

# Numerical Investigation on Microfluidic Integrated Side Polished Fiber to Detect Biological Analytes

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**Abstract**—To improve the integration and sensitivity of surface plasmonic resonance (SPR) detection systems, we have numerically investigate the SPR microfluidic integrated side polished fiber (SPF) sensor to detect the biological analytes. The proposed device consists of PDMS microfluidic channel and few layer MoS<sub>2</sub> nano coated SPF. Finally, the integrated microfluidic channel with SPF bio sensor evaluated by flow velocity of micro fluidic channel and SPR occurring wavelength.

**Index Terms**—Microfluidic Channel, Side polished fiber, Surface plasmonic resonance, MoS<sub>2</sub> nanosheets

## I. INTRODUCTION

The surface plasmon resonance (SPR) technique is fundamentally based on the evanescent field interaction at a metal-dielectric interface. These metallic structures possess a highly localized electromagnetic field, making them highly sensitive to the refractive indices changes of the surrounding medium [1]. This techniques can be extensively utilized for various bio-component detection such as biomarkers, toxins, pathogens and allergens. Microfluidic technology enables the analysis and manipulation of substances inside a microchannels with a requirement of minimum volume of samples [2]. The Properly designed microfluidic chips can improve the efficiency of substance detection. The hybrid system of optical fiber with microfluidics results in rapid detection, high sensitivity, good reproducibility and high integration. As a result, numerous researchers have combined fiber-based SERS technology with microfluidics to create various microfluidic fiber-based SERS

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probes. Recently, Fan et al. integrated end-face fiber SERS probes into a microfluidic chip, achieving a detection sensitivity of 10 mol/L for R6G [3]. In this work, We have integrated the microfluidics channel with SPF in order to improve resonance effect through controlled flow assistance.

## II. DESIGN

In section.II, we illustrate about the design of the microfluidic integrated MoS<sub>2</sub> nanocoated SPF to detect the bio-analytes which is shown in Fig.1(a). The proposed design

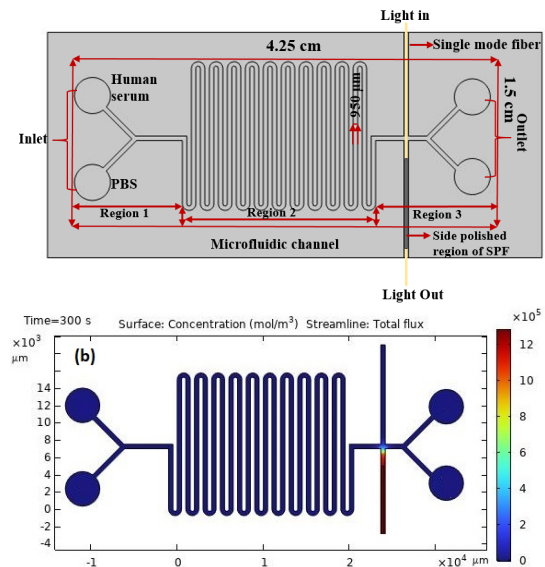


Fig. 1. (a) Schematic view of microfluidic integrated SPF Sensor. (b) Field profile of SPF sensor integrated with microfluidic channel.

consists of two major sections as microfluidic channel and MoS<sub>2</sub> nanocoated SPF. Firstly, we have design the microfluidic channel to utilize advantages of compact size, effecient flow control, requirement of low sample volume in order to improve the bio-substance detection of SPF. The proposed microfluidic channel has a length and height of 4.25 and 1.5 cm, respectively. The channel width of microfluidic channel is varied ranges from 1 to 1000  $\mu\text{m}$ . The proposed PDMS microfluidic channel comprises three regions. Region 1 and Region 3 has called as entrance and exit region, respectively, which consists of two inlets and outlets with a diameter of 0.5 mm. Inlets and outlets are used to insertion of human serum as well as PBS sloution and collection wastes, respectively. Region 2 is a middle region or mixing region of human serum and PBS solution. Further, the Region 2 supplies the mixed solution to the side polished MoS<sub>2</sub> coated sensing region. Further, MoS<sub>2</sub> nano coating is varied from monolayer to fewlayers(1 to 10) with a fixed single layer thickness as 0.625 nm. The considered SPF is regarded as a single-mode fiber, featuring core and clad diameters of 8  $\mu\text{m}$  and 125  $\mu\text{m}$ , respectively. The upper and above the core polished length of SPF is about 26 and 17 mm, respectively. The polished depth of SPF is about 58  $\mu\text{m}$ . Further, Fig.1(b) shows the field profile associated with SPF. In real time integration aspects, the microfluidic channel can be integrate with SPF by plasma bonding.

### III. RESULTS AND DISCUSSION

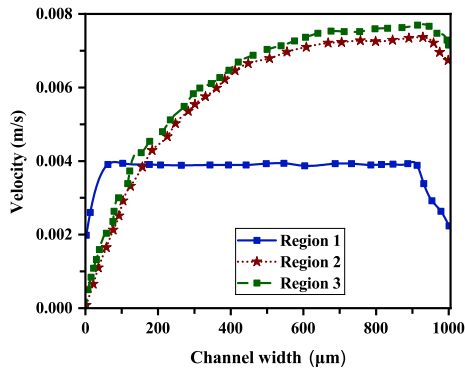


Fig. 2. Variation in velocity profile of microfluidic channel with respect to the variation in channel width.

Fig.2 depicts the velocity profile of proposed microfluidic channel with respect to the various channel width. Here, Region 1,2 and 3 represents the entrance,middle and exit region of microfluidic channel, respectively. one can see in Velocity profile that that the velocity of region 1 is same as  $3.9 \times 10^{-3} \text{ ms}^{-1}$  of region 2 and 3 at a channel with of 125 and 150  $\mu\text{m}$ , respectively. Initially,the velocity is set at  $2.5 \times 10^{-3} \text{ ms}^{-1}$  at the inlet of the microfluidic. The maximum velocity of region 2 (middle) and region 3 (exit) is attained  $7.3 \times 10^{-3} \text{ ms}^{-1}$  and  $7.7 \times 10^{-3} \text{ ms}^{-1}$ , respectively, at a channel width of 950  $\mu\text{m}$ . Further, Fig.2 indicates that the exit velocity get slightly exceeds the middle region velocity due to the reduction of cross sectional area of microfluidic channel

in the exit region. Hence, one can infer from the figure that constant velocity and enhanced velocity of entrance region and middle, exit region can be at microfluidic channel width of 950  $\mu\text{m}$ . Thus, we fix the channel width as 950  $\mu\text{m}$  through out studies.

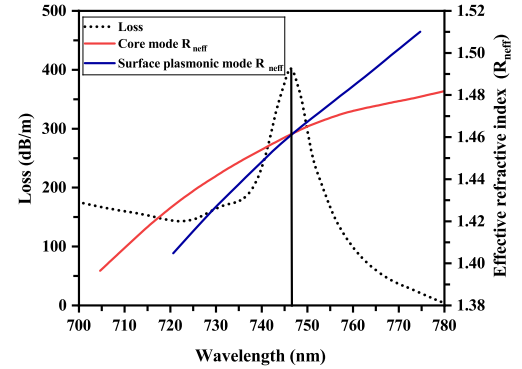


Fig. 3. Variation of loss as well as effective refractive index of core and SPR mode with respect to the wavelengths.

SPR is generally identified when the real effective indices and losses of the core and SPR mode indices match [4]. Resonance between core and evanescent field mode occurred only when the loss and real effective indices value of both modes gets matched. Fig.3 illustrates the real parts of the effective index and loss of both the core and the evanescent field mode as functions of wavelength. From Fig.3, it is clear that the real part of the both the mode indices of SPF match at 745 nm wavelength. On the same hand, the peak of the loss can also observed at 745 nm wavelength. Hence, the proposed SPF exhibits the SPR effect around the wavelength range of 745 nm.

### IV. CONCLUSION

In this work, the microfluidic channel integrated SPF has been proposed to improve the performance of SPR based bio analytes detection system. The proposed design exhibits the maximum velocity of  $7.3 \times 10^{-3} \text{ ms}^{-1}$  at 950  $\mu\text{m}$  of microfluidic channel width. Further, The SPF sensor exhibits the SPR at a wavelength of 745 nm for bio analytes detection.

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