

Supplementary Material

**Efficient sensing of nitrite by $\text{Fe}(\text{bpy})_3^{2+}$ immobilized Nafion
modified electrodes**

Uday Pratap Azad and Vellaichamy Ganesan*

Department of Chemistry, Faculty of Science

Banaras Hindu University, Varanasi-221 005, India.

E-mail: velganesh@yahoo.com and velgan@bhu.ac.in

Fax: + 91-542-2368127; Tel: + 91-542-2307321

Amount of $\text{Fe}(\text{bpy})_3^{2+}$ exchanged/adsorbed on the GC/Nf electrode:

The amount of $\text{Fe}(\text{bpy})_3^{2+}$ adsorbed on the GC/Nf electrode was calculated by measuring the difference in the absorbance of $\text{Fe}(\text{bpy})_3^{2+}$ solution before and after dipping the GC/Nf electrode.

Absorbance of measured volume of $\text{Fe}(\text{bpy})_3^{2+}$ solution before the dipping of GC/Nf electrode is measured. From the absorbance, amount of $\text{Fe}(\text{bpy})_3^{2+}$ (x_1 mol) present in the solution is calculated according to equation 1 (Reference A).

$$x_1 = (\text{Absorbance}/\epsilon l) * (V/1000) \text{ mol} \quad \dots(1)$$

where ‘ ϵ ’ is the molar extinction coefficient ($\text{L} \cdot \text{mol}^{-1} \cdot \text{cm}^{-1}$), ‘ l ’ is the path length (cm) ‘ V ’ is the volume of $\text{Fe}(\text{bpy})_3^{2+}$ solution (mL).

Similarly, after dipping the GC/Nf electrode for approximately 30 min. the absorbance of the solution is measured again and the amount of $\text{Fe}(\text{bpy})_3^{2+}$ (x_2 mole) remaining in the solution is calculated. The difference in amount of $\text{Fe}(\text{bpy})_3^{2+}$ ($(x_1 - x_2)$ mole) is exchanged/adsorbed on the GC/Nf electrode. As one molecule of $\text{Fe}(\text{bpy})_3^{2+}$ will exchange two protons from two $-\text{SO}_3\text{H}$ groups, one mol of $\text{Fe}(\text{bpy})_3^{2+}$ is considered as two equivalents.

Reference A:

D. A. Skoog, E. J. Holler and T. A. Nieman, Principles of Instrumental Analysis, Fifth edition (1998), Saunders College Publishing, Philadelphia, USA.

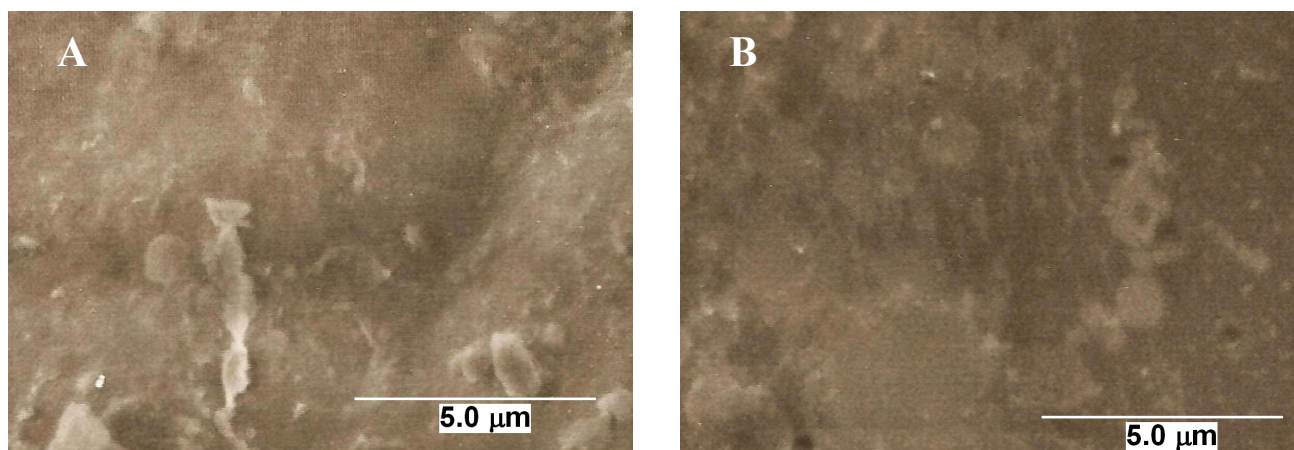


Fig. S1. SEM images of GC/Nf (A) and GC/Nf/Fe(bpy)₃²⁺ (B) films coated on a glass plate.

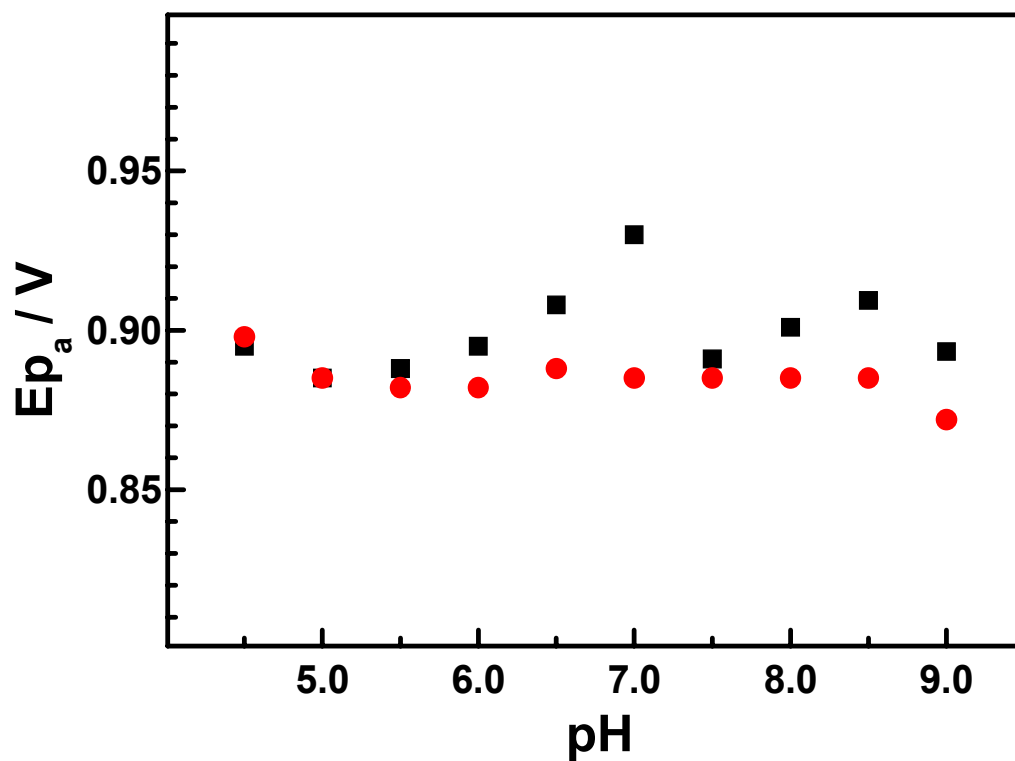
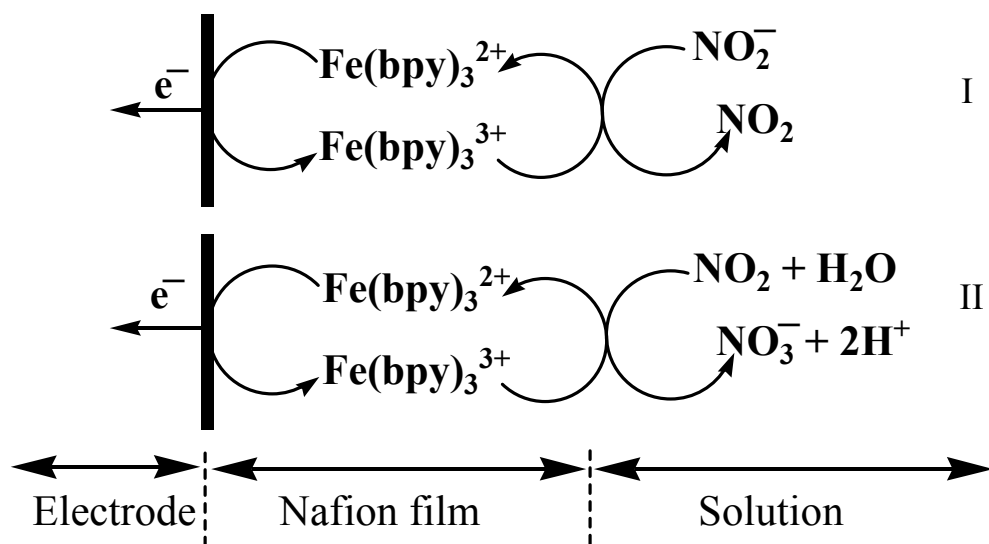


Fig. S2. Effect of pH on the oxidation peak potentials (Ep_a) of $Fe(bpy)_3^{2+}$ (●) and nitrite (■) at the GC/Nf/ $Fe(bpy)_3^{2+}$ electrodes. Note: ‘●’ are obtained in the absence of nitrite. ‘■’ are obtained in the presence of 5.0 mM nitrite.



Scheme S1. Schematic representation of electrocatalytic oxidation of NO_2^- at the GC/Nf/ $\text{Fe}(\text{bpy})_3^{2+}$ electrodes.

Table S1. Influence of interferences on the oxidation peak current of 1.0 mM NO_2^- at GC/Nf/ $\text{Fe}(\text{bpy})_3^{2+}$ electrodes.

Interference	Signal change %
Cl^-	+ 5.7
NO_3^-	+ 3.8
HCO_3^-	- 3.6
CH_3CO_2^-	- 1.2
Cu^{2+}	- 1.1
K^+	+ 3.2
O_2	+ 0.6
Glucose	+ 1.2
Urea	+ 5.5

Table S2. Recovery analysis for nitrite oxidation at GC/Nf/Fe(bpy)₃²⁺ electrodes.

Sample	Nitrite added (mM)	Nitrite found (mM)	Recovery (%)	Relative standard deviation (%)
Tap water	1.0	0.94	94	1.5 (n = 5)
Packaged drinking water (a popular brand)	1.0	0.96	96	0.9 (n = 6)