

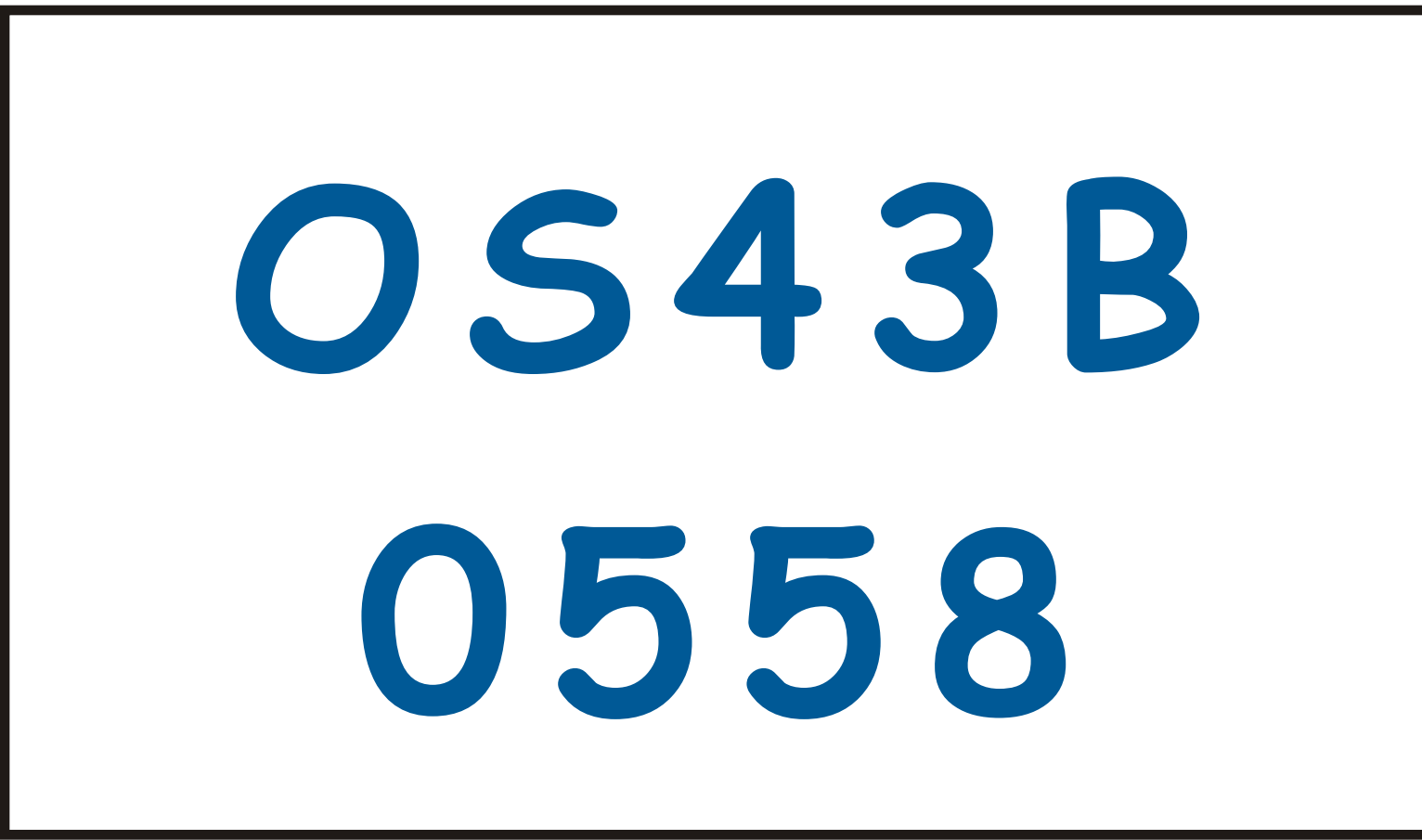


In Situ Raman Spectra from the SeaCliff Hydrothermal Field (Gorda Ridge)

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Abstract

MBARI's *in situ* laser Raman spectrometer (DORISS – Deep Ocean Raman *In Situ* Spectrometer) was deployed at the Sea Cliff Hydrothermal Field on the Gorda Ridge in July 2004. The first *in situ* Raman spectra of hydrothermal minerals and high-temperature fluid venting from the seafloor were obtained. These spectra are analyzed and compared to laboratory measurements of samples collected from the site.

Laser Raman spectroscopy is a proven, powerful geochemical technique for analyzing the chemical composition and molecular structure of solids, liquids, and gases. During an expedition to Gorda Ridge on the *R/V Western Flyer* in July 2004, DORISS was deployed successfully by the ROV *Tiburon* at hydrothermal vents on the seafloor (~2700 m depth). Data were collected from hydrothermal fluids, chimney minerals (e.g., anhydrite and barite), and bacterial mats using two types of sampling optics – an immersion optic, and a non-contact optic. To collect spectra from opaque mineral samples, a precision underwater positioner (PUP) was used to position the DORISS probe head. PUP is a stand-alone, three degree-of-freedom positioner capable of moving the DORISS probe head with a precision of 0.1 mm (required by the small focal volume of the sampling optic).

Raman spectra of ~300° C vent fluids were collected with both sampling optics. The Raman spectrum of seawater contains bands from the bending (~1640 cm⁻¹) and stretching (3000-3800 cm⁻¹) vibrational modes of the water molecule and a small peak from the S-O stretch of the sulfate ion (~981 cm⁻¹). Compared to ~2° C ambient seawater, vent fluid spectra show changes in the intensity ratios of the water bands due to the elevated temperature, and the sulfate peak is reduced. Additional components of hydrothermal fluid are present in such low concentrations that it is difficult to detect them with the current Raman system. The chimneys in the Sea Cliff field are primarily anhydrite, and debris in the area also contains barite. We were able to obtain quality spectra of both anhydrite (primary peak at ~1017 cm⁻¹) and barite (primary peak at ~987 cm⁻¹). In addition, we were able to detect elemental sulfur in the polycrystalline S₈ morphology which may have been produced by bacterial mats (peaks at ~218 and 473 cm⁻¹).

Geologic Setting

The Sea Cliff Hydrothermal Field is located on the northern end of the Gorda Ridge, an intermediate rate (~55 mm/yr) spreading segment in the northeastern Pacific Ocean (below). Hydrothermal activity is located approximately 2.6 km east of the spreading axis at a depth of 2675-2800 m (~300 m above the axial valley floor). The site was first identified by water column anomalies in 1986, and first visited by the U.S. Navy submersible DSV *Sea Cliff* (for which the field was named) in 1988.

Active venting is concentrated along ridges which trend perpendicular to the slope of the eastern axial valley wall (022°). The chimneys formed along these ridges are primarily composed of anhydrite and smectite with minor sulfide phases. The seafloor in the area is covered with a hydrothermal crust which consists of altered basalt, barite, amorphous silica, and sulfides. Large areas of the vent field between the chimney ridges are covered with blue mat-like material believed to be a type of ciliate. The vent fluids were essentially clear with temperatures up to ~300°C. The fluids have been found to be low in salinity and sulfide content. (Data from recent observations is presented on poster T21C-0538).

Gorda References

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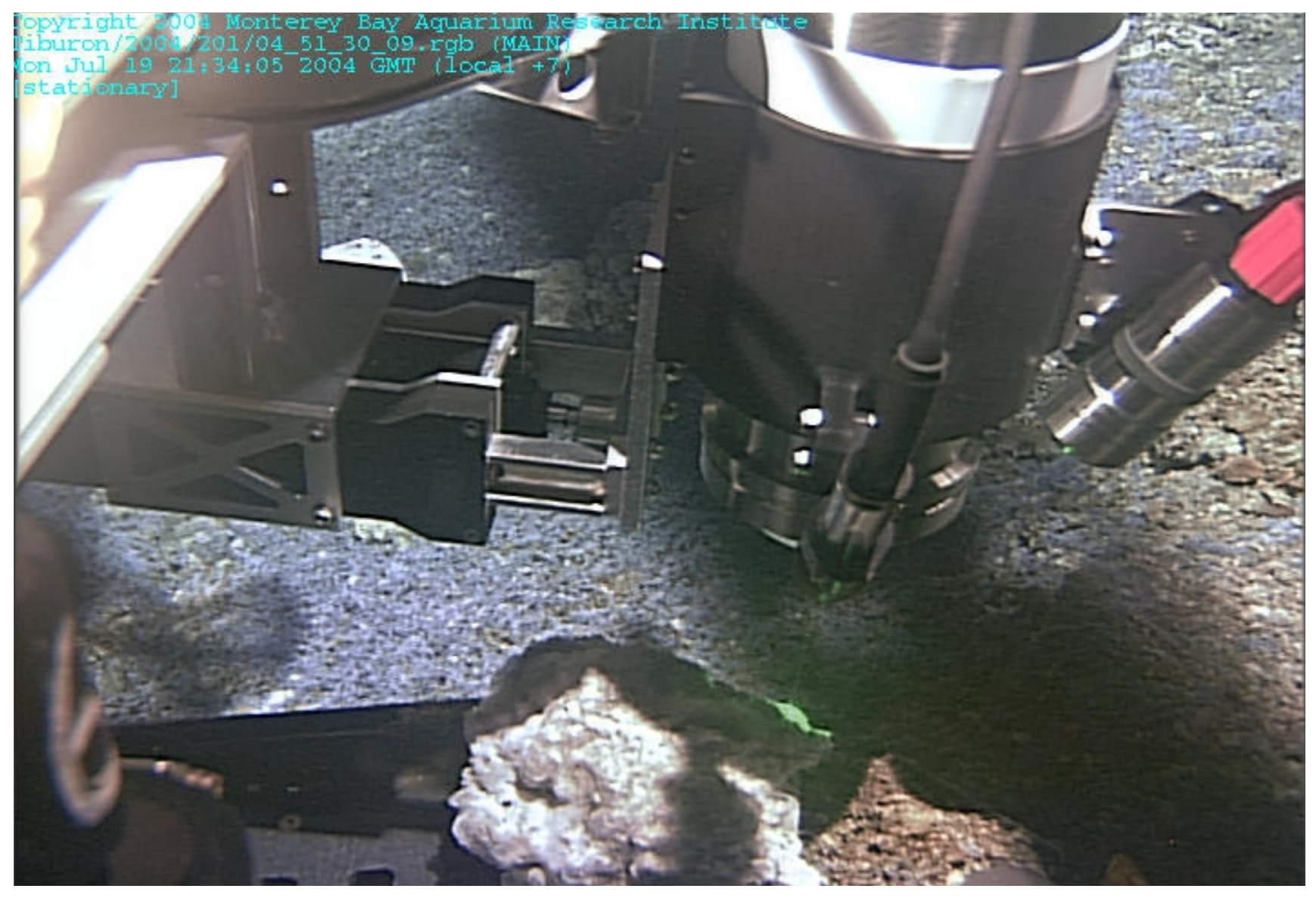


Background Picture: The DORISS probe head being positioned at a hydrothermal vent by the ROV *Tiburon* manipulator at the Sea Cliff Hydrothermal Field, Gorda Ridge.

Operations



The Precision Underwater Positioner (PUP - above) was used to position the DORISS probe head.

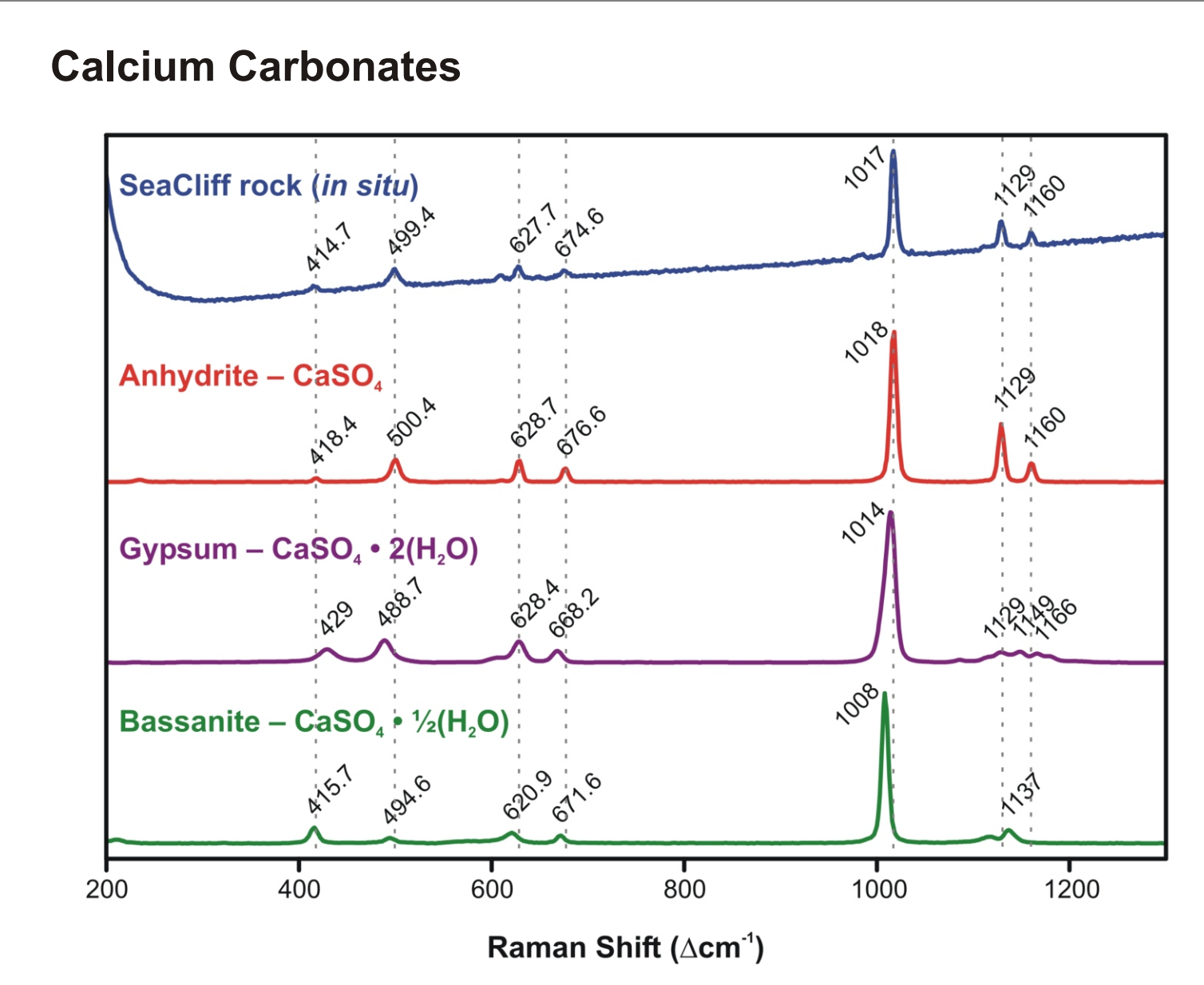
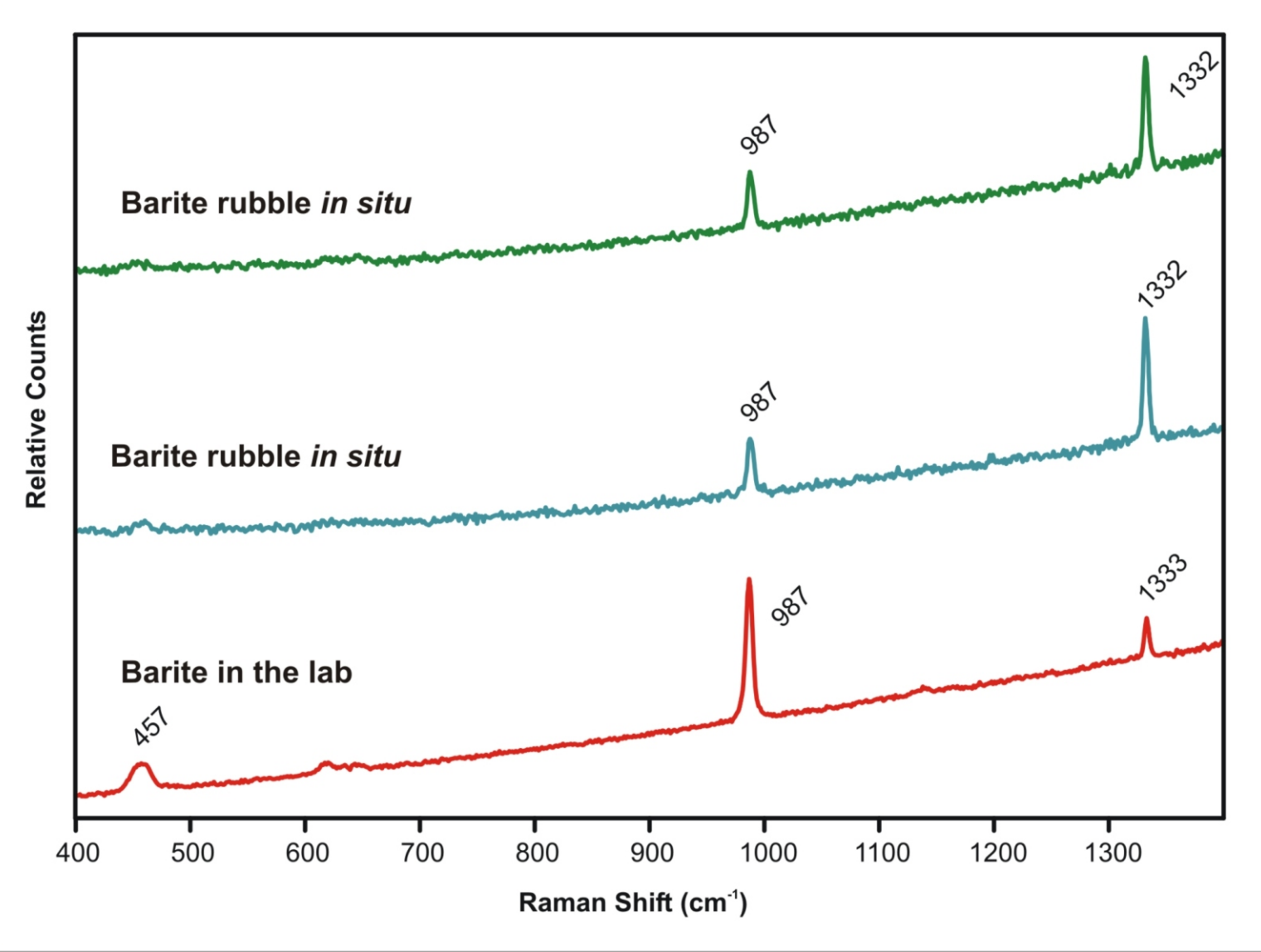


PUP (left) consists of a tripod frame and three actuators providing 3 degrees of movement – Z axis (15 cm up-and-down), R-axis (15 cm in-and-out), and theta axis (58° rotation about the Z axis). A camera, light, and red lasers that cross at the DORISS focal point assist the shipboard scientist in positioning the probe head.

In the picture above, the DORISS probe head is being positioned over a chunk of chimney material (the green laser spot is seen on the upper right of the rock). On the seafloor in the background is the blue mat-like material discussed in the Geological Setting section.

Hydrothermal Minerals

PUP was used to position the DORISS probe head at rock samples on the seafloor. In some cases, chimney material was broken off onto the ROV porch and analyzed there. Two minerals were primarily observed with the DORISS instrument: anhydrite and barite. When analyzing hydrothermal crusts and rubble in the vent field we detected barite (BaSO₄) (right). This mineral was mostly found as thin rinds or veins around darker material which exhibited significant fluorescence. Further analysis of the crust material will begin soon.



The chimneys are primarily anhydrite (CaSO₄), which is soluble in cold seawater. Thus anhydrite was not observed away from the high-temperature venting. There are two other common calcium sulfate minerals: bassanite (CaSO₄ · ½(H₂O)) and gypsum (CaSO₄ · 2(H₂O)). These three calcium sulfate species can be distinguished by their Raman spectra. The figure on the left compares the spectrum of a Sea Cliff sample to the calcium sulfate mineral reference spectra. The peak positions (indicated by dashed lines) and peak intensities of the Sea Cliff sample correspond well to anhydrite, and are clearly different from gypsum and bassanite by amounts greater than instrument error.

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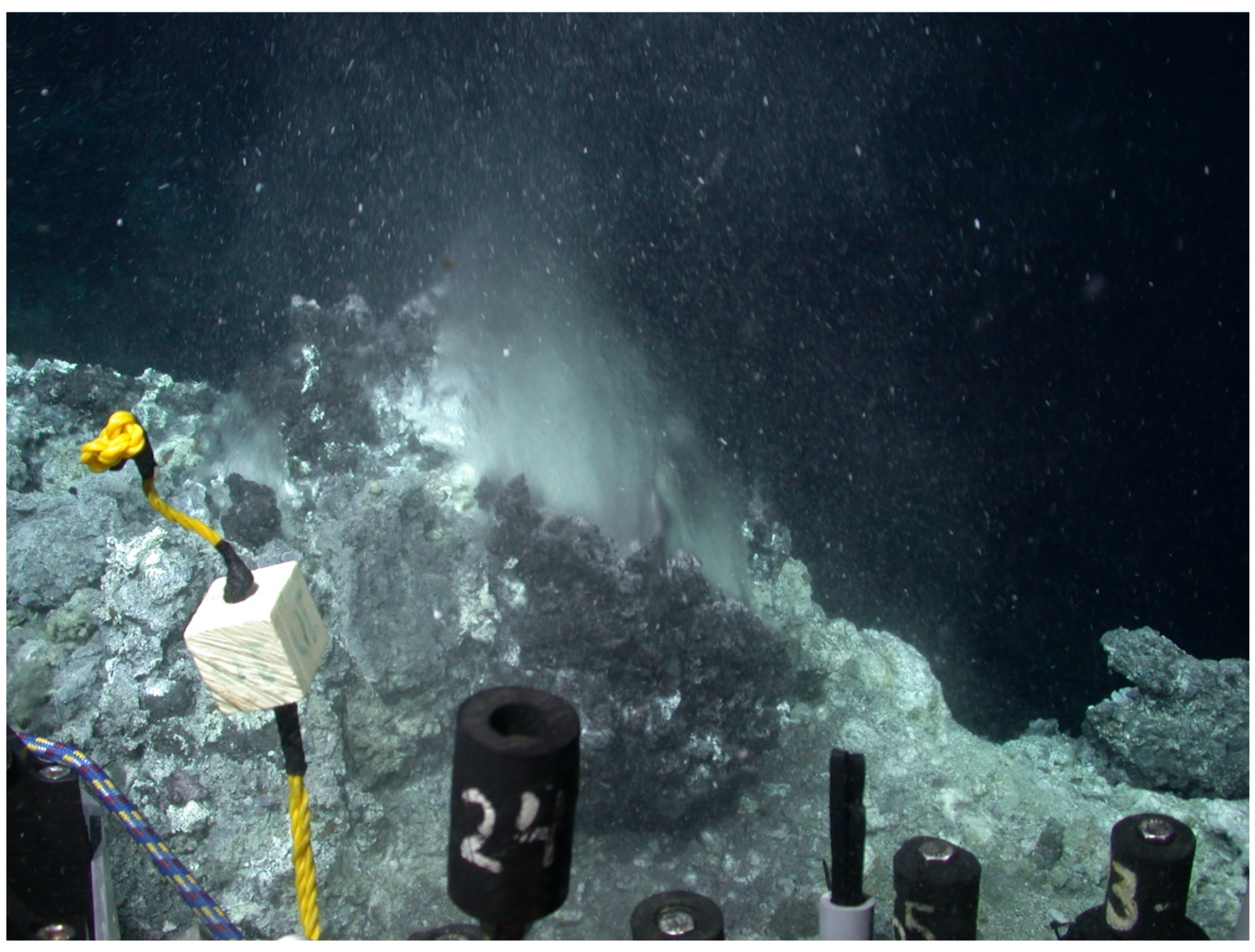
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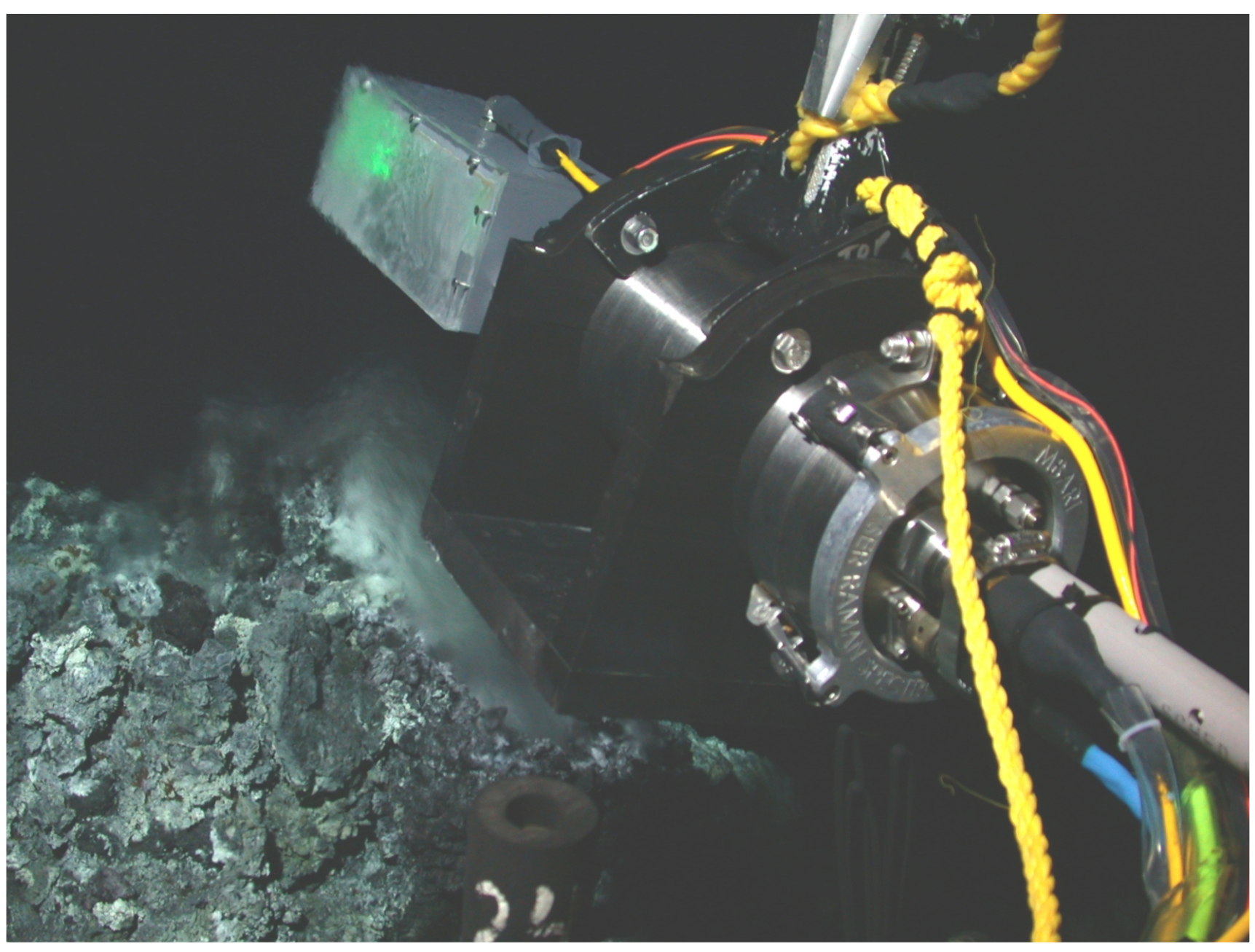
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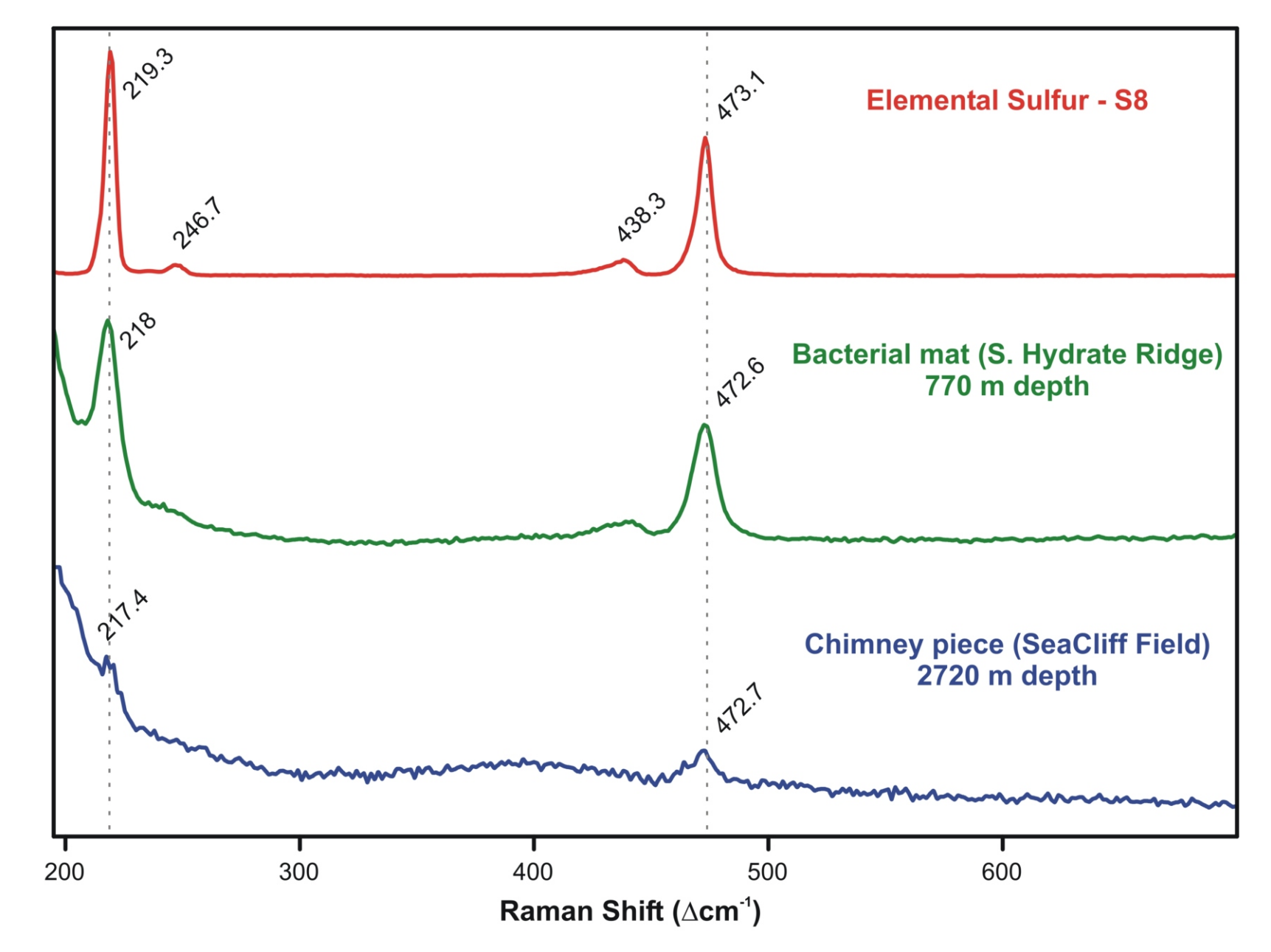


Vent fluid (above) was analyzed on two dives. On the first dive, a non-contact optic was used behind a dome window which was not in contact with the hydrothermal fluid; thus some ambient seawater was in the optical path. The focal point was positioned inside the volume of exiting fluid using the ROV manipulator. On the second dive, the immersion optic was used protruding into an open-bottomed box. The box was held over the vent to collect the buoyant hydrothermal fluid. In this case the optic window was immersed in the vent fluid.



Biological Material

Bacterial mat spectra were obtained at both the Sea Cliff Hydrothermal Field, and Hydrate Ridge. Previous laboratory studies of bacterial mat using Raman spectroscopy showed that the bacteria produced elemental sulfur in an S₈ morphology. *In situ* spectra collected from Gorda Ridge and Hydrate Ridge both correspond well to S₈ elemental sulfur (below). The spectrum from Hydrate Ridge (green) was collected using PUP to focus DORISS on seafloor bacterial mat. The spectrum from Gorda Ridge (blue) was serendipitously collected while trying to collect spectra from a chimney.



Hydrothermal Vent Fluids

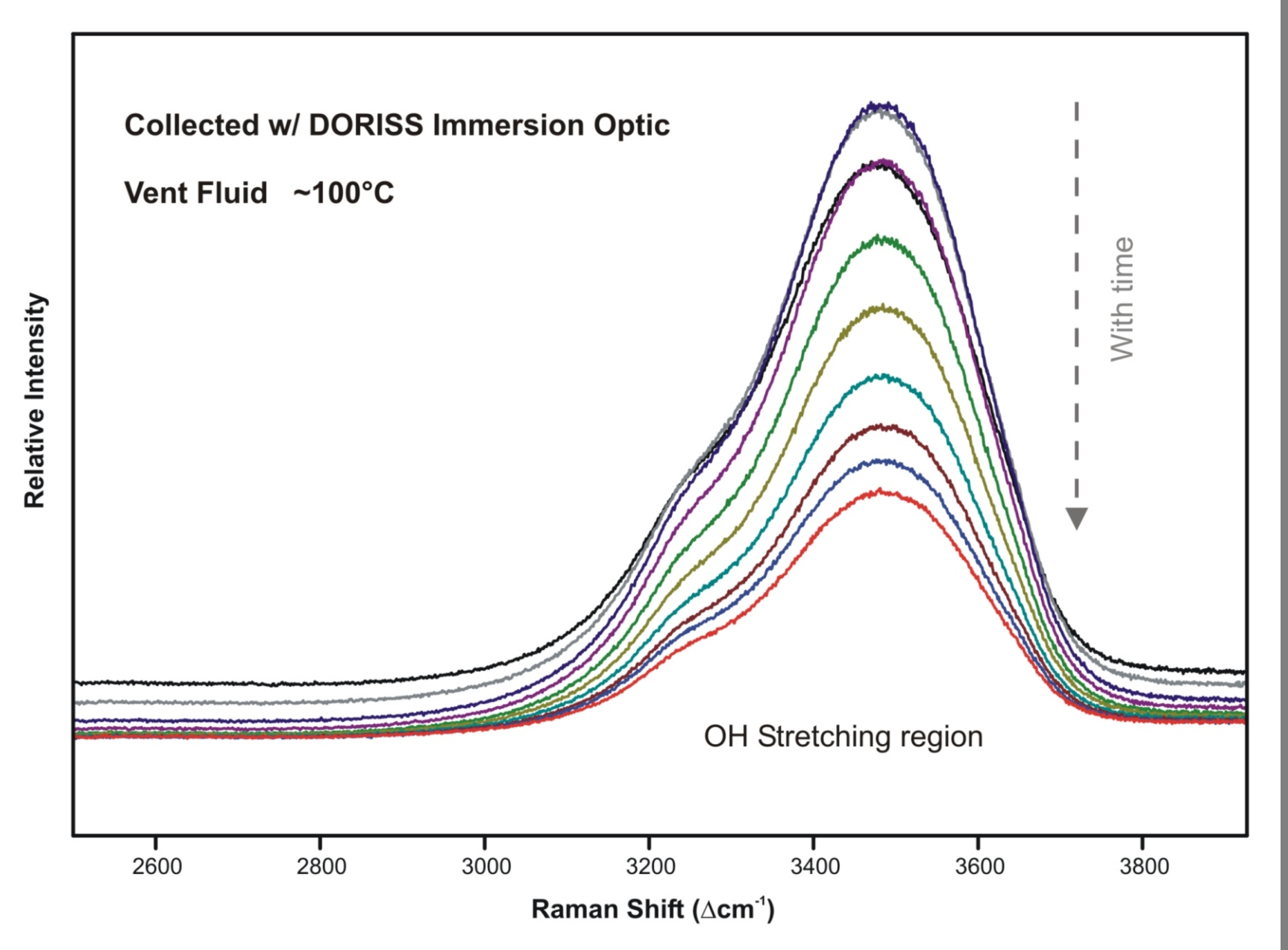
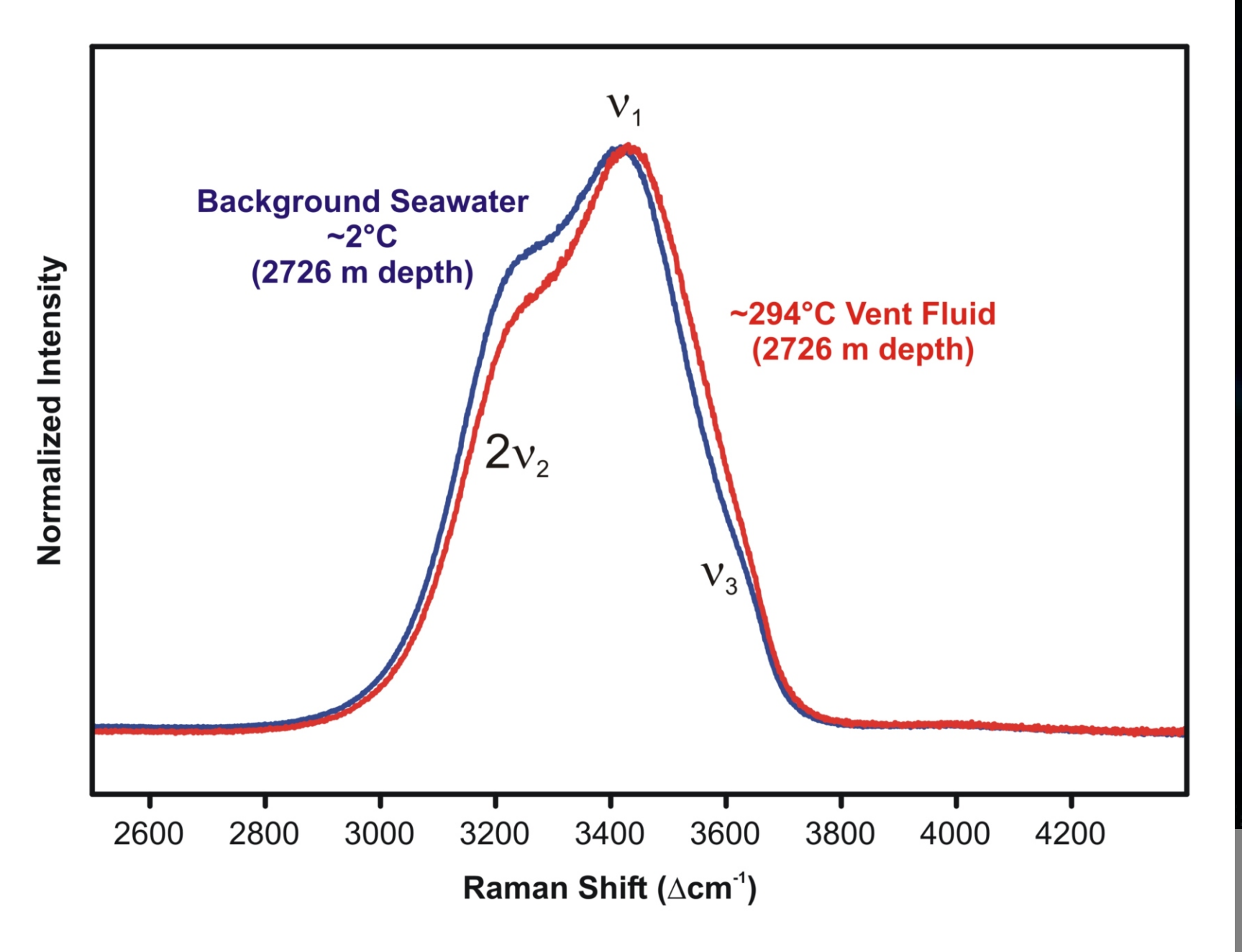
The primary purpose of our expedition to Gorda Ridge was to use Raman spectroscopy to detect the various constituents of hydrothermal vent fluid *in situ*. Gases such as CO₂ and H₂S are detectable by Raman spectroscopy even in a dissolved state.

The Raman spectrum of seawater is well-known and is composed of bands due to the bending (~1640 cm⁻¹ band) and stretching (~3000-3800 cm⁻¹ region) of the water molecule, and a peak at ~981 cm⁻¹ due to the sulfate ion. Data in the literature have shown that the stretching bands of the water molecule are affected by both temperature and pressure.

We obtained a number of spectra of vent fluid using two configurations (discussed at left). In all cases we observed changes in the region of the O-H stretching modes of water due to the elevated temperature of the hydrothermal fluid. The top right plot compares the spectrum of ~2°C ambient seawater to that of ~294°C vent fluid. (Note that this spectrum was collected using the non-contact optic, and thus the spectrum is a combination of the vent fluid being analyzed and ambient seawater in the optical path.) The ratio of the areas of the 2₂ band to the 2₁ band decrease as water temperature increases.

When using the immersion optic configuration (lower left picture) we found that the intensity of the Raman spectra decreased over time (lower right plot). This was due to the fact that the sapphire pressure window of the optic was immersed in the vent fluid and a light sulfide coating built up on the window. Although the fluid was ~294°C as it exited the chimney orifice, it was reduced to ~50-100°C in the box (as measured by a thermocouple in the top of the box) due to entrainment of ambient seawater. The change in the shape of the water stretching bands is more pronounced in the plot at right since no ambient seawater was in the optical path. For ambient seawater at 2627 m depth, the ratio of the areas of the 2₂ band to the 2₁ band is ~0.75. For hydrothermal fluid at ~100°C the ratio was ~0.30. The other major observation was the decrease in or lack of a signal from the sulfate ion. This is because vent fluid is stripped of sulfate in the subseafloor.

We did not see peaks characteristic of CO₂, CH₄, or H₂S. Based on previous dives, their concentrations were expected to be: CO₂ ~16 mmol/kg, CH₄ ~0.05 mmol/kg, H₂S ~0.06 mmol/kg. Vent fluid samples collected during the dive were analyzed (K. Von Damm, UNH) and found to be the same as those sampled in 2000 and 2002.



The plots above show seawater and high temperature vent fluid spectra in the O-H stretching region. Water bands present are the 2₂, 2₁, and 2₃ vibrational modes.

We attempted to collect Raman spectra from other biological targets as well. The blue mat-like material seen in the picture in the Operations section had no discernable Raman peaks but did exhibit strong fluorescence.

We also attempted to collect a Raman spectrum from a **tube worm** (*Ridgeia piscesae*) tube (below). We believe the tube to be chitinous. However no chitin Raman peaks were observed. We did observe two strong fluorescent bands. These bands were wide (~400-500 cm⁻¹), and the peak maximums were at wavelengths of 617 and 682 nm (excitation wavelength was 532 nm).

