Abstract: Toll-like receptors (TLR) play an eminent role in the regulation of immune responses to invading pathogens during sepsis. TLR genetic variants might influence individual susceptibility to developing sepsis. The current study aimed to investigate the association of genetic polymorphisms of the TLR2 and TLR4 with the risk of developing sepsis with both a pilot study and in silico tools. Different in silico tools were used to predict the impact of our SNPs on protein structure, stability, and function. Furthermore, in our prospective study, all patients matching the inclusion criteria in the intensive care units (ICU) were included and followed up, and DNA samples were genotyped using real-time polymerase chain reaction (RT-PCR) technology. There was a significant association between TLR2 Arg753Gln polymorphisms and sepsis under the over-dominant model ($p = 0.043$). In contrast, we did not find a significant difference with the TLR4 Asp299Gly polymorphism with sepsis. However, there was a significant association between TLR4 Asp299Gly polymorphisms and Acinetobacter baumannii infection which is quite a virulent organism in ICU ($p = 0.001$) and post-surgical cohorts ($p = 0.033$). Our results conclude that the TLR2 genotype may be a risk factor for sepsis in adult patients.

Keywords: TLR; polymorphism; infection; sepsis; septic shock

1. Introduction

Infection is one of the prominent causes of human morbidity and mortality, especially in patients requiring critical care [1,2]. Moreover, in intensive care units (ICUs), a serious complication of infection is sepsis and its maximal manifestation, septic shock [3]. Sepsis is an infection-induced life-threatening organ dysfunction with mortality rates reaching 20–70% [4,5].

Infectious diseases have been found to be a major selective pressure [6]. Despite the ambiguity of the precise etiology of sepsis, numerous studies have shown that gene polymorphisms have an important role in affecting individual susceptibility to sepsis [7]. Some polymorphisms of the innate immune system are supposed to mediate a predisposition to infectious complications including the outcome of patients with sepsis [8]. The
innate immune system is of crucial importance for both the direct defense against microorganisms and the activation of the adaptive immune system [9]. The innate immunity system is the main mediator of inflammation, and it recruits specific pattern recognition receptors (PRRs) capable of recognizing micro-organisms through identifying conserved pathogen-associated molecular patterns.

Toll-like receptors (TLR) are the most studied subtypes of pattern recognition receptors with their critical importance in the immune system [10,11]. Among the members of the TLR family, TLR2 and TLR4 are considered the most important PRRs that cover a wide range of antigenic determinants [12]. TLR4 has a distinctive ability to recognize a very wide range of microorganisms including Gram-negative bacteria through Lipopolysaccharide (LPS), in addition to many viruses and Fungi. Meanwhile, TLR2 is regarded as a key molecule in regulating our immune system with a crucial role in the recognition of Peptidoglycans of Gram-positive bacteria, in addition to different ligands of yeast, fungi, viruses, and parasites [12,13].

One of the most studied innate immunity polymorphisms is the TLR4 Asp299Gly (rs4986790) polymorphism, which interferes with TLR4 signal transduction; thus, it is supposed to affect host susceptibility to infections and microbial invasions [14]. Moreover, structural analysis of TLR4 Asp299Gly has revealed evidence of a resulted impairment in TLR4 binding to its ligands [15]. Meanwhile, one of the most important polymorphisms of TLR2 is Arg753Gln; the presence of this SNP was found to impair the signaling pathway of this key receptor [16], thus suggesting increased susceptibility to infections and sepsis.

Consequently, many studies have been conducted all over the world to reveal the prevalence of these SNPs and their impact on infection and sepsis susceptibility, but a varied pattern of prevalence was found for both TLR4 and TLR2 SNPs among different populations [13,17,18]. In addition, conflicting results were found regarding their impact on infection and sepsis susceptibility in different populations [19–21]. Therefore, a need was felt for further investigation on these issues. The usage of computational approaches in studying SNPs’ impact has gained momentum and importance in recent years [22–25]. Integrating the in silico approach with the experimental one provides great accuracy and depth to the analysis.

In this study, we aimed to investigate the possible role of TLR2 and TLR4 polymorphisms in affecting sepsis susceptibility and survival in critically ill patients in the Egyptian population using both in silico analysis and experimental methods.

2. Results

The study involved both a pilot study and in silico analysis. A scheme illustrating the layout of the study plan is shown in Figure 1.
2.1. In Silico Analysis

2.1.1. General Information: TLR2

TLR2 gene (ENSG00000137462) is a protein-coding gene located on 4q31.3. It is composed of five exons with a length of 26,564 nucleotides. It is located on Chromosome 4: 153684080-153710643 according to the Genome Reference Consortium Human Build 38 patch release 13 (GRCh38.p13) with NCBI Reference Sequence (NC_000004.12) (https://www.ncbi.nlm.nih.gov/gene/7097 (accessed on 29 August 2021)). There are eight transcripts for this gene (ensemble.org). This gene encodes Toll-like receptor 2 protein, a member of the Toll-like receptor (TLR) family. Figure 2A shows the subcellular localization of TLR2. The predicted network of protein–protein interactions of the TLR2 protein is shown in Figure 2B and its Gene Coexpression matrix (Figure 2C) shows coexpression with CD14, CLEC7A, and LY96 with scores of 0.611, 0.281, and 0.130, respectively (https://string-db.org (accessed on 29 August 2021)). Rs5743708 is an SNP located at Chromosome 4, position: 153705165 (forward strand) with two alleles (G and A). G is the ancestral allele and the minor allele frequency for A equals 0.01. This is a missense variant that causes the replacement of amino acid Arginine with amino acid Glutamine at position 753.

2.1.2. General Information: TLR4

TLR4 gene (ENSG00000136869) is a protein-coding gene located on 9q33.1. It is composed of four exons with a length of 20,333 nucleotides. It is located on Chromosome 9: 117704403-117724735 according to the Genome Reference Consortium Human Build 38 patch release 13 (GRCh38.p13) with NCBI Reference Sequence (NC_000009.12) (https://www.ncbi.nlm.nih.gov/gene/7099 (accessed on 29 August 2021)). There are four transcripts for this gene (ensemble.org). This gene encodes the Toll-like receptor 4 protein, a member of the Toll-like receptor (TLR) family as well. Figure 2D shows the subcellular localization of TLR4. The predicted network of protein–protein interactions of TLR4 protein is shown in Figure 2E and its Gene Coexpression matrix (Figure 2F) shows coexpression with LY86, CD14, and LY96 with scores of 0.301, 0.264, and 0.176, respectively (https://string-db.org (accessed on 29 August 2021)). Rs4986790 is an SNP located at Chromosome 9, position: 117713024 (forward strand) with three alleles (A, G, and T). A is the ancestral allele and the minor allele frequency for G equals 0.06. This is a missense mutation that causes the replacement of amino acid Aspartic acid with amino acid Glycine at position 299.

2.1.3. Predicting the Effect of SNPs on Protein Function

Five bioinformatics tools were used to predict the impact of rs4986790 and rs5743708 on the TLR4 and TLR2 proteins, respectively, to increase the accuracy of the results. For TLR4, all used bioinformatics tools predicted this variation to be neutral or benign as shown in Table 1. While for TLR2, all tools predicted this SNP to be damaging except SNPs and GO which predicted it to be neutral (Table 1). Figure 3 shows the structural and functional effects of SNPs.

Table 1. Predicting the effect of SNPs on protein function using bioinformatics tools.

| SNP     | Amino Acid Change | SIFT       | Polyphen2  | PANTHER          | PROVEAN   | SNPs and GO |
|---------|-------------------|------------|------------|------------------|-----------|-------------|
| rs4986790 | D299G             | Tolerated  | Benign     | probably benign  | Neutral   | Neutral     |
| rs5743708 | R753Q             | Deleterious| Probably Damaging | probably damaging | Deleterious | Neutral     |
Figure 2. Functional Analysis of TLR2 and TLR4 proteins. (A) Subcellular localization of the TLR2 protein. The gradient of green color indicates the degree of confidence (genecards.org) with (compartments.jensenlab.org) as the source of the image. (B) Predicted network of protein–protein interactions of the TLR2 protein.
Proteins are represented by nodes, while predicted associations are represented by edges that could be drawn with 7 colored lines that indicate different types of evidence. Redline fusion evidence, Light blue line—database evidence. Green line—neighborhood evidence. Blue line—co-occurrence evidence. Purple line—experimental evidence. Black line—coexpression evidence. Yellow line—text mining evidence. HMGB1: High mobility group protein B1, TIRAP: Toll/interleukin-1 receptor domain-containing adapter protein, VCAN: Versican core protein, HSPD1: 60 kDa heat shock protein, LY96: Lymphocyte antigen 96, CD14: Monocyte differentiation antigen CD14, TOLLIP: Toll-interacting protein, IRAK1: Interleukin-1 receptor-associated kinase 1, CLEC7A: C-type lectin domain family 7 member A, HSP90B1: Endoplasmin. STRING analysis (version 11.5). (C) TLR2 Gene Coexpression matrix. Predict association between protein functions, Color intensity shows the confidence level in the association between protein functions. TLR2 shows coexpression with CD14, CLEC7A, and LY96 with scores of 0.611, 0.281, and 0.130, respectively (https://string-db.org (accessed on 29 August 2021)). (D) Subcellular localization of TLR4 protein (genecards.org) with (compartments.jensenlab.org) as the source of the image. (compartments.jensenlab.org). (E) 1B predicted a network of protein–protein interactions of the TLR4 protein. TICAM1: TIR domain-containing adapter molecule 1, TICAM2: TIR domain-containing adapter molecule 2, TRAF6: TNF receptor-associated factor 6, LY86: Lymphocyte antigen 86. STRING analysis (version 11.5). (F) TLR4 Gene Coexpression matrix. TLR4 shows coexpression with LY86, CD14, and LY96 with scores of 0.301, 0.264, and 0.176, respectively (https://string-db.org (accessed on 29 August 2021)).
2.1.4. Identifying SNP Location on Protein Domains  
Using InterPro revealed that rs5743708 was found to be located on the Toll/Interleukin-1 Receptor Homology (TIR) Domain (InterPro entry: IPR000157) in the TLR2 protein which is an essential domain for protein function, while rs4986790 location was found to belong to a superfamily called Leucine-rich repeat domain superfamily (InterPro entry: IPR032675).

2.1.5. Prediction of Protein Stability with SNPs  
I-Mutant 2.0 web server analyzed the effects of rs5743708 and rs4986790 SNPs on the stability of TLR2 and TLR4 proteins, respectively, by calculating free energy change values (DDG) and the Reliability Index value (RI). For TLR4, rs4986790 was found to decrease stability with RI = 3 and DDG = 0.38 Kcal/mol. While for TLR2, rs5743708 was found to decrease stability with RI = 8 and DDG = −0.71 Kcal/mol.

![Figure 3](image_url)  
**Figure 3.** Functional and structural consequences of SNPs. (A) Predicting the impact of rs5743708 on TLR2 function—the score ranged from benign (0) to damaging (1) (B). Table showing the transcripts of rs5743708, allele (transcript allele), consequence type, amino acid fate, codons, and PolyPhen score. R: Arginine, Q: Glutamine (ensemble.org). (C) Predicting the impact of rs4986790 on TLR4 function the score ranges from benign (0) to damaging (1) (D). Table showing the transcripts of rs4986790, allele (transcript allele), consequence type, amino acid fate, codons, and PolyPhen score. D: Aspartate, V: valine, G: Glycine (ensemble.org).
2.1.6. Conservation Analysis

TLR2 and TLR4 proteins were analyzed by the ConSurf server to perform an evolutionary conservation analysis of their amino acid positions (Figures 4 and 5), respectively. In TLR2, position 753 (R753) was found to be an exposed and functional residue with high conservation. While in TLR4, position 299 (D299) was found to be an exposed and variable residue.

Figure 4. Evolutionary conservation analysis of TLR2 by Consurf.
Figure 5. Evolutionary conservation analysis of TLR4 by ConSurf.
2.1.7. Identifying the Structural Effects of SNPs

Using Project HOPE to analyze rs4986790 in TLR4, the new Glycine residue was found to differ in size and charge from the wild residue (Aspartic Acid) which could lead to a loss of interactions (Figure 6A). There was a difference in hydrophobicity too, which could cause loss of hydrogen bonds with possible disturbance of correct folding. Moreover, this replacement leads to an inability to form a Cysteine Bridge with its importance to protein stability, thus affecting the 3D structure of the protein and protein stability. In addition, Glycine flexibility affects the needed stability at that position. Meanwhile, analyzing rs5743708 in TLR2 revealed differences in size and charge between wild and mutant amino acids which could cause a loss of interactions (Figure 6B). Moreover, the different properties could lead to disturbance and elimination of (TIR) Domain function with its importance for protein function.

![Figure 6. HOPE illustration of mutation structural impacts](image)

2.2. Demographic and Microbiological Data

A total of seventy-five Egyptian unrelated patients were included in the study. All participants had developed an infection. The patients were followed up to assess sepsis and septic shock, and the demographic features and the clinical characteristics of ICU-admitted patients according to developing sepsis are presented in Table 2. The two groups had significant differences in age factor, APACHE score at admission, and some categories of admissions. Causative organisms are listed (Table 2). There was no statistically significant difference between any of the causative organisms and developing sepsis.

| Variables         | All     | No Sepsis | Sepsis | p-Value | OR (95% CI) |
|-------------------|---------|-----------|--------|---------|-------------|
| Number            | Number  | 75 (100%) | 48 (64.0%) | 27 (36.0%) |
| Age, years        | Mean ± SD | 60.0 ± 17.6 | 55.5 ± 18.9 | 68.0 ± 11.5 | 0.003       |
|                   | ≤40 years | 14 (18.7%) | 14 (29.2%) | 0 (0.0%) | 0.002       |
|                   | ≤60 years | 20 (26.7%) | 14 (29.2%) | 6 (22.2%) | 13.0 (0.67–252.6) |
|                   | >60 years | 41 (54.7%) | 20 (41.6%) | 21 (77.8%) | 30.41 (1.7–543.6) |
| Variables                        | All       | No Sepsis | Sepsis | p-value | OR (95% CI) |
|----------------------------------|-----------|-----------|--------|---------|-------------|
| **Sex**                          |           |           |        |         |             |
| Male                             | 47 (63.0%)| 31 (64.6%)| 16 (59.3%)| 0.64 | Reference   |
| Female                           | 28 (37.0%)| 17 (35.4%)| 11 (40.7%)| 1.25 | (0.48–3.30) |
| **Vital signs**                  |           |           |        |         |             |
| HR                               | 100.6 ± 21.3| 101.2 ± 20.8| 99.5 ± 22.4| 0.75 |             |
| MAP                              | 81.7 ± 25 | 85.4 ± 27.4| 75.2 ± 18.7| 0.09 |             |
| **Concomitant diseases**         |           |           |        |         |             |
| Diabetes                         |           |           |        |         |             |
| Positive                         | 29 (38.7%)| 17 (35.4%)| 12 (44.4%)| 0.44 | 1.46 (0.56–3.82) |
| Hypertension                     |           |           |        |         |             |
| Positive                         | 42 (56.0%)| 26 (54.2%)| 16 (59.3%)| 0.67 | 1.23 (0.47–3.20) |
| Vascular disease                 |           |           |        |         |             |
| Positive                         | 27 (36.0%)| 15 (31.3%)| 12 (44.4%)| 0.25 | 1.76 (0.66–4.66) |
| Chronic lung disease             |           |           |        |         |             |
| Positive                         | 6 (8.0%) | 5 (10.4%)| 1 (3.7%) | 0.41 | 0.33 (0.04–2.99) |
| Chronic liver disease            |           |           |        |         |             |
| Positive                         | 7 (9.3%) | 2 (4.2%) | 5 (18.5%)| 0.09 | 5.23 (0.94–29.10) |
| Chronic renal disease            |           |           |        |         |             |
| Positive                         | 17 (22.7%)| 10 (20.8%)| 7 (25.9%)| 0.61 | 1.33 (0.44–4.02) |
| **ICU assessment**               |           |           |        |         |             |
| APACHE score                     | Mean ± SD | 17.4 ± 8.3| 15.8 ± 6.0| 20.3 ± 10.8 | 0.024 |
| Glasgow scale                    | Mean ± SD | 9.8 ± 4.4 | 9.4 ± 4.1 | 10.5 ± 4.8 | 0.29 |
| Length of stay, days             | Mean ± SD | 19.7 ± 15.9| 17.6 ± 11.8| 23.4 ± 21.1 | 0.13 |
| **Consequence**                  |           |           |        |         |             |
| Discharge                        | 27 (36.0%)| 19 (39.58%)| 8 (29.63%)| 0.69 | Reference |
| Transferred                      | 5 (6.7%) | 3 (6.25%)| 2 (7.40%)| 1.58 (0.22–11.3) |
| Death                            | 43 (57.3%)| 26 (54.17%)| 17 (62.96%)| 1.55 (0.56–4.34) |
| **OS, days**                     | Mean ± SD | 19.6 ± 17.2| 17.5 ± 13.4| 22.8 ± 22 | 0.33 |
| **Admission category**           |           |           |        |         |             |
| Renal                            |           |           |        |         |             |
| Positive                         | 2 (2.7%) | 1 (2.1%) | 1 (3.7%) | 0.67 | 1.81 (0.11–30.1) |
| Cardiovascular                   |           |           |        |         |             |
| Positive                         | 3 (4%)  | 2 (4.2%) | 1 (3.7%) | 0.92 | 0.88 (0.08–10.2) |
| Infection                        |           |           |        |         |             |
| Positive                         | 21 (28%)| 7 (14.6%)| 14 (51.8%)| 0.001 | 7.23 (2.42–21.6) |
| Neurology                        |           |           |        |         |             |
| Positive                         | 20 (26.7%)| 17 (35.4%)| 3 (11.1%)| 0.022| 0.27 (0.07–1.03) |
| Post-surgical                    |           |           |        |         |             |
| Positive                         | 11 (14.7%)| 6 (12.5%)| 5 (18.5%)| 0.47 | 1.59 (0.44–5.80) |
| Respiratory                      |           |           |        |         |             |
| Positive                         | 10 (13.3%)| 10 (20.8%)| 0 (0.0%) | 0.011| 0.07 (0.00–1.19) |
| Trauma                           |           |           |        |         |             |
| Positive                         | 3 (4%)  | 3 (6.3%) | 0 (0.0%) | 0.54 | 0.24 (0.01–4.75) |
| Other causes                     |           |           |        |         |             |
| Positive                         | 5 (6.7%) | 2 (4.2%) | 3 (11.1%)| 0.24 | 2.88 (0.45–18.4) |
| **Variables**                    |           |           |        |         |             |
| Causative organism in culture    |           |           |        |         |             |
| Enterobacter spp.                | Positive  | 6 (6.3%) | 3 (6.3%) | 3 (11.1%)| 0.45 | 1.88 (0.35–10.01) |
| Acinetobacter baumannii          | Positive  | 11 (11.5%)| 7 (14.6%)| 4 (14.8%)| 0.97 | 1.02 (0.27–3.85) |
| Candida albicans                 | Positive  | 3 (3.1%) | 1 (2.1%) | 2 (7.4%) | 0.25 | 3.76 (0.32–43.53) |
| Escherichia coli                 | Positive  | 15 (15.8%)| 11 (22.9%)| 4 (14.8%)| 0.40 | 0.59 (0.17–2.06) |
| Gram negative bacilli            | Positive  | 3 (3.1%) | 2 (4.2%) | 1 (3.7%) | 0.92 | 0.90 (0.08–10.45) |
| Klebsiella pneumoniae            | Positive  | 20 (21.1%)| 13 (27.1%)| 7 (25.9%)| 0.91 | 0.94 (0.32–2.75) |
| Pseudomonas aeruginosa           | Positive  | 12 (12.6%)| 6 (12.5%)| 6 (22.2%)| 0.27 | 2.00 (0.57–6.96) |
| Staph spp.                       | Positive  | 17 (17.9%)| 12 (25.0%)| 5 (18.5%)| 0.52 | 0.68 (0.21–2.20) |
| Streptococcus spp.               | Positive  | 4 (4.2%) | 3 (6.25%)| 1 (3.7%) | 0.63 | 0.58 (0.06–5.84) |
| Aeromonas hydrophila             | Positive  | 1 (1.1%) | 1(2.1%) | 0 (0%) | 1.00 | 0.58 (0.02–14.6) |
| Proteus spp.                     | Positive  | 1 (1.1%) | 1 (2.1%) | 0 (0%) | 1.00 | 0.58 (0.02–14.6) |
| Citrobacter spp.                 | Positive  | 1 (1.1%) | 1 (2.1%) | 0 (0%) | 1.00 | 0.58 (0.02–14.6) |
| Serratia spp.                    | Positive  | 1 (1.1%) | 1(2.1%) | 0 (0%) | 1.00 | 0.58 (0.02–14.6) |

Data are shown as a number (percentage) or number ± standard deviation. HR: heart rate in beats per minute; MAP: mean arterial pressure in mmHg; OS: Overall survival. Chi-square ($\chi^2$) or Fisher’s exact tests were used for qualitative variables and student’s t-test was used for quantitative attributes. OR (95% CI), odds ratio, and confidence interval. Statistical analysis at p-value < 0.05.
2.3. Allele Frequencies of TLR2 and TLR4 Genes in the Study Population

Genotype and allele frequencies for TLR2 and TLR4 were detailed in Table 3. For TLR4, the frequency of wild-type genotype AA was 91%, while the heterozygous genotype AG was 8%, and the mutant genotype GG was 1%. The genotype frequencies followed the genotype frequencies expected by Hardy–Weinberg equilibrium ($p > 0.05$). For TLR2 the frequency of wild-type genotype GG was 92%, while the heterozygous genotype GA was 5%, and the mutant genotype AA was 3%. The genotype frequencies did not follow the genotype frequencies expected by Hardy–Weinberg equilibrium ($p < 0.05$).

Table 3. Genotype and allele frequencies of TLR2 and TLR4 genes in the study population according to developing or not developing sepsis.

| Variables | TLR2 (rs5743708) | TLR4 (rs4986790) |
|-----------|------------------|------------------|
| Genotype frequencies | | |
| All | Non-Septic | Septic | p-Value | All | Non-Septic | Septic | p-Value |
| A/A | 2 (3) | 1 (2) | 1 (4) | 0.22 | 68 (91) | 45 (94) | 23 (85) | 0.20 |
| G/A | 4 (5) | 1 (2) | 3 (11) | 6 (8) | 2 (4) | 4 (15) | |
| G/G | 69 (92) | 46 (96) | 23 (85) | 1 (1) | 1 (2) | 0 (0) | |

| Variables | Allele frequencies | | |
|-----------|-------------------|---|---|
| A | 8 (5) | 3 (3) | 5 (9) | 0.10 | 142 (95) | 92 (96) | 49 (91) | 0.39 |
| G | 142 (95) | 93 (97) | 49 (91) | 8 (5) | 4 (4) | 4 (7) | |
| $P_{HWE}$ | 0.009 | 0.032 | 0.180 | 0.180 | 0.063 | 1.00 | |

Data are shown as a number (percentage). Fisher’s Exact tests were performed. Statistical analysis at $p$ value $< 0.05$.

Genotype association models for the risk of sepsis were analyzed and a significant association was found between TLR2 Arg753Gln SNP and sepsis under the over dominant model ($p = 0.043$), but in the TLR4 polymorphism this difference did not reach statistical significance (Table 4).

Table 4. Genotype association models for sepsis risk assessment.

| Model | Genotype | Non-Septic | Septic | Adjusted OR (95% CI) * | p-Value |
|-------|----------|------------|--------|------------------------|--------|
| TLR2  | Codominant b | G/G | 46 (95.8%) | 23 (85.2%) | Reference |
|       |          | A/G | 1 (2.1%) | 3 (11.1%) | 11.42 (0.84–155.32) | 0.12 |
|       |          | A/A | 1 (2.1%) | 1 (3.7%) | 1.65 (0.09–29.49) | |
|       | Dominant | G/G | 46 (95.8%) | 23 (85.2%) | Reference | 0.07 |
|       |          | A/G-A/A | 2 (4.2%) | 4 (14.8%) | 5.34 (0.77–36.96) | |
|       | Recessive | G/G-A/G | 47 (97.9%) | 26 (96.3%) | Reference | 0.79 |
|       |          | A/A | 1 (2.1%) | 1 (3.7%) | 1.48 (0.09–25.46) | |
|       | Over-dominant | G/G-A/A | 47 (97.9%) | 24 (88.9%) | Reference | 0.043 |
|       |          | A/G | 1 (2.1%) | 3 (11.1%) | 11.27 (0.83–152.94) | |
|       | Log-additive | — | — | — | 2.42 (0.61–9.56) | 0.18 |
| TLR4  | — | — | — | — | — |

Int. J. Mol. Sci. 2022, 23, 10982
Table 4. Cont.

| Model       | Genotype | Non-Septic | Septic       | Adjusted OR (95% CI) a | p-Value |
|-------------|----------|------------|--------------|------------------------|---------|
| Codominant b | A/A      | 45 (93.8%) | 23 (85.2%)  | Reference              | 0.11    |
|             | A/G      | 2 (4.2%)   | 4 (14.8%)   | 7.23 (0.77–67.86)      |         |
|             | G/G      | 1 (2.1%)   | 0 (0%)      | 0.00 (0.00-NA)         |         |
| Dominant    | A/A      | 45 (93.8%) | 23 (85.2%)  | Reference              | 0.16    |
|             | A/G-G/G  | 3 (6.2%)   | 4 (14.8%)   | 3.68 (0.57–23.57)      |         |
| Recessive   | A/A-A/G  | 47 (97.9%) | 27 (100%)   | Reference              | 0.36    |
|             | G/G      | 1 (2.1%)   | 0 (0%)      | 0.00 (0.00-NA)         |         |
| Over-dominant | A/G      | 46 (95.8%) | 23 (85.2%)  | Reference              | 0.06    |
|             | A/G      | 2 (4.2%)   | 4 (14.8%)   | 7.49 (0.79–71.02)      |         |
| Log-additive | —        | —          | —           | 1.90 (0.45–8.04)       | 0.37    |

Values are shown as numbers (%). Chi-square ($\chi^2$) or Fisher’s exact tests were used. OR (95% CI), odds ratio, and confidence interval. a adjusted for confounding factors (age and sex). b represented both heterozygote and homozygote comparison models.

2.4. TLR2 and TLR4 Polymorphisms in Relation to Clinical and Laboratory Data

The association of single nucleotide polymorphisms (SNPs) with clinical and laboratory characteristics data is studied in Table 5. There was a statistically significant association between the TLR4 polymorphism (rs4986790) and infection with *Acinetobacter baumannii* ($p = 0.001$) and infection with undetermined Gram (−) bacilli. Moreover, a statistically significant association was found between the TLR4 polymorphism (rs4986790) and post-surgical patients’ admission category referred to ICU ($p = 0.033$). In addition, the TLR4 polymorphism (rs4986790) had a significant association with the selection of Azithromycin as an empirical antibiotic ($p = 0.003$), and Imipenem antibiotic ($p = 0.024$), while the TLR2 polymorphism (rs5743708) had an association with the selection of Teicoplanin ($p < 0.001$) and with Ampicillin + Sulbactam ($p = 0.022$). The selected empirical antibiotic depended on patient status and the severity of infection.

Table 5. Analysis for the association of variants with clinical and laboratory characteristics.

| Variables                  | TLR2 (rs5743708) p-Value | TLR4 (rs4986790) p-Value |
|----------------------------|--------------------------|-------------------------|
| Demographic                |                          |                         |
| Age, years                 | 0.84 (0.65–1.05)         | 0.99 (0.77–1.26)        |
| Sex                        | 0.27 (0.17–0.41)         | 0.71 (0.52–0.95)        |
| Vital signs                |                          |                         |
| HR, beats/min              | 0.27 (0.16–0.40)         | 0.47 (0.27–0.82)        |
| MAP, mm Hg                 | 0.70 (0.51–0.95)         | 0.84 (0.61–1.14)        |
| SBP, mm Hg                 | 0.70 (0.50–0.95)         | 0.86 (0.64–1.12)        |
| DBP, mm Hg                 | 0.80 (0.60–1.04)         | 0.92 (0.71–1.19)        |
| Concomitant diseases       |                          |                         |
| Diabetes                   | 0.80 (0.60–1.06)         | 0.08 (0.05–0.14)        |
| Hypertension               | 0.95 (0.74–1.22)         | 0.06 (0.03–0.19)        |
| Vascular dis               | 0.14 (0.07–0.28)         | 0.43 (0.24–0.73)        |
| Chronic lung disease       | 0.75 (0.55–1.01)         | 0.71 (0.50–1.00)        |
| Chronic liver disease      | 0.49 (0.33–0.71)         | 0.67 (0.45–0.99)        |
| Chronic renal disease      | 0.31 (0.19–0.52)         | 0.80 (0.57–1.13)        |
| Variables | TLR2 (rs5743708) | TLR4 (rs4986790) |
|-----------|-----------------|-----------------|
|           | p-Value         | p-Value         |
| ICU assessment |                 |                 |
| APACHE score | 0.75            | 0.70            |
| Glasgow scale | 0.89            | 0.24            |
| Length of stay | 0.84            | 0.36            |
| Sepsis | 0.22            | 0.20            |
| Septic shock | 0.74            | 0.44            |
| Death | 0.75            | 0.46            |
| Overall survival | 0.52            | 0.06            |
| Renal | 0.91            | 0.90            |
| Cardiovascular | 0.87            | 0.25            |
| Infection | 0.07            | 0.38            |
| Neurology | 0.30            | 0.77            |
| Post-surgical | 0.70            | 0.033           |
| Respiratory | 0.22            | 0.55            |
| Trauma | 0.87            | 0.85            |
| Other causes | 0.79            | 0.75            |
| Biochemical data |                 |                 |
| WBC, $\times 10^3$ cells/$\mu$L | 0.56 | 0.16 |
| HB, g% | 0.08            | 0.31            |
| Creatinine, mg/dL | 0.98            | 0.24            |
| Causative organism |             |                 |
| Enterobacter spp. | 0.07            | 0.05            |
| Acinetobacter spp. | 0.70            | 0.001           |
| Candida spp. | 0.08            | 0.85            |
| E. coli | 0.44            | 0.38            |
| Gram (-) bacilli | 0.87            | <0.001          |
| Klebsiella spp. | 0.30            | 0.69            |
| Pseudomonas spp. | 0.73            | 0.90            |
| Staph spp. | 0.63            | 0.80            |
| Streptococcus spp. | 0.83            | 0.80            |
| Aeromonas spp. | 0.95            | 0.94            |
| Proteus spp. | 0.95            | 0.94            |
| Citrobacter spp. | 0.95            | 0.94            |
| Serratia spp. | 0.95            | 0.94            |
| Type of culture |                 |                 |
| Blood | 0.68            | 0.76            |
| Sputum | 0.77            | 0.29            |
| Urine | 0.47            | 0.41            |
| Pus | 0.35            | 0.73            |
| CSF | 0.95            | 0.94            |
Table 5. Cont.

| Variables               | TLR2 (rs5743708) | p-Value | TLR4 (rs4986790) | p-Value |
|-------------------------|------------------|---------|------------------|---------|
| No of infections        | 0.48             |         | 0.94             |         |
| Empirical antibiotic    |                  |         |                  |         |
| No of antibiotics       | 0.05             |         | 0.77             |         |
| Cefoperazone            | 0.91             |         | 0.90             |         |
| Ceftazidime             | 0.57             |         | 0.90             |         |
| Levofloxacin            | 0.47             |         | 0.14             |         |
| Cefepime                | 0.16             |         | 0.23             |         |
| Ampicillin + sulbactam  | 0.022            |         | 0.70             |         |
| Imipenem                | 0.22             |         | 0.024            |         |
| Meropenem               | 0.95             |         | 0.94             |         |
| Ertapenem               | 0.75             |         | 0.69             |         |
| Azithromycin            | 0.95             |         | 0.003            |         |
| Rifampicin              | 0.95             |         | 0.94             |         |
| Teicoplanin             | <0.001           |         | 0.94             |         |
| Cefotaxime              | 0.57             |         | 0.90             |         |
| Piperacillin            | 0.95             |         | 0.94             |         |

Chi-square ($\chi^2$) or Fisher’s exact tests were used for qualitative variables and student’s t-test was used for quantitative attributes. Statistical analysis at $p$ value < 0.05.

2.5. Multivariate Analysis in Relation to Developing Sepsis

A multivariate analysis was performed to determine which variable was independently associated with the risk of sepsis (Table 6). Only age was found to be independently associated with the risk of sepsis with a $p$-value of 0.009.

Table 6. Multivariate analysis for the risk of sepsis in ICU-admitted patients.

| Risk Factors   | OR   | 95% CI (Lower) | 95% CI (Upper) | p-Value |
|----------------|------|----------------|----------------|---------|
| Age            | 0.940| 0.897          | 0.984          | 0.009   |
| Sex (female)   | 0.473| 0.116          | 1.925          | 0.30    |
| HR, beats/min  | 1.003| 0.974          | 1.033          | 0.83    |
| MAP, mm Hg     | 0.898| 0.654          | 1.233          | 0.51    |
| SBP, mm Hg     | 1.022| 0.915          | 1.142          | 0.70    |
| DBP, mm Hg     | 1.123| 0.901          | 1.400          | 0.30    |
| WBC, ×10^3 cells/µL | 0.926 | 0.854 | 1.003          | 0.06    |
| HB, g%         | 0.875| 0.666          | 1.150          | 0.34    |
| Creatinine, mg/dL | 0.938 | 0.724 | 1.217          | 0.63    |
| APACHE score   | 0.942| 0.815          | 1.088          | 0.42    |
| Glasgow scale  | 0.962| 0.800          | 1.156          | 0.68    |
| Length of stay | 0.962| 0.917          | 1.008          | 0.11    |
| TLR2 (A/G)     | 0.082| 0.001          | 6.474          | 0.26    |
| TLR2 (G/G)     | 1.939| 0.084          | 44.584         | 0.68    |
| TLR4 (A/G)     | 0.090| 0.005          | 1.785          | 0.11    |
| TLR4 (G/G)     | NA   | NA             | NA             | 1.00    |

OR: odds ratio; CI: confidence interval. Binary logistic regression analysis was performed.
2.6. Survival Analysis

Survival analysis was performed with the usage of Log-rank, Breslow, and Tarone–Ware tests which showed significance only with the length of stay (0.001, 0.001, and 0.001), respectively, and with the post-surgical category of admission with a log-rank test (0.03), as shown in Table 7.

Table 7. Survival analysis in ICU-admitted patients.

| Variables                  | Overall Comparisons | Log Rank | Breslow | Tarone–Ware |
|----------------------------|---------------------|----------|---------|-------------|
| Demographic data           |                     |          |         |             |
| Age                        | 0.44                | 0.36     | 0.35    |             |
| Sex                        | 0.23                | 0.50     | 0.36    |             |
| Vital signs                |                     |          |         |             |
| HR                         | 0.61                | 0.99     | 0.84    |             |
| MAP                        | 0.86                | 0.69     | 0.75    |             |
| SBP                        | 0.45                | 0.46     | 0.44    |             |
| DBP                        | 0.63                | 0.64     | 0.64    |             |
| Concomitant disease        |                     |          |         |             |
| Diabetes                   | 0.87                | 0.62     | 0.85    |             |
| Hypertension               | 0.12                | 0.26     | 0.17    |             |
| Vascular disease           | 0.39                | 0.28     | 0.28    |             |
| Chronic liver disease      | 0.58                | 0.50     | 0.50    |             |
| Chronic renal disease      | 0.42                | 0.55     | 0.48    |             |
| ICU assessment             |                     |          |         |             |
| APACHE score               | 0.81                | 0.84     | 0.76    |             |
| Glasgow scale              | 0.51                | 0.54     | 0.57    |             |
| Length of stay             | <0.001              | <0.001   | <0.001  |             |
| Sepsis                     | 0.91                | 0.82     | 0.78    |             |
| Septic shock               | 0.94                | 0.69     | 0.74    |             |
| No empirical drug          | 0.06                | 0.09     | 0.06    |             |
| Admission category         |                     |          |         |             |
| Renal                      | 0.53                | 0.54     | 0.54    |             |
| Cardiovascular             | 0.30                | 0.56     | 0.44    |             |
| Infection                  | 0.79                | 0.86     | 0.82    |             |
| Neurology                  | 0.33                | 0.29     | 0.31    |             |
| Post-surgical              | 0.030               | 0.09     | 0.06    |             |
| Respiratory                | 0.61                | 0.74     | 0.62    |             |
| Trauma                     | 0.32                | 0.39     | 0.37    |             |
| Other causes               | 0.26                | 0.59     | 0.40    |             |
| Lab data                   |                     |          |         |             |
| WBC, $\times 10^3$ cells/µL| 0.66                | 0.93     | 0.84    |             |
| HB, g%                     | 0.51                | 0.69     | 0.61    |             |
| Creatinine, mg/dL          | 0.71                | 0.42     | 0.55    |             |
| No of infection            | 0.48                | 0.47     | 0.54    |             |
| Molecular analysis         |                     |          |         |             |
| TLR2                       | 0.38                | 0.25     | 0.27    |             |
| TLR4                       | 0.63                | 0.39     | 0.43    |             |
| Combined                   | 0.12                | 0.35     | 0.22    |             |

Survival time is shown as mean and standard error, HR: Hazard ratio, CI; confidence interval. Log-rank, Breslow, and Tarone–Ware tests were used to find Kaplan–Meier estimates for survival. Quantitative variables were categorized by their medians.
In addition, Cox regression analysis was applied to the data to determine if any of these variables were independently associated with the duration of survival (Table 8). Hazard risk for TLR2 was 1.89 and hazard risk for TLR4 was 2.25 but these results did not reach significance, so the effect of TLR gene status during time remained constant.

Table 8. Multivariate analysis for the risk of mortality in ICU-admitted patients.

| Variables          | HR   | 95% CI      | p-Value |
|--------------------|------|-------------|---------|
| **Demographic data** |      |             |         |
| Age                | 1.90 | (0.47–7.57) | 0.36    |
| Sex                | 0.42 | (0.09–1.93) | 0.27    |
| **ICU assessment** |      |             |         |
| APACHE score       | 1.41 | (0.31–6.24) | 0.65    |
| Glasgow scale      | 1.86 | (0.38–8.89) | 0.44    |
| Septic shock       | 0.55 | (0.10–2.90) | 0.48    |
| No empirical drug  | 0.32 | (0.04–2.06) | 0.62    |
| No of infection    | 1.76 | (0.19–16.23)| 0.23    |
| **Molecular analysis** |     |             |         |
| TLR2               | 1.89 | (0.08–43.58)| 0.69    |
| TLR4               | 2.25 | (0.48–10.43)| 0.30    |

HR: hazard risk, CI: confidence interval. Cox Proportional Hazard Regression analysis was performed.

3. Discussion

The remarkable importance of TLR2 and TLR4 in our immune system and in modulating our response to infection suggested potential roles of their important variants, Arg735Gln and Asp299Gly, in increasing susceptibility to infection and sepsis as well. Different bioinformatics approaches were utilized in our analysis. Investigating the impacts of our variants depended on five various tools with various approaches to achieve a high robustness and effectiveness. While rs4986790 was predicted to possess a benign impact on TLR4 by all tools, rs5743708 was predicted by all tools except SNPs and GO to possess a damaging impact on TLR2. Moreover, the SNPs’ positions on the domains of their proteins were determined by InterPro, revealing the presence of rs5743708 on the important TIR domain. The TIR domain has a crucial role in the activation of TLR pathways [26]. Therefore, it is anticipated that this mutation could affect its protein function. In addition, since protein function and structure are critically dependent on its stability [27], the impacts of rs5743708 and rs4986790 on their proteins’ stability were investigated revealing how proteins’ stability was reduced by these SNPs. Furthermore, concerning the relationship between high scores of conservation and functionally significant residues [28], the conservation analysis was intended to anticipate those SNPs which could affect the significant functions. Rs5743708 of TLR2 was found to be a functional residue with high conservation. On the contrary, rs4986790 of TLR4 was found to be a variable residue. In addition, both rs5743708 and rs4986790 were anticipated to induce structural impacts on TLR2 and TLR4, respectively using the HOPE bioinformatics server.

In our prospective study, the genotype frequencies for TLR4 were in accordance with Hardy–Weinberg equilibrium. On the contrary, the genotype frequencies for TLR2 were not in accordance with the Hardy–Weinberg equilibrium, and this aberrant result was also found by Saleh et al. in the Egyptian population, in his study about Toll-like receptor-2 polymorphisms and the susceptibility to pulmonary and peritoneal tuberculosis [29] which may require further investigation. The different prevalence of these SNPs between different populations have been steadily observed by different researchers [13,17,18] with obvious differences between Asian, African, and European ethnicities for both SNPs. These different distribution patterns between populations were suspected to be responsible for different susceptibility patterns to infectious diseases and other serious diseases such as coronary artery disease and type 2 Diabetes as well [18,30].
In our study, there was a significant association between TLR2 Arg753Gln polymorphism and sepsis under the over-dominant model \( (p = 0.043) \), while the TLR4 polymorphism did not show such significance. Some other investigators reached the same results in some populations despite the observed conflict between studies. A meta-analysis study conducted by Gao and colleagues found an association in this study between Arg753Gln SNP and the risk of sepsis among critically ill adult patients in Europe. Meanwhile, this study also shed light on the issue of the conflicting results regarding the TLR2 polymorphism and developing sepsis [21]. The TLR4 polymorphism studies also showed conflicting results; a study conducted in France by Lorenz et al. found that the Asp299Gly and Thr399Ile polymorphisms of TLR4 might potentially be linked to Gram-negative septic shock [19]. On the contrary, some studies showed an absence of association between TLR4 SNP and sepsis; a study conducted by Kumpf et al. found no association between Asp299Gly and Thr399Ile polymorphisms of TLR4 and the incidence of sepsis syndrome or the type of organisms causing surgical infection in German adults [20]. In addition, another study by Shan Xo et al. in Wenzhou found that the Asp299Gly and Thr399Ile polymorphisms may not correlate with susceptibility to sepsis in Chinese Han children [31]. These conflicting results can be seen frequently among different ethnic groups in these types of genetic association studies investigating diseases that depend on several genetic factors [32], as sepsis is believed to be initiated and augmented by multiple genes and there is no full control over sepsis by a single gene [33,34]. Consequently, the various frequency of different SNPs in different ethnic groups, and the difference in the penetration and the effect of SNPs because of other factors such as gender or age variations in different studies could explain these conflicting results among different populations.

Developing infection with *Acinetobacter baumannii* was found to have a statistically significant association with the TLR4 polymorphism \( (p = 0.001) \). This finding is in agreement with a recent study conducted by Chatzi et al. who found that the Asp299Gly and Thr399Ile polymorphisms of TLR4 could play an essential role in developing multidrug resistance to *Acinetobacter baumannii* in CNS infections [35]. In addition, other researchers have confirmed the role of TLR4 in *Acinetobacter baumannii* infection in vitro and in vivo and found that the production of IL-8 by epithelial A549 cells in the human lung as a response to *Acinetobacter baumannii* required both TLR2 and TLR4 [36]. However, other studies showed that the recognition of *Acinetobacter baumannii* depends on TLR4 rather than TLR2, as TLR4 is the dominant receptor in this type of recognition. Knapp et al. found that TLR4-deficient mice, not TLR2-deficient mice (with intranasal inoculation of *Acinetobacter baumannii* Lipopolysaccharides) showed the impaired production of TNFα in bronchi alveolar lavage fluid and the impaired recruitment of polymorph nuclear cells, compared with Wild Type mice [37]. Moreover, Kim et al. found that the production of *Acinetobacter baumannii*-induced cytokines was impaired with TLR4-deficient bone marrow-derived macrophages or dendritic cells, while it was not the case with TLR2-deficient macrophages [38]. Besides, Erridge et al. found that the activation of human monocytes (resulting from phenol water re-extracted Lipopolysaccharides from *Acinetobacter baumannii*) was the responsibility of the TLR4 signaling pathway [39]. This association between the TLR4 polymorphism and this virulent bacterium could allow proper management and prevention measures where high rates of *Acinetobacter baumannii* infection are found. This could be an important step towards the individualization of host susceptibilities towards virulent microorganisms in intensive care units.

Our study also found a significant association between the TLR4 polymorphism (rs4986790) and the post-surgical category among patients referred to ICU. This role of the TLR4 polymorphism in post-surgery was investigated by a clinical study conducted by Koch and colleagues who found that the presence of a TLR4 polymorphism influenced the immune–endocrine stress response which resulted from the systemic inflammation caused by major surgery. They found decreased serum concentrations of ACTH, IL-8, IL-10, and GM-CSF postoperatively in those surgical patients who carried that polymorphism [40]. This might explain this significant association found in our study.
The multivariate analysis was also performed to analyze the effects of our variables on the development of sepsis syndrome, but it was only the age factor that was found to have an independent association with the risk of sepsis in our study group. The age factor is a well-identified risk factor for developing this syndrome [41].

Survival analysis found that the length of stay and the surgical category of admission had a significant association with time of survival in intensive care units. Our results are in agreement with several studies that found an association between prolonged ICU stay and higher hospital mortality as well. Those patients, with an ICU length of stay of 14 days or longer, were found to have a mortality rate of more than 50% [42,43].

Overall, our study is characterized by the usage of both experimental and in silico methods. Our investigation showed promising results regarding the analysis of the role of TLRs variants in infection and sepsis. However, our study had its limitations as in most genetic polymorphism studies, as the number of patients carrying the variant alleles was relatively small due to the small number of these polymorphisms in the general population. As a result, there is a need for multi-center studies conducted on a larger scale to validate these findings.

4. Materials and Methods

4.1. Ethics Statement

The study protocol was approved by Scientific Research Ethics Commission at Suez Canal University (reference No. 201709MH1). All subjects or their next of kin gave informed consent before inclusion in the study.

4.2. In Silico Analysis

4.2.1. General Information

National Center for Biotechnology Information (NCBI) and Ensembl databases were used to retrieve general information about TLR2 and TLR4 genes. Subcellular localization was retrieved from compartments.jensenlab.org mainly and geneCards.org. Gene coexpression and predicted protein–protein interactions were obtained from the String Biological database. General information about rs5743708 and rs4986790 were brought from the dbSNP and Ensembl databases. (https://web.expasy.org (accessed on 29 August 2021)) was used for retrieving data about the variants' effect on sequences of our proteins with these data gained from UniProtKB/Swiss-Prot databases.

4.2.2. Predicting the Effect of SNPs on Protein Function

Five bioinformatics tools were used to predict the effect of SNPs on protein function to increase the strength and accuracy of results; 1-SIFT (Sorting Intolerant from Tolerant) (https://sift.bii.a-star.edu.sg/ (accessed on 30 August 2021)). SIFT depends on sequence homology in addition to the physical properties of amino acids to predict the effect of missense mutations on protein function [44]. 2-PolyPhen-2 (Polymorphism Phenotyping v2) (http://genetics.bwh.harvard.edu/pph2 (accessed on 30 August 2021)). PolyPhen-2 uses comparative and physical approaches to predict the effect of amino acid substitution [45]. 3-PANTHER (Protein Analysis Through Evolutionary Relationship) (http://www.pantherdb.org/tools/csnpScoreForm.jsp (accessed on 30 August 2021)). This method depends on calculating the evolutionary preservation of an amino acid to predict the likelihood that a nonsynonymous SNP could cause a functional impact on the protein [46]. 4- PROVEAN (Protein Variation Effect Analyzer) (http://provean.jcvi.org/seq_submit.php (accessed on 30 August 2021)). PROVEAN uses blast hits to calculate the delta alignment score and computes the PROVEAN score finally with a cutoff at −2.5 [47]. 5-SNPs and GO (https://snps.biofold.org/snps-and-go/snps-and-go.html (accessed on 30 August 2021)). SNPs and GO depend on protein functional annotation to predict the impact of variations [48].
4.2.3. The Identification of SNP Location on Protein Domains

The locations of SNPs on conserved domains on TLR2 and TLR4 proteins were identified using the InterPro bioinformatics tool (https://www.ebi.ac.uk/interpro/ (accessed on 30 August 2021)), a bioinformatics tool that could perform functional analysis of protein and identify domains and functional sites [49].

4.2.4. The Prediction of Protein Stability with SNPs

We used I-Mutant 2.0 (https://folding.biofold.org/i-mutant/i-mutant2.0.html (accessed on 30 August 2021)) to predict the stability of the TLR2 and TLR4 proteins with rs5743708 and rs4986790 SNPs, respectively [50]. I-Mutant 2.0 is considered a support vector machine that was tested depending on the ProTherm database which contained the largest experimental data about stability changes with protein mutations [51].

4.2.5. The Identification of Evolutionarily Conserved Positions in a Protein Sequence

This identification was performed using the ConSurf server (https://consurf.tau.ac.il (accessed on 30 August 2021)) which depends on phylogenetic relations between homologous sequences to identify the evolutionary conservation of amino acids in protein sequences [28,52].

4.2.6. The Identification of Structural Effects of SNPs

Structural effects of rs5743708 and rs4986790 SNPs on TLR2 and TLR4, respectively, were analyzed using HOPE (https://www3.cmbi.umcn.nl/hope/ (accessed on 30 August 2021)) which is a mutant analysis server that could analyze the effects of SNPs on protein structure [53].

4.3. The Study Design

This was a prospective observational study that was conducted in intensive care units in Suez Canal University Hospitals, Ismailia, Egypt, for seven months. All ICU Patients who contracted infections with a positive culture or a chest X-ray were included in the study group. All included patients were Egyptian adults of both sexes. Exclusion criteria were patients younger than 18 years old, pregnancy, immune suppression, and patients with radiation therapy or chemotherapy.

Once admitted, general examination and clinical status were assessed for patients; both Acute Physiology and Chronic Health Evaluation (APACHE II) scores and sequential organ failure assessment (SOFA) scores were measured. In addition to vital signs check (blood pressure, heart rate, respiratory rate, central venous pressure, and temperature) and laboratory analyses such as complete blood count, blood sugar, CRP, blood urea nitrogen, serum calcium, potassium, sodium, aspartate aminotransferase, alanine aminotransferase, and arterial blood gas analysis were carried out.

The patients were further followed up to assess infection, sepsis, and septic shock. Routine cultures of sputum, blood, urine, and pus were collected to determine the presence of infection and identify the causing organism. Assessment of sepsis and septic shock was performed by daily evaluation for sepsis or septic shock. Sepsis and septic shock were defined and diagnosed according to “The Third International Consensus Definitions for Sepsis and Septic Shock (Sepsis-3)” [4].

4.4. Samples Collection

Two milliliters of venous blood sample were collected into EDTA tubes from all admitted patients in the study group under complete aseptic conditions and stored at −80 °C until processed for DNA extraction.

4.5. Genotyping

Genomic DNA was extracted from venous blood with a QIAamp DNA Blood Mini kit (Cat. No. 51104; QIAGEN, Hilden, Germany) according to the manufacturer’s protocol. The
measurement of both the concentration and purity of the extracted DNA was performed by NanoDrop ND-1000 (NanoDrop Tech., Inc., Wilmington, DE, USA).

Genotyping for the TLR4 gene polymorphism (Asp299Gly; rs4986790) and TLR2 gene polymorphism (Arg753Gln; rs5743708) was performed using real-time polymerase chain reaction technology using TaqMan allelic discrimination assay. The required reagents for the TaqMan assay including TaqMan genotyping assay and TaqMan genotyping master mix were brought from Applied Biosystems (Foster City, CA, USA). The assay ID for rs5743708 is C_27860663_10 and for rs4986790 is C_11722238_20. PCR was run with a total reaction volume of 25 µL reaction volume. The components of PCR reaction were 12.5 µL TaqMan genotyping master mix; No AmpErase UNG (2 ×), genomic DNA (20 ng) diluted to 11.25 µL with DNase-RNase free water, and 1.25 µL TaqMan SNP genotyping assay mix (Cat. No. 4351379, Applied Biosystems, Foster City, USA). Nuclease-free water was used as a negative control.

The PCR amplification was carried out in a StepOne™ real-time PCR system (Applied Biosystems, Foster City, CA, USA) according to the following conditions: a hold cycle (95 °C for 10 min) followed by a 40-cycle PCR consisting of 95 °C for 15 s and 60 °C for one minute. SDS software version 1.3.1 (Applied Biosystems) was used for allelic discrimination.

Genotyping was performed with blindness to sepsis/non-sepsis status.

### 4.6. Statistical Analysis

Statistical analysis was carried out using Microsoft® Excel 2010 and the “Statistical Package for the Social Sciences (SPSS) for windows” software, version 24. Odds ratios (OR) with a 95% confidence interval (CI) were calculated. Descriptive statistics were expressed as percentages for qualitative variables and mean ± standard deviation (SD) for quantitative variables. Testing differences between septic patients and no septic patients were performed using Student’s t-test, Chi-square ($\chi^2$) test, or Fisher’s exact tests. $p$-value was considered statistically significant below 0.05. The Hardy–Weinberg equilibrium (HWE) was calculated by the Online Encyclopedia for Genetic Epidemiology (OEGE) software (http://www.oeg.org/software/hwe-mrcalc.shtml (accessed on 10 March 2019)). The relationship between the risk factors including our polymorphisms and the development of sepsis was further determined using logistic regression after adjustment of factors. Survival analysis was performed as well. Log-rank, Breslow, and Tarone–Ware tests were used to find Kaplan–Meier estimates for survival. Cox regression analysis was applied to the data to determine if any of the variables were independently associated with the duration of survival.

### 5. Conclusions

Rs5743708 was predicted by nearly all used bioinformatics tools to possess a damaging impact on TLR2 which was not the case with rs4986790 of TLR4. Meanwhile, the conducted pilot study concluded that the TLR2 genotype may be a risk factor for sepsis in adult patients. Moreover, our study showed that Asp299Gly polymorphism in TLR4 may be associated with an increased risk of Acinetobacter baumannii infection. In addition, a significant association was found between the TLR4 polymorphism and the post-surgical category of patients admitted to intensive care units. Identification of the role of TLR2 and TLR4 polymorphisms in developing infection and sepsis could allow early prediction, prevention, and management of these serious diseases.

**Author Contributions:** M.Y.B. (Investigation, Methodology, Conceptualization, Formal Analysis, Writing–Original Draft Preparation); M.M.A. (Supervision, Conceptualization, Formal Analysis, Methodology, Writing–Original Draft Preparation); E.E.-D.A.I. (Supervision, Methodology, Writing–Review and Editing); E.A.T. (Supervision, Methodology, Formal Analysis, Writing–Review and Editing); A.A.A. (Supervision, Methodology, Conceptualization, Writing–Original Draft Preparation); H.R.H. and A.A.S. (Supervision, Formal Analysis, Writing–Review and Editing). All authors have read and agreed to the published version of the manuscript.
22. Hosseinkhani, M.S.; Roy, A.S.; Islam, M.S. In silico analysis predicting effects of deleterious SNPs of human RASSF5 gene on its structure and functions. *Sci. Rep.* 2020, 10, 14542. [CrossRef] [PubMed]

23. Behairy, M.Y.; Abdelrahman, A.I.A.; Abdallah, H.Y.; Ibrahim, E.E.-D.A.; Sayed, A.A.; Azab, M.M. In silico analysis of missense variants of the ClqA gene related to infection and autoimmune diseases. *J. Taibah Univ. Med. Sci.* 2022. Available online: https://www.sciencedirect.com/science/article/pii/S1658361222000890 (accessed on 4 August 2022).

24. Behairy, M.Y.; Soltan, M.A.; Adam, M.S.; Refaat, A.M.; Ezz, E.M.; Albogami, S.; Fayad, E.; Alothbaiat, F.; Gouda, A.M.; Sileem, A.E.; et al. Computational Analysis of Deleterious SNPs in NRAS to Assess Their Potential Correlation with Carcinogenesis. *Front. Genet.* 2022, 13, 872845. Available online: https://www.frontiersin.org/articles/10.3389/fgene.2022.872845 (accessed on 19 August 2022).

25. Behairy, M.Y.; Abdelrahman, A.A.; Abdallah, H.Y.; Ibrahim, E.E.-D.A.; Hashem, H.R.; Sayed, A.A.; Azab, M.M. Role of MBL2 Polymorphisms in Sepsis and Survival: A Pilot Study and In Silico Analysis. *Diagnostics* 2022, 12, 460. [CrossRef] [PubMed]

26. Takeda, K.; Akira, S. TLR signaling pathways. *Semin. Immunol.* 2004, 16, 3–9. [CrossRef] [PubMed]

27. Deller, M.C.; Kong, L.; Rupp, B. Protein stability: A crystallographer’s perspective. *Acta Cryst. Sect. F Struct. Biol. Commun.* 2016, 72 Pt 2, 72–95. [CrossRef]

28. Berezin, C.; Glaser, F.; Rosenberg, J.; Paz, I.; Pupko, T.; Fariselli, P.; Casadio, R.; Ben-Tal, N. ConSeq: The identification of functionally and structurally important residues in protein sequences. *Bioinformatics* 2004, 20, 1322–1324. [CrossRef]

29. Saleh, M.A.; Ramadan, M.M.; Arram, E.O. Toll-like receptor-2 Arg753Gln and Arg677Trp polymorphisms and susceptibility to pulmonary and peritoneal tuberculosis. *APMIS* 2017, 125, 558–564. [CrossRef]

30. Liu, F.; Lu, W.; Qian, Q.; Qi, W.; Hu, J.; Feng, B. Frequency of TLR 2, 4, and 9 gene polymorphisms in Chinese population and their susceptibility to type 2 diabetes and coronary artery disease. *J. Biomed. Biotechnol.* 2012, 2012, 373945. [CrossRef]

31. Shan, X.; Wu, Y.; Ye, J.; Ding, Z.; Qian, C.; Zhou, A. Gene polymorphisms of Toll-like receptors in Chinese Han children with sepsis in Wenzhou. *Zhonghua Er Ke Za Zhi = Chin. J. Pediatr.* 2010, 48, 15–18.

32. Fawzy, M.S.; Hussein, M.H.; Abdelaziz, E.Z.; Yamany, H.A.; Ismail, H.M.; Toraih, E.A. Association of MicroRNA-196a2 Variant with Response to Short-Acting β2-Agonist in COPD: An Egyptian Pilot Study. *PloS ONE* 2016, 11, e0152834. [CrossRef]

33. Tumangger, H.; Jamil, K.F. Contribution of genes polymorphism to susceptibility and outcome of sepsis. *Egypt. J. Med. Hum. Genet.* 2010, 11, 97–103. [CrossRef]

34. David, V.L.; Ercisli, M.F.; Rogobete, A.F.; Boia, E.S.; Horhat, R.; Nitu, R.; Diaconu, M.M.; Pirtea, L.; Ciua, I.; Horhat, D.I.; et al. Early Prediction of Sepsis Incidence in Critically Ill Patients Using Specific Genetic Polymorphisms. *Biochem. Genet.* 2017, 55, 193–203. [CrossRef]

35. Chatzi, M.; Papanikolaou, J.; Makris, D.; Papathanasiou, I.; Tsezou, A.; Kariou, M.; Zakyntinos, E. Toll-like receptor-2 and -4 polymorphisms and susceptibility to type II diabetes and coronary artery disease. *Int. J. Mol. Sci.* 2013, 14, 2267–2280. [CrossRef] [PubMed]

36. Fawzy, M.S.; Hussein, M.H.; Abdelaziz, E.Z.; Yamany, H.A.; Ismail, H.M.; Toraih, E.A. Association of MicroRNA-196a2 Variant with Response to Short-Acting β2-Agonist in COPD: An Egyptian Pilot Study. *PloS ONE* 2016, 11, e0152834. [CrossRef] [PubMed]

37. Knapp, S.; Wieland, C.W.; Florquin, S.; Fantophlet, R.; Dijkstra, M.; Tshimbalanga, N.; Akira, S.; van der Poll, T. Differential roles of CD14 and toll-like receptors 4 and 2 in murine Acinetobacter pneumonia. *Am. J. Respir. Crit. Care Med.* 2006, 173, 122–129. [CrossRef] [PubMed]

38. Kim, C.-H.; Jeong, Y.-J.; Lee, J.; Jeon, S.-J.; Park, S.-R.; Kang, M.-J.; Park, J.-H.; Park, J.-H. Essential role of toll-like receptor 4 in Acinetobacter baumannii-triggered inflammatory response. *PLoS ONE* 2010, 5, e10033. [CrossRef] [PubMed]

39. Knapp, S.; Wieland, C.W.; Florquin, S.; Fantophlet, R.; Dijkstra, M.; Tshimbalanga, N.; Akira, S.; van der Poll, T. Differential roles of CD14 and toll-like receptors 4 and 2 in murine Acinetobacter pneumonia. *Am. J. Respir. Crit. Care Med.* 2006, 173, 122–129. [CrossRef] [PubMed]

40. Kim, C.-H.; Jeong, Y.-J.; Lee, J.; Jeon, S.-J.; Park, S.-R.; Kang, M.-J.; Park, J.-H.; Park, J.-H. Essential role of toll-like receptor 4 in Acinetobacter baumannii-induced immune responses in immune cells. *Microb. Pathog.* 2013, 54, 20–25. [CrossRef]

41. Erridge, C.; Moncayo-Nieto, O.L.; Morgan, R.; Young, M.; Poxton, I.R. Acinetobacter baumannii lipopolysaccharides are potent stimulators of human macrophage activation via Toll-like receptor 4 signalling. *J. Med. Microbiol.* 2007, 56 Pt 2, 165–171. [CrossRef] [PubMed]

42. Koch, A.; Hamann, L.; Schott, M.; Boehm, O.; Grottemeyer, D.; Kurt, M.; Schwenke, C.; Schumann, R.R.; Bornstein, S.R.; Zacharowski, K. Genetic variation of TLR4 influences immunoenocrine stress response: An observational study in cardiac surgical patients. *Crit. Care.* 2011, 15, 109. [CrossRef]

43. Mayr, F.B.; Yende, S.; Angus, D.C. Epidemiology of severe sepsis. *Virology* 2014, 5, 4–11. [CrossRef] [PubMed]

44. Wang, D.T.; Gomez, M.; McGuire, G.P.; Kavanagh, B. Utilization of intensive care unit days in a Canadian medical-surgical intensive care unit. *Crit. Care Med.* 1999, 27, 1319–1324. [CrossRef]

45. Abela, F.J.; Castro, M.A.; Landeiro, N.M.; Neves, A.M.; Santos, C.C. Mortality and length of stay in a surgical intensive care unit. *Rev. Bras. Anestesiol.* 2006, 56, 34–45. [CrossRef] [PubMed]

46. Sim, N.L.; Kumar, P.; Hu, J.; Henikoff, S.; Schneider, G.; Ng, P.C. SIFT web server: Predicting effects of amino acid substitutions on proteins. *Nucleic Acids Res.* 2012, 40, W452–W457. [CrossRef] [PubMed]

47. Choi, Y.; Chan, A.P. PROVEAN web server: A tool to predict the functional effect of amino acid substitutions and indels. *Bioinformatics* 2016, 32, 2230–2232. [CrossRef] [PubMed]

48. Capriotti, E.; Calabrese, R.; Fariselli, P.; Martelli, P.L.; Altman, R.B.; Casadio, R. WS-SNPs&Go: A web server for predicting the deleterious effect of human protein variants using functional annotation. *BMC Genom.* 2013, 14 (Suppl. S3), S6.
49. Blum, M.; Chang, H.Y.; Chuguransky, S.; Grego, T.; Kandasaamy, S.; Mitchell, A.; Nuka, G.; Paysan-Lafosse, T.; Qureshi, M.; Raj, S.; et al. The InterPro protein families and domains database: 20 years on. *Nucleic Acids Res.* 2021, 49, D344–D354. [CrossRef]

50. Capriotti, E.; Fariselli, P.; Casadio, R. I-Mutant2.0: Predicting stability changes upon mutation from the protein sequence or structure. *Nucleic Acids Res.* 2005, 33, W306–W310. [CrossRef]

51. Bava, K.A.; Gromiha, M.M.; Uedaira, H.; Kitajima, K.; Sarai, A. ProTherm, version 4.0: Thermodynamic database for proteins and mutants. *Nucleic Acids Res.* 2004, 32, D120–D121. [CrossRef]

52. Ashkenazy, H.; Abadi, S.; Martz, E.; Chay, O.; Mayrose, I.; Pupko, T.; Ben-Tal, N. ConSurf 2016: An improved methodology to estimate and visualize evolutionary conservation in macromolecules. *Nucleic Acids Res.* 2016, 44, W344–W350. [CrossRef]

53. Venselaar, H.; Te Beek, T.A.; Kuipers, R.K.; Hekkelman, M.L.; Vriend, G. Protein structure analysis of mutations causing inheritable diseases. An e-Science approach with life scientist friendly interfaces. *BMC Bioinform.* 2010, 11, 548. [CrossRef]