

Characterization of PVA-Enzyme Coated Indicator Electrodes GA coated again with PVC-KTpClPB-o-NPOE SEM-EDS, FTIR and XRD analysis

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Submission date: 09-Jan-2022 09:06PM (UTC-0600)

Submission ID: 1739336954

File name: 1265-Article_Text-6198-1-10-20220108.pdf (416.9K)

Word count: 3783

Character count: 19538



Characterization of PVA-Enzyme Coated Indicator Electrodes GA coated again with PVC-KTpCIPB-o-NPOE SEM-EDS, FTIR and XRD analysis

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DOI: [10.29303/jppipa.v8i1.1265](https://doi.org/10.29303/jppipa.v8i1.1265)

Article Info

Received: November 30, 2021

Revised: December 27, 2021

Accepted: January 8, 2022

Published: January 31, 2022

Abstract: This study aims to characterize the tungsten-urea analyte indicator electrode. The method used is biosensor potentiometry with urease enzyme immobilization technique. This indicator electrode was coated with PVA-enzyme coated with glutaraldehyde (GA) 2.9% coated with PVC-KTpCIPB- o-NPOE with o-NPOE variation of 61% and 66%. Characterization of coated indicator electrodes using SEM-EDS, FTIR and XRD analysis. A1-4 61% indicator electrode sample coated PVA-enzyme 1x coated with glutaraldehyde (GA) 2.9% 1x coated PVC-KTpCIPB- o-NPOE 1x, with o-NPOE 61%, A3-4 61% indicator electrode sample coated PVA-enzyme 3x coated with glutaraldehyde (GA) 2.9% 1x coated PVC-KTpCIPB- o-NPOE 1x, with o-NPOE 61%. Likewise, the reasoning of samples A1-4 66% and A3-4 66%. There are four indicator electrodes made with the notation A1-4 61%, A1-4 66%, A3-4 61% and A3-4 66%. The best results were obtained at the indicator electrode sample A1-4 61%, contributing to the urea sensor of the potentiometer cell

Keywords: Tungsten indicator electrodes; GA and o-NPOE; SEM-EDS; FTIR and XRD; Biosensor Potentiometry; Immobilization.

Citation: Hakim, S.A. (2022). Characterization of PVA-Enzyme Coated Indicator Electrodes GA coated again with PVC-KTpCIPB-o-NPOE SEM-EDS, FTIR and XRD analysis. *Jurnal Penelitian Pendidikan IPA*, 8(1), 103-108. <https://doi.org/10.29303/jppipa.v8i1.1265>

Introduction

Research has been carried out on indicator electrodes coated with PVA-enzyme coated with PVC-KTpCIPB with a detection range of 10-5-10-4M (Hakim, 2019). This electrode analysis has been characterized using SEM-EDS (Hakim, 2018). According to (Rahman, 2008) the range is 5.10-7-10-2M. On this basis the researchers increased the detection range with GA and o-NPOE. Experimental research on indicator electrodes B1-4 and B3-4 has been carried out. B1-4 is an indicator electrode coated with PVA-Enzyme 1x coated GA 1x coated again with PVC-KTpCIPB 1x. B3-4 is an indicator electrode coated with PVA-Enzyme 3x coated GA 1x coated again with PVC-KTpCIPB. B1-4 and B3-4 analyzed by SEM-EDS can be seen in Table 1.

Research experiments have been carried out with indicator electrodes A1-4 61%, A1-4 66%, A3-4 61% and A3-4%. The results can be seen in Table 2. Indicator electrodes A1-4 61% were indicator electrodes coated with PVA-Enzyme 1x coated with GA 1x coated again with PVC-KTpCIPB-o-NPOE 1x with o-NPOE 61%. Indicator electrodes A1-4 66% are indicator electrodes coated with PVA-Enzyme 1x coated with GA 1x coated again with PVC-KTpCIPB-o-NPOE 1x with o-NPOE 66%. Indicator electrode A3-4 61% is indicator electrode coated with PVA-Enzyme 3x coated GA 1x coated again with PVC-KTpCIPB-o-NPOE 1x with o-NPOE 61%. Indicator electrode A3-4 66% is indicator electrode coated with PVA-Enzyme 3x coated GA 1x coated again with PVC-KTpCIPB-o-NPOE 1x with o-NPOE 66%.

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To coat the PVA indicator electrode, enzyme, PVC-KTpCIPB, o-NPOE in solution form. PVA 0.0350 g dissolved in 10 mL of hot water to cool in a glass tube. PVC 0.0350 g + KTpCIPB 0.0500 g dissolved in 10 mL THF in a glass tube and covered with a plastic cover/aluminum foil. 2.9% GA solution is 0.29 g of GA in 10 mL of distilled water. The o-NPOE solution consisted of 5 mL, 61% (0.0647 gr = 0.0622 mL) and 66% (0.0700 gr = 0.0673 mL).

Based on Table 1. the weight percent of tungsten sample B1-4 is 73.68%, B3-4 is 76.24%. Likewise, Table 2 weight percent of tungsten samples A1-4 61% is 91.91%, A3-4 61% is 95.98%, A1-4 66% is 84.93% and A3-4 66% is 79.81%. There was an increase in weight percent of tungsten with the addition of o-NPOE 61% and 66% through EDS analysis. The morphology of the indicator electrodes A1-4 61%, A3-4 61%, A1-4 66% and A3-4 66% can be seen in Figure 1. The best four samples have not been selected, for this reason, FTIR and XRD analysis are carried out.

An ion selective electrode (ISE) has been used (Huang, 2014) with 2-nitrophenyloctyl ether (NPOE) and mixed in poly(vinyl chloride) (PVC). The obtained sensitivity is 19.7 mV/decade with a range of 5.10-7-10-2M. According to (Rahman, 2008) the optimum sensitivity is 59 mV/decade, for a reaction to be reversible (in practice the difference is typically between 70-100 mV).

The immobilization technique in the construction of the biosensor involves the urease enzyme and a polymer consisting of PVC for the urea bioser (Ulianas, Heng and Ahmad, 2011). Poly(vinylchloride) (PVC) is a polymer matrix commonly used in ISE (Mir, Lugo, Tahirbegi and Josep Samitier, 2014).

The permissible limit of PVC on the membrane electrode has the composition of PVC: o-NPOE: DPA: NaTPB as 33:61:3:3 (w/w) as the optimum composition with the closest Nernstian response (Kaur, Chhibber and Mittal, 2017). The composition according to (Badakhshan, 2019) is 30:65.5:0.5 mg of PVC: o-NPOE: KTpCIPB, as a composition of PVC, plasticizer and lipophilic additive. Ionophores (I) 1.10% (w/w), lipophilic salt (KTpCIPB) 0.25% (w/w), plasticizer (o-NPOE) 65.65% (w/w), and PVC 33.00 % (w/w) dissolved in tetrahydrofuran (Lenar, 2020; Hasan, 2019); 0.05 wt% KTpCIPB as membrane additive, and 1 wt% ionophore, 33 wt% PVC and 66 wt% plasticizer (o-NPOE) (Rizk, et al, 2009); 62.5 mg PVC, 130 mg o-NPOE, 1% by weight s prepared containing 62.5 mg PVC, 130 mg o-NPOE, 1% by weight ionophore and 80 to 140 mol% KTpCIPB (relative to the ionophore). The membrane components were dissolved in 1.5 mL of THF (Pomečko, R., Luboch, E., and Jeszke, M., 2021).

Method

The materials used in the manufacture of the indicator electrodes were tungsten with a diameter of 1.0 mm 99.99% Aldrich 267562, PVA [-CH₂CHOH-]_n, enzyme EC 3.5.1.5 (Urease) U4002, 50 - 100 ku type ix, glutaraldehyde (GA); PVC (CH₂CHCl)_n, potassium tetrakis (4-chlorophenyl) borate (KTCIPB), o-nitrophenyloctylether (o-NPOE), Tetrahydrofuran C₄H₈O, all from Sigma-Aldrich.

The indicator electrodes were made by using the potentiometric biosensor of the urease enzyme immobilization technique in two notations B1-4 2.9% GA and B3-4 2.9% GA. Then analyzed by SEM-EDS. From the two samples above, it was continued to manufacture four indicator electrodes with notation A1-4 61%, A3-4 61% and A1-4 66%, A3-4 66%. Then the four were analyzed by SEM-EDS, FTIR and XRD.

Result and Discussion

EDS and SEM analysis results

Analysis of indicator electrodes B1-4 and B3-4 using EDS and SEM has not been able to ensure the best indicator electrode samples. For strong conclusions, further FTIR and XRD analyzes are needed.

Table 1. Weight percent, atomic and atomic ratio of Tungsten without o-NPOE.

Indicator electrode (%)	Element	Weight %	Atomic %	Ratio Atomic
B1-4	w	73.68	80.30	2.41
B3-4	w	76.24	87.78	2.49

After UV-Vis and SEM-EDS analysis, samples of indicator electrodes B1-4 and B3-4 without o-NPOE were further analyzed by cell potentiometer, to characterize their sensitivity and detection range. The best results were obtained at the indicator electrode B1-4.

Table 2. Weight percent, atomic and atomic ratio of Tungsten Using o-NPOE.

Indicator electrode (%)	Element	Weight %	Atomic %	Ratio Atomic
A1-4 61	w	91.91	91.80	2.93
A3-4 61	w	95.98	92.78	3.04
A1-4 66	w	84.93	86.27	2.74
A3-4 66	w	79.81	87.45	2.59

However, samples B1-4 and B3-4 continued to the next step in the manufacture of indicator electrodes A1-4 61%, A3-4 61%, A1-4 66% and A3-4 66%, the

results of the SEM analysis can be seen in Figure 1 and EDS Table 1.

SEM-EDS analysis for indicator electrodes having o-NPOE can be seen in table 2. The weight percent of tungsten A1-4 61% is 91.91% and the weight percent of tungsten A3-4 61% is 95.98%. The weight percent of tungsten A1-4 66% was 84.93% and the weight percent of tungsten A3-4 66% was 79.81%. The largest weight percentage of tungsten was at the best 61% o-NPOE content at the indicator electrodes A1-4 61% and A3-4 61%. A good SEM morphological analysis of the membrane was also found at the indicator electrodes A1-4 61% and A3-4 61%, the pores slightly increased the weight percent of tungsten. At the indicator electrodes A1-4 66% and A3-4 66% the number of pores reduces the weight percent of tungsten.

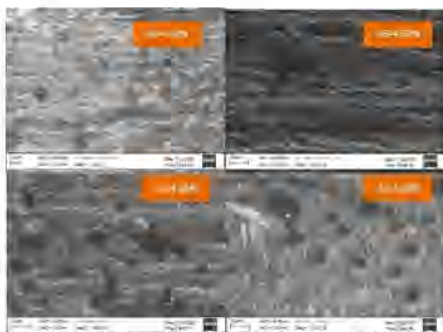


Figure 1. Morphology of sample size 10 m voltage 20kV magnification 600 x (A1-4 61%) and 100 x (A3-4 61%, A1-4 66% and A3-4 66%).

Morphology of indicator electrodes A1-4 61% EDS analysis peak intensity 4.3 keV weight percent tungsten 91.91%; morphology of indicator electrodes A1-4 66% EDS analysis peak intensity 2.5 keV weight percent tungsten 84.93%; morphology of indicator electrode A3-4 61% EDS analysis peak intensity 4.3 keV weight percent tungsten 95.98%; morphology indicator electrode A3-4 66% EDS analysis peak intensity 4.5 keV weight percent tungsten 79.81%. This confirmed the high purity of the prepared tungsten (Shawky, 2021).

FTIR Characterization Results using o-NPOE

The characterization of indicator electrodes A1-4 61%, A3-4 61%, A1-4 66% and A3-4 66% can be seen in Figure 2. The FTIR spectrum pattern for A1-4 61% indicator electrodes has a transmittance percentage between 85-100. The spectrum pattern of A1-4 is 61% similar to (Badakhshan, 2019), the transmittance percent is between 90 -100. There is an increase in %Transmittance of the FTIR spectrum pattern by an increase in o-NPOE. Meanwhile, A1-4 66% and A3-4

66% decreased the transmittance of the FTIR spectrum pattern by increasing o-NPOE.

(Elbehery, 2019) plasticizer (either DOP or o-NPOE, 133 mg), and PVC (66 mg) in THF (3 mL), (Hassan, 2019) The composition of the membrane was 32.0 wt.% PVC, 65.0 wt. % plasticizer, and 3.0 wt.% ionophore. (Badakhshan, 2019) 30.0 mg of PVC, 65.5 mg of o-NPOE, and 0.5 mg of KTpCIPB were dissolved in 3 ml of THF to make a polymer membrane. (Kaur, Chhibber and Mittal, 2017, 2017) The variation of plasticizers, lipophilic salts, and PVC within the permissible limits makes the membrane electrodes have a PVC composition: o-NPOE: DPA: NaTPB as 33:61:3:3 (w/w). (Amr, 2021) 2 mg potassium tetrakis (4-chlorophenyl) borate, 63.0 mg o-NPOE, and 34.0 mg PVC; all in 2-mL THF. (Alarfaj, N.A. and El-Tohamy, M. F., 2020) PVC 190 mg, plasticizer (o-NPOE, 0.35 mL) in 5 mL THF.

1.66.5 mg PVC, 127 mg plasticizer (o-NPOE) and 2.2 mg potassium tetrakis (3,5-bis (trifluoromethyl) phenyl) borate (KTFPB) were thoroughly mixed in a 3 cm diameter petri dish and dissolved in 3 mL (Kamel, et al, 2020).

Although the dielectric constant = 24 of o-NPOE is the highest value, it does not significantly affect the membrane potentiometric detection limit in the case of drugs (Hassan, 2020).

The observed absorption bands at 745 cm⁻¹ and 1081 cm⁻¹ associated with the C-Cl and B-C stretches, respectively confirmed the presence of the lipophilic additive KTpCIPB. The presence of PVC as an inert membrane matrix was confirmed by the gauche absorption band observed in the 669 cm⁻¹ region (Badakhshan, 2019).

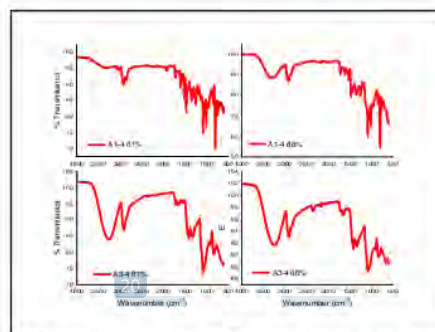


Figure 2. Spectrum pattern % Transmittance A1-4 61%, A1-4 66%, A3-4 61% and A3-4 66%.

The ratio of plasticizer/PVC plays a major role to obtain optimal response. o-NPOE was chosen as a plasticizer because of the good solubility of the membrane components and high dielectric constant. Meanwhile, NaTPB is used as a lipophilic additive

because of its important role in increasing the sensitivity and selectivity of the electrode and reducing anionic interference (Alharthi, S.S., Fallatah, A.M. and Al-Saidi, H.M., 2021). The lipophilic additive in this study was KTpCIPB.

UV-Vis spectroscopy is the most useful and reliable technique for confirming the main characterization of the size, shape and stability of the nanostructures of metal oxides synthesized in liquid suspension. According to the analysis (Shawky, 2021; Al-Mohaimeed, et al, 2021) the increase in the absorption peak with respect to the wavelength has an effect on the increase in % Transmittance of the wave number.

According to (Alharthi, S.S., Fallatah, A.M. and Al-Saidi, H.M., 2021), the increase in % Transmittance with respect to the wavelength of the FTIR, was followed by an increase in the peak intensity with respect to the atomic elements of the EDS. If you observe Figure 2 of the FTIR spectrum pattern, compared to Table 2 of the high peak intensity of EDS, there is only one best sample, namely A1-4 61%. It can be seen in Table 3. the following is the relationship between the composition of o-NPOE in PVC with weight percent and peak intensity height and the pattern of the FTIR spectrum.

Table 3. The relationship between tungsten weight percent, EDS intensity peak height and FTIR spectrum pattern

Sample (%)	Weight percent Wolfram (%)	High Peak intensity (EDS) keV	FTIR Spectrum Pattern
A1-4 61	91.91	4.30	Increase
A1-4 66	84.93	2.50	Decrease
A3-4 61	95.98	4.30	Decrease
A3-4 66	79.81	4.50	Decrease

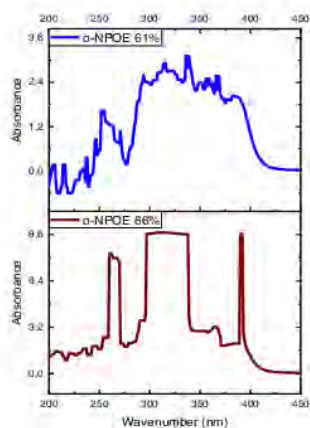


Figure 3. Absorbance spectrum pattern of PVC-KTpCIPB-o-NPOE at o-NPOE 6 and 66%.

Research experiments have been carried out with UV-Vis analysis, the absorbance spectrum pattern of PVC-KTpCIPB-o-NPOE at o-NPOE 61% two absorption peaks at 250 nm and 337 nm; at o-NPOE 66% three absorption peaks of 262, 320 and 390 nm. Meanwhile, the best PVA-Enzyme-GA absorbance spectrum pattern was at GA 2.9%, with a peak absorbance height of 300 nm. There is an increase in the absorbance peak with the addition of the concentration of o-NPOE (Hassan et al., 2020), and an increase in the absorbance of ionopores with the addition of metal salts (Golcs et al., 2018).

XRD Characterization Results using o-NPOE

XRD is an analytical method for estimating and measuring different crystal forms in the tested samples [Shawky, 2021]. (Alarfaj, N.A. and El-Tohamy, M. F., 2020) have used the characterization of UV-Vis, FT-IR, XRD, and EDX spectroscopic methods in determining the calibration curve of the potentiometric system of the o-NPOE modified coated wire membrane sensor. There was a change in the XRD spectrum pattern (Hakim et al, 2021) of KTpCIPB as shown in Figure 4.

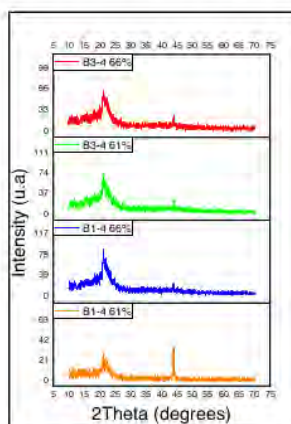


Figure 4. XRD diffraction spectrum patterns A1-4 61%, A1-4 66%, A3-4 61% and A3-4 66%.

(El-Naby, E.H., 2019) CE NaTPB ionophore based polymer membrane sensor with incorporation of anionic additives; potassium tetrakis(p-chlorophenyl)borate (KTpCIPB) produces an excellent "CE NaTPB/KTpCIPB" sensor. High sensitivity with a response slope closer to the ideal Nernstian value and a wider linear range and anionic additives cause a dramatic increase with the o-NPOE plasticiser in the lower limit of detection 5.10⁻⁵ M (with a DOP limit of detection of 7.5.10⁻⁷ M), also improves the interfacial ion exchange kinetics.

(Kaur, Chhibber and Mittal, 2017, 2017), Response time is an important parameter to evaluate

the working efficiency of the electrode, the best content of o-NPOE in PVC is 61%.

Conclusion

Based on the data above, the manufacture of tungsten indicator electrodes with the best membrane modifications at B1-4 and A1-4 66%. B1-4 membrane consists of 1x PVA-E coated GA 1x coated PVC-KTpCIPB 1x. A1-4 66% membrane consists of 1x PVA-E coated GA 1x PVC-KTpCIPBo-NPOE coated 1x.

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