

Prospects of nanotechnology for detection of water borne pathogen

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In recent scenarios, the development of nanotechnology with novel size and shape dependent properties has shown remarkable potential for fabrication of devices like biosensor for environmental and biomedical applications. These nanostructured materials form an interface with biomolecules, cell-lines, microorganisms etc that creates a channel for fast communication in nanoscale devices that can be utilized for diverse biomedical applications including biosensors, drug delivery, hyperthermia treatment, cell separation etc.¹ In this context, we have synthesized different nanostructured materials including iron oxide, zirconium oxide, nickel oxide and their composites with biopolymers like gelatin and chitosan.²⁻⁵ These functionalized nanostructured materials have been utilized for detection of water borne pathogens.

Water borne pathogen can cause severe infectious disease including diarrhea, gastrointestinal infection and systematic illnesses. Thus, it is important to monitor these pathogens in real samples to minimize these types of diseases. So immunosensing platform has been fabricated using different nanostructure materials and these deposited onto indium-tin-oxide (ITO) coated glass substrate. These materials provide a suitable environment for immobilization of specific biorecognition elements (antibodies) for detection of *Vibrio cholerae* (*Vc*) utilizing electrochemical technique. It has been found that these nanoparticles provide the improved biosensing properties including sensitivity, selectivity and detection range for detection of pathogens. Figure 1 shows the stepwise fabrication of the sandwich electrode based on the CH–NiO/ITO electrode by the functionalization of BSA and Ab-*Vc* onto the CH–NiO surface for the detection of *Vc* concentration.⁴ For the confirmation of interaction of *Vc* onto BSA/ Ab-*Vc*/CH–NiO/ITO, the secondary labelled antibodies (HRP–Ab-*Vc*) are specific to the other epitopes of *Vc*, which do not interact with primary

antibodies. The change in the electrochemical response of the HRP–Ab-*Vc*/ *Vc*/BSA/Ab-*Vc*/CH–NiO/ITO immune-electrode was monitored as a function of H₂O₂ concentration using electrochemical as well as optical techniques. This BSA/Ab-*Vc*/CH–NiO/ ITO immunoelectrode provides an excellent detection range of 20–700 ng mL⁻¹, a low detection limit of 0.108 ng mL⁻¹ and a sensitivity of 0.644 mA ng mL⁻¹ cm⁻². The electrochemical based HRP–Ab-*Vc*/ *Vc*/BSA/Ab-*Vc*/CH–NiO/ ITO sandwich immunoelectrode exhibits a higher sensitivity of 2.95 mA mM mL⁻¹ cm⁻² and a K_m value of 19.96 mM. However, the optical based HRP–Ab-*Vc*/ *Vc*/BSA/Ab-*Vc*/CH–NiO/ ITO sandwich immune-electrode reveals a sensitivity of 0.064 A mM⁻¹ cm⁻² with a low K_m value of 1.0 mM.

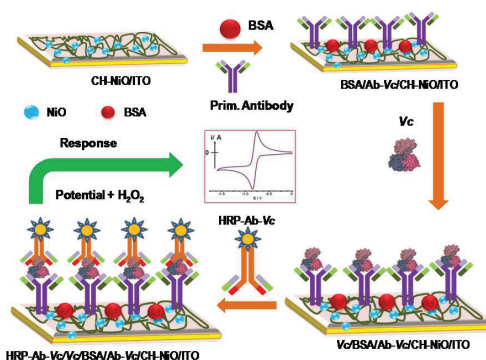


Figure 1: The preparation of the HRP–Ab-*Vc*/ *Vc*/BSA/Ab-*Vc*/CH–NiO/ITO immunoelectrode

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