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# Milled Microchannel-Assisted Open D-Channel Photonic Crystal Fiber Plasmonic Biosensor

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**ABSTRACT** A surface plasmon resonance (SPR) based photonic crystal fiber (PCF) sensor having a milled microchannel, and an open D-channel has been proposed in this paper. The sensor shows good functionality in the wide sensing range of 1.14-1.36 Refractive Index Units (RIU) of the analyte, having the capability to detect low refractive index (RI). The Finite Element Method (FEM) based numerical investigations dictate that the proposed sensor has been able to gain a maximum wavelength sensitivity of 53,800 nm/RIU according to the wavelength interrogation technique. The amplitude interrogations show that the sensor has the highest amplitude sensitivity of 328 RIU<sup>-1</sup>. The highest FOM (Figure of Merit) has been found to be 105 RIU<sup>-1</sup>. The sensor evinces a minimum wavelength resolution of  $1.86 \times 10^{-6}$  RIU, which secures high detection accuracy. A circular perfectly matched layer (PML) is implemented in the sensor's outermost layer as a boundary condition to absorb surface radiations. Gold is the plasmonic metal, while TiO<sub>2</sub> acts as the adhesive layer for gold attachment on silica. Due to the high sensitivity with a broad range of analyte detection, the sensor is well suited for practical biochemical detection purposes.

**INDEX TERMS** Broad sensing range, low analyte refractive index, plasmonic oscillations, wavelength sensitivity.

## I. INTRODUCTION

A photon coupling to an exciton gives rise to a polariton at the right frequency. The plasmonic oscillations occurring on a metal surface produce surface plasmonic waves (SPW). SPWs can occur in a metal-dielectric interface when photons with the right energy are absorbed under resonance conditions. This optical phenomenon has been engineered to act like a biosensor for biochemical analyte detection purposes. The SPR technology has shown good promise as a new optically advanced sensing scheme for its dynamic applications. These applications include temperature sensing [1], chemical sensing [2], label-free applications [3], and many more. With the advancement of the fabrication

technology, porous silica formation has been employed for sensing purposes [4]. SPR based PCF has found its way to widespread applications in biological implications. It has been reported that cancer cell detection is possible with this SPR based PCF biosensor [5]. Early detection of blood cancer is also reported in [6]. Recently, it has been reported that using plasmonic photothermal effect (PPT) in combination with localized SPR (LSPR), the genetic and nucleic acid testing of the coronavirus is possible [7]. It indicates Covid19 detection with more precision than the routinely utilized RT-PCR (Reverse Transcription Polymerase Chain Reaction) technique. The LSPR peak shifts have been reported to be utilized previously to detect the neurotoxin, ADDL (Amyloid  $\beta$ -derived diffusion ligand) related to Alzheimer's disease [8]. Refractive index detection of low carbon alcohols, such as methanol, ethanol, butanol, propanol, and phenol is possible

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using PCF SPR sensors [9]. In this work, the specific alcoholic analytes that are mentioned here are sensed based on SPR such that, the analytes are detected solely based on their refractive index. The highest wavelength sensitivity of the lattice design is 40,000 nm/RIU. However, the transmittance profile depicts that, there has been multiple resonance in the same wavelength for the same analyte, which puts forward a detection hindrance.

Plasmonic metals like gold and silver are most commonly used in designing biosensors with the SPR technology, because of the inter-band transitions as the metals are the noble ones. Gold is more preferred due to its higher chemical stability and higher resonance wavelength peak shifts. One of the most sensitive photonic crystal designs with only circular arrangements has been proposed in [10], where the highest wavelength sensitivity,  $S_\lambda$  is 28,000nm/RIU and the maximum amplitude sensitivity  $S_A$  is 6829 RIU<sup>-1</sup>. The RI sensing range of the analyte lies in 1.33-1.42 RI. However, with the advancement of the fabrication technology, different D-shaped open channel PCF-SPR sensors have been designed, which show higher sensitivity with a broad analyte sensing range. A novel D-shape design has been proposed that shows the highest  $S_\lambda = 11055$  nm/RIU within 1.20-1.29 RI of analytes [11]. Another team of researchers modified the airhole orientation of the design in recent years to gain a maximum  $S_\lambda = 20,000$ nm/R.I.U [12]. in the broad analyte sensing range 1.18-1.36 RI, capable of detecting low RI of analytes. However, their sensitivities are moderate in comparison to the most recent advancements in PCF SPR sensor designs. For accelerating the enhancement of the analyte's coupling more with the incident light wave guided through the core region, dual D-shape open-channel designs have also been proposed [13]. However, the wavelength sensitivity is quite low in the analyte sensing range 1.33-1.39 RI. Several dual D-shape open channels have been modeled, but the sensitivity did not reach moderately high values with respect to the different sensing ranges [14]–[16]. A circularly slotted multiple D-channel design has been proposed, which shows the highest  $S_\lambda = 16,000$ nm/RIU in the sensing range 1.40-1.46 RI [17]. Though the D-shape designs have introduced more volume of the analyte to interact with the core, the photonic crystal orientation did not support the lattice design to reach moderately high sensitivity values. All these lattice designs have been modeled, and the numerical explorations have been conducted using the FEM. Very few works have been conducted previously, which show good sensitivity with analyte RI changes. Similar work with a sensor design has been proposed and fabricated with moderate sensitivity [18].

The PCF sensor design introduced in this work utilizes an open D-channel along with a U-shaped deep groove channel to form the lattice design. Airhole arrangements have been carefully designed such that the periodic variations of the refractive index of airholes result in good coupling. Gold is the plasmonic metal with TiO<sub>2</sub> as the adhesive layer, which attaches gold with silica and enhances the

evanescent field's effect on the surface plasmons. The sensor has a broad refractive index sensing range of analytes ranging from 1.14-1.36 RI, capable of detecting lower refractive index analytes. Any type of optically isotropic liquid analyte with refractive index within our detectable sensing range 1.14-1.36 RI is possible to be detected and sensed with our lattice design structure with high sensitivity. The highest wavelength sensitivity has been ascertained to be 53,800 nm/RIU, with the highest  $S_A = 328$  RIU<sup>-1</sup>. This is the highest spectral sensitivity that has been achieved in this broad analyte sensing range to the best of our knowledge, capable of detecting very low refractive indices of analytes.

## II. THEORETICAL CONSIDERATIONS

As mentioned earlier, a photon coupling to an exciton produces a polariton, which is a part-light part-matter quasi-particle. This light-matter interactive optical phenomenon is responsible for the optical energy to be absorbed to create electronic oscillations in a metal. The metallic nanoparticles act as the quantum well, where the photon energy is utilized to create an electron-hole pair, which again gives up its energy to convert back to photons. In the presence of a metal-dielectric interface, when an electromagnetic wave falls on it, maximum optical energy is transferred from the incoming light to the surface electronic oscillations on the metal, known as surface plasmons at resonance condition. SPR can be achieved with the right wavelength of the electromagnetic (EM) wave entering the rightly suited crystal orientation of the PCF waveguide. SPWs give rise to the electric field generated at the interface. SPWs travel forward along with the interface. This corresponds to propagation constant [19]

$$\beta = \frac{\omega}{c} \sqrt{\frac{\epsilon_M \epsilon_D}{\epsilon_M + \epsilon_D}} \quad (1)$$

Here,  $\omega$  is the angular frequency,  $c$  is the velocity of light. The metal and the dielectric medium have permittivity  $\epsilon_M$  and  $\epsilon_D$ .

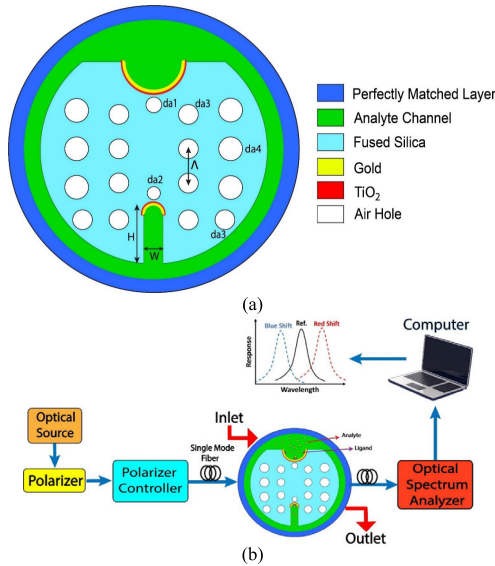
The phases coincide when the propagation constants of the incident light wave and the SPWs coincide. This leads to the matching of the frequency and the momentum of both the waves. Maximum modal power is delivered at this resonance condition. This causes the propagation loss to get elevated at resonant wavelengths due to the leaky gaussian nature of the core mode under such conditions. This confinement loss has a linear relation with effective mode index imaginary part  $\text{Im}(n_{\text{eff}})$  [10]

$$CL = 8.686 \times k_0 \times \text{Im}(n_{\text{eff}}) \times 10^4 \text{ (dB/cm)} \quad (2)$$

where,  $k_0 = \frac{2\pi}{\lambda}$  is the wave number in the free area.

## III. MODELLING AND NUMERICAL ANALYSIS

The representational model of the sensor is illustrated in Figure 1(a). The combination of a D-shape channel and a milled deep groove microchannel placed at the top and bottom of the sensor forms up the sensor design. Airholes have been arranged with four different diameters marked as  $da_1$ ,  $da_2$ ,  $da_3$ , and  $da_4$  with a square lattice around the center



**FIGURE 1. (a) Geometry and (b) experimental setup for making the sensor functional.**

of the sensor. For better internment of the electric field in the core, air holes have been placed incrementally from inwards to outwards. Two different airholes with diameters of  $da_1$  and  $da_2$  are placed separately just under the D shape channel and above the U-shaped deep groove, respectively. These airholes assist in creating a stable core mode with respect to  $\lambda$  alterations of the incoming EM wave. The D-channel and the groove's apex are coated with gold, which is the plasmonic metal, along with an adhesive layer of  $TiO_2$ . Gold shows poor attachment with silica. So, we have introduced the adhesive layer.  $TiO_2$  helps gold to boost up the sensing performance by elevating the surface plasmonic excitation. Because of  $TiO_2$  having higher RI, when it is allocated on the sensor surface, it can create better coupling between photons and polaritons. The ideal airhole parameters found after proper anatomization of the design are  $da_1 = 1.27\mu m$ ,  $da_2 = 1.20\mu m$ ,  $da_3 = 1.65\mu m$ , and  $da_4 = 2.2\mu m$ . The distance between two airholes is marked as pitch,  $\Delta$  which is kept at  $2.9\mu m$ . The suitable thickness of gold and  $TiO_2$  for both the D channel and the deep groove is kept at 80nm and 9nm, respectively. The D channel radius is  $2.40\mu m$ , and the U-shaped deep groove has a height of  $4.83\mu m$  and a width of  $1.67\mu m$ . The stack and draw technique supports the formation of the square lattice airholes [20], where pulsed lasers can be utilized to form the microchannels [21]. The side polishing method is an effective way of fiber polishing [22]. Chemical vapor deposition and atomic layer deposition allows the formation of nano-metallic layers with minimized surface roughness [23]. A femtosecond laser can create microchannels in optical fibers [24]. Focused ion beaming to create deep groove milled microchannels has been reported earlier [25].

Fused silica is employed as the base material because of its low absorption loss as well as high mechanical strength.

The Sellmeier's equation dictates the fused silica RI,  $\eta_{si}$  [26]:

$$\eta_{si}^2(\lambda) = 1 + \frac{B_1\lambda^2}{\lambda^2 - C_1} + \frac{B_2\lambda^2}{\lambda^2 - C_2} + \frac{B_3\lambda^2}{\lambda^2 - C_3} \quad (3)$$

where  $B_1, B_2, B_3, C_1, C_2, C_3$  are the Sellmeier constants for silica. Numerical values of each constant are  $B_1 = 0.696163$ ,  $B_2 = 0.4079426$ ,  $B_3 = 0.8974794$ ,  $C_1 = 4.67914 \times 10^{-3}\mu m^2$ ,  $C_2 = 1.3512 \times 10^{-2} \mu m^2$  and  $C_3 = 97.934 \mu m^2$  [5].

The RI of gold can be calculated as [10]:

$$n_{Au}^2(\lambda) = \epsilon_d = \epsilon_\infty - \frac{\omega_D^2}{\omega(\omega + j\gamma D)} - \frac{\Delta\epsilon\Omega_L^2}{(\omega^2 - \Omega_L^2) + j\Gamma_L\omega} \quad (4)$$

where  $\epsilon_\infty = 5.9673$  is the permittivity of gold at a high frequency.  $\omega_D, \gamma D, \Gamma_L, \Omega_L$  are the plasma frequency, damping frequency, spectral width, and oscillator strength of Lorentz oscillators, respectively. All the parameters for the above equation are taken from [10].

A thin  $TiO_2$  layer is placed between gold and silica for strengthening the attachment of gold with silica.  $TiO_2$  also raises the interaction between surface plasmons and its joining material, gold. The RI of titanium oxide can be expressed as [10]:

$$n_{TiO_2}^2 = 5.913 + \frac{2.441 \times 10^7}{(\lambda^2 - 0.803 \times 10^7)} \quad (5)$$

where  $n_{TiO_2}$  is the wavelength-dependent RI of  $TiO_2$ .

The whole numerical analysis has been carried out using Comsol Multiphysics as a simulation tool. The total fiber radius is kept at  $12.2\mu m$  with a non-physical PML having a thickness of  $1.2\mu m$ . The PML acts like a boundary condition which absorbs surface radiations and reflected light at the boundary. The PML's physical and the optical properties are kept the same as the base fiber material of the sensor to avoid a significant effect on sensing performance.

An experimental setup for the sensor's practical understanding is depicted in Figure 1 (b) as a resemblance. Broadband or super-continuum light source can be utilized and passed through a polarizer. A polarizer controller can also be used to have linearly polarized light. The polarized light can enter into single-mode fiber (SMF) via the splicing technique. An analyte channel has been introduced in our sensor at the outer layer, which eases the analyte's inlet and outlet. A pump can control the inlet and outlet. SPR establishes a binding environment between the ligands of the plasmonic metal and the analyte, which gives relocation of resonance peak resulting in either a blue or redshift of the response curve. An optical spectrum analyzer (OSA) can be used to sense the wavelength shifts, and a computer can be utilized to analyze the output.

## IV. SIMULATIONS AND RESULTS

### A. WAVELENGTH INTERROGATION METHOD

The core and the SPP mode's field profiles are shown in Fig. 2 (a) and (b). Because of the lattice structure's precise

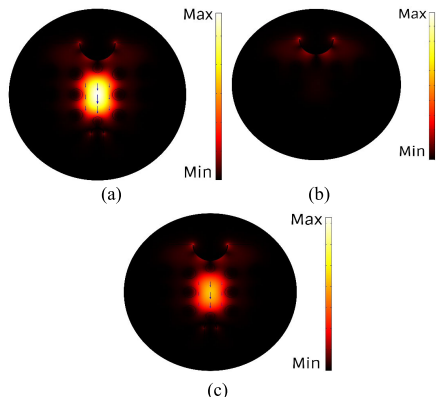


FIGURE 2. Electric field profile of the (a) core mode (b) SPP mode and (c) the resonance coupling.

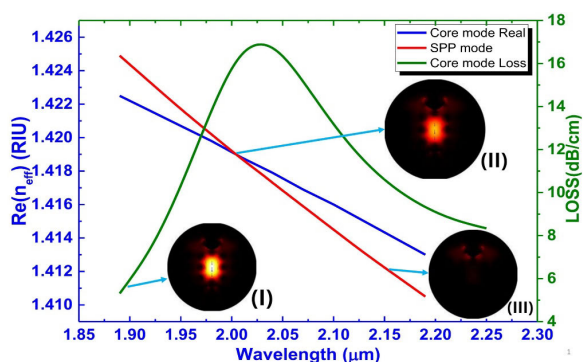


FIGURE 3. The real part of the effective index of the core mode (blue), SPP mode (red) and confinement loss of the core mode (green) with respect to wavelength alterations.

photonic crystal orientation, the incident light is guided through the core. The evanescent field of the core guided mode causes the energy leakage from the guided EM wave to the SPWs. At the phase-matching condition, the highest energy content is transferred from the core mode to the SPP mode, which is portrayed by the resonance coupling mode in Fig. 2(c). The square lattice design with increasing diameter from inwards to outwards with a D-channel and a deep groove at the bottom better interlinks the evanescent waves from the core to the polaritons. This enhances the coupling for a broad range of refractive indexes of the analytes as more analyte volume can interact with the core guiding light. The plasmonic metal introduced at the deep groove’s apex lets the evanescent field interact with lesser damping. Fig. 3 shows the dispersion relations, the real part of the refractive indices of the core and the SPP mode, along with the confinement loss curve. The crossing of the real parts of both the modes dictates that incomplete coupling has taken place for the lattice design. The crossing ensures the phase matching of the EM wave with the SPW. The wavelength of the incident EM wave at the point of the crossing is the resonant wavelength. Any kind of minuscule changes in the analyte causes the optical properties of the analyte to change as well, particularly, the refractive index. Hence, the RI alteration of the analyte causes the overall effective mode index to change, which

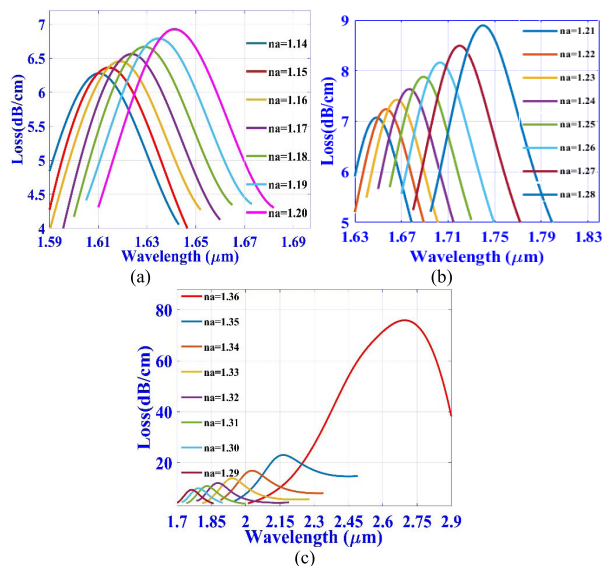


FIGURE 4. Loss curves for analyte refractive index variations (a) 1.14-1.20 (b) 1.21-1.28 and (c) 1.29-1.36.

causes the crossing to take place at different wavelengths. That is, the resonant wavelength shifts with the analyte RI alterations. From Fig. 4, it is evident that the loss curves show red shifting with the increment in the analyte RI. The probable cause for the red shifting of the resonant wavelength,  $\lambda$  is that higher RI of analyte on the periphery causes the damping of the plasmonic waves. This causes the propagation constants to decrease, resulting in the refractive index matching at higher wavelengths.

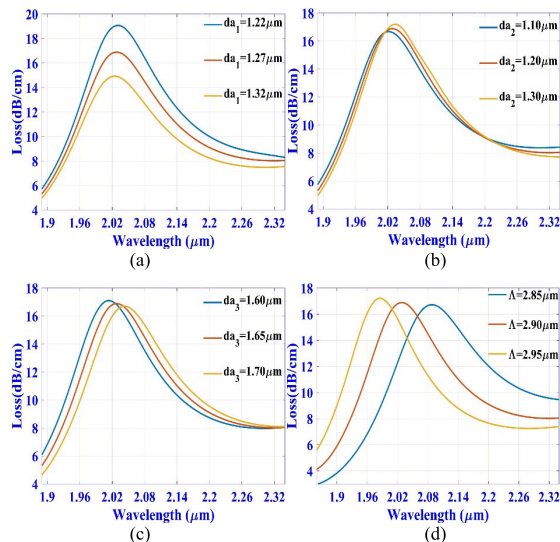


FIGURE 5. Loss curves for diameter variations (a)  $da_1$  (b)  $da_2$  (c)  $da_3$  and pitch (d) variations at 1.34RI of the analyte.

The fabrication tolerance has been studied on the lattice design numerically. The airhole diameter  $da_1$ ,  $da_2$ , and  $da_3$  variations are elucidated in Fig. 5 (a), (b), (c). The tolerance of  $da_1$  has a greater effect on the sensor response. As it directly lies in between the core and the metal-dielectric interface,



decreasing this airhole diameter causes greater energy transfer between photon and polariton, resulting in better coupling.  $da_2$  variations have a lesser effect on the loss curves as the surface plasmons are lesser in the groove's apex. D-shaped channel diameter is kept at  $4.8\mu\text{m}$  and U-shaped groove width is kept at  $1.67\mu\text{m}$  in our work. From these two values, it is evident that generation of the evanescent field and the amount of surface plasmons are much higher in the D-shaped dielectric interface than that of the U-shaped groove dielectric interface because of the D-shaped channel having much higher area.  $da_2$  airhole is placed just above the U-shaped groove, where the amount of surface plasmons and their coupling with the generated evanescent field is not high. For this reason, we have not seen any significant changes in the loss curves in Fig. 5(b) although we have varied  $da_2$  by almost  $\pm 8.35\%$ . On the contrary,  $da_1$  airhole is placed just under the D-shaped channel where the amount of surface plasmons and generation of evanescent field is high. Hence, we saw a significant change in loss curves in Fig. 5(a) although we varied  $da_1$  only by  $\pm 3.9\%$ . It can be added that the presence of  $da_1$  and  $da_2$  comes off with great significance for confining light in the core. Without these airholes, a stable core mode cannot be obtained with respect to sweeping the wavelength of the incoming light.

The increment of  $da_3$  causes a slight red-shifting of the curve. Pitch variations have a negligible effect on the energy transfer between the two modes, as depicted in Fig.5 (d). However, the resonant wavelength shows blue shifting with an increment in the pitch,  $\Lambda$ . As the gap between the airholes is increased, the confinement of light in the core decreases, which excites the surface plasmons a little lesser. The performance of the sensor corresponding to analyte RI variations has been tabulated in Table 1. According to the wavelength interrogations, the wavelength sensitivity parameter evaluates the sensing response of the sensor, which is primarily dependent on the resonance  $\lambda$  peak shifts. This shifting of the loss curve is peaked by a spectrometer at the output to detect the analyte changes. The wavelength sensitivity,  $S_\lambda$  is defined [10]

$$S_\lambda [nm/RIU] = \frac{\Delta\lambda}{\Delta n_a} \tag{6}$$

where,  $\Delta\lambda$  is the peak wavelength shifts and  $\Delta n_a$  denotes to successive analyte RI changes. The sensor's highest  $S_\lambda$  for the analyte changes from 1.35-1.36RI, which is  $53,800\text{nm}/\text{RIU}$ . The  $S_\lambda$  increases non-linearly from lower analyte RI to the higher ones.

For the least possible detection of spectral shifting, the sensor wavelength resolution evaluates the sensor response with respect to the analyte alterations [27]

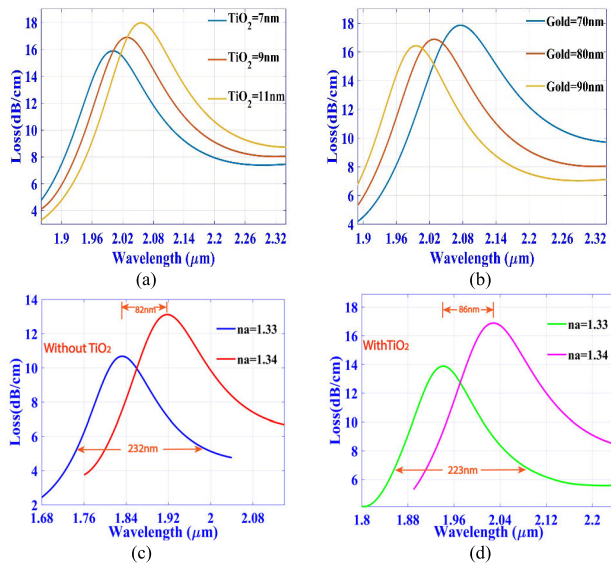
$$R = \Delta n_a \Delta\lambda_{min} / \Delta\lambda_{peak} RIU \tag{7}$$

where,  $\Delta\lambda_{min} = 0.1\text{nm}$  is considered as the minimum spectral resolution [10]. The R is inversely proportional to  $S_\lambda$ . The proposed sensor exhibits an R of  $1.859 \times 10^{-6}$  RIU.

TABLE 1. Sensor performance within sensing range.

ANALYTE INDEX	Loss Peak $\alpha$ (dB/cm)	$\lambda_{peak}$ ( $\mu\text{m}$ )	Resonant Peak Shift (nm)	$S_\lambda$ (nm/RIU)
1.14	6.277	1.610	4	400
1.15	6.362	1.614	5	500
1.16	6.456	1.619	4	400
1.17	6.562	1.623	6	600
1.18	6.669	1.629	6	600
1.19	6.790	1.635	7	700
1.20	6.926	1.642	7	700
1.21	7.074	1.649	8	800
1.22	7.239	1.657	9	900
1.23	7.425	1.666	11	1100
1.24	7.637	1.677	12	1200
1.25	7.880	1.689	14	1400
1.26	8.161	1.703	17	1700
1.27	8.494	1.720	20	2000
1.28	8.896	1.740	24	2400
1.29	9.393	1.764	30	3000
1.30	10.030	1.794	37	3700
1.31	10.890	1.831	47	4700
1.32	12.080	1.878	63	6300
1.33	13.880	1.941	86	8600
1.34	16.880	2.027	136	13,600
1.35	23.06	2.163	538	53,800
1.36	75.92	2.701	N/A	N/A

Fig. 6 (a) and (b) show the effect of metal layer variations on the sensor response's loss curves. An increase in the gold layer dimensions causes the coupling to decrease by a small amount. Also, the resonant  $\lambda$  is blue-shifted as the propagation constant of the plasmonic wave increases due to greater plasmonic oscillations. The increment of the titanium oxide layer enhances the coupling, followed by red-shifting. This will keep on increasing with the  $\text{TiO}_2$  layer thickness. A very high thickness will cause high losses, which limit the sensing length, causing fabrication impediments. Hence, we prefer an ideal thickness of around 9nm. Fig. 6(c) and (d) portray the loss curve shifting for the analyte RI change from 1.33 to 1.34 RI. The two figures reveal the  $S_\lambda = 8200\text{nm}/\text{RIU}$  and  $S_\lambda = 8600\text{nm}/\text{RIU}$  without using  $\text{TiO}_2$  and with using  $\text{TiO}_2$  along with gold respectively. That is, usage of  $\text{TiO}_2$  ameliorates the sensitivity. In addition to this, the detection accuracy increases for using  $\text{TiO}_2$  as the Figure of Merit (FOM) is improved. This occurs for the reduction of the Full Width Half Maximum (FWHM) of the Gaussian loss curves as the two quantities are reciprocal to each other.



**FIGURE 6.** Loss curves for metal layer thickness variations at analyte index 1.34RI for (a) TiO<sub>2</sub> (b) gold (c) loss curve shifting for the analyte index alteration from 1.33 to 1.34RI, without TiO<sub>2</sub> and (d) loss curve shifting for the analyte index alteration from 1.33 to 1.34RI, with TiO<sub>2</sub>.

The same is true for the whole sensing range of the sensor. TiO<sub>2</sub> is significant as an adhesive layer for gold attachment on silica. It is important that this additional adhesive layer does not intervene with the sensing potential of the sensor. The didactic inference from Fig. 6(c) and (d) suggests that TiO<sub>2</sub> further enhances the sensitivity instead of impeding it. Hence, we promote the usage of this adhesive layer along with the plasmonic metal.

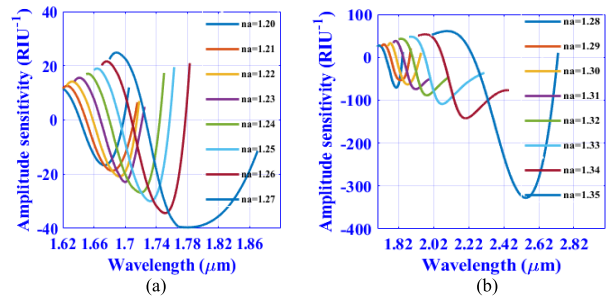
**B. AMPLITUDE INTERROGATION METHOD**

Amplitude sensitivity, S<sub>A</sub> is another significant cost-effective means of judging the sensor performance. An OSA measures the resonant wavelength shifts. However, the output light intensity measured at different input EM wavelengths varies with the analyte RI alterations. The output intensity shows dips with the subsequent rise in the confinement loss. This output intensity is measured by a photodetector, which is a cheap means of getting viable output for detecting analytes. Good S<sub>A</sub> readings of any sensor pave the way for the sensor to be used with a photodetector to get good transmittance curve dips at the resonant λ. Fig. 7 (a) and (b) show the S<sub>A</sub> shifts with RI variations. Increasing the analyte RI causes the S<sub>A</sub> curves to be red-shifted with a non-linear rise in the sensitivity. The S<sub>A</sub> is stated as [28]

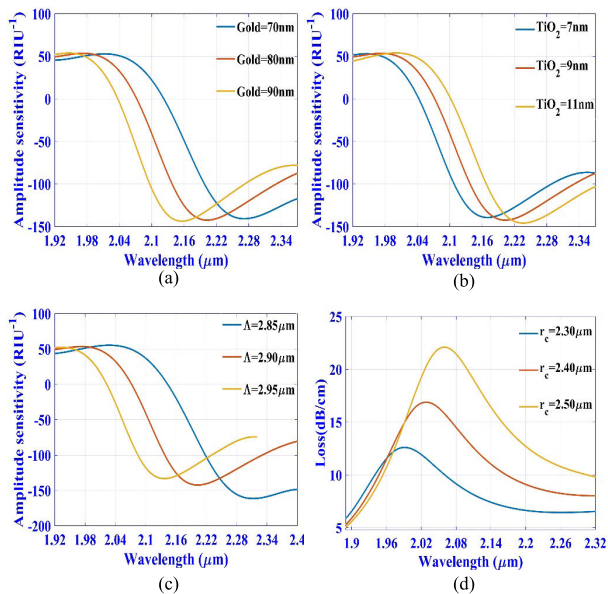
$$S_A(\lambda) \left[ RIU^{-1} \right] = -\frac{1}{\alpha(\lambda, n_a)} \frac{\delta\alpha(\lambda, n_a)}{\delta n_a} \quad (8)$$

where, α(λ, n<sub>a</sub>) is the confinement loss for any analyte RI, δn<sub>a</sub> is the successive analyte RI difference and δα(λ, n<sub>a</sub>) is the difference of the loss obtained for consecutive analyte RI. The highest S<sub>A</sub> of the sensor is found to be 328 RIU<sup>-1</sup>.

The effect of metal layer thickness variations has been studied on the S<sub>A</sub> and depicted in Fig. 8 (a) and (b).



**FIGURE 7.** Amplitude Sensitivity curves for analyte index sensing range (a) 1.20-1.27 RI and (b) 1.28-1.35 RI.



**FIGURE 8.** Amplitude Sensitivity curves with respect to (a) gold layer thickness variations (b) TiO<sub>2</sub> layer thickness variations, (c) pitch variations and (d) channel variations on the loss curve.

The increment of the gold layer, and the TiO<sub>2</sub> layer causes the resonance λ to shift, causing the amplitude sensitivity curves to shift. Increasing the gold layer causes the blue shifting of the amplitude sensitivity curve and increasing the TiO<sub>2</sub> layer causes red-shifting. As mentioned earlier, an increment of the pitch results in the blue shifting of the resonance λ. This results in the blue shifting of the S<sub>A</sub> curve as shown in Fig. 8(c).

Fig. 8(d) elucidates the effects of the channel area variations on the sensor response. It is evident that, with the increase of channel radius the loss increases, followed by slight redshifts. As the channel area increases, more analyte volume can interact with more surface plasmons on the metal. This causes more energy transfer at the resonance conditions.

**C. COMPARATIVE DISCUSSION**

Table 2 comes forward with a relativistic study of the proposed sensor with respect to the other proposed designs in terms of the S<sub>λ</sub> and the analyte RI sensing range. All the other

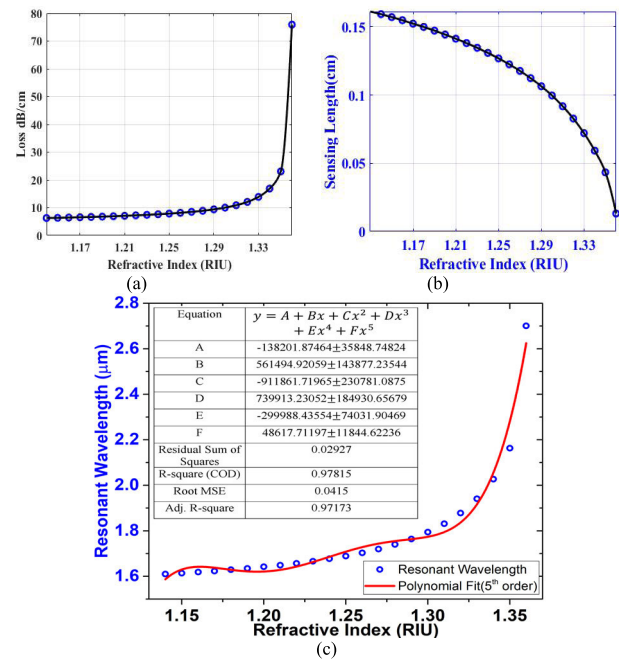
**TABLE 2.** Comparison of the sensor performance with other recent publications.

Ref.	Structure	WS (S <sub>i</sub> ) (nm/RIU)	RI Sensing Range
[10]	Hexagonal lattice	28,000	1.33-1.42
[17]	Circular slotted D-channel	16,000	1.40-1.46
[29]	Concave shaped	9314.28	1.33-1.38
[30]	Spiral lattice	23,000	1.32-1.40
[31]	H-shaped	25,900	1.33-1.49
[32]	Square lattice	25,000	1.33-1.38
[13]	Two-sided open channel	5,000	1.33-1.39
[33]	Hollow core	21,000	1.33-1.42
[34]	Series-wound	17,500	1.13-1.35
[12]	D-shaped	20,000	1.18-1.36
[35]	Concave shaped	10,700	1.19-1.29
[36]	D-shape	15,000	1.22-1.33
[9]	Square lattice	40,000	1.33-1.53
Proposed design	D-channel deep groove	53,800	1.14-1.36

references are proposed designs of PCF SPR sensors, based on FEM, whose results have been numerically investigated. Our proposed lattice design has been able to gain a very high sensitivity in terms of wavelength interrogations for a very wide analyte sensing range. The wide analyte sensing range includes any liquid analyte having a refractive index range of 1.14-1.36 RI. References [12], [34]–[36] illustrate sensor works targeted for low analyte RI detections. However, with our non-linear rise of wavelength sensitivity, the sensing readings are much higher in a broader sensing range of analytes. A higher maximum sensitivity leads to rising slopes when going from any low RI analyte to the higher one. This indicates the improvement of the average sensitivity of the sensor for the whole sensing range. The comparative analysis with the other published lattice design works shows different lattice shapes with various photonic crystal arrangements. However, such types of proposed models lack the precise arrangement for the guided photons to couple with the polaritons for a broad sensing range, such that the sensitivity to the analyte RI alteration is high enough within the working range. The milled channel introduced with the open D-channel and the enhanced form of the well-calibrated photonic crystal lattice design have produced the desired results that have been necessary for the domain of PCF-SPR sensor works.

**D. SENSOR EVALUATION**

The sensor shows a non-linear rise in the resonant wavelength loss values for the increment of the incident wavelength. This has been shown in Fig. 9(a). Fig. 9(b) portrays the sensing length of the sensor with respect to the analyte index alterations.



**FIGURE 9.** Response of the sensor for analyte RI alterations in terms of (a) confinement loss at resonant  $\lambda$  (b) sensing length of the sensor and (c) resonant  $\lambda$ .

The actual sensing length for such sensors has a reciprocal relationship with the confinement loss at the resonant  $\lambda$  [32].

$$L = \frac{1}{\alpha(\lambda, n_a)} \tag{9}$$

Greater confinement loss impedes the practical realization of the sensor as it decreases its sensing length. This makes it harder for the practical analysis of the sensor when splicing with fibers at the input and the output. The proposed sensor shows a maximum CL = 75.92 dB/cm, which is reasonably low in comparison to the other sensor works previously done. The sensor shows a non-linear rise in the resonant  $\lambda$  with increasing RI of analytes. A polynomial is fitted to the 5<sup>th</sup> order for the resonant  $\lambda$  increment with the analyte alterations in the increasing order, as illustrated in Fig. 9(c). This gives the characteristic polynomial equation for the sensing performance. Polynomial fitting accuracy largely depends on two factors; 1. Adjusted R<sup>2</sup> value and 2. Root mean square error (Root MSE). A polynomial fit is considered to be the most accurate when the adjusted R<sup>2</sup> value is close to 1 and the Root MSE is close to 0. A 5<sup>th</sup> order polynomial fit gives the most accurate fitting in terms of these two factors. It has been also highlighted in Fig. 9(c).

A better Figure of merit (FOM) can improve the sensor response in terms of realizing the output response from the

unwanted noise signals. The FOM is reciprocal to the full width of the half maximum (FWHM) of the gaussian sensor response [10].

$$FOM = \frac{S_\lambda}{FWHM} \quad (10)$$

The highest figure of merit for the sensor design is 105 RIU<sup>-1</sup>. The figure of merit improves for the increase in wavelength sensitivity at higher analyte RI. However, the FWHM consequently decreases with analyte RI increments.

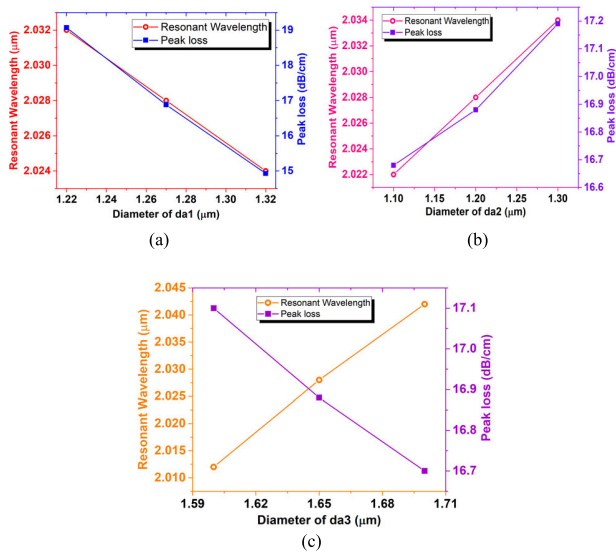


FIGURE 10. Resonant wavelength and corresponding loss as a function of (a)  $da_1$ , (b)  $da_2$  and (c)  $da_3$ .

The study of the optical physics behind the loss curve shifts in Fig. 5 and 6, as mentioned earlier, can directly lead to Fig. 11 and 12. Fig. 10(a) reveals that the sensor shows almost a linear response with equal negative slopes for the resonant  $\lambda$  and the corresponding loss for increasing the diameter  $da_1$ . The same goes for  $da_2$  with rising slope, depicted in Fig. 10(b). Fig. 10(c) shows a reciprocal relationship between resonant  $\lambda$  and loss for  $da_3$ . Fig. 11(a) focuses the same for the gold layer thickness increments. Increasing gold film thickness results in the change of resonant  $\lambda$  and loss with negative slope. TiO<sub>2</sub> thickness changes them with positive slopes, the slope of resonant  $\lambda$  being a bit higher, as shown in Fig. 11(b). Fig. 11 (c) illustrates that there exists almost a reciprocal relationship between the two variables under study for pitch variations. Fig. 11(d) elucidates the changes on them having almost equal positive slopes for the increasing D-channel radius. These relationships give an intuitive idea of the sensor response with respect to the parameters under study, revealing more about the fabrication tolerance.

### V. ANECDOTE FOR FUTURE IMPROVEMENTS

As inferred previously, the quality simulation works and their numerical investigations give a broad spectrum of data and knowledge about how to go about with the geometrical features, such that the experimental researchers can correlate more about the geometry of the design and the

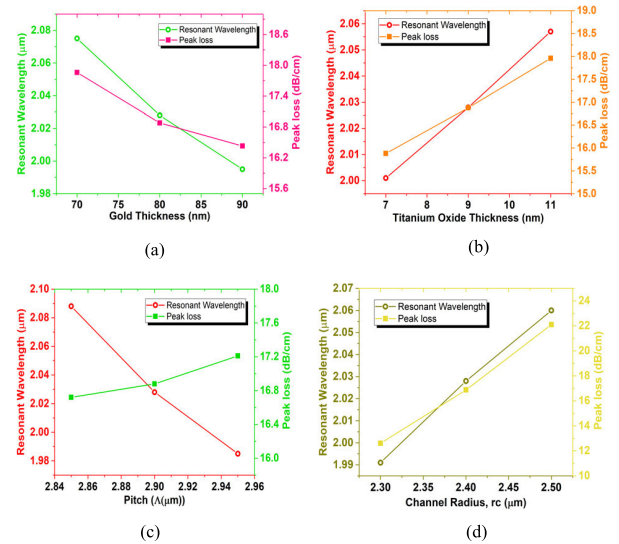


FIGURE 11. Resonant wavelength and corresponding loss as a function of (a) gold thickness, (b) TiO<sub>2</sub> thickness, (c) pitch and (d) channel radius.

sensor response. With a view to achieving that, wavelength sensitivity and the amplitude sensitivity have been extensively studied. Lower birefringence peak shift study is another way to go about judging the sensor response [27]. However, the sensitivity enhancement factor (SEF) comes forward from a pragmatic point of view in judging the sensor response [37]. It gives a better estimation of the content of the target molecules required for obtaining a detectable signal by selective localization in the strong fields. Besides, in surface enhanced Raman scattering sensors, substrates with high SEFs are desired [38]. Analysis of a PCF SPR sensor in terms of SEF has not been yet carried out. But it is matter of concern to study in our future works.

### VI. CONCLUSION

In this work, a milled deep groove has been introduced into the fiber sensor design with an open D-channel. This results in more volume of the analyte to interact with the core mode. This broadens the analyte sensing capability of the sensor, ranging from 1.14-1.36 RI. The plasmonic oscillations occurring in very close proximity of the stable core mode, due to the lattice design's geometric orientations result in improved spectral sensitivity. The highest wavelength sensitivity is 53,800 nm/RIU with the highest amplitude sensitivity of 328 RIU<sup>-1</sup>. High sensitivity having such a broad sensing range makes the sensor design suitable for dynamic biochemical detections.

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