

Diagnosis of Toxoplasmosis Using Surface Antigen Grade I Detection by ELISA, Nano-Gold ELISA, and PCR in Pregnant Women

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Introduction: The accurate diagnosis of toxoplasmosis has critical importance in pregnant women. Nanotechnology and molecular biology are making possible opportunities for accurate and rapid diagnosis of many infectious diseases.

Aim and Methods: The aim of our study was to compare nano-gold ELISA with ELISA and PCR for diagnosis of toxoplasmosis using *Toxoplasma* surface antigen grade 1 (SAG1) in pregnant women seeking antenatal care in outpatient clinics.

Results: PCR showed the highest diagnostic values than nano-gold ELISA and ELISA regarding sensitivity (97.3% versus 89.2% and 83.8%); specificity (100% versus 94% and 88%); and diagnostic accuracy (98.9% versus 91.95% and 86.2%), respectively. There is no statistical difference between PCR and nanogold ELISA results.

Discussion: Nano-gold ELISA had a significant improvement in diagnosis than the traditional ELISA method. Most likely with the assistance of nanoparticles, more antibodies enter the antigen-antibody complex because of the considerable improvement in the surface area of nano-gold particles.

Conclusion: Although PCR had higher diagnostic values than nano ELISA, nano ELISA is cheaper and easier than PCR. We recommend nano-gold ELISA with SAG1 as a promising technique in the diagnosis of toxoplasmosis and survey studies.

Keywords: toxoplasmosis, SAG 1, immunodiagnosis, nanogold, ELISA, PCR

Introduction

Toxoplasma gondii (*T. gondii*) is a worldwide obligate intracellular parasite.¹ According to estimates, toxoplasmosis has infected one-third of the world's population.^{2,3} *T. gondii* IgG antibody prevalence in pregnant women was surveyed in several governorates of Egypt, ranging between 44.0 to 67.5% and 23.0% of aborted women.⁴ Also, the prevalence among dogs (23.3%) and cats (9.8%) in Egypt was reported.⁵ About 0.8% of pregnant females were confirmed to be at hazard of acute seroconversion. Pregnant women with toxoplasmosis run the risk of miscarriage, stunted growth, early delivery, or congenital anomalies (encephalitis, hydrocephalus, microcephalus, retinochoroiditis, blindness, mental retardation epilepsy).^{4,6} The severity of toxoplasmosis relies on the host's immune response and the *T. gondii* strain.⁷

Toxoplasmosis can be diagnosed by various methods, including antibody detection using serological tests, and parasite DNA determination by polymerase chain reaction (PCR).

The most widely utilized technique nowadays is antibody detection, which provides serological proof of immunoglobulin (Ig)G and IgM antibodies specific to *T. gondii* antigens. These tests could have sensitivity and specificity problems, which could produce false-positive or false-negative results.⁸ The emergence of molecular biology and serological techniques coincided with the emergence of many *T. gondii* stage-specific genes. One of

these, surface antigen grade 1 (SAG1), which is specific to tachyzoites, makes up 3.0–5.0% of the surface membrane proteins. It is essential for parasite adhesion and host-cell penetration. It also has an important role in immune modulation.⁹ First trials of PCR detection based on *SAG1*-specific primers showed that they were able to amplify DNA from seven test *T. gondii* strains with no cross-reactivity with other parasites or microbes. Hence, these primers are highly specific.¹⁰ The sensitivity of PCR is not only affected by the genomic copy number of the target gene but is also affected by the infection stage. PCR detection using primers specific to several target genes including SAG 1 is often very effective.¹¹

Nano diagnostics is the application of nanotechnology in clinical diagnosis to solve the need for increased sensitivity, specificity, and early detection. The nanomaterial helps many target molecules bind more easily for ultrasensitive detection due to their large surface area.¹² Different materials were used for nanoparticle synthesis and used in medical research to cure and diagnose various illnesses, such as gold, chitosan, and silver. Nowadays, using nano-gold particles as a weapon against various parasites has become a common practice in research and medicine.¹³ The currently available and frequently used toxoplasmosis diagnostic procedures can be improved by nanomaterials or perhaps replaced by them.¹⁴ We used pregnant women sera in our study due to the availability of routine *Toxoplasma gondii* screening tests during prenatal care. The aim of our study was to compare a new nano-gold ELISA with ELISA and PCR for diagnosis of toxoplasmosis using SAG1.

Patients and Methods

Sample Collection

Samples of blood were collected from pregnant women seeking antenatal care in the outpatient clinic of the Gynecology and Obstetrics Department of Benha University Hospital, Benha Teaching Hospital, and also from different laboratories using commercial IgG and IgM ELISA kits (Sigma, St. Louis, MO, USA). In the period between October 2017 and December 2020, the study protocol complies with the Declaration of Helsinki¹⁵ and was revised and approved by the Medical Research Ethical Committee at Benha Faculty of Medicine, Benha University (Approval no. MD-5-11-2017). The objective of the study was explained to all pregnant females participating in this work and informed consent was taken. The study was conducted on 87 pregnant females, and they were grouped as follows: Group I: Thirty-seven *T. gondii* infected patients (presence of IgM, seroconversion, or rising IgG antibodies titer in their serum). Group II: Thirty patients having other parasites than *T. gondii*: ten patients infected with *Cryptosporidium parvum* (*C. parvum*), 10 patients infected with *Giardia lamblia* (*G. lamblia*), and 10 patients infected with *Entamoeba histolytica* (*E. histolytica*). The idea behind including patients harboring other parasites than *T. gondii* in our study was to avoid cross-reactivity in the results. Group III: Twenty apparently healthy pregnant women free from parasitic infections served as noninfected control (negative control). Blood samples were stored at -20°C till required for the study tests.

Animals

Three Swiss Albino male mice, weighting roughly 20–30 grams, were used in our study in the preparation of Toxoplasma surface antigen grade I (SAG1). Mice were obtained from the animal house of Theodore Bilharz Research Institute (TBRI). They were housed in plastic cages with white wood chips for bedding, fed by commercial complete food mixture and tap water for drinking and maintained temperature (22 ± 2). Also, two New Zealand white male rabbits, weighing approximately 1.5 Kg and about two months age, were used in the production of polyclonal antibodies (pAb). They were purchased from the Rabbit Research Unit (RRU), Agriculture Faculty, Cairo University. The rabbits were housed for four weeks (experiment duration) in the animal house in TBRI under standard laboratory care at 21°C , 16% moisture, filtered drinking water with additional salts 1cm/5 liters and vitamin 1cm/10 liters. A diet containing 15% protein, 3% fat, and 22% fiber was purchased from RRU. All procedures related to experimental animals in the present study approved by TBRI and met the International Guiding Principles for Biomedical Research Involving Animals, as issued by the International Organizations of Medical Sciences (NRC, 2006).¹⁶

ELISA Techniques

Preparation of SAG I Antigen

T. gondii tachyzoites (RH strain), kindly provided by Parasitology Department of (TBRI), were used for the preparation of *Toxoplasma* surface antigen grade I (SAG1).¹⁷ Briefly, *T. gondii* RH strain tachyzoites were grown and maintained in three Swiss Albino male mice by intraperitoneal inoculation. The peritoneal fluids from infected mice were pooled, centrifuged, and then cleaned three times with phosphate buffer saline every four days after infection, disrupted by sonication then centrifuged at 12,000 rpm for 1 hour, and the supernatant collected and the purified target antigen was obtained by DEAE-Sephadex A-50 ion exchange chromatography as a soluble antigen.^{18,19} The concentration of protein was determined dependent on Coomassie brilliant blue G-250 dye color change and the soluble antigen stored at -20°C till use. Assessment of reactivity and specificity of *Toxoplasma* surface antigen grade I (SAG1) was done by indirect ELISA.²⁰

Production of Polyclonal Antibodies (pAb)

By immunizing one of the New Zealand white rabbits with SAG1, through intramuscular injection (i.m.) of a priming dose at two sites, 1mg *Toxoplasma* SAG1 mixed 1:1 in complete Freund's adjuvant (CFA) (Sigma), this was followed by 3 booster doses, each was 0.5 mg antigen emulsified in equal vol. of incomplete Freund's adjuvant (IFA) (Sigma). Two weeks after the priming dose, the first boosting was given. The following boosting doses were given at weekly intervals.²¹ Purification of the rabbit's anti-*Toxoplasma* serum by ammonium sulfate (50%) precipitation method was done as 50% Saturated ammonium sulfate solution was added dropwise to rabbit anti-*Toxoplasma* serum to reach 50% saturation, with continuous stirring and then centrifuged for 20 min at 3500 rpm in a cooling centrifuge (Heraeus) at 4°C . The supernatant was discarded, and the ammonium sulfate precipitation steps were repeated several times. The final precipitate was dissolved in a suitable amount of 0.01 M PBS, pH 7.2. The ammonium sulfate was removed by dialysis against 0.01 M PBS, pH 7.2 for 72 hrs at 4°C . Then, purification by caprylic acid (7%) method was done, caprylic acid was added dropwise on the precipitated whole Igs after 50% ammonium sulfate precipitation, with slow magnetic stirring for 30 min at 4°C then the mixture was centrifuged at 1000 rpm for 30 minutes. Precipitation was discarded, while the supernatant contained nearly pure IgG.²²⁻²⁴

One fraction of polyclonal antibodies was conjugated with Horseradish peroxidase (HRP). This was done on two successive days. On the 1st day, (A) Five mg of horseradish peroxidase (HRP) (Sigma) was suspended in 1.2 mL of dist. H₂O, then 0.3 mL (300ul) of freshly prepared sodium periodate was added and incubated at room temperature for 20 min. The HRP solution was dialyzed against 1 mM sodium acetate buffer. (B) Five mg/mL of polyclonal antibody was dialyzed against 0.02M carbonate buffer. On the 2nd day, A and B were mixed and then incubated for two hours at room temperature. 100 μL of freshly prepared Na borohydride was added to the mixture of A +B and then incubated for two hours at 4°C .²⁵ The second fraction of polyclonal antibodies remained without conjugation. The third fraction was conjugated with nanogold particles (AuNPs-MUA) and is mentioned in the next section. The three fractions were stored at -20°C till required.

Nano-Gold Particles (AuNPs)

AuNPs were purchased from Nanotech Company (6th October City, Egypt) and scanned by transmission electron microscope. AuNPs functionalized with mercaptoundecanoic acid (AuNPs-MUA) by adding 30 mL of the prepared AuNPs to 45 μL of 1 mM mercaptoundecanoic acid (MUA) in ethanol solution (Sigma Aldrich, Germany). The solution was mixed and left overnight at 4°C . The concentration was determined the next day by UV/vis spectrophotometry using Beer's law.²⁶

The 3rd fraction of polyclonal antibodies was conjugated with nanogold particles (AuNPs-MUA). Two mg of anti-*Toxoplasma* pAb was added to 5 mL of AuNPs-MUA (2 nM; pH = 7.4). To obtain more robust AuNPs-pAb conjugates, the covalent linkage between pAb and AuNPs-MUA was done through the use of N-hydroxy succinimide/1-Ethyl-3-[3-dimethylaminopropyl] carbodiimide (NHS/EDC) cross-linkers (NHS: Fluka, Germany). Five mL of a mixture of 5 mM sodium phosphate buffer (pH = 7), 1.2 mM NHS, and 2.8 mM EDC was added to 5 mL of AuNPs-pAb conjugates. The mixture was mixed and left to incubate overnight at 4°C .^{27,28}

ELISA Procedure

Microtiter plates were coated with 100 μL /well of purified pAb in a concentration of 20 $\mu\text{g}/\text{mL}$ as a capture antibody diluted in 0.8 M carbonate buffer, pH 9.6 (1/20). The plates were washed with a washing buffer [0.1 M PBS/0.05% Tween 20, PH 7.4 (PBS/T)] three times. The free sites were blocked with 200 μL /well of 0.1% BSA/PBS/T and incubated for 1 h at 37°C. The plates were washed with 0.1 M PBS/T five times. Pregnant women's sera (100 μL /well), in dilution 1/50, were added and incubated for 2 h at 37°C, and the plates were washed 3 times. Peroxidase-conjugate pAbs (100 μL /well) in concentration 20 $\mu\text{g}/\text{mL}$ diluted in PBS (1/80) were distributed, and plates were incubated for 2 h at 37°C. The reactions were visualized by the addition of 100 μL /well of O-phenylenediamine substrate solution for 30 min at room temperature. The reaction was stopped by adding 50 μL /well of 8 N H_2SO_4 , and the plates were read at optical density (OD) 490 nm with an ELISA microplate reader (Bio-Rad). The cutoff point was 0.16 in ELISA.^{29,30}

Nano-Gold ELISA

Microtiter plates were coated with 100 μL /well of nano-gold conjugated pAb in concentration of 10 $\mu\text{g}/\text{mL}$ as a capture antibody diluted in 0.8 M carbonate buffer, pH 9.6 (1/40). The plates were washed with a washing buffer [0.1 M PBS/0.05% Tween 20, PH 7.4 (PBS/T)] three times. The free sites were blocked with 200 μL /well of 0.1% BSA/PBS/T and incubated for 1 h at 37°C. The plates were washed with 0.1 M PBS/T five times. Pregnant women's sera (100 μL /well), in dilution 1/50, were added and incubated for 2 h at 37°C, and the plates were washed 3 times. Peroxidase-conjugate pAbs (100 μL /well) in concentration 15 $\mu\text{g}/\text{mL}$ diluted in PBS (1/80) were distributed, and plates were incubated for 2 h at 37°C. The reactions were visualized by the addition of 100 μL /well of O-phenylenediamine substrate solution for 30 min at room temperature. The reaction was stopped by adding 50 μL /well of 8 N H_2SO_4 , and the plates were read at optical density (OD) 490 nm with an ELISA microplate reader (Bio-Rad). The cutoff point was 0.203 in nano-gold ELISA.^{29,30}

Polymerase Chain Reaction (PCR)

Human DNA was extracted from blood samples by using QIAmpDNA Blood Mini Kit supplied by (Qiagen, Germany) according to the manufacturer's instructions. After extraction, DNA samples were measured using NanoDrop 1000 Spectrophotometer (Thermo Scientific, USA). The eluted DNA was then preserved at -20°C until further processing. *T. gondii* major surface antigen (SAG1) gene (Accession NO. MK250980.1) was amplified using two oligonucleotide primers: F: 5'-GCTGTAACATTGAGCTCCTTGATTCCTG-3' and R: 5'-CCGGAACAGTACTGATTGTTGTCTTGAG-3' and Maxima Hot Start Green Master Mix (Fermentas, Thermo Fisher Scientific Co., USA). PCR reaction was conducted in a total volume of 50 μL using 25 μL of Taq PCR Master Mix, 2.5 μL forward and reverse primers, 10 μL of template DNA, 10 μL of nuclease-free. An initial denaturation step at 94 °C for 5 min, then 35 cycles consisting of 94 °C for 30 sec, 62 °C for 30 sec, 72 °C for 30 sec, and a final extension step at 72 °C for 10 min. The amplified DNA of the SAG1 gene was electrophoresed with 1% agarose gel.^{31–33}

Statistical Analysis

The collected data were analyzed by SPSS program (SpssInc, Chicago, ILL Company) version 16 software. Chi-square (χ^2) and Fisher's exact tests were used. Receiver operator characteristic (ROC) curves were constructed to assess the performance of PCR and ELISA in the diagnosis of toxoplasmosis. MedCalc Software (Mariakerke, Belgium) was used to compare areas under the curve of the 3 parameters.^{34,35} The data were considered significant if *p*-values were equal to or less than 0.05.

Results

Toxoplasma SAG I Antigens

The SAG1 antigen was purified using DEAE Sephadex A-50 ion exchange chromatography, and the OD₂₈₀ profile is obtained and shown in Figure 1. A single peak with a maximum OD value of 1.2 at fraction number 10 represents the eluted antigen (represented by the fraction with the highest protein content). We collected fraction no. 10 as SAG 1 antigen.

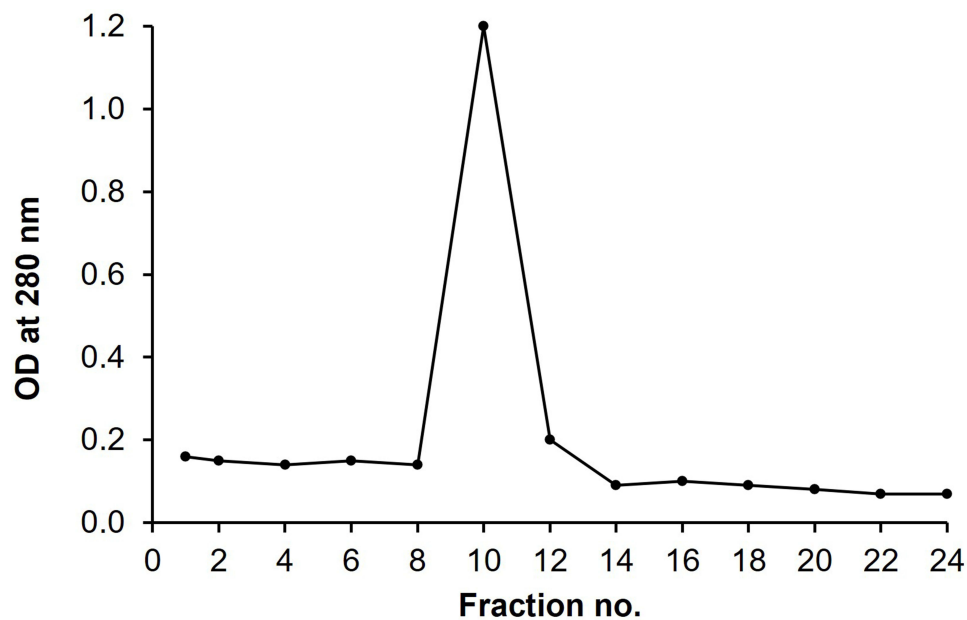


Figure 1 Purification of the SAG1 antigen by DEAE-Sephadex A-50 ion exchange chromatography. The OD₂₈₀ profile of the antigen fractions obtained following purification by DEAE Sephadex A-50 ion exchange chromatography. The eluted antigen is represented by a single peak with maximum OD value equal to 1.2 at fraction number 10 (represented the fraction with the highest protein content).

Anti-Toxoplasma pAb

Figure 2 shows increasing antibody level in the immunized rabbit started one week after the first booster dose. With the 3rd booster dose, the immune sera gave a high titer against *Toxoplasma* antigen with an OD of 1.8 at 1/250 dilution. The non-immunized rabbit showed no increase in antibody level. Table 1 shows that the produced anti-*Toxoplasma* antibodies diluted 1/250 in PBS/T gave strong reactivity to *Toxoplasma* antigen. The OD reading at 490 nm for *T. gondii* was 1.82 compared to 0.28, 0.2, and 0.22 for *C. parvum*, *E. histolytica*, and *G. lamblia*, respectively.

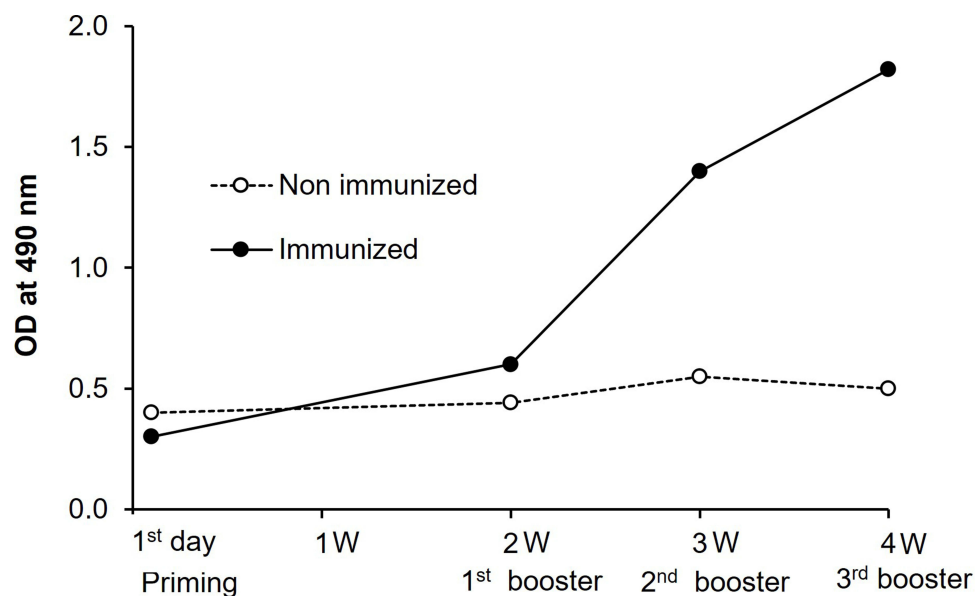


Figure 2 Reactivity of immunized rabbit anti-*Toxoplasma gondii* IgG polyclonal antibodies (diluted 1/250) against *Toxoplasma* SAG1 by ELISA. The figure showing increasing antibody level in the immunized rabbit started one week after the first booster dose. With the 3rd booster dose, the immune sera gave a high titre against *Toxoplasma* antigen with OD 490 of 1.8 at 1/250 dilution. The non-immunized rabbit showing no increase in antibody level.

Table 1 Reactivity of Purified Target Antigen by ELISA

Serum Samples	OD Readings at 490 nm
<i>T. gondii</i>	0.95 ± 0.22
<i>C. parvum</i>	0.21 ± 0.02
<i>E. histolytica</i>	0.2 ± 0.01
<i>G. lamblia</i>	0.18 ± 0.07

Note: Data represented as M±SD value.

Abbreviation: OD, optical density.

Nano-Gold Particles (AuNPs)

Transmission electron microscopy (TEM) was used to get direct information about the size of the produced nano-gold particles. TEM images showed that most particles (66%) were approximately about 40 nm (Figure 3).

ELISA

As regards detecting *Toxoplasma* (SAG1) by ELISA in the serum, the results were positive in 31 cases (83.8%) of group 1 (*T. gondii* infected cases), while 6 cases (16.2%) were negative. In group II, 6 cases (20%) were detected as false positive (1 with *C. parvum*, 3 with *G. lamblia* and 2 with *E. histolytica*), while the other 24 cases (80%) were negative. All healthy control samples were negative (Table 2).

Nano-Gold ELISA

The results of nanogold ELISA were positive in 89.2% of group 1, while 10.8% were negative. In group II (patients with other parasites) 10.0% were detected as false positive (2 cases with *G. lamblia* and 1 case with *E. histolytica*), while the other 90.0% were negative. All healthy control cases (group III) were negative. There was a statistically significant difference among the studied groups regarding positive and negative results (Table 2).

PCR

When detecting *Toxoplasma* SAG1 gene in the blood, 36 cases (97.3%) were positive in group I (*T. gondii* infected cases). In group II (patients infected with other parasites) and group III (healthy control group), all samples were

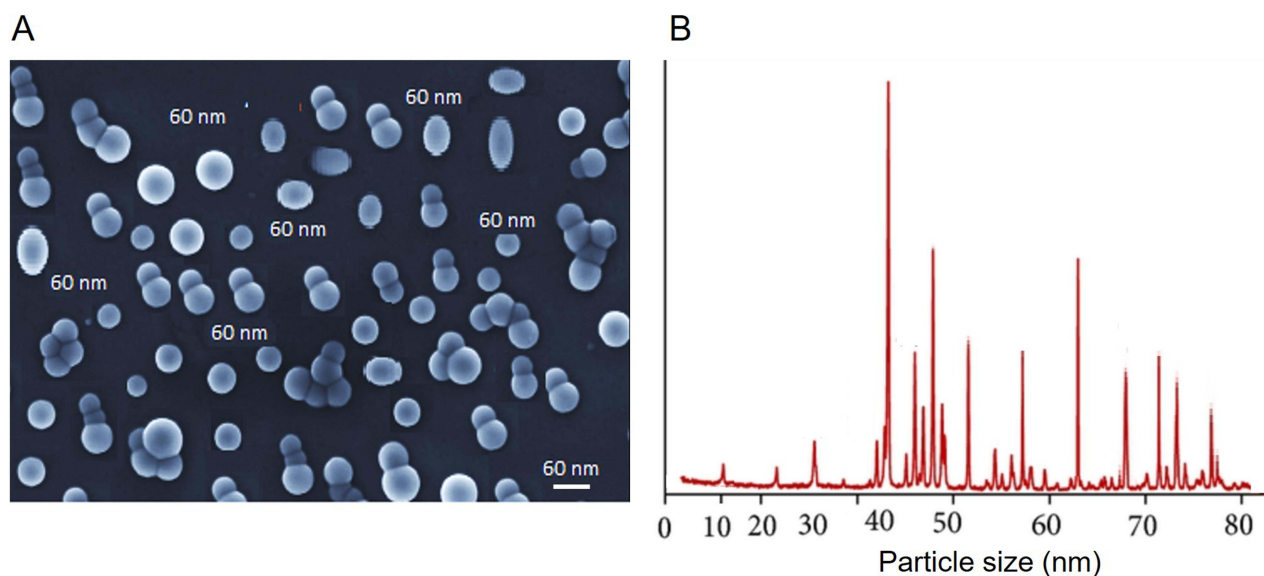


Figure 3 Nano-gold particles size. (A) Transmission Electron microscopy images of gold nanoparticles. The mean size 40–60 nm. (B) Histogram showing the size distribution of gold nanoparticle. White line indicated scale bare of 60 nm.

Table 2 SAG I Detection Using PCR, Nano-Gold ELISA, and ELISA Techniques

Groups		PCR*		Nano-Gold ELISA*		ELISA*	
		Positive No. (%)	Negative No. (%)	Positive No. (%)	Negative No. (%)	Positive No. (%)	Negative No. (%)
GI (n=37)	<i>T. gondii</i> (n=37)	36 (97.3)	1 (2.7)	33 (89.2)	4 (10.8)	31 (83.8)	6 (16.2)
GII: Other parasites (n=30)	<i>C. parvum</i> (n=10)	0 (0)	10 (100)	0 (0)	10 (100)	1 (10.0)	9 (90.0)
	<i>G. lamblia</i> (n=10)	0 (0)	10 (100)	2 (20.0)	8 (80.0)	3 (30.0)	7 (70.0)
	<i>E. histolytica</i> (n=10)	0 (0)	10 (100)	1 (10.0)	9 (90.0)	2 (20.0)	8 (80.0)
GIII: Negative control (n=20)	Healthy control (n=20)	0 (0)	20 (100)	0 (0)	20 (100)	0 (0)	20 (100)

Note: *Statistically significant results between positive and negative cases ($p < 0.001$).

negative. No false-positive results were detected. Electrophoresis bands of PCR are shown in Figure 4, group I showed a negative sample in lane 1 and positive samples in lanes 2, 3, and 4 at 356 bp. Other parasites and negative control groups showed no PCR bands. There was a significant difference among the different studied groups between positive and negative results (Table 2).

Comparison Between Different Techniques

PCR results were higher than nano-gold ELISA and ELISA regarding sensitivity (97.3% versus 89.2% and 83.8% respectively); specificity (100% versus 94% and 88% respectively); PPV (100% versus 91.7% and 83.8% respectively); NPV (98% versus 92.2% and 88% respectively) and diagnostic accuracy (98.9% versus 91.95% and 86.2% respectively) (Table 3). ROC curve (Figure 5) was used for comparing the performance of PCR, nano-gold ELISA, and ELISA in the diagnosis of toxoplasmosis. There was no significant difference between nano-gold ELISA and PCR results, while there was a significant difference between ELISA and PCR (Table 4).

Discussion

Toxoplasmosis in pregnancy is a serious problem that necessitates the development of a more appropriate diagnostic method.^{36,37} SAG I is stage-specific, detected only in the tachyzoite stage. It is one of the most immunogenic *T. gondii* antigens and could represent an important indicator of active infection.^{38,39} The entire *Toxoplasma* soluble extract is used as the antigen in the majority of commercially accessible assays. However, due to the absence of a purified standardized

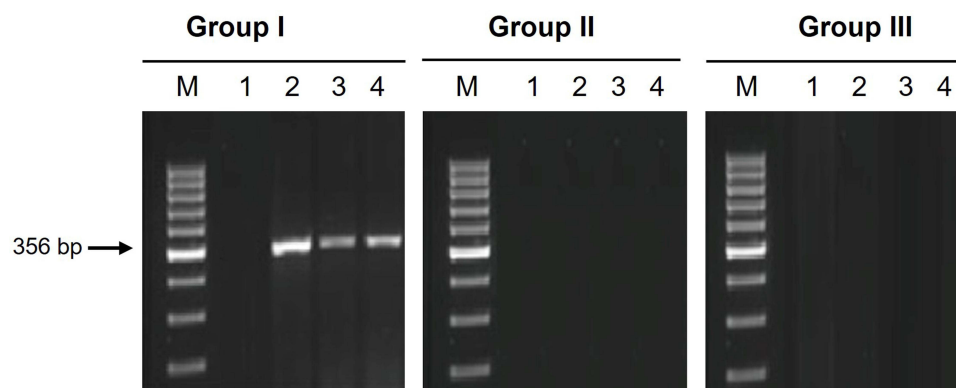


Figure 4 Molecular results of amplification of SAG I obtained from DNA isolated from blood samples by PCR in the different studied groups. M: molecular weight marker. G I: lane 1 was negative lanes 2, 3, 4 were positive. GII and III: All samples showing negative results.

Table 3 Diagnostic Values of PCR, Nano-Gold ELISA, and ELISA

	PCR	Nano-Gold ELISA	ELISA
Sensitivity%	97.3%	89.2	83.8
Specificity%	100%	94	88
PPV%	100%	91.7	83.8
NPV%	98%	92.2	88
Accuracy	98.9%	91.95	86.2
AUC	0.986	0.916	0.859
95% CI	0.96–1.0	0.85–0.98	0.77–0.95

Abbreviations: PPV, positive predictive value; NPV, negative predictive value; AUC, the area under the ROC curve; CI, coefficient interval.

antigen, the tests currently in use for the detection of specific anti-Toxoplasma antibodies differ in their abilities to detect anti-Toxoplasma immunoglobulins. We tried to improve the sensitivity and specificity of toxoplasmosis diagnosis. SAG 1 was used instead of the whole *Toxoplasma* antigen.

In the current work, the higher improvement in gold nanoparticle surface area, which encourages more antibodies to enter the antigen–antibody complex, is likely the source of the enhanced diagnostic values of ELISA using nanoparticles. Additional factors may be small size, high-binding capacity and rapid reaction. These properties allow nano-gold-based assays to be highly sensitive.⁴⁰ The use of nanomaterials is capable of assisting and improving the detection of antibodies, antigens, and DNA of *T. gondii*, with quick and easy handling techniques, presenting good sensitivity and specificity, indicating a potential to act in the screening and diagnosis in different stages of toxoplasmosis.⁴¹

Jalallou et al used ELISA for the detection of SAG1 *T. gondii* using specific IgG in human sera. Sensitivity was 88.4%, and specificity was 88%.⁴² They did not use IgM positive sera, while in our study this antigen was surveyed with IgM-positive sera. Another survey could detect 83% of positive IgG sera by SAG1 ELISA.⁴³ Differences in insensitivity

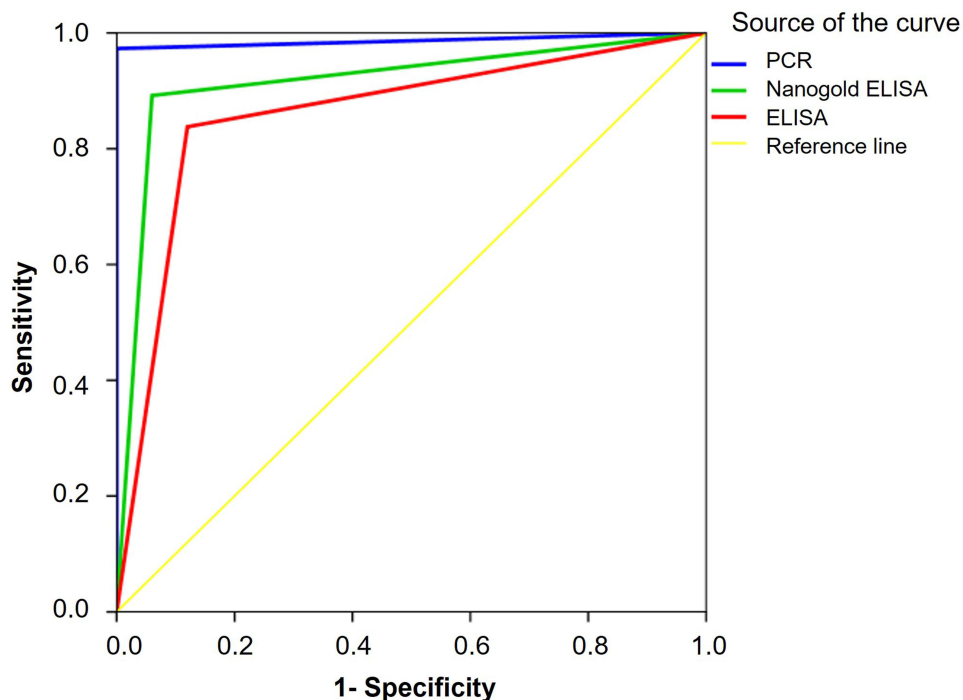


Figure 5 ROC curve for the performance of PCR, nano-gold ELISA and ELISA in diagnosis of toxoplasmosis.

Table 4 Comparing Areas Under the ROC Curve

Statistics	PCR ≠ Nano-Gold ELISA	PCR ≠ ELISA
Difference	0.07	0.127
Standard Error	0.0394	0.0468
z test	1.777	2.713
p-value	0.076	0.007

rates between our study and other research might be related to the purification of antigens.⁴⁴ Other nanoparticles (chitosan and silica gel) were used in the diagnosis of toxoplasmosis and the authors demonstrated that nano-particle ELISA outperformed traditional ELISA in terms of sensitivity and specificity.^{13,45}

Although serological testing has been the main diagnostic for toxoplasma infection, it has many drawbacks. DNA in blood and body fluids minimizes the problems faced when using serodiagnostic assays and makes diagnosis in challenging cases easier.⁴⁶ The use of PCR has made a significant improvement in both detection of acute disease in immunocompromised patients and the prenatal diagnosis of congenital toxoplasmosis.^{47,48} Another study reported that in asymptomatic newborns, postnatal PCR testing of the placental tissues for *Toxoplasma* may provide the only sign of infection. To increase the accuracy of detecting recent infection and/or fetal transmission, serological and molecular approaches should be combined.^{4,49} In our study, one sample was negative with PCR. Since quick processing was not possible, DNA samples were kept at -20°C before the amplification was carried out. This may be the cause of some test sensitivity lowering. On the other hand, the presence of inhibitors in the blood samples cannot be excluded despite the effective DNA extraction technique. Our results of SAG1 detection had some limitations; first, the study of SAG1-based nano-gold ELISA needs experimental skills; second, small numbers of samples were tested. Further study is needed to include a large number of confirmed *T. gondii* samples.

Our results showed that PCR and nano ELISA had a high specificity for *T. gondii* detection than traditional ELISA. Blood from people who were not infected with *T. gondii*, including those who had other parasitic infections, did not trigger a PCR reaction. SAG1 does not share any characteristics with other parasite antigens, which eliminates false-positive PCR results.

In conclusion, there was no significant difference between PCR and nano-gold ELISA using SAG1 regarding sensitivity and specificity. PCR and nano-gold ELISA using SAG1 appear to be effective methods for *T. gondii* diagnosis. We recommend nano-gold ELISA as a promising technique in the diagnosis of toxoplasmosis and survey studies. Also, using the polyclonal antibodies conjugated with gold nanoparticles as coating antibodies and conjugate antibodies in nano-gold ELISA may achieve better sensitivity and specificity.

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Disclosure

The authors report no conflicts of interest in this work.

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