

A White Light Reflectance Spectroscopy Label-Free Biosensor for the Determination of Fungicide Carbendazim [†]

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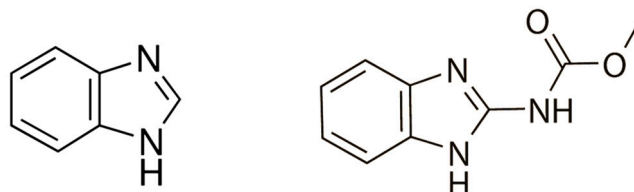
Abstract: Carbendazim is a broad-spectrum benzimidazole-type fungicide effective against fungi that compromise the safety/quality of food products. Despite its potential usefulness, carbendazim constitutes a major environmental pollutant, being hazardous for humans and animals; therefore, reliable determination of carbendazim levels in water, soil, and food samples remains a highly desirable analytical goal. Herein, an optical (white light reflectance spectroscopy, WLRS) label-free biosensor for fast and sensitive determination of carbendazim is presented. The transducer is a SiO₂/Si chip, on which a suitable benzimidazole-conjugate has been immobilized; determination is based on the competitive immunoassay format: A mixture of an in-house developed anti-carbendazim antibody with the calibrators/samples is pumped over the chip, followed by biotinylated secondary antibody and unlabeled streptavidin. The WLRS platform allows for real-time monitoring of biomolecular interactions carried out onto the SiO₂/Si chip by transforming the shift in the reflected interference spectrum caused by the immunoreaction to effective biomolecular adlayer thickness. The sensor is capable of detecting carbendazim levels within 28 min (LoD: 20 ng/mL; intra- and inter-assay CVs: ≤6.9% and ≤9.4%, respectively). Excellent analytical characteristics and short analysis time combined with its small size render the proposed WLRS biosensor ideal for future point-of-need determination of carbendazim in food and environmental samples.

Keywords: white light reflectance spectroscopy; label-free immunosensor; pesticides; carbendazim

1. Introduction

Fungal pathogens cause significant damages to the food crops every year resulting in poor yield, deficient food quality, and huge economic loss; therefore, the use of fungicides gains ground to circumvent these problems [1]. Carbendazim (methyl 1*H*-benzimidazol-2-ylcarbamate) is a low molecular weight (191.19), systemic, broad-spectrum, benzimidazole-type fungicide (Scheme 1), used worldwide as pre- and post-harvest treatment to control fungi that compromise the quality of various vegetables, fruits, cereals, and seeds [2,3]. Despite the unquestionable benefits provided to crop yield,

carbendazim is a major pollutant, which induces acute and late effects on humans and livestock. In this context, carbendazim has been documented to induce infertility, embryotoxicity, teratogenicity, hepatocellular dysfunction, endocrine-disrupting effects, disruption of hematological functions, and mutagenicity [2]. Additionally, the World Health Organization (WHO) has classified carbendazim in group 2B, as a possible human carcinogen [4]. Due to its aforementioned severe toxicities and its persistence in food and the environment, carbendazim has been officially banned in most of European Union, USA, and Australia. However, some countries, such as UK, Portugal, India, China, and Brazil, are still permitting the production and use of carbendazim in various formulations [5], raising a growing concern for the health and safety of humans and animals.



Scheme 1. Chemical structures of benzimidazole core-moiety (**left**) and carbendazim (**right**).

Analytical methodologies that ensure fast and accurate determination of carbendazim are a prerequisite to guarantee protection of public health. Determination of carbendazim is routinely performed by instrumental analysis techniques, mainly high-performance liquid chromatography (HPLC) coupled to mass spectroscopy (MS) or ultraviolet spectroscopy (UV) [6,7]. Alternatively, immunochemical techniques have been developed and used, offering low-cost and short-time analyses, simple assay protocols, and minimum sample pre-treatment, as well as high-throughput sample capacity. In this context, classic enzyme-linked immunosorbent assays (ELISA) and immunochromatographic strips have appeared in the literature [8,9], while the modern technologies of biosensors have lately attracted much attention due to their simplicity, rapidity, portability, and potential of point-of-need application [10].

In this work, a label-free sensor based on White Light Reflectance Spectroscopy (WLRS) is presented for the fast and sensitive determination of carbendazim. The transducer employed is a Si chip with a 1 μm thick thermal SiO_2 on top, which is transformed to a biosensing element through immobilization of a suitable benzimidazole conjugate on the SiO_2 surface. Carbendazim determination is based on the competitive immunoassay format, i.e., a mixture of the antibody with the calibrators or the samples is pumped over the chip surface followed by reaction with biotinylated secondary antibody/streptavidin for signal enhancement. The WLRS biosensing platform allows for the label-free, real-time monitoring of biomolecular interactions carried out onto the SiO_2/Si chip by transforming the shift in the reflected interference spectrum caused by the immunoreaction to effective biomolecular adlayer thickness. The developed sensor is capable of real-time detection of carbendazim with LoD: 20 ng/mL, within 28 min total analysis time.

2. Materials and Methods

2.1. Reagents and Chemicals

Four-inch Si wafers (<100>), purchased from Si-Mat Germany (Kaufering, Germany), were cleaned with acetone and isopropanol and then a 1- μm thick SiO_2 layer was grown on them by thermal oxidation at 1100 $^\circ\text{C}$ in the clean room facility of the Institute of Nanoscience and Nanotechnology of NCSR “Demokritos”. Rabbit polyclonal antibody against carbendazim (anti-carbendazim antibody) and benzimidazole-conjugate were in-house developed, as previously described [11]. Carbendazim pestanal[®], biotinylated goat anti-rabbit IgG (biotinylated secondary antibody), streptavidin, highly pure ethanol, acetone, isopropanol, and (3-aminopropyl) triethoxysilane (APTES) were obtained from Sigma-Aldrich (St. Louis, MO, USA). Bovine serum albumin (BSA) was purchased from Acros Organics (Geel, Belgium). IgG Elution buffer was from

Thermo FisherScientific Inc. (Waltham, MA, USA). Salts and other chemicals were purchased from Sigma or Merck. The water used throughout the study was distilled.

2.2. WLRS Instrumentation

In the present work, an optical label-free sensor is presented based on White Light Reflectance spectroscopy (WLRS). WLRS methodology involves a visible/near infra-red light source (ThetaMetrisis SA) and a miniaturized USB controlled spectrometer (Ocean Optics Inc., Dunedin, FL, USA), operating in the corresponding spectral range. The white light emitted from the light source is guided to a reflection probe (ThetaMetrisis SA), which delivers the incident light to the biomodified surface and collects the reflected light directing it to the spectrometer. The surface consists of a transparent SiO₂ film over a Si reflecting substrate covered by a custom designed microfluidic cell (Jobst Technologies GmbH) providing the fluidic connections to the solutions. The particular sensor allows the label-free and real-time monitoring of the biomolecular reactions by recording the shift in the interference spectrum caused by the increase of the effective thickness of the biomolecular layer on the sensing surface during immunoreaction.

2.3. Preparation of Calibrators

Carbendazim stock solution of 5 mg/mL was prepared in pure ethanol, aliquoted, and stored at -30 °C. Calibrators with concentrations ranging from 20 µg/mL to 20 ng/mL were prepared in 10 mM phosphate buffer, pH 7.4, containing 0.9% NaCl, 0.4% (*w/v*) BSA and 5% (*v/v*) ethanol (assay buffer).

2.4. Biochip Preparation and Assay Performance

Chips were first cleaned and hydrophilized by O₂ plasma treatment (10 mTorr) for 30 s in a Reactive Ion Etcher. Then, they were immersed for 20 min in a 2 % (*v/v*) aqueous APTES solution, gently washed with distilled water, dried under a nitrogen (N₂) stream and cured by heating at 120 °C for 20 min. Chips were kept at room temperature (RT) in a desiccator for at least 48 h prior to use. For the chip biofunctionalization, a suitable benzimidazole conjugate was deposited on the chips and incubated overnight at RT. The following day, chips were rinsed with phosphate buffer (washing buffer), blocked with immersion for 3 h in 2% (*w/v*) BSA solution in washing buffer, rinsed with washing buffer and distilled water, dried with N₂, and used for the assay.

Prior to assay, each biofunctionalized chip was assembled with the fluidic module, placed on the docking station and equilibrated with assay buffer to acquire a stable baseline. For the assay, 1:1 *v/v* mixtures of calibrators (0.02–20 µg/mL in assay buffer) with the rabbit anti-carbendazim antibody (2 µg/mL in assay buffer) were passed over the chip for 18 min (50 µL/min flow rate), followed by a biotinylated anti-rabbit IgG antibody (7 min), and streptavidin (3 min). Finally, the biochip was regenerated by passing IgG elution buffer, followed by re-equilibration with assay buffer. The calibration curve was created by plotting the effective thickness of the built-up biomolecular layer (signal) corresponding to different calibrators, S_x (expressed as percentage of the zero-calibrator signal -maximum signal; S_0), for each one analyte against the respective analyte concentration in the calibrator solution.

3. Results and Discussion

3.1. Assay Optimization

In order to achieve a fast and sensitive immunochemical detection of carbendazim, many assay parameters had to be optimized. Due to the competitive nature of the assay, the two most crucial parameters to optimize were the concentration of benzimidazole-conjugate used for coating and the concentration of anti-carbendazim antibody, since their combination determines the highest signal (zero calibrator signal) and the assay sensitivity. Hence, based on preliminary titration experiments using different concentrations of the benzimidazole-conjugate (100–1000 µg/mL) in combination with different concentrations of the anti-carbendazim antibody (0.5–5 µg/mL), the highest analytical signal

was obtained when 500 µg/mL benzimidazole-conjugate was employed for surface functionalization in combination with 4 µg/mL concentration of anti-carbendazim antibody. Nevertheless, an antibody concentration of 2 µg/mL was adopted in the final protocol, since this resulted in an adequately high zero-calibrator signal along with significantly improved assay sensitivity. Another parameter to optimize was the duration of the whole assay, in order to achieve a fast and reliable analysis. As shown in Figure 1a, a total analysis time of 28 min was adopted, consisting of 18 min for the primary immunoreaction, 7 min for the secondary immunoreaction, and 3 min for the biotinylated secondary antibody-streptavidin reaction.

3.2. Assay Characteristics

A typical calibration curve obtained from the WLRs biosensing system for carbendazim is shown in Figure 1b.

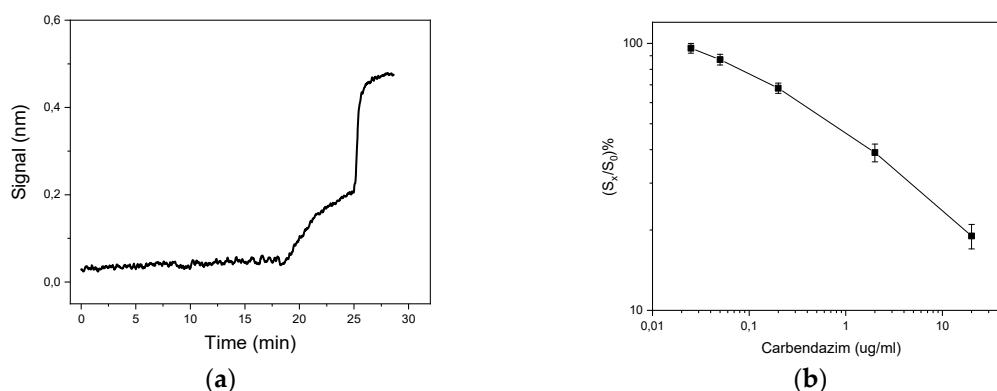


Figure 1. (a) Real-time response curve obtained from a biochip in the presence of zero calibrator, applying the optimized immunochemical protocol (b) Typical calibration curve obtained with carbendazim calibrators prepared in assay buffer. Each point represents the mean value of three independent runs ± SD. S_0 = zero-calibrator signal; S_x =calibrator signal.

The main analytical characteristics of the immunochemical detection of carbendazim using the WLRs sensor are summarized in Table 1. As shown, the analytical sensitivity of the proposed immunosensor was evaluated through calculation of the assay limit of detection (LoD) corresponding to a mean value -3SD of 15 replicate measurements of zero calibrator, and was found to be 20 ng/mL, whereas the dynamic range extended up to 20 µg/mL. The reproducibility of the assay was assessed by running three control samples in triplicate within the same day, and also in seven different days, in order to calculate the intra- and inter-assay coefficients of variation (CV), respectively. The intra-assay CV values were ≤6.9%, while the inter-assay CVs ≤9.4%.

Table 1. Immunochemical detection of carbendazim using the white light reflectance spectroscopy (WLRs) biosensing platform: Assay characteristics.

Assay Characteristic	Value
LoD (ng/mL)	20
Dynamic range (ng/mL)	up to 20,000
Intra-assay CVs (%)	≤6.9
Inter-assay CVs (%)	≤9.4
Duration	28 min

4. Conclusions

Herein, a WLRS-based biosensing platform for the label-free and real-time immunochemical determination of carbendazim was successfully developed. The proposed sensor allowed for the sensitive and fast quantification of carbendazim levels down to 20 ng/mL within less than 30 min. The proposed biosensor is planned to be applied for the determination of carbendazim in samples of interest, e.g., citrus fruit juices. Taken altogether, excellent analytical characteristics and short analysis time combined with small size of the analytical device render the proposed WLRS biosensor ideal for future point-of-need determination of carbendazim in food and environmental samples.

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