses the FT signals as the fluorescence threshold, and compares results to what was seen in the Hernandez et al., 2016 publication, however this publication uses the default FT + 3 \( \sigma \) threshold. Response: The paper has been rewritten to correct any ambiguities within regard to the threshold used for the work. The default mean Forced Trigger + 3 \( \sigma \) threshold was used manuscript changes: Page 9 Line 11: The WIBS fluorescence data obtained in the experiments were filtered using thresholds most commonly utilized in the literature (ie the mean of forced trigger mode values + 3\( \sigma \) method) (Hernandez et al., 2016):

Comment: laboratory and field study should be more explicit (e.g. size calibration information, fluorescence calibration information, the fluorescence threshold chosen-whether it is the average, median, etc.). Size calibrations were carried out using several PSL sphere ranges 0.5, 0.82, 1, 2, 4, 10, 12 microns Response: At the time of the work the Robinson fluorescence calibration paper had not been published. Thus a fluorescence calibration was not under taken. Manuscript changes: Page 9, Line 23: For the WIBS-4A instrument, size calibrations were carried out using Polystyrene Latex Spheres (PSL) with diameters 0.5, 0.82, 1, 2, 4, 10, 12 \( \mu \)m. The internal photomultipliers for each WIBS were not measured at the time.

Comment: Section 2.2 Field Instrumentation: On page 6 lines 21-24, the author states that both the WIBS-4 and the WIBS-4A units were identical in terms of functionally-this is strong statement. Response: We agree with the reviewer and have modified
this point significantly. **Manuscript changes:** The section has been edited to read 

*Page 6, Line 31:* Both instruments display similarities in terms of sampling methods and build but have a few distinctions such as the WIBS-4 dual gain detection approach and the WIBS-4A double threshold system. The WIBS-4A has a slightly higher flow rate at 2.5 L/min and 300 ml/min (flow velocity of 18 m/s) compared to the WIBS-4 at 2.4 L/min and 230 ml/min (flow velocity of 12 m/s). Similarly, variation in fluorescent intensity between WIBS instruments for identical particles is a potential problem in such studies a problem which has been discussed previously in the literature. (Robinson et al., 2017, Savage et al., 2017, and Könemann et al., 2018)

**Comment:** Can the author please comment on whether such calibration (fluorescence) was done? Where the PMT voltages measured for each WIBS unit?

**Response:** No as explained in response to reviewer 1, the PMT voltages were not measured

**Comment:** Page 11, lines 5- 20: In general, I think this section needs more discussion in regards to the suggested publications and their analysis strategies - Gabey et al., 2010, Perring et al., 2015, Savage et al., 2017 and Savage et al., 2018. It is not clear what the author means by stating “Unusually, fluorescence signals were mainly measurable in the FL1 channel. FL2 registered little emission above threshold as illustrated in Figure 4, which shows plots of size/AF data as a function of the FL1 and FL2 channels. (FL3 showed no fluorescence). The larger size feature (2-6 um) consisting of highly fluorescent solid particles/droplets but only in the FL1 channel represents a behaviour that has not been observed previously in any WIBS field campaign. Hence fungal spores, certain pollen and bacteria as large as 2 um (Hernandez et al., 2016) can be found in the 2-6 um size regime but are fluorescent in all channels because of their amino acid, tryptophan and NAD(P)H contents”. **Response:** We agree with the reviewer and the section has been rewritten to reflect this as follows. **Manuscript**
changes: Page 11, Line 19: The work by Hernandez et al (2016) suggests that some fungal spores show fluorescent characteristics that are present in FL1 but not FL3. However, the conditions on site would not favour spore release as the island has very little soil-based vegetation with only sea kelp present. Very low wind speeds were recorded during the measurement periods (<2.5 m/s during the 15th – 16th July and <5 m/s during the 26th-28th July). Therefore, it was highly unlikely that material could be carried on to the island from the mainland. In any case, the particles are less likely to be bacteria or pollen because of size constraints. Hence bacteria sizes are found at the lower limit (and below) of WIBS detection. Pollen sizes are often measured at much higher than the upper limit of WIBS detection and so are generally captured by the particle trap. During the summer period, the dominant fungal spore in locations close to but not at the Cork Harbour coastline, is known to be Cladosporium which is generally released during the day time (10:00 am - 12:00 pm onwards) and under dry conditions (O’Connor et al., 2015, Healy et al., 2014). By contrast in this study WIBS particle detection was found in the night-time period between 00:00 – 08:00 am.

Page 12, line 11: Ascospores are linked with rain releases but only 0.2 mm of rainfall was recorded after 09:00 am on the 16th July, after which the WIBS signal is seen to decease (O’Connor et al., 2015).

Comment: Several studies suggest there are non-biological, fluorescent particles that may be interferences when discriminating between bio vs. non-biological particles, and even different particle types (Huffman et al., 2010, Pohlker et al., 2015, and Savage et al., 2017, and references there-in). Can the author comment on these possible interferences, and if these substances were taken into consideration during their field analysis? Manuscript changes: The manuscript has been updated to reflect the comments of the reviewer. Page 20, Line 12: It should be not that other non-biological particles have been seen to be fluorescent. Mineral dust was also considered as a potential source of fluorescent particles. In fact, studies have shown that fluorescent mineral dust can contribute up to 10% of the total measured (Toprak and Schnaiter., C11
However in this study, using the FL1 and FL3 channel filters this dust artefact was removed. A lack of FL3 fluorescence signals in this study rules out the presence of mineral dust because it weakly fluoresces in the FL1 and FL3 detection ranges and therefore is considerably weaker than biofluorophore signals (Toprak and Schnaiter, 2013; Pohlker et al., 2012). Polycyclic Aromatic Hydrocarbons (PAH’s) could also be considered a potential interference to the measurements made here due to their highly fluorescent nature but since they are largely present on the surface of soot particles which generally exist in submicron sizes, detection by the WIBS is unlikely unless oil droplets were present. The complex chemical environments associated with soot particles can also lead to fluorescence quenching of PAH’s (Pohlker et al., 2012). Humic-like substances (HULIS) and secondary organic aerosols (SOA) have also been indicated as potential interference signals in the literature (Pohlker et al., 2012).

References added:


Interactive comment on Atmos. Chem. Phys. Discuss., https://doi.org/10.5194/acp-2018-673,
Fig. 1. UV Absorption spectrum of iodine in milliQ water with a range from 250 – 600 nm
Fig. 2. UV Absorption spectrum of iodine in saltwater mimic
Fig. 3. UV Absorption spectrum of iodine in a saltwater mimic after 4 days

- 288nm $I_2$ or $I_3$
- 352nm $I_3$ (Trace)
- 450nm $I_2$