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## Infrared Degenerate Four-Wave Mixing with Upconversion Detection for Quantitative Gas Sensing

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Dr. Jialan Zhang  
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*Jove*

2018-09-03

Dear Dr. Jialan Zhang:

I am pleased to submit an original research article entitled “Infrared Degenerate Four-Wave Mixing with Upconversion Detection for Quantitative Gas Sensing” by Rasmus Pedersen and Zhongshan Li for consideration for publication by Jove. We previously shown the advantages of combining upconversion and DFWM, and this manuscript aims to make the technique more approachable.

In this manuscript, we describe the procedure for aligning the degenerate four-wave mixing spectroscopy setup, as well as the procedure for setting up the upconversion detector for consistent, optimal detection.

We believe that this manuscript is appropriate for publication by Jove because it is focused on the practicalities of the technique, which matches the scope of Jove.

This manuscript has not been published and is not under consideration for publication elsewhere. We have no conflicts of interest to disclose. If you feel that the manuscript is appropriate for your journal, we suggest the following reviewers:

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Thank you for your consideration!

Sincerely,

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**TITLE:**

Infrared Degenerate Four-wave Mixing with Upconversion Detection for Quantitative Gas Sensing

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**KEYWORDS:**

gas spectroscopy, photon upconversion, upconversion detection, infrared, degenerate four-wave mixing, intracavity upconversion

**SUMMARY:**

Here, we present a protocol to perform sensitive, spatially resolved gas spectroscopy in the mid-infrared region, using degenerate four-wave mixing combined with upconversion detection.

**ABSTRACT:**

We present a protocol for performing gas spectroscopy using infrared degenerated four-wave mixing (IR-DFWM), for the quantitative detection of gas species in the ppm-to-single-percent range. The main purpose of the method is the spatially resolved detection of low-concentration species, which have no transitions in the visible or near-IR spectral range that could be used for detection. IR-DFWM is a nonintrusive method, which is a great advantage in combustion research, as inserting a probe into a flame can change it drastically. The IR-DFWM is combined with upconversion detection. This detection scheme uses sum-frequency generation to move the IR-DFWM signal from the mid-IR to the near-IR region, to take advantage of the superior noise characteristics of silicon-based detectors. This process also rejects most of the thermal background radiation. The focus of the protocol presented here is on the proper alignment of the IR-DFWM optics and on how to align an intracavity upconversion detection system.

**INTRODUCTION:**

IR-DFWM provides the capability of measuring concentrations of IR active species down to the ppm level<sup>1</sup>, with spatial resolution. IR-DFWM has several advantages that make it an attractive technique for combustion research. Flames can be changed drastically by the insertion of probes, but IR-DFWM is nonintrusive. It has a spatial resolution, so species' concentrations at different points in the flame structure can be measured. It provides a coherent signal, which can be isolated from the thermal emission of the flame. Additionally, DFWM is less sensitive to the collision environment than, for instance, laser-induced fluorescence (LIF), which can be difficult to determine in a flame. The technique also provides access to molecular species that are IR

active but lack visible or near-visible transitions that can be used to measure them with other techniques.

While DFWM has a number of advantages, alternative techniques might be preferable if one or more of these advantages are not needed. If spatial resolution is not needed, absorption-based techniques will be both simpler and more accurate. If the molecular species in question have transitions in the visible or near-IR region, LIF might be preferable, as LIF can provide spatially resolved information from a plane rather than just a single point. Under the right conditions, nonlinear methods, such as DFWM and PS, can also be used for single-shot 2D measurements<sup>2</sup>. The signal of these nonlinear methods is proportional to the probe beam intensity cubed, and as the pump beam must be expanded to cover the area of the 2D measurement, this requires either very high pulse energies or a combination of high third-order susceptibility, high concentrations, and low background noise to work. Therefore, it mostly depends on the molecular species whether this is a possibility.

In a more direct competition with DFWM, there are the other four-wave-mixing-based spectroscopic techniques: coherent anti-Stokes Raman spectroscopy (CARS), laser-induced grating spectroscopy (LIGS), and polarization spectroscopy (PS). CARS is a well-established technique for measuring the temperature and major species in combustion environments. However, it lacks the sensitivity to detect minor species, as the detection limit is usually approximately 1%<sup>2</sup>. PS and DFWM have previously been shown to have similar sensitivity and detection limits<sup>3</sup>; however, the signal-to-noise ratio of DFWM has been shown to increase by factor 500 when combined with upconversion detection<sup>4</sup>, while PS has shown only a 64-fold increase<sup>5</sup>. LIGS has the advantage of inducing a grating, using mid-IR light, but measures the effect by the refraction of a probe laser of this grating, and the wavelength of this probe laser can be chosen freely<sup>6</sup>. The wavelength of the probe laser can, therefore, be in the visible region, where fast, low-noise silicon-based detectors are available. This is the same advantage that is achieved by using upconversion. LIGS does have the drawback that it is very sensitive to collision<sup>2</sup>, which means that the concentration of major gas species must be known for precise concentration or temperature measurements with LIGS. If that issue is overcome, LIGS does have a similar sensitivity to both DFWM and PS at atmospheric pressure<sup>3</sup>, but where the LIGS signal increases with increasing pressure, the signal from DFWM and PS increases at lower pressures, which means the preferred technique will depend on the pressure environment.

Upconversion detection is the technique of converting a signal from long wavelengths to shorter ones by using sum-frequency generation. The advantage of this is that the detectors in the visible or near-infrared range have lower noise and a higher sensitivity than their counterparts in the mid-IR region. This was first investigated five decades ago<sup>7</sup>, but saw very little attention and use since then, due to the low conversion efficiencies. However, with advances in the production techniques for periodically poled lithium niobate (PPLN) and other materials with high nonlinear coefficients, as well the increased availability of high-power laser diodes (LDs), the technique has attracted increased attention in the last decade, with applications covering areas such as mid-IR single-photon detection<sup>8-11</sup>, IR lidar<sup>12,13</sup>, and hyperspectral imaging<sup>14,15</sup> and microscopy<sup>16</sup>. The main advantage of combining upconversion detection with IR-DFWM is that the phase-match

condition has a narrow angular and spectral acceptance band, which discriminates heavily against the thermal background, allowing the detection of weaker signals.

## PROTOCOL:

The setup of the upconversion detector is shown in **Figure 1**; mirrors, lenses, or other optics referenced in the protocol are identified here or in the diagram of the IR-DFWM setup shown in **Figure 2**. The protocol section mainly deals with aligning the optical setup used for this method, and the process can be paused at any point by turning off all running equipment. All mirrors are adjusted manually. The software used here to control the camera and LD was delivered together with the upconversion detector. The use of the software is described at the end of the protocol.

### 1. Upconversion

1.1. Place the end mirror of the alignment cavity, UH, as indicated in **Figure 1**.

1.2. Remove the PPLN crystal from the crystal mount.

1.3. Place an IR-sensitive card (sensitive at 1,064 nm) at position A, see **Figure 1**.

1.4. Turn the angle of the kinematic mount holding UH to the extreme position in both the horizontal and vertical direction. Then, turn on the LD at approximately 1/3 of the maximum output.

1.5. Align the alignment cavity as follows.

1.5.1. Change the angle of UH by  $+0.2^\circ$  in the horizontal direction.

1.5.2. Sweep the vertical angle of UH from one extreme to the other, while watching the IR card for a beam from the alignment cavity.

1.5.3. Repeat steps 1.5.1 and 1.5.2 until the cavity starts lasing.

1.5.4. When the alignment cavity is lasing, switch between adjusting the angle of UH for a higher power and reducing the LD drive current. The LD is dimensioned to drive the full cavity, which has much higher losses than the alignment cavity. Keep the power where the beam leaving UH is easily visible with the IR card but no more than that.

1.6. Remove the IR card.

1.7. Adjust the angle of U2, so the alignment beam is reflected in the center of U3 (**Figure 1**).

NOTE: The beam from the alignment cavity should hit U2 in the center.

1.8. Adjust the angle of U3 so the beam continues to U4, U5, and U6 and is reflected from U6 to

U7.

1.9. The beam must pass through the PPLN mount at the height of the middle of the channels of the PPLN crystal, and it must enter the crystal perpendicular to the surface. Use U2 to correct the height and angle, while adjusting U3 to keep the beam level and centered through the holes x and y.

1.10. Remove the germanium window and place the IR card behind U7, so that an IR beam leaving the cavity will hit the card, and the fluorescence will be visible to the person aligning the cavity.

NOTE: The alignment beam will now be passing through the PPLN mount and hit U7.

1.11. Adjust the angle of U7, so that the reflection from U7 passes back along the path of the alignment beam. While adjusting the angle of U7, watch for a beam on the IR card. When a beam is seen, adjust the angle of U7 to maximize the output.

1.12. Mount the PPLN in the mount. Make sure the mount is placed so that the beam goes through one of the channels in the crystal.

1.13. Continue with the substep (step 1.13.1, 1.13.2, or 1.13.3) matching the current situation.

1.13.1. If an IR beam is still visible exiting U7, adjust U7 to maximize the output, and continue with the next step.

1.13.2. If the IR beam exiting U7 is no longer visible, increase the LD current to 1/3 of the maximum output, and check if the IR beam can be seen. If a beam is visible, go to step 1.13.1; otherwise, go to step 1.13.3.

1.13.3. Reduce the LD current to the previous level and trace the guide beam to see if it passes through the PPLN in the center of one of the channels. If it does not, repeat from step 1.7, but with the PPLN in the mount.

1.14. Turn off the LD, remove UH, and attach the LP750 filter at position B (see **Figure 1**).

1.15. Place the power meter behind U7 but leave space for checking the beam with an IR card. Then, turn on the LD at full power.

1.16. If no signal is seen on the power meter, make small angle changes to U7, while watching for a signal on the power meter. If a signal is found, continue to the next step; otherwise, return to step 1.1.

1.17. Optimize the cavity alignment by adjusting the angles of U2 and U7 to maximize power, while using a high-power IR card to check that the cavity is running in the fundamental Gaussian

mode.

NOTE: While it might be possible to get higher power in a higher order mode, it is essential for the conversion efficiency that the laser is running in the fundamental mode.

1.18. If the cavity is not running in the fundamental mode, it will be running in a higher order mode where multiple lobes are visible on the IR card. Turn U7 so that the lobes are brought closer together on the IR card, until they merge.

1.19. Record the power output at U7. Use this and the transmission of U7 to calculate the intracavity field. Compare this value to the calibration curve in **Figure 6**.

1.20. When the cavity has been optimized, remove the LP750 filter, and reattach the germanium window.

## 2. IR-DFWM Alignment

NOTE: See **Figure 2** for a diagram of the DFWM setup.

2.1. Align the HeNe laser beam (the guide beam) with M3 and M4 to hit L1 in the center, going horizontally from M4 to L1.

2.2. Insert BOXCARS plate 1 at a 45° angle to the beam (vertical direction) and ensure the beam goes through, producing two output beams.

2.3. Insert BOXCARS plate 2 at a 45° angle to the beams (horizontal direction) and ensure the beam goes through, producing four output beams. Adjust the angles of the plates so the beams are spaced like the corners in a square.

2.4. Adjust the position of L1 until the beams are equally spaced around the center of the lens.

2.5. Leave the signal beam, which will be generated along the path of the beam blocked by the beam block, unblocked for now, so it can be used to align the rest of the setup. Place the iris so it blocks the three pump beams but allows the fourth beam, the signal beam, to pass through.

2.6. Align L2 so the signal beam is collimated. This must be done using the focal lengths at the wavelength of the pulsed laser and not by visual inspection, as the focal lengths will be different for the wavelength of the guide beam and the mid-IR.

2.7. Place M5 and M6 so that the guide beam is centered on the input window of the upconversion detector and perpendicular to the input window.

2.8. Place L3 one focal length optical distance from the center of the PPLN. Take into account the refraction of the germanium window, the cavity mirror, and the PPLN itself.

221  
222 2.9. Set up the upconversion module and turn it on (see section 1).  
223

224 2.10. Remove the germanium window of the upconversion detector. This will allow a 1064 beam  
225 to exit the upconversion module.  
226

227 2.11. Overlap the HeNe laser beam and the 1064 beam from the upconversion detector by using  
228 M6 to move the 1064 beam onto the signal beam so that they overlap at L2, and by using M5 to  
229 move the guide beam onto the 1064 beam at L3. Alternate between the two mirrors until the  
230 guide beam and the 1064 follow the same path.  
231

232 2.12. Reattach the germanium window.  
233

234 2.13. Place several ND filters in the beam path, in front of the upconversion detector. Take great  
235 care to never let an unattenuated beam from the pulsed laser into the upconversion detector, as  
236 the high energy will likely damage the detector.  
237

238 2.14. Turn on the pulsed laser and ensure it is running stable and at an appropriate energy per  
239 pulse.  
240

241 2.15. Overlap the pulsed laser and the guide beam as follows.  
242

243 2.15.1. Adjust the angle of M1 till the pulsed laser overlaps the guide beam at the beam combiner  
244 (M2).  
245

246 2.15.2. Adjust the angle of the M2 so that the pulsed laser is reflected in the propagation  
247 direction of the guide beam.  
248

249 2.15.3. Check that the beams are overlapping at the beam combiner and at distances of 1 m, 2  
250 m, and 3 m.  
251

252 2.16. Find the focal point of the beams after L1. Place the gas flow or flame to be measured so  
253 that the measurement point is at the focal point of the beams.  
254

255 2.17. Connect the trigger signal from the pulsed laser to the upconversion detector to time gate  
256 the detection. If the time delay and gate time are not known, start with a long duration of the  
257 time gate and narrow it down when the signal is found.  
258

259 2.18. Search the setup, especially the BOXCARs plates, for errant reflections and ensure they are  
260 blocked.  
261

262 2.19. Optimize the alignment of the signal beam in the upconversion detector as follows.  
263

264 2.19.1. If a signal is visible on the detector, adjust M5 and M6 to maximize the signal.



2.19.2. If no signal is visible on the detector, reduce the ND filtering by one order of magnitude. Repeat until a signal is seen.

2.19.3. If the signal on the detector is saturated, increase the ND filtering by one order of magnitude. Repeat until the signal is no longer saturated.

2.19.4. Go through steps 2.19.1–2.19.3 until the signal can no longer be increased by adjusting M5 and M6.

2.20. Place the beam block so that it blocks the signal beam, as indicated in **Figure 2**. Then, remove the ND filters.

2.21. Adjust the position of the beam block to reduce any scattering (background noise) seen on the detector. Take great care not to unblock the beam accidentally and expose the detector to direct light from the pulsed laser.

2.22. Prepare the gas flow or flame to be measured. Then, scan the pulsed laser across the wavelength range of interest, while recording the signal from the detector. This will generate a spectrum matching the gas composition at the overlap of the beams.

### 3. Laser Diode Software

3.1. Run the LabVIEW program **AuroraOne control.vi**.

3.2. Click the **Laser TEC enable** button to the **on** position and click the **RW/TW safety** button **off**.

3.3. Set the laser current by entering the desired value in microwatts in the **TA set point** field. Entering a new value while the laser is running will adjust the current.

3.4. Click the **TA enable** button to the **on** position to turn the laser diode current on.

3.5. Turn off the laser diode by clicking the **TA enable** and the **Laser TEC enable** to the **off** positions.

### 4. IDS Imaging Development Systems

4.1. Run the LabVIEW program **UpconversionControl.vi**.

4.2. Under the tab **Settings**, set the shutter speed to 8  $\mu$ s by typing the value in the field labeled **Exposure Time (secs)**.

4.3. Under the tab **Settings**, set the shutter type to **Global** in the field labeled **IDS Shutter**.

- 4.4. Under the tab **DBG**, set the trigger type to **Lo\_Hi** in the **IDS trigger** field.
- 4.5. Under the tab **DBG2**, set the trigger delay in the field labeled **IDS Trigger Delay ( $\mu$ s)**. This will depend on the delay between the trigger pulse and the laser pulse from the laser.
- 4.6. Under the tab **Settings**, set the **off set x** and **off set y** to 480 pixels and the **Width** and **Height** to 96 pixels.
- 4.7. Under the tab **Settings**, set the frame rate to 0 in the **Framerate** field; this sets the camera to take one frame per trigger signal.
- 4.8. Turn on the camera by pressing the **Start Acquisition** button.
- 4.9. When a signal is entering the upconversion detector, the signal will be visible as a bright spot in the middle of the image shown on the right in the LabVIEW program. Use the **Rect** function on the left bar next to the image to draw a 6 x 6 pixel rectangle around the signal.
- 4.10. View the average intensity from the selected pixels as a function of time under the **History** tab. If necessary, the graph can be cleared by right-clicking it and selecting **clear**.
- 4.11. Press the **Stop Acquisition** button to stop the acquisition of new images from the camera.
- 4.12. Export the data by right-clicking the intensity plot, select **copy data to clipboard**, and paste the data into a **.txt** file.
- 4.13. Turn off the camera and control program by pressing the **Shut down** button.

#### **REPRESENTATIVE RESULTS:**

**Figure 3** shows the signal from different concentrations of HCN in  $N_2$ , averaged over three scans for each concentration. The mixture was prepared by mixing 300 ppm HCN in  $N_2$  with pure  $N_2$  using mass flow controllers and heating it to 843 K. The central peak is the P(20) line of the  $v_1$  vibrational band of HCN. The inset in **Figure 3** shows the peak value of the signal from this line for each concentration, with a second-degree polynomial fit. The concentration dependency of the signal can be described by  $S = ax^2 + b$ , where  $S$  is the signal and  $a$  and  $b$  are fitting constants<sup>17</sup>. Absolute concentration measurements in a flame require a calibration measurement as shown here, at a known temperature, to determine the constant  $a$ . The temperature in the measurement volume in the flame must also be measured as the constant  $a$  scales with temperature; a full discussion of this has already been published<sup>17</sup>. The poling period used for this measurement was 21.5  $\mu$ m, with a crystal temperature of 104.5 °C.

**Figure 4** presents raw data from a premixed flame. It shows five consecutive scans over the range 3229.5–3232  $cm^{-1}$ , each scan taking approximately 65 s. These cover three groups of waterlines, used for temperature measurements. Ideally, when working with a stable system, each scan over the same range should be identical, as the concentration, pressure, and temperature should be

unchanged. The intensity of the lines seen here changes drastically from scan to scan, which is because the laser pulse mode and energy is not stable from scan to scan. Results like these are unusable unless the laser pulse energy has been recorded and can be used to sort measurements with sufficient laser pulse energy from the rest. The poling period used for this measurement was 21.5  $\mu\text{m}$ , with a crystal temperature of 123  $^{\circ}\text{C}$ .

In **Figure 4**, the background scattering is not seen because an ND2 filter was used to reduce the signal, to avoid saturating the detector. For weaker signals, it was found that the background scattering is on the order of 5 pJ per pulse, which corresponds to the signal generated from the P(20) line of the  $\nu_1$  vibrational band of 100 ppm HCN at room temperature.

#### FIGURE AND TABLE LEGENDS:

**Figure 1: Diagram of the upconversion detector.** U1–U7 and UH are mirrors, highly reflective (HR)-coated for 1,064 nm. All the mirrors are planes, except for U3, which has a 200 mm radius of curvature. Mirrors U1–U5 were made to be transmissive at the wavelength of the laser diode, to ensure that the LD light does not reach the detector. U6 is transmissive for the upconverted signal, 650–1,050 nm. U7 is transmissive for the mid-IR signal. UH is 95% reflective for 1,064 nm and 5% transmissive. The path length from U1 to U3 is 156 mm, and the path length from U3 to U7 is 202 mm. L4 and L5 are achromatic lenses with 60 mm and 75 mm focal lengths, respectively. Both are transparent for 650–1,050 nm. The camera used as detector is placed 75 mm from L5. The cavity field is vertically polarized. The PPLN used here has poling periods of 21.0  $\mu\text{m}$ , 21.5  $\mu\text{m}$ , 22.0  $\mu\text{m}$ , 22.5 mm, and 23.0  $\mu\text{m}$ , and the crystal length is 20 mm. The visible and near-infrared detector used is a UI-5240CP-NIR-GL camera from IDS Imaging Development Systems.

**Figure 2: Diagram of the DFWM setup.** M1 is a dielectric mirror highly reflective (HR) at the wavelength of the pulsed laser. M2 is a dielectric mirror coated to be HR at the wavelength of the pulsed laser and transmissive for the HeNe guide beam. M3–M6 are protected gold mirrors. B.C.1 and B.C.2 are BOXCARS plates 1 and 2. L1 is a 500 mm focal length  $\text{CaF}_2$  lens with a 5.1 cm diameter. L2 is a 500 mm focal length  $\text{CaF}_2$  lens with a diameter of 2.54 cm. L3 is a 100 mm focal length  $\text{CaF}_2$  lens. The pulsed laser is vertically polarized.

**Figure 3: Signal from different concentrations of HCN in  $\text{N}_2$ .** The central peak is the P(20) line of the  $\nu_1$  vibrational band of HCN. The inset shows the peak signal from each concentration (diamond markers), with a second-order polynomial fit.

**Figure 4: Five consecutive scans of ca. 65 s in duration per scan, done in a premixed flame.** The laser was scanned over the range of 3229.5–3232  $\text{cm}^{-1}$ . The peaks seen here are the signal from several collections of  $\text{H}_2\text{O}$  transition lines. The signal was reduced with an ND1 and an ND0.6 filter, to avoid saturating the detector.

**Figure 5: A BOXCARS plate seen from the side.** It is a block of transparent material. On the input side, it is coated with an antireflective coating on half of the surface. The laser beam enters here and reaches the output side, where half the surface is coated for a 50% transmission. The light

reflected internally in the plate is then refracted to the part of the input side coated for high reflection and is reflected through the top half of the output side. This splits one beam into two parallel beams. The same effect could be achieved with a beam splitter and a mirror, but a beam splitter would have some reflection from the rear surface, which could increase the background noise. Also, the BOXCARS plate requires no alignment to ensure the two beams produced are parallel.

**Figure 6: Intracavity power as a function of pump laser diode current for the upconversion module.** Each point is an average of the power measured from three separate alignments of the cavity and the error bars indicate the spread between the separate alignments. The deviation from the ideal laser behavior is caused by thermal effects in the laser crystal and the PPLN crystal.

#### DISCUSSION:

The precision of the alignment of the pulsed laser beam is critical to the sensitivity of the method. Special care must be taken to ensure that the beams are separated by equal distance after the BOXCARS plates and that the beams are equally spaced around the center of L1. Deviation from this will lead to a significant drop in signal intensity and, therefore, sensitivity. Likewise, care must be taken that the upconversion module cavity is running in the fundamental mode and that the signal beam is aligned for an optimal overlap with the upconversion pump. The signal can easily be reduced by one or two orders of magnitude if the upconversion cavity is running in a wrong mode or the signal beam overlap with the cavity field is suboptimal. This includes placing L3 with millimeter precision so that the signal beam focal point is in the middle of the PPLN crystal. With an optimal overlap and 80 W of cavity power, a 6% quantum efficiency of the SFG stage is possible. With the detector and wavelength used here, the total detection efficiency is 3%. The maximum intracavity power that can be reached is 120 W, but 80 W can be achieved reliably. The conversion efficiency is proportional to the intracavity power, so signals recorded with a different intracavity power can be compared if the intracavity power is recorded.

The main limiting factor for the sensitivity of this method is the background scattering, which drowns weak signals. To limit this scattering, it is critical that the optics are kept dust-free, especially lens L1. Care must also be taken that the position of the beam block minimizes the background noise. The beam block should be placed on an xy-stage so that it can be moved in a controlled manner in both the horizontal and vertical plane, perpendicular to the direction of the beams.

The scanning discussed here is done with the PPLN at a constant temperature. The conversion efficiency is proportional to  $\text{sinc}(\Delta k L / 2\pi)^2$ , where  $\Delta k$  is the phase mismatch and  $L$  is the crystal length. The full width half maximum (FWHM) of this function is the bandwidth of the detector at a constant PPLN crystal temperature. The FWHM of this function changes with the crystal temperature and wavelength but is generally on the order of  $5 \text{ cm}^{-1}$  in the mid-IR, for a 20 mm long crystal. The exception is near 4,200 nm, where the width increases greatly<sup>18</sup>.

No scaling optics have been included in the setup diagram in **Figure 2**, because there are a number of issues to consider before deciding what, if any, scaling is needed. For the setup

described here, the pulsed laser beam is collimated at a beam diameter of ca. 2 mm when reaching L1. This gives a beam waist at the focal point of approximately 400  $\mu\text{m}$ , using a wavelength of 3  $\mu\text{m}$ . When implementing this technique, it might be desirable to change the focal length of L1, either because more space is needed between L1 and the focal point for practical reasons, or to shorten the measurement volume by increasing the convergence angles, which can be done by using a shorter focal length. In this case, the beam waist at the focal point should be kept at ca. 400  $\mu\text{m}$  and the collimated beam should be scaled to match. It should, however, be taken into account that increasing the beam diameter without increasing the spacing of the beams will increase the scattering from the beam block edges. The spatial resolution is given by the overlap of the pump beams. For the setup described here, the overlap is 6 mm long, so the measurement volume is a cylinder of 6 mm long, with a radius of 0.4 mm.

To achieve quasi-phase-matching in the PPLN crystal, both the mid-IR signal and the intracavity field of the upconversion cavity must be extraordinarily polarized in the PPLN crystal. The upconversion cavity should be built so that the polarization of the intracavity field is automatically right. If the mid-IR laser does not already match this, a waveplate can be inserted at the mid-IR laser output to turn the polarization.

IR-DFWM requires relatively high energy pulses, 1–4 mJ, combined with a narrow enough laser linewidth to resolve molecular lines, which are on the order of 0.1  $\text{cm}^{-1}$ . Lasers that match these criteria generally have low repetition rates, and as data acquisition with DFWM is generally done by scanning the wavelength of the laser, this limits measurements speed. This means the method is most readily applied to measurements where the subject does not change over time, although it has also been applied to temporally resolved measurements<sup>17</sup>. Another limitation is that, because of the sensitivity to scattered light, particles in or near the measurement volume will create scattering events that completely drown the signal<sup>17</sup>. The phase-match condition of the upconversion process is spectrally narrow, which helps eliminate noise from the thermal background radiation, but it does make scans over broad wavelength ranges more time-consuming as the PPLN temperature must be tuned to keep the signal wavelength phase-matched.

Future uses of IR-DFWM are planned for the detection of  $\text{NH}_3$  in flames, or to continue the work with HCN in more practical environments. The most obvious means for improving the method is to further reduce the background from scattered light. This might be done using spatial filtering of the signal beam after the signal is collected by L2.

#### **ACKNOWLEDGMENTS:**

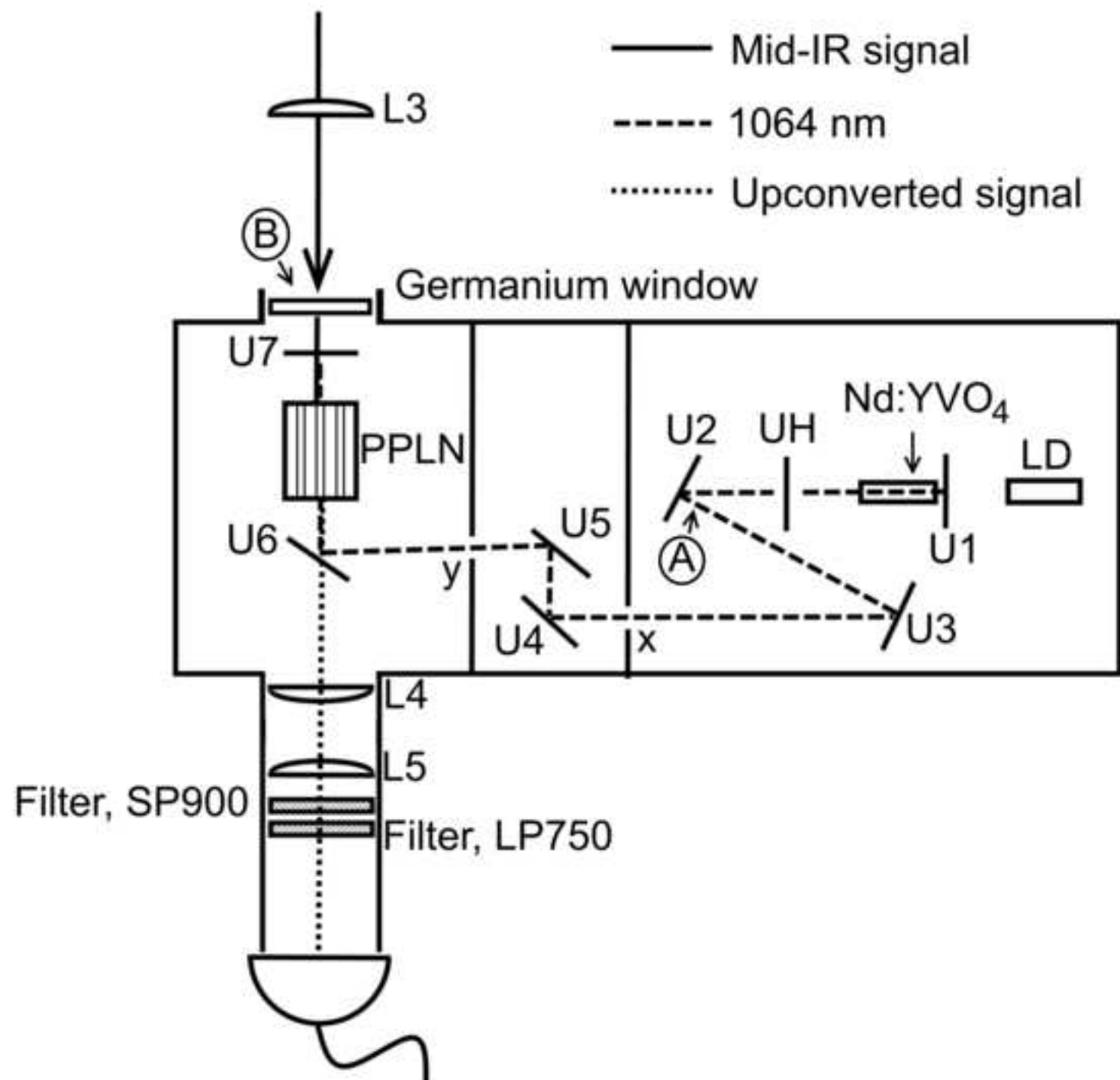
The funding received by the authors within the scope of Horizon 2020 by the European Union is highly appreciated. This work was conducted as part of the Mid-TECH Marie Curie innovative training network [H2020-MSCA-ITN-2014-642661].

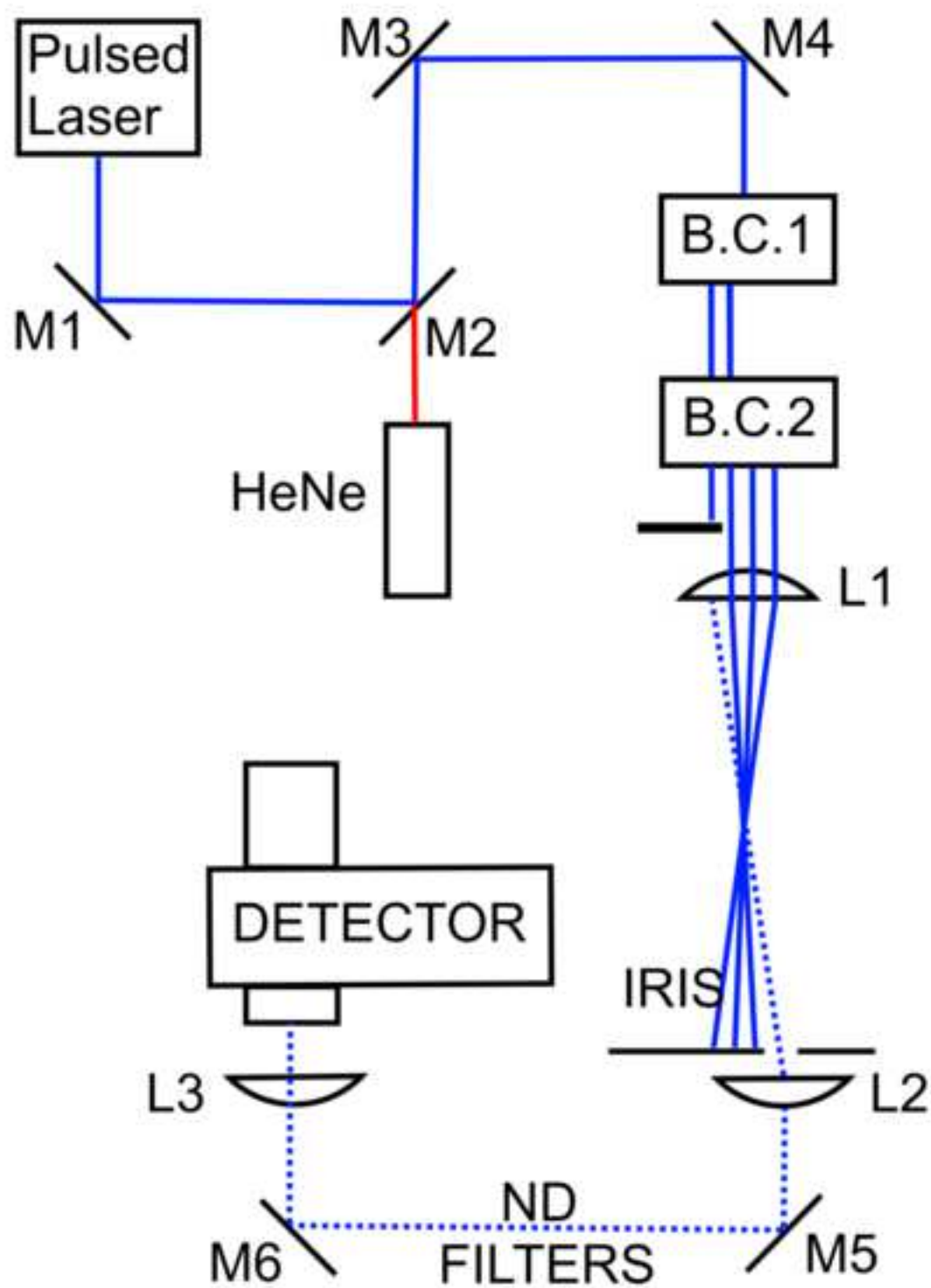
#### **DISCLOSURES:**

The authors have nothing to disclose.

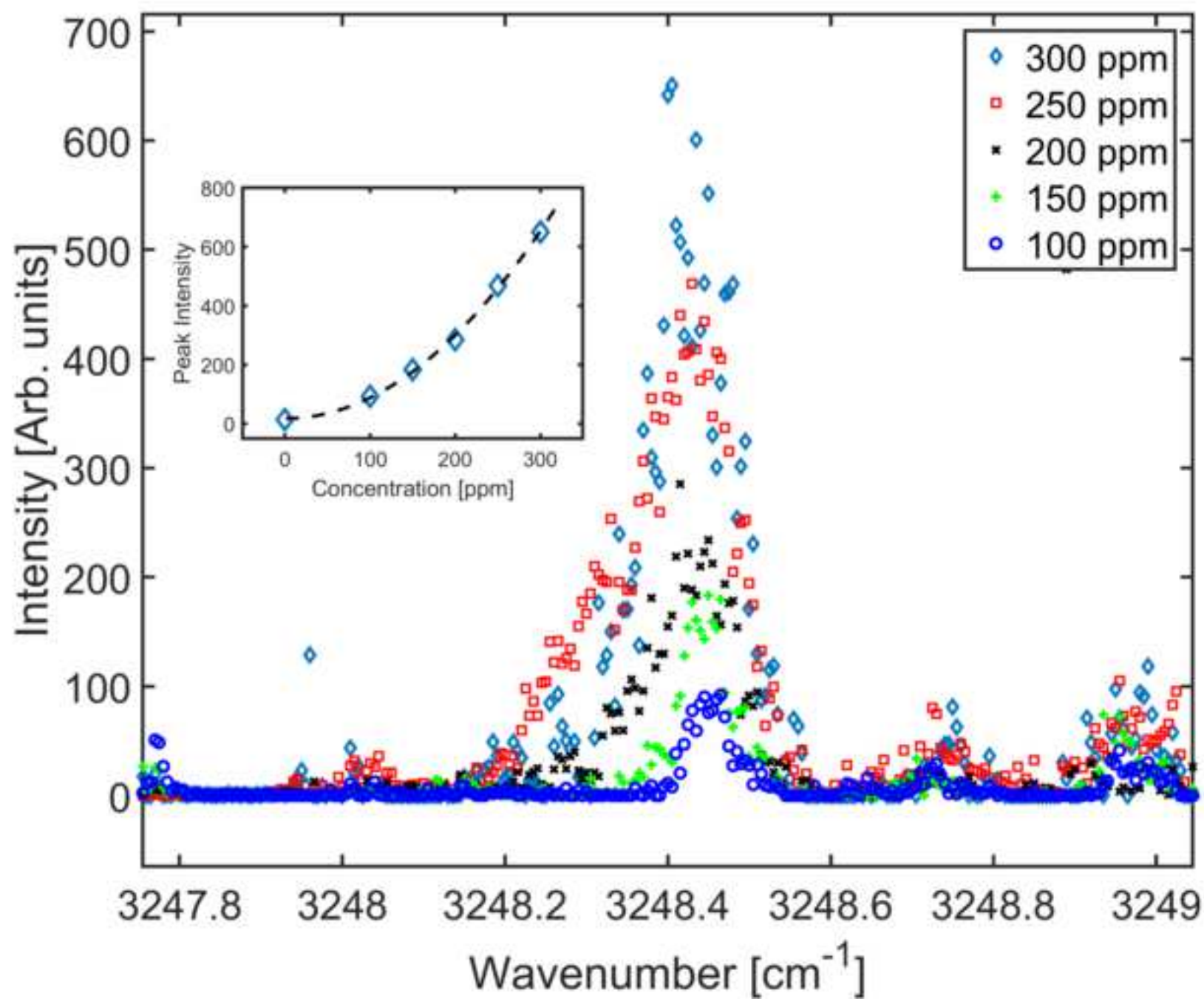
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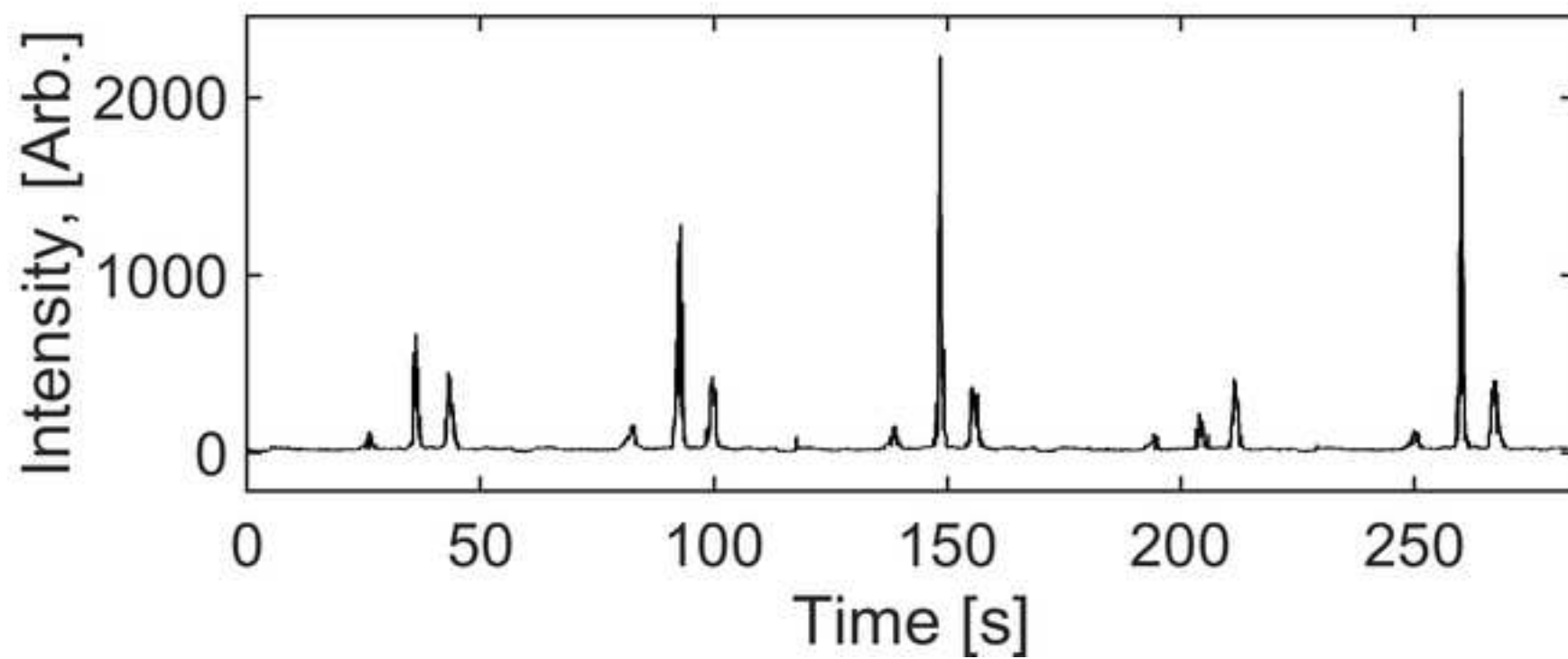
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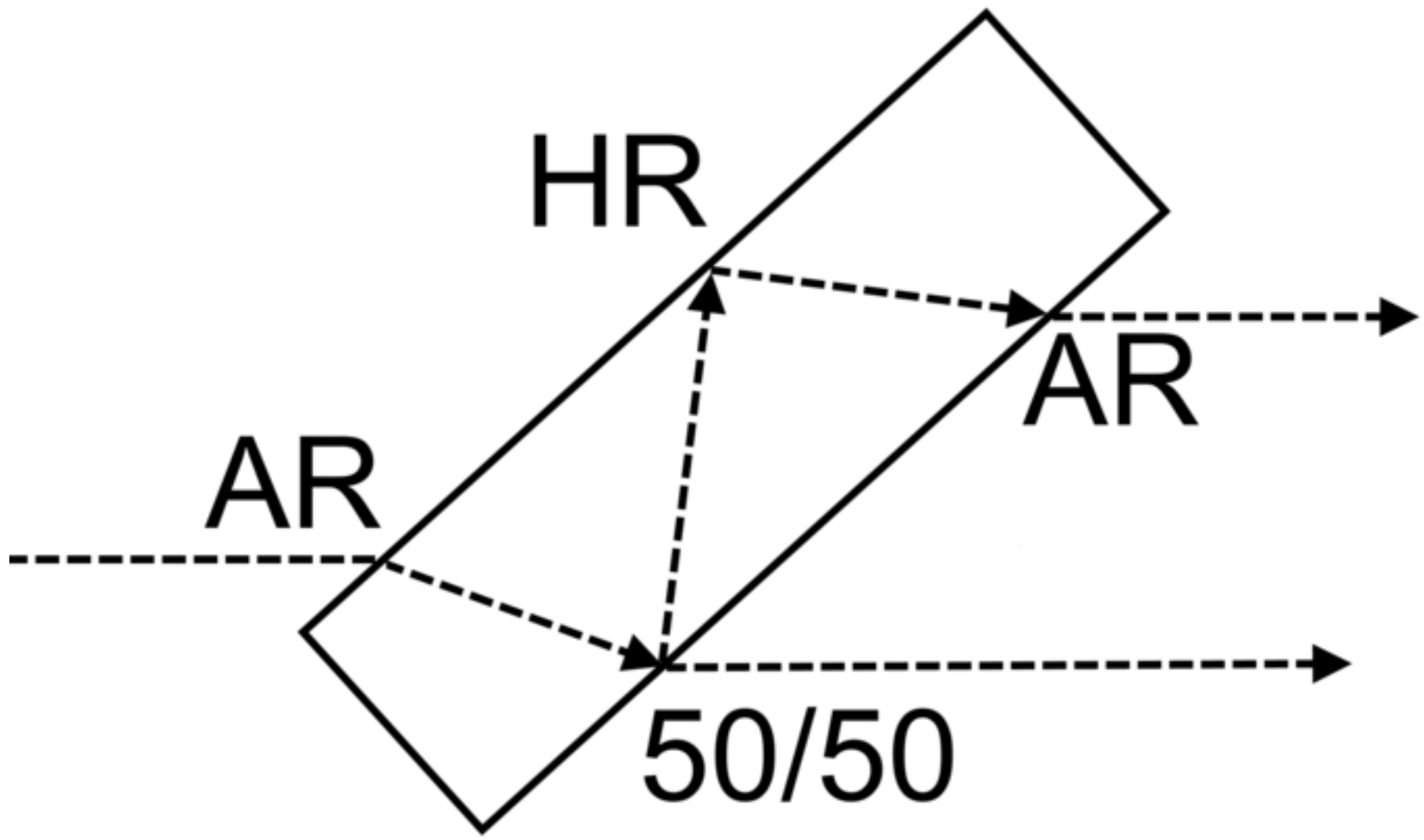


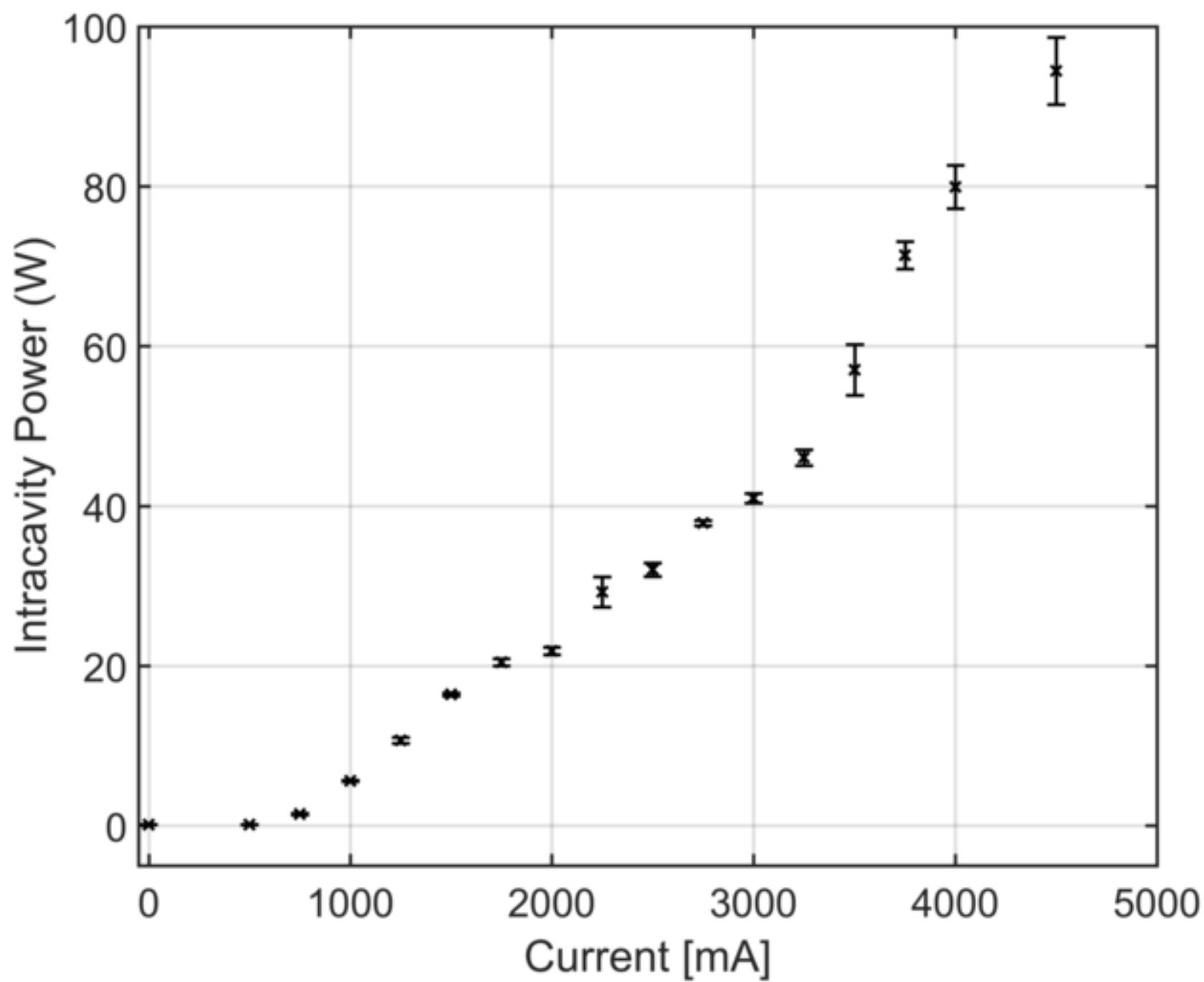












Name of Material/ Equipment	Company	Catalog Number	Comments/Description
Nd:YAG laser, pulsed	Spectra Physics	Quantarau Pro-290-10	Quantity: 1 (For pumping the mid-IR laser)
Nd:YAG laser, injection seeding system	Spectra Physics	6350	Quantity: 1
NIR Dye laser - OPA system	Sirah	OPANIR	Quantity: 1
HeNe laser	Thorlabs	HNL100LB	Quantity: 1
Dichroic mirror	LASEROPTIK	NA	Quantity: 1, custom order: HR for the mid-IR, transparent for 632 nm
Protected Gold Mirrors	Thorlabs	PF10-03-M01	Quantity: 5
BoxCars Plate	LASEROPTIK	NA	Quantity: 2, Custom order
xy-stage	Thorlabs	DTS25/M	Quantity: 2
500 mm focal length CaF2 lens, Ø2"	Eksmaoptics	110-5523E	Quantity: 1
500 mm focal length CaF2 lens, Ø1"	Thorlabs	LA5464	Quantity: 1
100 mm focal length CaF2 lens, Ø1"	Thorlabs	LA5817	Quantity: 1
Iris, Ø50 mm	Thorlabs	ID50/M	Quantity: 1
ND1 IR filter (ZnSe)	Thorlabs	NDIR10B	Quantity: 1
ND2 IR filter (ZnSe)	Thorlabs	NDIR20B	Quantity: 1
ND3 IR filter (ZnSe)	Thorlabs	NDIR30B	Quantity: 2
Upconversion Detector	NLIR	NA (closest equivalent: U3055 3.0-5.5µm)	Quantity: 1, Custom order
VIS/NIR Detector Card	Thorlabs	VRC2	Quantity: 1, (low intensity)
NIR Detector Card	Thorlabs	VRC4	Quantity: 1, (high intensity)
MIR Detector Card	Thorlabs	VRC6S	Quantity: 1
Thermal Power Sensor Head	Thorlabs	S302C	Quantity: 1
Power meter console	Thorlabs	PM100D	Quantity: 1



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INFRARED DEGENERATE FOUR-WAVE MIXING WITH  
UPCONVERSION DETECTION FOR QUANTITATIVE GAS SENSING

Author(s):

RASMUS L. PEDERSEN, ZHONGSHAN LI

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### Revisions of manuscript JoVE59040R1

We thank the editor for the comments. We have taken into account all comments and deal with them point by point below. The revisions were made with the 'track changes' function on, to make them visible in the revised document.

- 1) *"Please take this opportunity to thoroughly proofread the manuscript to ensure that there are no spelling or grammar issues."*

**Our reply:** This has been done.

- 2) *"The highlighted protocol steps are over 2.75 page limit (including headings and spacing). Please highlight fewer steps for filming."*

**Our reply:** We apologise for the mistake, the length have now been reduced.

- 3) *"Please combine some of the shorter Protocol steps so that individual steps contain 2-3 actions and maximum of 4 sentences per step."*

**Our reply:** This has been done.

- 4) *"Step 2.1: Please write this step in imperative tense"*

**Our reply:** Step 2.1 has been changed to the imperative tense.

- 5) *"Are you adjusting the mirrors manually or using a software? Is there a software used in the protocol? If yes, please provide a step-wise description of software usage. Please mention what button is clicked on in the software, or which menu items need to be selected to perform the step. If you are adjusting the mirrors manually, please specify that in the protocol."*

**Our reply:** The mirrors are adjusted manually, this is now mentioned in the protocol.

Software is used to control the laser diode and the camera. Two sections have been added to the protocol describing the use of the software.