Tip-Enhanced Raman Spectroscopy and Microscopy on Single Dye Molecules with 15 nm Resolution

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Our recently developed approach of UHV-tip-enhanced Raman spectroscopy permits us to acquire Raman spectra of a few single brilliant cresyl blue (BCB) molecules and even a single one adsorbed on an Au(111) surface. This is substantiated by simultaneously recorded STM images. Furthermore, because of the reduced photobleaching in UHV, the time frame for spectral acquisition is sufficiently extended to allow tip-enhanced Raman imaging of a single BCB molecule with a lateral resolution of 15 nm.

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With the invention of scanning tunneling [1] and atomic force microscopy [2] the characterization of surfaces with atomic resolution has become feasible. In the form of inelastic electron tunneling spectroscopy (IETS) the former technique permits chemical identification of molecules and even single molecules [3–5]; this can, however, only be achieved using cryogenic temperatures; moreover, at present IET spectra have a rather poor fingerprint quality. Optical spectroscopies, on the other hand, can provide detailed and rich chemical information about an adsorbate in a wide temperature range. However, for studies in the nanometer region one has to overcome limitations of the low sensitivity of standard Raman spectroscopy and the diffraction limited resolution of light. Both of these restrictions can be surmounted by generating and using enhanced, narrow-sized electromagnetic near fields in the vicinity of a metallic tip, which enable both ultrahigh sensitivity and spatial resolution to be achieved. In the first papers on tip-enhanced Raman spectroscopy (TERS) [6–8], enhancement factors of about \(10^4\) were reported. More recently, enhancement factors of \(10^6–10^9\) have been observed [9–11]. With such high sensitivities the spectroscopical detection of individual molecules seems feasible. Indeed, there are recent reports on single-molecule TERS [10–13], based, however, only on indirect evidence such as spectral fluctuations.

For 1D objects such as carbon nanotubes, Anderson et al. demonstrate enhanced Raman spectroscopy and microscopy with a lateral resolution of \(\sim 15\) nm [14]. Theoretical calculations indicate a lateral resolution of \(\sim 8\) nm for a gold tip of 20 nm radius above a gold substrate [15]; other authors even consider molecular resolution possible [16].

In this Letter we present a tip-enhanced Raman spectroscopic and microscopic study on brilliant cresyl blue (BCB) adsorbed on gold single-crystal surfaces performed in ultrahigh vacuum (UHV). Because of substantially reduced photobleaching under UHV conditions, we are able not only to record TER spectra of single dye molecules, but also Raman images of a single dye molecule with a resolution of a few nm.

The UHV-TERS and TERS imaging system employed here has been described recently [17]. It mainly consists of a DME Polyscope 500 UHV-STM and an optical setup employing a parabolic mirror with a numerical aperture of \(\lambda/4\) made by Rodenstock Präzisionsoptik, Germany, for focusing the laser and collecting the Raman light. The laser source is a HeNe-laser with a wavelength of 633 nm and a linear polarization mode which is converted into a pseudoradial polarization mode by an arrangement of four \(\lambda/2\) retarders [18]. Radial polarization combined with a high numerical aperture parabolic mirror leads to a sharp focus (\(\sim \lambda/2\)) [19,20] as well as to an increased longitudinal electric field component along the tip axis [21,22]; such focal properties are essential for a high field enhancement [23,24].

For spectral analysis a Spex Triplamate spectrophotograph equipped with a Princeton Instruments Spec-10 liquid nitrogen cooled charge coupled device was used. Spectra were acquired with a laser power of 0.4 mW and an integration time of 1 s and images with 0.5 s per data pixel. Tips were etched from a gold wire of 0.25 mm diameter in a solution of 1:1 ethanol/fuming HCl [25]. This yields smooth tips with radii of 15–20 nm as indicated by scanning electron microscopy images (not shown). After each experiment the tip was checked for cleanliness. The bias was +150 mV and the tunnel current was set to 1 nA. Au single crystals were obtained from Mateck. They were cleaned in piranha solution and annealed by inductive heating in an Ar atmosphere. BCB (Fluka) was adsorbed from a \(10^{-6}\) M or, for lower coverages, \(10^{-8}\) M solution in Millipore water. The crystals were subsequently rinsed with pure Millipore water to remove excess BCB molecules. The UHV experiments were performed at pressures in the \(10^{-10}\) mbar range. For measurements in an oxygen atmosphere the chamber was vented with oxygen 5.5.

With this experimental setup a focal spot with a diameter of 300 nm is achieved [17]. The tip radius of 15 nm leads to a diameter of the TERS region of 15 nm [26].

BCB is an organic dye that has electronic transitions near to the resonance of the localized surface plasmon modes of the tunneling gap between the gold tip and...
gold sample, making it an ideal model system. Its resonant excitation with a 633 nm wavelength laser produces a weak, but detectable resonance Raman signal (far-field), while its adsorption at a metal surface effectively quenches the dye fluorescence. Because of our preparation procedure, on atomically flat Au surfaces about a 1/2 monolayer of BCB is adsorbed, corresponding to about 36 000 molecules in the focal area [27]. Upon tip approach, the signal intensity of the strongest band at 570 cm⁻¹ raises to a 3900-fold level corresponding to a Raman enhancement of 1.6 × 10⁶. The corresponding spectra show strong intensities though the TER signal stems from only ~90 molecules located within the TERS region (see Fig. 4 below). In this experiment we are still about 2 orders of magnitude short of the detection of single molecules; this indicates single-molecule TERS is in reach.

In the experiments shown in Figs. 1(a)–1(d), regions of a gold surface covered with either five BCB molecules or only a single, well separated one, were imaged by scanning tunneling microscopy. The STM height profiles [Figs. 1(b)–1(e)] display BCB molecules with a lateral size of 1.4 × 0.7 nm² and an effective height of 120–140 pm. TER spectra were recorded with the molecules located exactly in the gap between gold tip and substrate [Figs. 1(c)–1(f)]. For 90 molecules a Raman intensity of 38700 counts/s was recorded with an integration time of only 1 s, the size of 1.4 nm² corresponding to 110 counts/s. The local signal intensity consists of a constant background that only a single Raman scatterer is placed. Thus, the sensitivity also allows Raman imaging of single molecules on a time scale in which the probed surface section is stable.

For the single molecule in Fig. 1(c), Fig. 2(a) displays the local intensity of the band at 570 cm⁻¹ during a scan across the surface section. Under the present experimental conditions, i.e., a well-defined hot spot and a large suppression of diffusion by adsorption on a metal surface, probably on a specific site, there is no spectral blinking that is often observed in surface-enhanced single-molecule Raman spectra. After performing the Raman mapping, a STM image of essentially the same area was recorded, still showing the same single molecule (not shown). Figure 2(b) shows the horizontal and vertical TER intensity cross sections through the center of the molecule as well as a theoretical curve of the Raman enhancement profile along the substrate surface following a simple model. It describes the enhancement as the near-field interaction of an oscillating dipole in the center of a spherical particle with a molecule [28]. The electromagnetic coupling of the tip and the substrate is neglected which could result in an even more confined near field. Any deviation from a perfect shape of the tip cone or apex may result in a contrary effect. Therefore, the simplified model probably yields a realistic distance dependence of TERS as the r⁻¹⁵ dependence dominates all other effects [26]. The Raman intensity I(r) is given by

\[ I(r) = I_0 \left(1 + \frac{r^2}{(R_s + d)^2}\right)^{-6} + I_{bg}, \]

where r is the lateral distance from the molecule to the central point underneath the tip. A tip radius of \( R_s = 20 \) nm and a tunneling gap width of \( d = 1 \) nm were assumed. The values for \( I_0 \) and \( I_{bg} \) were chosen in correspondence with the spectra of the single BCB molecule and the tip above the uncovered gold sample in Fig. 1(f). Note that the probe is scanned across a homogeneous surface on which only a single Raman scatterer is placed. Thus, the local signal intensity consists of a constant background caused by the probe itself and the contribution of the scatterer. The latter is determined by the tip-scatterer distance dependent Raman enhancement, which exhibits an

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**FIG. 1 (color).** STM images of five (a) and a single BCB molecule (d) adsorbed on Au(111), the corresponding height cross sections (b),(e) and the UHV-RR and UHV-TERR spectra (c),(f).
dependence \[26\]. Consequently, the Raman image displays the above TERS profile along the surface. The agreement of the intensity cross sections with the theoretical curve confirms the validity of the simplified model employed here. Based on the same model the resolution of TERS imaging can be estimated by calculation of the Raman intensity map of two equal point scatterers placed on a plane surface \[29\]. If we assume that the intensity at the minimum along the line between the two scatterers should not exceed the maximum intensity of a single separated scatterer, we can postulate a simple rule for the image resolution of TERS: The minimum distance that can still be resolved, is approximately 70% of the sum of tip radius \(R_s\) and tunneling gap width \(d\).

Another example for tip-enhanced Raman mapping is presented in Fig. 3. The STM image shows a Au(111) surface with a few monoatomic steps at which BCB molecule clusters are preferentially adsorbed. ATER scan of the intensity at 570 cm\(^{-1}\) clearly resolves the position of these clusters that are separated by distances of 15–20 nm.

Above we presented single-molecule Raman imaging, which requires the dye to remain stable during the exposure of the enhanced field; i.e., the photobleaching must be substantially reduced under UHV conditions. However, for resonantly excited dye molecules, the huge electromagnetic enhancement also boosts the rate of photobleaching of dye molecules, with bleaching time constants (in air) down to subseconds \[9,26\] often yielding deteriorated spectra due to photodegradation of the dye. Under vacuum conditions, Raman studies on dye molecules without near-field enhancement yield a delayed photobleaching \[30\]. That this also holds for strongly enhanced near fields, will be shown in the following experiment.

Besides these benefits, the pronounced resonance of the system also causes a rapid photobleaching of the dyes in the vicinity of the tip. To determine the rate of photobleaching under UHV conditions, 30 subsequent spectra of a BCB monolayer on Au(111) were recorded first. The next experiment was performed with a freshly prepared sample and the same tip, but after filling the chamber with 1 atm of pure oxygen. Both series of spectra are shown in Figs. 4(a) and 4(b). In Fig. 4(c) the signal and background intensities at about 570 cm\(^{-1}\) are plotted vs the time of illumination showing a very different bleaching behavior in the presence and absence of oxygen. In UHV, the band intensity only drops to \(<0.0024\) within 30 s and all spectral features remain clearly visible. By contrast, the adsorbate is strongly and rapidly bleached in an \(O_2\) atmosphere which already affects the first spectrum. Almost the complete spectral information is lost during further illumination of the sample; after 30 s the band intensity is reduced to \(<15\%\) of the initial value. Figure 4(c) shows an initial time constant of photobleaching in an oxidative atmosphere of only 2.9 s. Another interesting aspect is the different evolution of the background intensity in UHV and in \(O_2\). In UHV, both band and background intensity decrease in almost equal measure. In \(O_2\), the rapid decrease of the signal intensity is accompanied by a strong, but temporary increase of the background around 250–1000 cm\(^{-1}\). It reaches a maximum value in approximately 3 s after which it is also bleached. The absence of this rise of the background under UHV conditions indicates that this effect must be associated with decomposition products of BCB accumulated during rapid bleaching, which are then themselves exposed to a further photodecomposition pro-
cess. Under vacuum conditions the rate of photobleaching is too low to visibly affect the background intensity.

Because of the extremely sharp focus of the parabolic mirror with high numerical aperture as well as the use of smooth and sharp gold tips, an excellent contrast between the far-field and near-field signal of \( \frac{2500}{3900} \) is achieved with the present UHV-TERS setup, employing monocrystalline Au samples as substrates. The strong contrast indicates an electromagnetic Raman enhancement of \( \frac{2400}{10^6} \) which is sufficient for the detection of single resonant dye molecules even within integration times as short as 1 s. The Raman spectra can be assigned to their source via the topographic images of the STM. The Raman mapping yields chemical information with a lateral resolution of \( \frac{24}{15} \) nm as proven by TERS imaging of a single BCB molecule as well as of larger molecular ensembles. We showed that, under UHV conditions, the adsorbed dye molecules are much more stable against photobleaching than in an oxygen atmosphere, even if very strong near-field intensities are present. Thus, in UHV a substantially extended measurement time is available, which permits Raman mapping (TER microscopy) on a sample.

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