On-chip surface enhanced Raman spectroscopy using ALD grown plasmonic nanotrenches integrated with a silicon nitride slot waveguide

(Student Paper)

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ABSTRACT

We present an enhanced Raman spectroscopy using sub 10 nm plasmonic nanotrenches directly grown on a silicon nitride slot waveguide using atomic layer deposition (ALD). A novel ALD process for gold deposition at 100°C is used, the precursor and reactant used for this process are Me3AuPMe3 and H2 plasma, respectively. The fabricated Raman sensor exhibits ∼ 1.5 × 10−8 pump to Stokes conversion efficiency for a monolayer of 4-Nitrophenol. This is at least an order of magnitude higher than the state of art nanoplasmonic waveguide based Raman sensors.

Keywords: Nanoplasmonics, Integrated Optics, Surface enhanced Raman spectroscopy, Waveguide enhanced Raman spectroscopy, Monolayer sensing, Atomic layer deposition.

1 INTRODUCTION

Surface-enhanced Raman spectroscopy (SERS) is a sensing technique that uses a nanoplasmonic structure to enhance the light in a nanometer-sized gap hence producing high electric field hot spots [2]. The analyte molecules present in these hot spots are then excited and the scattered Raman signal is collected either by a high NA microscope objective or a higher index contrast waveguide. The former technique is called free space SERS and the latter is known as waveguide based SERS [5]. Various nanoplasmonics structures i.e. nanobowties [5], [6] and nanotriangles [10] etc. have been integrated on a photonic platform to demonstrate waveguide based SERS. The fabrication of these plasmonics structures demands high resolution and alignment dependent patterning techniques. Recently, enhanced Raman spectroscopy using a hybrid nanoplasmonic waveguide [1], [7] has been demonstrated whereby mass scalable UV-photolithography is used. Despite all these design and technological advancements, the detection of an ultra-small concentration of weak Raman scatterers, demands further sensitivity improvement. One of the key applications in this regard is the single molecule spectroscopy [12] on a photonic chip.

In this work, we present an on-chip Raman sensor that uses plasmonic nanotrenches directly defined on a single mode silicon nitride Si3N4 slot waveguide using atomic layer deposition. Highly conformal ALD of Al2O3 and Au allows defining the nanotrenches in a more controlled and uniform fashion all along the waveguide width and length. Furthermore, the gold layer is deposited by a novel gold ALD process. The SERS performance of the resulting sensor is characterized using a monolayer of 4-Nitrophenol.

2 METHODS

2.1 Fabrication

The sensor is fabricated in three steps. In the first step, Si3N4 slot waveguides are fabricated using 193 nm optical lithography and subsequently etched by fluorine based inductive coupled plasma-reactive ion-etch process [8] to get the final structure as depicted in Fig. 1. Then a 40 nm layer of Al2O3 is deposited using ALD. Finally, a 100 μm window is defined using DUV lithography followed by an ALD deposition of a 24 nm thick gold layer composed of 7.9 nm wide nanotrenches. A home built ALD reactor with a base pressure of 2×10−6 mbar is used to deposit the Al2O3 and Au layers [3]. Both layers are deposited at a substrate temperature of 100°C. The Al2O3 layer is deposited using TMA and H2O as the precursor and reactant, respectively. The used pressure for both gases is 6x10−7 mbar. Au is deposited with a new ALD process, using Me3AuPMe3 and H2 plasma (20% H2 in Ar) as the precursor and reactant. A pressure of 1 mbar is used for the Au reactant and
Figure 1. The SEM images of the fabricated on-chip SERS device. a) Top view. Inset shows the Au island directly grown on a flat Si substrate. b) The SEM cross section of the waveguide showing the stack of the material layers.

6x10^{-3} \text{ mbar} for the hydrogen gas. In order to characterize the free space SERS strength of the ALD gold islands, a nanostructured gold layer is also grown on a flat silicon substrate as shown in Fig. 1.a (inset).

After the patterning, the chip was thoroughly cleaned with Acetone/IPA/DI water and dried with \( \text{N}_2 \) gun. For the monolayer coating, the chip was immersed in 1mM solution of 4-Nitrophenol (pNTP) in Ethanol. pNTP selectively binds to the gold using a thiol–gold bond.

2.2 Experiment

A confocal Raman microscope (WITEC Alpha 300R+) equipped with 100x/0.9 objective is used to characterize the Raman sensor. We acquire Raman spectra in a back-reflection geometry. A 785 nm laser and -70°C cooled Andor IDUS 401 CCD camera is used. All spectra are recorded using 1 mW pump power and a 1 sec integration time. For on-chip Raman measurements, the light is edge coupled to the waveguide and the polarization is aligned to the TE-mode of the waveguide. Also, same experimental conditions are used for free space SERS substrates. Further details about the experiment are reported in [7].

3 RESULTS AND DISCUSSION

In the first set of experiments, free space SERS performance of ALD gold layers is characterized. In order to optimize the gold thickness, four different thicknesses of ALD gold are deposited on a flat Si substrate using 100, 200, 400 and 800 ALD cycles. The measured Raman spectra are shown in Fig. 2.a (inset). Each spectrum is recorded by optimizing the CCD counts for the 1339 cm\(^{-1}\) (\(\nu_s(NO_2)\)) [4] Raman mode of pNTP. Very weak pNTP peaks were measured with the gold substrate where 100 ALD cycles are used. On the other hand, the stronger pNTP Raman modes appear in a Raman spectrum measured from substrate with 800 ALD gold cycles. This is attributed to the formation of nanotrenches \(\sim 7.9 \text{ nm} \) size that lead to a huge field enhancement.

Based on the 1339 cm\(^{-1}\) Raman mode, the pump to Stokes conversion efficiency \(\frac{\text{P}_s}{\text{P}_p}\) is measured using the method reported in [6]. The measured values of \(\frac{\text{P}_s}{\text{P}_p}\) are shown in Fig. 2.a. A highest value of \(\frac{\text{P}_s}{\text{P}_p} = (8.7 \pm 0.4) \times 10^{-9}\) is recorded for the 800 ALD cycles SERS substrate. This is almost one order of magnitude weaker than state of art free space SERS substrates such as a nanodome substrate [11] with \(\frac{\text{P}_s}{\text{P}_p} \sim 7 \times 10^{-8}\). However, it is worth mentioning that in the present case, a nanoplasmonic gap of 7.9 nm is achieved without any patterning technique and with further optimization of ALD gold process, \(\frac{\text{P}_s}{\text{P}_p}\) can be greatly enhanced.

After the free space SERS characterization, the Raman spectrum is measured using an on-chip SERS sensor. The spectrum is shown in Fig. 2. b. In order to reduce the silicon nitride background [9], the length of the access waveguide is limited to 1 \(\mu\text{m}\). The measured spectrum contains all the pNTP Raman modes. The measured \(\frac{\text{P}_s}{\text{P}_p}\) measured for the 1339 cm\(^{-1}\)mode is \(\sim 1.5 \times 10^{-8}\). Note that a coupling loss of 4.2 dB/facet has been taken into account for the on-chip \(\frac{\text{P}_s}{\text{P}_p}\) value. This is \(\sim 2\) times stronger than its free space counterpart. This is attributed to the long interaction length (few microns), leading to an efficient excitation and collection of Raman signal. Also, this is an order of magnitude stronger than the previously reported nanoplasmonic waveguide based on-chip Raman sensor where \(\frac{\text{P}_s}{\text{P}_p} = (1.0 \pm 0.057) \times 10^{-9}\) [7]. This substantial improvement is ascribed to the formation of 7.9 nm wide nanotrenches inside \(\text{Si}_3\text{N}_4\) slot waveguide. This leads to an efficient modal overlap between a photonic and a plasmonic mode hence producing a huge Raman enhancement \(|E|^4\).

CONCLUSION

We have reported a SERS sensor for an on-chip Raman spectroscopy using a novel ALD process of gold. Me\(_3\)AuPMe\(_3\) and H\(_2\) plasma are used as the precursor and reactant, respectively. The formation of 7.9 nm wide nanotrenches along the width and length of the \(\text{Si}_3\text{N}_4\) slot waveguide leads to a large pump to Stokes
conversion efficiency of $1.5 \times 10^{-8}$. This is the highest on-chip pump to Stokes conversion efficiency known to the date.

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