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Original Article

Effect of BaTiO3 and Chitosan Composite Material on the Enhancement of the Sensitivity and Limit of Detection of the Surface Plasmon Resonance **Sensor-Divergent Beam Based**

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Divergence Beam, SPR Sensor, Limit of Detection (LOD). Chitosan and Sensitivity.

The SPR sensor's poor sensitivity and low limit of detection (LOD) limit its use for tiny contamination detection when the measured sample ranges from 1.3317 to 34. Thus, study examined the combined influence of CaF2/ZnO/TiO2/Silver/BaTiO3/Chitosan on the performance of an SPR biosensor employing the Kretschman configuration. The proposed SPR sensor use the Kretschmann configuration, which employs the divergence beam produced by the Powell lens to avoid mechanical rotation. Through the use of an STM32F401RE controller with 12-bit ADC, it was determined that the proportionate alloying of BaTiO3 and chitosan with optimal thickness could enhance the designed SPR sensor's theoretical sensitivity to 480.45°/RIU and limit of detection (LOD) to. Thus proposed SPR sensor showed the improvement in sensitivity and LOD by 16.67% and 33.69%, respectively, as compared to the SPR sensor without the combined effect of optimal chitosan and BaTiO3 thickness.

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INTRODUCTION

The optical technique of surface plasmon resonance spectroscopy operates on the principle of attenuated total internal reflection (ATR). When P-polarized light illuminates the prismmetal interface at an angle greater than the critical angle, the entire light is reflected, a phenomenon known as total internal reflection [1, 2]. Even if all the light is reflected, a propagating wave known as an evanescent wave will occur at the metaldielectric interface, which is extremely sensitive to slight changes in the refractive index of the dielectric material [1, 2]. Consequently, surface plasmon resonance occurs when the evanescent wave's propagation constant equals that of the surface plasmon wave [2]. As a result, when the refractive index at the metal-dielectric interface changes, total internal reflection disappears. This is known as attenuated total internal reflection (ATR). The resonance angle is the reflected angle at which the reflected intensity is nearly zero [1, 2]. The detected intensity is indicated by the dark band at this location. When the detected intensity zero signify optimal photon energy transformation have been achieved.

Researcher have used SPR sensor in many fields like detecting waterborne and airborne harmful bacteria and virus which are dangerous for public health [3, 4]. Quantization of kinetic and equilibrium characterization of biomolecular interactions are easily measured by SPR sensor [5]. Despite the researcher's efforts to improve sensitivity and limit of detection (LOD) in conventional prism based SPR sensors (single metal layer system) that work on angular interrogation, the sensor still suffers from insufficient sensitivity and LOD, despite the fact that high sensitivity and LOD is the primary goal of the researcher. Therefore, recently researchers have shifted their focus to multilayer SPR based as an alternative mechanism of improving the sensitivity and LOD of the SPR sensor.

The designed SPR sensor used Kretschmann based configuration with six layers. Silver as a metal was selected because it has high detection accuracy in comparison to other metals like gold

[6]. The adhesion between glass and metal is typically poor because of this, chromium (Cr) and titanium (Ti) are frequently used to increase the adhesion between metal and glass [7]. These materials' low optical transmission and metal interdiffusion limitations are a drawback due their refractive index in visible range has real and imaginary, imaginary part absorb fraction of the photon energy resulting to decrease in sensitivity and widening of the reflected SPR reflectivity curve [7].

The use of the adhesion layer with the real refractive index was an option for improving the sensitivity and detection accuracy [8]. Many studies have proposed the use of Si, SiO₂, ZnO and TiO₂ which gives better SPR performance than those with complex refractive index in the visible [8-10]. However, it has been experimented that the combination of low and high purely real refractive index makes the composite layer of ZnO and TiO₂ produce better sensitivity than single adhesion layer [9, 11].

When silver used in SPR sensor gives better performance than gold in terms of sensitivity and detection accuracy because the ratio of real part to imaginary part of silver is higher than other metals like gold, copper, aluminium and palladium [12]. However, silver has a detrimental effect of easily oxidized when encounter with water, it is hardly to be reused. The researchers have addressed the problem by adding transition metal dichalcogenides (TMDCs) monolayers such as graphene, molybdenum disulfide (MoS₂) [13] with the sensitivity of 230.66°/RIU, MoSe₂ [14] with sensitivity of (215.5°/RIU), MXene [15, 16] which are resistant to oxidation, meanwhile the sensitivity is improved.

BaTiO₃ is widely used due high purely real refractive index when compared to other transition metal dichalcogenides (TMDCs. Recently research have shown that when transition metal dichalcogenides (TMDCs) combined perovskite material like barium titanate (BaTiO₃) the sensitivity of the SPR sensor will be highly improved. It has been observed that BaTiO₃ is widely used compared to other

perovskite material like, lead titanate (PbTiO₃) [17], strontium titanate SrTiO₃ and potassium niobate (KNbO₃) [18] this is because of due high purely real refractive index [19].

Wang et al. proposed an SPR sensor with a maximum sensitivity of 294°/RIU in a (prism/Ag/Au/BaTiO₃/Graphene) configuration structure [20]. Wang et al. [21] present another work in which they design an SPR sensor with a multilayer structure of (Ag/BaTiO₃/Graphene) and a sensitivity of 257°/RIU. Saharia et al [22] proposed an SPR sensor sandwiched with (BaTiO₃) and graphene that has a sensitivity of 116.67 °/RIU. Pal and Jha [23] proposed SPR multilayer (Ag/BaTiO₃/Ag/Graphene) sensor with sensitivity of 280°/RIU. Imam et al [24] proposed calcium fluoride prism/BaTiO₃/Blue Phosphorus/WS₂ multilayer structure obtain maximum sensitivity of 413.7 °/RIU.

In this article the SPR sensor was configured as calcium fluoride prism CaF₂/ZnO/TiO₂/Ag/ BaTiO₃/chitosan. Prism CaF₂ was opted since it has the lowest refractive index which results in high sensitivity and improved LOD compared to the prism with higher refractive index. Apart from prevention of oxidation on the silver, this study has found that optimum thickness of BaTiO₃ enhances the sensitivity by the factor of 1.7 in comparison to convection SPR sensor [25]. For the purposes of increasing absorption chitosan film was added on top to BaTiO₃ which has high absorption capability [26]. Chitosan is the copolymer which can be formed naturally in some microorganisms such as fungi and yeast [27, 28]. It is biodegradable and harmless to human health. It has numerous applications in food and agriculture technology, biomedicine and cosmetic [27, 29]. Moreover, it can be derived from alkaline N-deacetylation which is mostly found in shells of shrimps, krill, lobsters, and crabs [27]. This signify that chitosan is healthy and easily found. Henceforth, it has low cost. Chitosan has reasonable absorption capacity due to the presence of amine group and hydroxyl group [28, 29]. Therefore, the presence of amine group is helpful for immobilizing the surface which increases carboxylic bonding [30].

Significant effort has been dedicated to improving the sensitivity and limit of detection (LOD) of the SPR sensor using an additional multilayer approach, which are essential attributes for identifying tiny changes in the measured sample. Nevertheless, no theoretical studies have employed a BaTiO₃ and Chitosan alloy with appropriate thickness to enhance the sensitivity and detection limit of the SPR sensor. Thus, study concluded that the integration of an optimal chitosan thickness, alongside a suitable BaTiO₃ thickness, significantly improved sensitivity and limit of detection (LOD), exceeding the findings of the latest published research in these metrics. The results were realized by using transfer matrix method with help of MATLAB software [2].

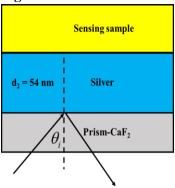
THEORETICAL DEVELOPMENT SPR SENSOR

The evanescent wave's vector along the horizontal axis can be represented as follows.

$$K_{x} = \frac{2\pi}{\lambda} n_{p} \sin \theta_{i} \tag{1}$$

where n_p is the prism RI, λ is the wavelength of the light source and θ_i in the incidence angle at prism/metal interface

Figure 1: Conventional SPR sensor



When there is an oscillation of surface charges on the gold layer, the surface plasmon wave vector K_{sp} [8] can be expressed as

$$K_{sp} = \frac{2\pi}{\lambda} \sqrt{\frac{n_{silver}^2 n_{sample}^2}{n_{silver}^2 + n_{sample}^2}},$$
 (2)

Here, n_{silver} and n_{sample} are refractive indices of the silver and dielectric sample respectively. Therefore, (2) show how the horizontal wave vector of metal oscillating electron charges is impacted by changes in refractive index. For the plasmons to resonate and produce the resonance angle, the two wave vectors must coincide.

$$\theta_{spr} = \sin^{-1} \left[\frac{1}{n_p} \sqrt{\frac{n_{silver}^2 n_{sample}^2}{\left(n_{sillver}^2 + n_{sample}^2\right)}} \right]$$

(3)

SPR sensor design considerations, mathematical models' performance measurement parameters

SPR sensor design considerations

Figure 2 illustrates a six-layers structure of the SPR when the p-polarized light with wavelength λ incident on the prism. To prevent mechanical translation or movement, the collimated source is routed through the Powell lens, which diverges the laser beam so that the beam incident on the prism-ZnO contact covers the RI measurement range of 1.3317 to 1.34. The redesigned structure of the SPR sensor is proposed by stacking the

layers of different materials along the z-axis. This work's CaF₂ prism has a refractive index of 1.4329, which is lower than BK7 and FK51a's refractive indexes of 1.5151 and 1.4853, respectively. It has been demonstrated that SPR sensors that use prisms with low RI have higher sensitivity. The RI of CaF₂ is calculated using [31]

$$n_{CaF_2} = \left(1 + \frac{0.567588\lambda^2}{\lambda^2 - 0.050263^2} + \frac{0.4710914\lambda^2}{\lambda^2 - 0.10039^2} + \frac{3.8484723\lambda^2}{\lambda^2 - 34.64904^2}\right)$$
(4)

The stacked layers consist of $CaF_2/ZnO/TiO_2/Silver/BaTiO_3/Chitosan/sensing$ sample as shown in the **Figure 2**. The thickness of ZnO, TiO₂, silver, BaTiO₃ and Chitosan is d_2 , d_3 , d_4 , d_5 and d_6 and their corresponding refractive index shown in **Table 1**. The reflected beam from prism – ZnO interface is reflected to other side of the prism. The refractive index of ZnO can be computed as [32]

$$n_{ZnO} = \sqrt{\left(2.81418 + \frac{0.87968\lambda^2}{\lambda^2 - 0.3042^2} - 0.00711\lambda^2\right)}$$
(5)

Therefore, the reflected intensity can be captured by either a charge couple device (CCD) or complementary metal-oxide (CMOS) sensor. From composite structure **Figure 2** in which an angle θ_i is higher than a critical angle this ensures total internal reflection occurs hence the evanescent wave is generated at prism/ZnO interface which responsible for detecting changes in the sample at chitosan/sample interface. The beam used is diverged beam by using Powell lens

which cover entire measurand (1.3317 - 1.34) without any mechanical scanning.

Figure 2: Proposed of a six-layers SPR sensor.

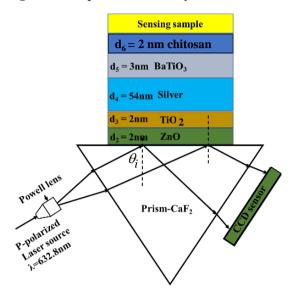


Table 1 shows optical properties material used in the proposed SPR sensor as per provided references.

Table 1: Optical and physical parameters used in the designing of the proposed SPR sensor.

Number of layers	Used material	The refractive index of the material at 632.8 nm	Thickness (nm)	References	
1	Prism (CaF ₂)	1.4329	-	[31]	
2	ZnO	1.98	$d_2 = 2$	[32]	
3	TiO_2	2.5837	$d_3 = 2$	[33]	
4	Silver	0.05626+4.2776i	$d_4 = 54$	[34]	
5	BaTiO ₃	2.4042	$d_5 = 3$	[35]	
6	Chitosan	1.54+0.015i	$d_6 = 2$	[36]	
7	Analyte	1.3317 - 1.34	-		

SPR sensor mathematical model for multilayers

The SPR reflectivity curve was calculated in this study using the matrix method [2] on MATLAB software. According to the boundary condition, the electric and magnetic fields, U_1 and V_1 , at the first layer's boundary are related to those U_6 and V_6 at the last layer's interface, respectively, by

$$\begin{bmatrix} U_1 \\ V_1 \end{bmatrix} = M \begin{bmatrix} U_6 \\ V_6 \end{bmatrix}, \tag{6}$$

where M is the matrix representing the wave propagation response through different layers by

$$M_{ij} = \prod_{k=2}^{6} \begin{pmatrix} \cos \beta_k & -(i/q_k) \sin \beta_k \\ -iq_k \sin \beta_k & \cos \beta_k \end{pmatrix}_{ij}$$
$$= \begin{pmatrix} M_{11} & M_{12} \\ M_{21} & M_{22} \end{pmatrix}. \tag{7}$$

Here, the factor q_k is defined as

$$q_{k} = \frac{\left(n_{k}^{2} - \left(n_{p} \sin \beta_{i}\right)^{2}\right)^{1/2}}{n_{k}^{2}},$$
(7)

while β_k is the phase shift of k^{th} layer with a thickness d_k

$$\beta_k = \frac{2\pi d_k}{\lambda} \left(n_k^2 - n_p^2 \sin^2 \theta_i \right)^{1/2} \tag{8}$$

As the result of the wave propagation through multilayer media, the total reflection coefficient for the transverse magnetic field is expressed by

$$r^{TM} = \frac{\left(M_{11} + M_{12}q_{sample}\right)q_{1} - \left(M_{21} + M_{22}q_{sample}\right)}{\left(M_{11} + M_{12}q_{sample}\right)q_{1} + \left(M_{21} + M_{22}q_{sample}\right)}$$
(9) with

$$q_{sample} = \frac{\left(n_{sample}^2 - \left(n_p \sin \theta_i\right)^2\right)^{1/2}}{n_{sample}^2} \tag{10}$$

Finally, the reflectivity power can be calculated as

$$R = \left| r^{TM} \right|^2. \tag{11}$$

SPR sensor measurement parameters

The SPR sensor measurement parameters were extracted from the resonance curve, in which sensitivity, detection accuracy and quality factor.

Sensitivity

Sensitivity of the SPR biosensor is measured as change in resonance position in the reflectance curve ($\Delta\theta res$) corresponding to the small change in the refractive index (Δns) of the analyte. The sensitivity is given as[37, 38]

$$S = \frac{\Delta \theta_{res}}{\Delta n_s} \tag{12}$$

Detection Accuracy

DA is inversely proportional to the full width at half maxima (FWHM) of the reflectance curve and is given by[37, 38],

$$DA = \frac{1}{FWHM} \tag{13}$$

Quality Factor (QF)

QF is defined as the ratio of S to the FWHM [37-39], and has unit of RIU⁻¹

$$Q = \frac{S}{FWHM} \tag{14}$$

Limit of detection (LOD)

LOD is the minimum value in which CCD sensor can detect the change of the sample. This parameter depends on angular resolution and the sensitivity of the SPR sensor. Consider the theoretical use imbedded linear array

TCD1304AP CCD number of pixel $\binom{N_{pixel}}{of}$ of 3648 pixels. The controller for the CCD sensor is STM32F401RE microcontroller which has 12-bit ADC converter.

$$LOD(\sigma_{RI}) = \frac{\sigma_{s}}{Sensitivity}$$
(15)

Where σ_s is the angular resolution of the SPR sensor. σ_s depends on the pixel range covered by resonance angle and the ADC conversion resolution $1/(2^n-1)$ is multiplied by the resolution of the CCD. The resolution of the CCD sensor is the minimum angle at which CCD sensor can track changes in one pixel is computed as

$$\sigma_{s} = \frac{\Delta \theta_{res}}{N_{pixel}} \times \frac{1pixel}{\left(2^{n} - 1\right)}$$
(16)

 N_{pixel} is the total number of pixels of CCD sensor and n is the number of bit of ADC converter

Sensor merit

Sensor merit is the parameter which relates the broadening and minimum dip of the SPR reflectivity curve. The broadening of SPR reflectance dip, is quantified by combining the effect of the resolution of the sensor and FWHM and shape of resonance damping in the

neighborhood of the minimum. Therefore, sensor merit is expressed as [40-43],

$$SM = \frac{R_{\text{max}} - R_{\text{min}}}{FWHM} \tag{17}$$

Combined sensitivity factor

Combined sensitivity factor (CSF), is the product of the sensitivity sensor merit (SM) in this work [41, 42]. In [43], it was shown that the aforementioned CSF is inversely proportional to the refractive index resolution and can be written Finally, the CSF is given by S multiplied by SM [40, 43] where the unit is RIU⁻¹ and is expressed as:

$$CSF = S \times SM \tag{18}$$

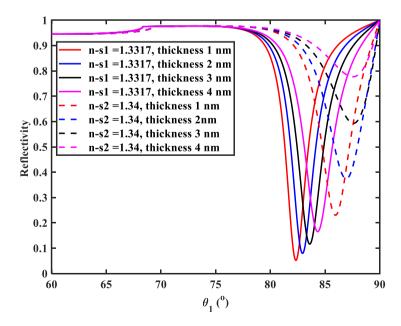
The performance parameters indicate how good the performance of the SPR biosensor is. Note that the behavior of the performance parameters is analyzed with respect to the variation of the chemical potential of graphene

RESULTS AND DISCUSSIONS

Figure 3 depicts reflectance curves for an SPR sensor plotted against incidence angle for various

refractive indices (n-s1 = 1.3317 and n-s2 = 1.34) and chitosan thicknesses (1, 2, 3, and 4 nm). The solid lines reflect the case where n-s1 = 1.3317, whereas the dashed lines represent n-s2 = 1.34. As the refractive index grows from 1.3317 to 1.34, the reflectivity dip goes towards higher angles, suggesting a shift in the resonance angle caused by the change in refractive index. [44, 45]. As the thickness increases from 1 nm to 4 nm for both refractive indices, the resonance dip becomes shallower, while the reflectivity curve broadens and shifts slightly. Thicker layers provide broader reflectivity minima, which affects resolution (detection accuracy). All curves show a characteristic drop in reflectivity, corresponding to the resonance state where SPR occurs. This dip is more evident for thinner layers and gets less sharp as the thickness grows, showing a trade-off between detection accuracy and layer thickness [46]. The plot highlights the importance of layer thickness and refractive index in determining the performance of an SPR sensor, with significant implications for the sensitivity and resolution of detection [45].

Figure 3: Shows the SPR reflectivity with variation of the chitosan thickness by 1 nm, 2 nm, 3 nm, and 4 nm on

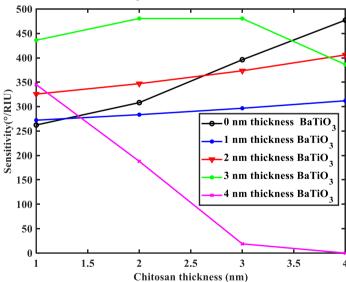


The Figure 4 shows the sensitivity of a SPR sensor as a function of the chitosan layer thickness for different thicknesses of a BaTiO₃ layer (1 nm, 2 nm, 3 nm, and 4 nm). The **Figure 4** shows that when the thickness of the chitosan increases the sensitivity increases only when thickness of BaTiO₃ is less than 3 nm. As it is clearly shown that when the chitosan thickness of increase from 1 nm to 2 nm the sensitivity also increases from 436.02 to 480.48 \(^{\text{RIU}}\), which implies that the sensitivity improved by 10.20 %. However, when the chitosan thickness increases to 3 nm the sensitivity slightly dropped to 480.36 RIU, this signify the optimum thickness of chitosan is 2 nm in which high highest sensitivity is achieved. For 1 nm thickness of BaTiO₃ (black line) the sensitivity remains fairly constant around 300 °/RIU as the chitosan thickness increases from 1 to 4 nm. For 2 nm thickness of BaTiO₃ (red line), the sensitivity gradually increases from about 350 °/RIU to around 400 °/RIU as the chitosan thickness increases. This suggests that an intermediate chitosan thickness, in conjunction

with a moderate BaTiO₃ layer, provides the best sensitivity for the SPR sensor.

For 4 nm thickness of BaTiO₃ (purple line) the sensitivity sharply decreases from around 350 °/RIU at 1 nm of chitosan to almost 0 °/RIU at the range 3 to 4 nm of chitosan, indicating a rapid loss of sensor sensitivity with increasing chitosan thickness for this configuration due to the damping effect of the chitosan as it becomes thicker [46, 47]. This suggests that when the BaTiO₃ layer becomes too thick, the sensor's ability to detect changes in refractive index is diminished, possibly due to the reduced interaction of the evanescent field with the sensing medium [46, 47]. The impact of chitosan thickness in most configurations (except for the 4 nm BaTiO₃ increasing the chitosan thickness enhances the sensitivity, up to a certain limit. This indicates that the chitosan layer plays a significant role in enhancing the sensor's plasmonic response, but beyond a certain thickness, its benefit may diminish or even become detrimental, as seen in the 4 nm BaTiO₃.

Figure 4: Shows the effects of increasing the thickness of the BaTiO₃ layer by 1 nm, 2 nm, 3 nm, and 4 nm on sensitivity



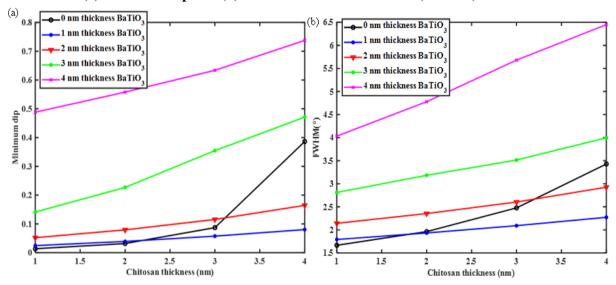
The **Figure 5** shows the minimum dip and FWHM of a SPR sensor as a function of the chitosan layer thickness for different thicknesses of a BaTiO₃ layer (1 nm, 2 nm, 3 nm, and 4 nm). The **Figure 5(a)** shows that when the thickness of the chitosan

and $BaTiO_3$ increases the minimum dip is getting shallow and shallow decreases only when thickness of $BaTiO_3$ is less than 3 nm. The similar was followed by FWHM as shown in **Figure 5(b)**. The broader FWHM leading to poorer detection

accuracy for the sample which are very close to each other in terms of refractive index. In which at the thickness of 2 nm the FWHM is around 3.18°. This yielded as the results of decreasing the evanescent field strength hence reduction in the penetration depth to the measured sample. When the thickness of the chitosan is 2 nm the minimum

dip is around 0.22 which implies that optical coupling efficiency is around 78 % which is acceptable value where detection of the minimum dip is possible. Also, at that thickness the maximum sensitivity of 480.4819 is achieved (see **Figure 4**).

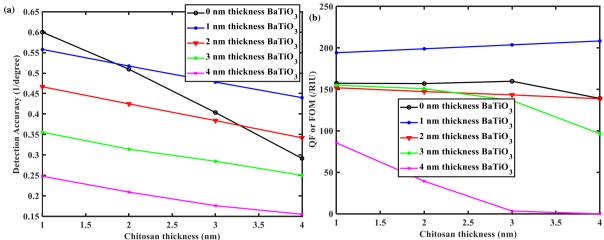
Figure 5: Shows the effects of increasing the thickness of the BaTiO₃ layer by 1 nm, 2 nm, 3 nm, and 4 nm on (a) Minimum dip and (b) full width at half maximum (FWHM)



The Figure 6 shows the detection accuracy and quality factor (QF) of a SPR sensor as a function of the chitosan layer thickness for different thicknesses of a BaTiO₃ layer (1 nm, 2 nm, 3 nm, and 4 nm). The variation trend for DA and QF is similar because DA is proportional to QF, since $DA = QF \times \Delta n_s$ in which Δn_s is constant for which is 0.0083 (1.34 - 1.3317). When the thickness of BaTiO₃ is less than 3 nm, it is observed that as the chitosan increases up to 3 nm the both DA and QF increases. However, beyond 3 nm of the chitosan and when the thickness of BaTiO₃ is more than 3 nm it clearly seen that the both DA and QF decreases. The Figure 6(a) shows that when the thickness of the chitosan increases the detection accuracy (DA) decreases since DA is inversely proportional to FWHM while the quality factor (QF) is decreases which the thickness of BaTiO₃ is higher than 2 nm. While

when the thickness of BaTiO₃ is below 2 nm the QF increase with the increase of the thickness of chitosan. When the thickness of BaTiO₃ is more than 3 nm the evanescent field strength penetrated into the sample is reduced as results of the thickness. But when there is no it is seen that the detection accuracy and QF is decreases this implies the proper interfacial bonding between the chitosan and BaTiO3 layers plays great role in determining detection accuracy and QF. When the thickness of BaTiO₃ is 1 nm it is observed that the both detection accuracy and QF increases which signifies the right proportion for bonding between the chitosan and BaTiO₃ adversely the sensitivity is low for thickness. In comparison with other related work the results for DA of our proposed sensor is higher than those published recently [48] [49] [50] and QF (150.88 RIU⁻¹) is higher than [51] which is 118.4 RIU⁻¹ and [52] which is 88.84 RIU⁻¹.

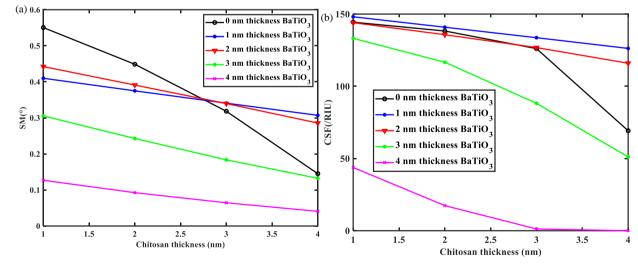
Figure 6: Shows the effects of increasing the thickness of the BaTiO₃ layer by 1 nm, 2 nm, 3 nm, and 4 nm on (a) the detection accuracy (DA) and (b) quality factor (QF).



The **Figure 7** shows the SM and CSF of a SPR sensor as a function of the chitosan layer thickness for different thicknesses of a BaTiO₃ layer (1 nm, 2 nm, 3 nm, and 4 nm). The variation trend for SM varies in inversely proportional FWHM and proportional to power photon transfer efficiency $\left(R_{\text{max}}-R_{\text{min}}\right)$, which implies that the dipper the minimum the higher SM, while the narrower the FWHM the higher SM. From **Figure 7** is clearly seen that as the thickness of chitosan increases the lower the value of SM and CSF.CSF is

proportional to the product of sensitivity and SM, thus the higher SM and S the better CSF. When the thickness of BaTiO₃ is less than 3 nm, it is observed that as the chitosan increases up to 3 nm the both DA and QF increases. Decrease in both SM and CSF as the chitosan increases caused by reduction of the penetration depth of evanescent was as the chitosan becomes thicker. Also, it is observed that as BaTiO₃ thickness increases SM and CSF decreases because of the reduction of evanescent field strength.

Figure 7 Shows the effects of increasing the thickness of the BaTiO₃ layer by 1 nm, 2 nm, 3 nm, and 4 nm on (a) Sensor merit (SM) and (b) Combined sensitivity factor (CSF)



In general, the optimum performance of the SPR sensor was obtained when the thickness of BaTiO₃ is 3 nm while the thickness of chitosan is 2 nm. The overall results performance can be

summarized as shown in the **Table 2.** The theoretical resolution of the proposed SPR sensor is computed by considering the range of the

resonance angle is 3.988° which means from 82.939 to 86.927° as shown in **Figure 3.**

$$\sigma_s = \frac{3.988^{\circ}}{3648 \, pixel} \times \frac{1 \, pixel}{4095} = \frac{3.988^{\circ}}{14938560} = 2.6696 \times 10^{-7\circ}$$
(19a)

$$LOD(\sigma_{RI}) = \frac{2.6696 \times 10^{-7}}{480.482^{\circ} / RIU} = 5.5561 \times 10^{-10} RIU$$
(19b)

Therefore, the limit of detection LOD of the proposed SPR sensor has improved from $8.3790\times10^{-10}RIU$ [25] to $5.5561\times10^{-10}RIU$. Thus, the limit of detection LOD of the proposed SPR sensor has improved by 33.69 %.

Table 2: An overview of the optimal performance parameters of the proposed SPR sensor.

Parameters	S	FWHM	Minimum	DA	SM	CSF	QF
	(/ RIU)	(degree)	dip	(1/degree)	(1/degree)	(1/RIU)	(1/RIU)
1	480.482	3.18	0.227	1.25	0.24	116.67	150.88

Comparison of the Proposed SPR Sensor with Recent Publications Pertaining to BaTiO₃

Table 2 contrasts the effectiveness of the proposed SPR sensors with that of recently published research. The findings demonstrate that the suggested SPR sensor surpasses a recently published study on sensitivity (S), detection accuracy (DA), and figure of merits (FOM), all of which are essential attributes for detecting analytes with refractive index values ranging from

1.3317 to 1.34. The sensitivity of the proposed SPR sensor is improved by the factor of 1.2 when chitosan of 2 nm was added on top of BaTiO₃ [25]. Meanwhile the DA and FOM of the proposed SPR sensor is lowered by the factor of 1.47 and 1.2, respectively, compared to the to the SPR structure without chitosan layer [25]. However, in this application the most important parameters are sensitivity and limit of detection of the SPR sensor which outperform the recently published paper.

Table 2: A comparative study of the proposed SPR sensor and most recently published research work in terms of sensitivity (S), detection accuracy (DA) and Quality factor (QF).

Reference	Reported Year	Structure	Sample range	S (deg./RIU)	DA	FOM or QF (deg./RIU)
[24]	2022	CaF ₂ /BaTiO ₃ /Blue Phosphorus/WS ₂	1.365 - 1.388	413.7 °	-	-
[49]	2023	BK7/Ag/BaTiO ₃ /Ag/Graphene	1.33 - 1.39	220.83	0.819	102.45
[53]	2023	Ag/BaTiO ₃ / Ag/ WS ₂	1.368 -1.392	253.5	0.576	146.02
[54]	2023	$Ag/BaTiO_3/Ag_1/MoS_2$	1.368 -1.392	290.71	0.346	100.59
[51]	2023	BK7/Ag/ BaTiO ₃ /Blue P/MoS ₂ /WS ₂	1.33 -1.335	326	-	-
[48]	2023	CaF ₂ /Copper /BaTiO ₃ /Graphene	1.33 - 1.34	219	0.47	102.93
[55]	2023	BK ₇ /Ag/BaTiO ₃ /Graphene	TB	300. 83	0.249	74.91
[50]	2024	BK7/Ag/ BaTiO ₃ / MoSe ₂	1.371, 1.41,1.378	247.813	0.17437	43.211
[25]	2024	CaF ₂ /TiO ₂ /Ag/BaTiO ₃	1.3317 -1.34	399.45	0.4606	183.98
	Proposed	CaF ₂ /ZnO/TiO ₂ /Ag/BaTiO ₃	1.3317 -1.34	480.48	0.314	150.88

CONCLUSION

The designed SPR sensor, enhanced by a chitosan layer above the BaTiO₃, has demonstrated improvements in sensitivity and detection accuracy by factors of 16.67 % and 33.69 %, respectively. The enhancement resulted from the precise optimization of the thickness of BaTiO₃ and chitosan, with optimal sensitivity and limit of detection (LOD) achieved at a BaTiO₃ thickness of 3 nm and a chitosan thickness of 2 nm. The suggested SPR sensor can detect minute variations in the measuring sample within the range of 1.3317 to 1.34 without mechanical rotation. The high sensitivity and low limit of detection (LOD) of 480.45°/RIU $5.5561 \times 10^{-10} RIU$, respectively, results in the ability to detect even little variations in measuring samples.

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