Hexaferrocenium Tri [Hexa(Isothiocyanato)Iron(III)] trihydroxonium Complex as a New DNA Intercalator for Electrochemical DNA Biosensor

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Abstract

Ferrocene or ferrocenium has been widely studied in the field of organometallic complexes because of its stable thermodynamic, kinetic and redox properties. Novel Hexaferrocenium Tri[Hexa(Isothiocyanato)Iron (III)]Trihydroxonium (HexaFc) complex was the product from the reaction of ferrocene, maleic acid and ammonium thiocyanate and it was confirmed by elemental analysis CHNS, FTIR and single crystal x-ray crystallography. In this study, HexaFc was used for the first time as an electroactive indicator for DNA hybridisation biosensor. The UV-Vis DNA titrations with this compound showed hypochromism and redshift at 250 nm with increasing DNA concentrations. The binding constant (K_b) for HexaFc complex towards CT-DNA (calf-thymus DNA) was $3.1 \times 10^4 \text{ M}^{-1}$. These results indicated intercalator behaviour of the complex. To test the usefulness of this complex for DNA biosensor application, a porcine DNA biosensor was constructed. The recognition probes were covalently immobilised onto silica nanospheres (SiNS) via glutaraldehyde linker on a screen-printed electrode (SPE). After intercalation with the complex, the biosensor response to complementary porcine DNA was measured using differential pulse voltammetry. The DNA biosensor demonstrated a linear response range to the complementary porcine DNA from 1×10^{-6} to $1 \times 10^{-3} \mu\text{M}$ ($R^2 = 0.9642$) with a limit detection of $4.83 \times 10^{-8} \mu\text{M}$. The results indicated that HexaFc complex is a feasible indicator for DNA hybridisation evaluation without the need of chemical attachment as a label.

Introduction

Ferrocene is a stable metal complex and consists an Iron(II) atom sandwiched between two cyclopentadienyl ligands¹. Ferrocene has been commonly studied due to its properties and applications in research comprising organic synthesis, catalyst and materials science². Ferrocene when loses an electron, ferrocenium species are formed resulting in the oxidation state to increase from 2+ to 3+. Ferrocenium is a free radical species of good stability as it has an unpaired electron in one of the nonbonding orbitals. Hence, because of all of these features, ferrocene and ferrocenium complexes are a good agent that is often used in biomedical fields³.

One of the usefulness of ferrocenium complexes and derivatives in biomedical fields is as an indicator for DNA detection. This characteristic is attributed to the electrochemical properties of the remarkable and straightforward redox-active unit of the ferrocene⁴. In addition, it has attracted a lot of attention of many researchers because it is an inexpensive but stable compound for a DNA detection system. The differential of the indicator's concentration before and after DNA hybridisation is often related to the voltammetry peak current. Liu et al.⁴ had employed the interaction between DNA and ferrocenium complex interaction based on the Langmuir-Blodgett film to develop a DNA electrode. Ferrocenium complex was used as an electroactive indicator to detect dengue DNA⁵. Ju et al.⁶ have reported a DNA hybridisation indicator using ferrocene derivatives to label the yeast DNA chain.

Ferrocenylnaphthalenediimide was used as DNA hybridisation intercalator to detect Sus scrofa mtDNA in food adulteration⁷. Sato and Takenaka⁸ studied electrochemical DNA detection using

ferrocenylnaphthalenediimide. In most of the reported work involving ferrocene/Fe(III) compounds of DNA hybridisation indicator, they were used as labels that needed to be chemically attached to the DNA probes to detect a hybridisation event^{9,10}.

In this work, a novel compound from a hexaferrocenium complex was used for the development of a DNA electrochemical biosensor. The use of a combination of hexaferrocenium cation and hexa(Isothiocyanato)Iron(III) anion for binding to DNA simultaneously after the hybridisation process on the electrode surface was attempted here for the first time. Both complex ions are expected to interact with the DNA hybridisation via intercalation mode during the hybridisation event. Hence the use of a combination of hexaferrocenium cation and hexa(Isothiocyanato)Iron(III) anion will allow direct detection of DNA hybridisation without the need of any further procedure of label attachment to the DNA probes. This is the uniqueness of using this complex ligand for DNA hybridisation.

Pork is typically involved in meat adulteration for example it is added to beef because pork is cheaper than beef. Pork is also frequently added into beef and chicken meatballs as additive¹¹. It is difficult and impossible in differentiating between beef and pork by visual inspection¹². Fish breeders also use boiled pork intestines and internal organs to feed their fish¹³. Some religious laws strictly- forbidden the existence of pork and its by-products in foods¹⁴. It is difficult for a consumer to identify between halal and haram meat where the presence of pork is the criteria for assessment. The assessment of the possible appearance of pork meat in food is thus a major consumer's demand for protection against falsely labelled food. A porcine DNA biosensor is one way to overcome the detection of pork in food as DNA biosensors are easily manageable, transportable, rapid detection, and a verification method for pork detection.

Thus, to develop a porcine DNA biosensor, the porcine DNA recognition probes were covalently immobilised onto silica nanospheres (SiNS) via glutaraldehyde linkers. Gold nanoparticles (AuNPs) were used to improve the overall electrochemical response of the DNA biosensor¹⁵. Differential pulse voltammetry was used to measure the current changes in response to complementary porcine DNA in the presence of HexaFc complex as DNA intercalator after the hybridisation process.

Methods

Apparatus. Field emission scanning electron microscope (FESEM) images were recorded in FESEM model Zeiss SUPRA 55VP with Energy-dispersive X-ray Analysis (EDX). The crystalline structure was assessed by X-ray diffraction (Cu K_α radiation, $\lambda=0.15416$ nm; $0.025^\circ.0.1\text{ s}^{-1}$) (model D8 Advance, brand Bruker AXS Germany). DNA titration were recorded using Varian Cary® 50 UV-Vis Spectrophotometer. Differential pulse voltammetry (DPV) measurements were carried out using AUTO-LAB PGSTAT and NOVA software package. A three-electrode system using Ag/AgCl as a reference electrode (Metrohm), platinum electrode as the counter electrode and screen-printed electrode (SPE) as the working electrode. The electrochemical measurements were carried out in 0.05 M potassium phosphate buffer solution (PBS) at pH 7 and were scanned from -0.4 V to 0.4 V with a step potential of 0.01 V.

Reagents and Materials. Synthetic oligonucleotides, calf-thymus DNA (CT-DNA) and the reagents (3-Aminopropyl)triethoxysilane (APTES, 99 %), glutaraldehyde, DNA, gold nanoparticles (AuNPs, <100 nm diameter), potassium chloride (KCl), ferrocene, maleic acid, ammonium thiocyanate) were supplied by Sigma Aldrich. The dihydrogen phosphate (KH_2PO_4) and potassium ferricyanide ($\text{K}_3[\text{Fe}(\text{CN})_6]$) chemicals were received from Merck, tetraethyl orthosilicate (TEOS, 98 %) was obtained from Fluka, ammonia (25 %) and ethanol (95 %) from System. Besides, dipotassium hydrogen phosphate (K_2HPO_4) was supplied by BDH Laboratory Supplies. MilliQ-deionized water was used to prepare all standard buffer and chemical solutions. The DNA oligonucleotide sequences used in the present research:

Porcine DNA probe (100 OD): 5'-CTG ATA GTA GAT TTG TGA TGA CCG TAG [AmC3]

Complementary Porcine DNA: 5'-CTA CGG TCA TCA CAA ATC TAC TAT CAG

Beef taurusa cytb: 5'-CAT AGC AAT TGC CAT AGT CCA CCCTA

Gallus gallus cytb: 5'-CGC AGG TAT TAC TAT CAT CCA CC

Synthesis and DNA binding of Hexaferrocenium Complex

An ethanolic solution of ferrocene (1.864 g; 0.01 mol) was added into 50 mL solution mixture of maleic acid (1.260 g; 0.01 mol) and ammonium thiocyanate (1.5224 g; 0.02 mol). The mixture was heated and stirred for 30 minutes. The excess ferrocene was filtered out, and the mixture was left to crystallise at room temperature. Bluish-black shiny needle-like crystal was formed after one day. Yield: 29 %. Melting point: 171-172 °C. Analysis: Calcd (%): C 39.17, H 3.00, N 10.54, S 24.11. Found (%): C 40.89, H 2.98, N 8.19, S 23.02 IR (cm^{-1}): 3099 $\nu(\text{C-H})$, 1413 $\nu(\text{C-C})$, 1007 $\delta(\text{C-C})$, 852 $\delta(\text{C-C})$, 2050 $\nu(\text{NCS})$.

5 mM of Tris-HCl buffer at pH 7.1 was used to prepare CT-DNA stock solution. UV-Vis spectra were recorded in 1 cm path length of quartz cuvettes using a Shimadzu UV-1800 spectrophotometer. The ratio of the absorbance of the CT-DNA solution at 260 nm and 280 nm was 1.87, which was more than 1.8, indicating that CT-DNA was not contaminated by proteins. A molar absorption coefficient value of 6600 $\text{M}^{-1} \text{cm}^{-1}$ was applied to calculate the concentration of CT-DNA from its absorption intensity at 260 nm¹⁴.

Fabrication and Characterisation of DNA biosensor

Silica nanospheres were prepared as per-described by Sani et al.¹⁶. The SPE was deposited with 10 μL of colloidal AuNPs (0.005 mg/ μL) and dried at room temperature. Then, 5 μL from 2 mg SiNSs suspended in 300 μL of ethanol was deposited onto the AuNPs-modified SPE and air-dried. Next, the SPE electrode was immersed in 10 % glutaraldehyde for 1 hour, followed by immersion in 300 μL of porcine DNA probe solution (1 μM) for 24 hours. The biosensor was then rinsed with 0.05 M potassium phosphate buffer, pH 7 and hybridised in 300 μL of complementary porcine DNA solution for 1 hour. The DNA was rinsed again with 0.05 M potassium phosphate buffer, pH 7 followed by dipping it in 2×10^{-5} M HexaFc solution for 1 hour and the response of the DNA biosensor was scanned using DPV. The fabrication of porcine DNA

biosensor using HexaFc as a DNA hybridisation marker is shown in Figure 1. The morphology of the SiNSs-DNA was determined by FESEM and FESEM-EDX (energy-dispersive X-ray) elemental mapping with platinum coating by sputtering. Meanwhile, the X-ray diffraction study of the DNA biosensor was performed using powder X-ray diffraction (XRD). The diffraction angle was measured with the X-ray radiation of Cu K α .

Performance Evaluation of Electrochemical Porcine DNA Biosensor

The scan rate study was carried out to understand the nature of the reaction. Scanning rate studies of modified SPE electrodes were performed in K₃[Fe(CN)₆] at 1 mM using cyclic voltammetry (CV) techniques at scan rates of 10 mVs⁻¹ to 300 mVs⁻¹. The deposited layer on the electrode surface was characterised by CV at a scan rate of 100mVs⁻¹ by using 2×10^{-4} M HexaFc complex as a redox label. The linear range of the porcine DNA biosensor was obtained by using a series of complementary porcine DNA concentration between 1×10^{-7} μ M until 1×10^{-2} μ M with constant HexaFc complex concentration at 2×10^{-5} mM. DNA biosensor selectivity with complementary DNA and non-complementary DNA was examined at the same concentration. The new porcine DNA biosensor was tested with raw pork samples. Pork DNA from raw meat was extracted according to the Spin-Column Protocol by using a DNeasy kit. The DNA obtained was hybridised with the immobilised DNA sequence of porcine on the biosensor in the presence of the new DNA redox indicator. The signal received was compared to the unhybridised response of the biosensor.

Results And Discussion

Infra-red Spectroscopy and Crystal Structure. Five significant peaks of ferrocenium compound at 2050.63, 3099.75, 1413, 1007 and 852 cm⁻¹ can be observed. The presence of a sharp stretching peak at frequency 2050.63 cm⁻¹ is owing to the N=C=S. Meanwhile the peak at 3099.75 cm⁻¹ is a characteristic for CH stretching of the cyclopentadienyl rings for ferrocenium group. The peak at 1413 cm⁻¹ corresponds to the antisymmetrical C-C stretching while the peak at 852.17 cm⁻¹ represents a CH out of plane mode for ferrocenium. The presence of all these significant peaks indicates that this ferrocenium compound has successfully formed¹⁷. The compound crystallized in trigonal crystal system with space group P-3, $a = b = 18.1606(12)$ Å, $c = 8.9808(6)$ Å, $\alpha = \beta = 90^\circ$, $\gamma = 120^\circ$, $Z = 1$ and $V = 2565.1(4)$. CCDC number for this compound is 1914296. From X-ray crystallography investigation, the titled compound consisted of six ferrocenium moiety, three hexa(isothiocyanato)iron(III) complexes and three hydronium group (Figure 2). The positive charge was stabilised by the presence of three hexa(isothiocyanato)iron(III) complexes and three hydronium group. All hexa(thiocyanato)iron(III) complexes in the compound adopt octahedral structure.

DNA binding of Hexaferrocenium Complex. The binding mode and the binding affinity between Hexaferrocenium complex and CT-DNA were performed using UV-Vis spectroscopic titrations. The UV-Vis absorption spectra of the title compound show two intense bands at 250 nm and 280 nm, respectively

(Figure 3). The absorption band at 250 nm is assigned to the π - π^* transition, which attributes to the localisation of molecular orbitals on the C=C group of hexaferrocenium cation. The absorption band at 280 nm is referring to the n - π^* transition of C=N of hexa(isothiocyanato)iron(III) anion. Absorption measurements were carried out by using constant HexaFc complex concentration (3×10^{-5} M) while increasing the concentration of DNA until no changes could be seen on the UV-Vis spectrum. The spectrum of the HexaFc complex was recorded after each addition of the DNA.

Changes that could be observed in the spectrum were either hyperchromism (increased in absorption) or hypochromism (decreased in absorption). Hyperchromism occurs due to the secondary damage of the double helix structure of the DNA, causing the DNA to be single-stranded^{18,19}. The occurrence of hypochromism is caused by the contraction of DNA in the helical axis. It is also affected by the transformation in DNA conformation²⁰. Besides, the change in the wavelength of either the redshift (the absorption to the longer wavelengths) or the blue shift (the absorption change to the shorter wavelength) can also be observed.

The percentage of hypochromicity was calculated following Equation (1):

$$\%Hypochromicity = \frac{\epsilon_{free} - \epsilon_{bound}}{\epsilon_{free}} \times 100(1)$$

Based on the UV-Vis DNA titrations, this compound exhibited a change in hypochromism and redshift at 250 nm when DNA concentration increased. The hypochromicity shown was 40 %, and the redshift was 2.5 nm and in agreement with the established intercalators^{21,22}. According to Wu et al.²³ and Shahabadi et al.²⁴, hypochromism event showed the binding strength of compounds towards DNA through intercalation mode. DNA binding through intercalation causes redshift. This involves strong overlapping interactions between the aromatic ligand chromophores of the metal complex with DNA bases^{25,26}. Meanwhile, the effects of hyperchromism or hypochromism can be observed when the occurrence of electrostatic interaction or groove binding followed by a blue shift (hypsochrome effect) or a minor change in the wavelength absorption of the UV-Vis spectrum²⁷. The intrinsic binding constant K_b of hexaferrocenium salt–DNA was determined according to Equation (2):

$$\frac{[DNA]}{(\epsilon_A - \epsilon_F)} = \frac{[DNA]}{(\epsilon_B - \epsilon_F)} + \frac{1}{K_b(\epsilon_B - \epsilon_F)}(2)$$

Where the apparent molar extinction coefficients, $\Delta\epsilon_{ap} = |\epsilon_A - \epsilon_F|$, $\epsilon_A = A_{observed}/[Complex]$, $\Delta\epsilon = |\epsilon_B - \epsilon_F|$. ϵ_F and ϵ_B represent molar extinction coefficients for the free hexaferrocenium salt and the DNA bound hexaferrocenium complex, respectively. To interpret the binding affinity of the compound to DNA, the intrinsic binding constant K_b was discovered by recognising the changes of maximum absorption bands centred at around 250 nm region. From the plotted graph of $[DNA]/|\epsilon_A - \epsilon_F|$ versus $[DNA]$, the y-intercept is equal to $1/(|\epsilon_B - \epsilon_F| \times K_b)$ whereas the slope is equal to $1/|\epsilon_B - \epsilon_F|$. K_b values can be determined by dividing

the slope value by the y-intercept. The binding constant (K_b) for HexaFc complex towards CT-DNA was $3.1 \times 10^4 \text{ M}^{-1}$. Also, this compound exhibited the same approximation value of binding constant with other reported ferrocene derivatives towards CT-DNA^{28,29}.

Characterisation of DNA Biosensor. We observed the characteristic modification when SiNSs immobilised porcine DNA probe and interacted with complementary DNA molecules. The XRD patterns of layer-by-layer DNA biosensor are shown in figure 4. The diffractogram of SPE exhibited a band at 25.6° . The intensity band and the percentage of crystallinity decreased after AuNPs and SiNSs were deposited on SPE. But the intensity band and the percentage of crystallinity were slightly increased after treatment with glutaraldehyde (GA)³⁰. The intensity band and the percentage of crystallinity continued to decrease with the addition of the porcine DNA probe and the complementary DNA. After the addition of HexaFc complex, the intensity and the percentage of crystallinity began to increase due to the presence of iron metal from the complex.

FESEM examined the morphology of the DNA biosensor. FESEM can provide a clear view of the structure of the DNA biosensor. Mapping with EDX made it possible to see the specifics of the DNA biosensor behaviour. Particularly after immobilisation of the aminated probes and hybridisation with complementary DNA. Figure 5 shows the morphology of the stepwise fabrication of the DNA biosensor. From the FESEM micrograph (Figure 5a), SiNSs were spherical with diameter between 20 to 200 nm while AuNPs with diameter less than 100 nm. Both SiNSs and AuNPs were dispersed with high homogeneity. More DNA probes could be immobilised on silica nanospheres than flat surface silica, hence an increase the biosensor sensitivity³¹. The DNA hybridisation in the presence of complementary DNA involving the intercalation with hexaferrocenium complex can be confirmed from FESEM studies (Figures 5). Thus, a noticeable change on the biosensor surface before and after interaction with a complementary porcine DNA was observed.

The dispersive energy X-ray (EDX) elementary mapping analysis was employed to detect the distribution of the different elements present in the biosensor³². This biosensor consists of carbon, oxygen, silicon and aurum elements. The elemental mappings of Au (yellow), C (red), Si (blue) and O (green) were observed and have shown to be well distributed in the biosensor (Figure 5a and 5b). As can be seen, nitrogen element was appeared after DNA immobilisation (Figure 5c) and it is derived from the aminated DNA probe (N-H). The presence of the nitrogen atom on the biosensor caused a higher attenuation of X-ray and gave a better contrast on the image. The N element increases in figure 5d due to the formation of double-stranded DNA³³. The process of DNA hybridisation is successful and was proved by the addition of a nitrogen element in figure 5d. Meanwhile, Figure 5e shows the presence of Fe element after HexaFc complex intercalation is performed. The combination of FESEM micrograph and EDX mapping supports the observation of stepwise fabrication of DNA biosensor, including DNA immobilisation and DNA hybridisation process.

Electrochemical Studies. The electrochemical study of HexaFc complex solution was inspected by cyclic voltammetry (CV) to determine their electron transfer properties. Figure 6 shows every layer of modified SPE that were tested by CV with HexaFc complex solution as a redox-active test probe. The sigmoid curve from the biosensor response proposed that the majority of radial diffusion occur on the electrode surface³⁴. Table 1 shows the peak separation between anodic and cathodic peak potential (ΔE_p) and anodic and cathodic peak current ratio (I_{PA}/I_{PC}). The separation between anodic and cathodic peak relates to ion resistance involved in the redox reaction³⁵.

Table 1 display the potential difference (ΔE_p) increase in the electrode order of AuNPs/SiNSs-SPE<AuNPs-SPE<bare SPE<SiNSs-SPE as the electron transfer rate decreased at the electrode surface in the electrode order of AuNPs/SiNSs-SPE>AuNPs-SPE>bare SPE>SiNSs-SPE. As the electron transfer rate decreased, CV became more expansive, and ΔE_p value increased. According to Monk³⁶, the reversible redox system can be determined by peak potential differences and I_{PA}/I_{PC} values ~ 1 . The bare SPE and the SiNSs-SPE depict greater peak separations (Figure 6). This may be attributed to the sluggish electron conductivity of the bare electrode, and the non-conductive properties of the SiNSs, which resulted in low electron transfer on the electrode surface. Therefore, the difference between anodic and cathodic peaks can be an indication of the resistance of electron transfer of the electrode³⁵. The value of ΔE_p generally decreases due to the presence of AuNPs of good electrical properties because AuNPs enhanced the electron transfer rate³⁶. The current oxidation-reduction peak (I_{PA}/I_{PC}) ratio of AuNP / SiNSs-SPE was close to 1. It shows that this system is still reversible even though the electrode has been modified.

Table 1. Electrodynamics data about anodic peak potential (E_{PA}), cathodic peak potential (E_{PC}), potential difference (ΔE_p), anodic peak current (I_{PA}), cathodic peak current (I_{PC}) and anodic to cathodic peak current ratio (I_{PA}/I_{PC}) of HexaFc with different surface-modified working electrodes.

	E_{PA} (V)	E_{PC} (V)	ΔE_p (V)	I_{PA} (A)	I_{PC} (A)	I_{PA}/I_{PC}
Bare SPE	0.26566	0.18524	0.08042	4.5673×10^{-6}	-3.0541×10^{-6}	1.4955
AuNPs- SPE	0.26566	0.18997	0.07569	6.4508×10^{-6}	-7.0527×10^{-6}	0.9147
SiNSs-SPE	0.26093	0.17578	0.08515	5.9079×10^{-6}	-3.0966×10^{-6}	1.9079
AuNPs/SiNSs-SPE	0.27039	0.19943	0.07096	6.191×10^{-6}	-6.1335×10^{-6}	1.0094

The scan rate study was conducted from 0.008 until 0.1 Vs⁻¹ in the K₃[Fe(CN)₆] redox indicator (Figure 7a). The K₃[Fe(CN)₆] system is an appropriate and valued tool for monitoring the characteristics of the modified electrode³⁷. According to figure 7a, the peak current for oxidation and reduction increases proportionally with the scan rate from 0.008 until 0.1V/s. Figure 7b demonstrates the cyclic voltammetry plot at different scan rates. Polarisation increase with the increasing of scan rate, and it develops wide and distorted oxidation and reduction peak. Other than that, the increasing scan rate will lead to the increasing of voltage change among anodic and cathodic peaks³⁸.

Scan rate study was also performed in HexaFc complex from 0.008 until 0.08 Vs⁻¹ to investigate the electrochemical process of the complex. Figure 7c displays that the redox peak current increases with the scan rate from 0.008 until 0.08 Vs⁻¹. However, there is no much difference in the redox peak current for the scan rates within 0.09 and 0.1 Vs⁻¹. This result suggests that HexaFc complex is irreversible at the higher scan rate. The scan rates from 0.008 until 0.08 Vs⁻¹ were more reversible, proof by the linear increase in the current redox (Figure 7d). According to the larger peak area of the CV curves, more material was set based on the Faraday Law³⁹.

Analytical Performance. SiNSs is a non-conductive material and has been used as DNA probe immobilisation sites. The low conductivity of SiNSs reduces biosensor performance. AuNPs have therefore been used to overcome this problem. AuNPs can enhance the potential of electron transfer from the redox indicator and improve the biosensor sensitivity³⁷. Glutaraldehyde has been utilised as a link between porcine DNA probe and to the amine groups of SiNSs. The pH buffer also plays an essential role in producing DNA biosensor because it provides a suitable environment for DNA hybridisation process^{31,40}. Generally, an acidic or basic environment can cause DNA damage³¹. All the parameters in the optimised condition were used to fabricate porcine DNA biosensor and shown in Table 2.

Table 2. Optimised parameter of porcine DNA biosensor

Parameters	Optimum amount
Amount of gold nanoparticles	0.05 mg
Amount of silica nanospheres	0.04 mg
% of glutaraldehyde	10
Potassium phosphate buffer concentration	0.05 M
Potassium phosphate buffer pH	pH 7
Ionic strength	1 M
Concentration of DNA probe	1 μ M
Immobilization time	24 hours
Concentration of HexaFc complex	2×10^{-5} M

The performance of the biosensor was tested in a sodium phosphate buffer solution, 0.05 M containing complementary porcine DNA with different concentration and 1 M Na⁺ ion at pH 7. Figure 8 shows a good linearity response of complementary porcine DNA from 1×10^{-6} μ M until 1×10^{-3} μ M with the correlation coefficient, $R^2=0.9642$. Limit of detection (LOD) for complementary porcine DNA was determined at 4.83×10^{-8} μ M. The LOD of the porcine DNA biosensor was calculated following the three times of the standard deviation of the biosensor response in the linear range divided by the direct calibration code⁴¹. The average reproducibility relative standard deviation for each calibration point of this biosensor is excellent with most values below 4.5 % (n=3).

Figure 9a exhibits the DPV response before and after the probe-complementary DNA hybridisation process. The DNA electrode based on AuNPs/SiNSs-SPE showed a 100 % matching of the complementary-porcine DNA sequences. The strong and sharp of the DPV peak at a potential ~ 0.22 V revealed that the HexaFc complex had been successfully intercalated into the porcine DNA double-stranded and it suggested that DNA probe is immobilised and hybridised with complementary-DNA. DNA probe covalently immobilised with SiNSs. No DPV peak was found at potential ~ 0.22 V for single-stranded porcine DNA probe, and only a small peak was obtained after this biosensor exposed to beef and chicken DNA. This new research will be beneficial for determining porcine DNA in food products by applying new HexaFc complex as an electrochemical porcine DNA indicator.

This DNA biosensor was tested with raw pork meat from several markets and supermarkets around Bangi, Malaysia. It was discovered that the signal from raw pork was found to have a high current value compared to a DNA probe (Figure 9b).

Performance Comparison with Other Reported Porcine DNA Biosensors

Table 3 displays a performance comparison between the constructed porcine DNA biosensor and the other electrochemical porcine DNA biosensors. This comparison is needed for the validation of the constructed biosensor. The constructed porcine DNA biosensor showed substantial improvements in high linear range response and low detection limit compared to other DNA immobilisation matrixes, such as disposable electrochemical printed chips⁴², gold nanoparticles⁴³, AuNP/NBA-NAS⁴⁴ and graphene biochips⁴⁵. This outcome is attributed to the large surface area of silica nanospheres (SiNSs) for DNA immobilisation sites. When compared with DNA hybridisation indicator that employed ferrocenium compounds^{9,10}, the biosensor developed here has comparable LOD, but it did not require any procedure for ferrocene labelling, i.e. chemical attachment of ferrocene compounds to the DNA. This advantage is because the ferrocenium complex used in this work detects DNA hybridisation by intercalation.

Table 3. Comparison of performance between the DNA biosensor reported here with other biosensors using different ferrocene and non-ferrocene based compounds as an indication of hybridisation

DNA electrode materials	DNA label	Linear range (μM)	LOD (μM)	Reference
SPE/AuNPs/SiNSs	HexaFc complex	1×10^{-6} - 1×10^{-3}	4.83×10^{-8}	This work
Fc-acid-OMPA	Ferrocene oligomer	1.0×10^{-9} - 1.0×10^{-4}	2.0×10^{-9}	[9]
Graphene	Tetraferrocene	2.0×10^{-8} - 2.0×10^{-3}	8.2×10^{-9}	[10]
Gold nanoparticles-DNA bioconjugate	Methylene blue	7.35×10^{-3} - 3.68×10^{-1}	4.26×10^{-2}	[43]
SPE/AuNP/NBA-NAS	Ruthenium(II) complex	1.0×10^{-7} - 1.0×10^{-2}	-	[44]
Graphene biochips	Ruthenium hexamine	-	6.89	[45]
Disposable electrochemical printed chips	Hoechst 33258	-	1.34	[42]

Conclusion

The new HexaFc complex has demonstrated good DNA binding and selectivity from spectrophotometric studies. It was then successfully used as a new redox indicator for an electrochemical DNA biosensor to determine porcine DNA. Most importantly, the DNA biosensor with this ferrocenium indicator showed a good response towards complementary porcine DNA. It is potentially an easy and rapid method for the determination of porcine DNA in food products.

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Declarations

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Author contributions

E.Y.A. conceived and designed the experiment except for synthesis and DNA binding of Hexaferrocenium complex, performed the experiment, analysed the data and wrote the paper except the DNA binding of Hexaferrocenium complex part; F.R. synthesised Hexaferrocenium complex; E.I.Z. performed the DNA binding experiment and wrote the DNA binding part; M.K., B.M.Y., L.Y.H., and S.A.H. together conceived

and designed the experiments, contributed the reagents/materials/analysis tools and proof-read the manuscript.

Figures

Figure 1

Fabrication of porcine DNA biosensor using HexaFc as DNA hybridisation marker

Figure 2

HexaFc crystal structure with 30% ellipsoid probability at c-axis.

Figure 3

Absorption spectral changes of HexaFc complex (30 μM) in Tris- HCl buffer (pH 7.1) in the absence and presence of increasing concentrations of CT-DNA. Inset: Fitting of the absorbance data that were used to obtain the binding constant at 250 nm.

	Crystallinity (%)	Amorphous (%)
SPE	87.4	12.6
SPE-SiNSs/AuNPs	73.9	26.1
SPE-SiNSs/AuNPs-GA	83	17
SPE-SiNSs/AuNPs-GA-DNA probe	81.7	18.3
SPE-SiNSs/AuNPs-GA-DNA probe-complementary DNA	58.1	41.9
SPE-SiNSs/AuNPs-GA-DNA probe-complementary DNA-complex HexaFc	82.9	17.1

(b)

Figure 4

(a) XRD diffractogram and (b) crystallinity table of layer-by-layer DNA biosensor.

Figure 5

EDX elemental mapping analysis. Inset: FESEM micrograph of DNA biosensor; (a) silica nanospheres (SiNSs) with gold nanoparticles (AuNPs); (b) SiNSs, AuNPs, glutaraldehyde; (c) after immobilization with DNA probe; (d) after hybridisation with complementary DNA; (e) after intercalation with HexaFc.

Figure 6

Cyclic voltammograms of 2×10^{-4} M HexaFc complex from bare SPE (SPE), gold nanoparticles modified SPE (AuNPs-SPE), gold nanoparticles/silica nanospheres modified SPE (AuNPs/SiNSs-SPE), and silica nanospheres modified SPE (SiNSs-SPE) at a scan rate of 80 mVs⁻¹.

Figure 7

(a) Scan rate study of DNA biosensor in $K_3Fe(CN)_6$; (b) cyclic voltammetry plot at different scan rate in $K_3Fe(CN)_6$; (c) scan rate study of DNA biosensor in HexaFc complex solution and (d) cyclic voltammetry plot at different scan rates in HexaFc complex solution.

Figure 8

Linear response range for porcine DNA biosensor

Figure 9

(a) Porcine DNA biosensor selectivity on complementary porcine DNA, chicken DNA and beef DNA and (b) Real samples measured with porcine DNA biosensor.