

Vaccine design against *Toxoplasma gondii* in ovines using surface antigen-1 (SAG1) and dense granule protein-6 (GRA6) antigens through immunoinformatic approach

Thabile Madlala, Dr M Okpeku, Dr SI Tshilwane And Prof MA Adeleke

Discipline of Genetics, School of Life Sciences,
College of Agriculture, Engineering and Science,
University of KwaZulu-Natal, Westville P/Bag X54001,
Durban 4000 South Africa



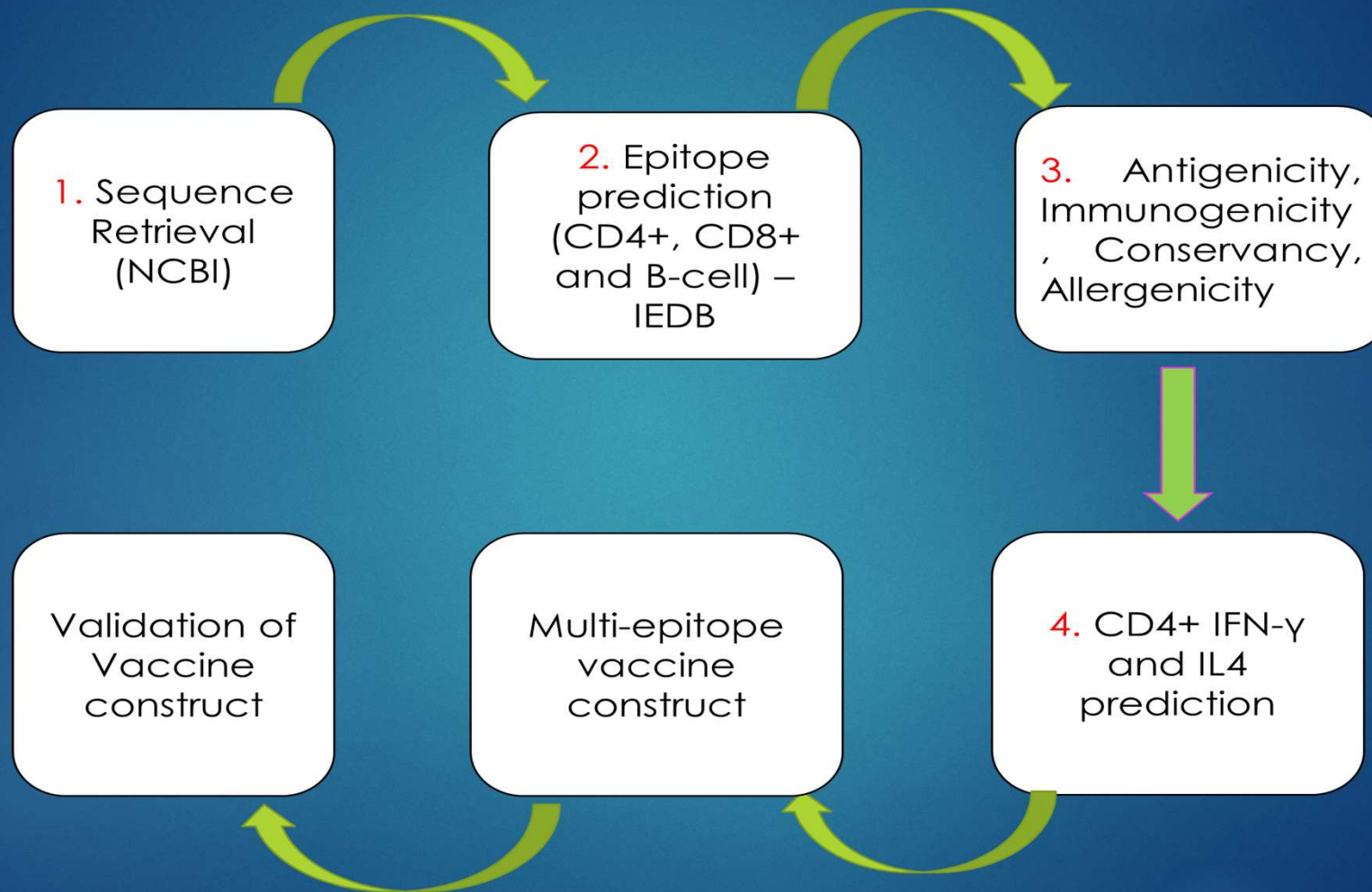
Introduction

- ▶ Toxoplasmosis: Zoonotic disease known to infect almost all warm-blooded animals (including humans).
- ▶ Caused by *Toxoplasma gondii* resulting in congenital infections and abortion in humans and livestock.
- ▶ Primary carrier/host: Cats (Felidae)
- ▶ Intermediate hosts: Humans and livestock
- ▶ Threatens public health due to increased zoonotic risk of transmission of infections to humans
- ▶ Mode of transmission: Consumption of contaminated food, water and dust.
- ▶ In humans: Poorly cooked meat containing *T. gondii* tissue cysts or accidental ingestion of contaminated water or vegetables with infected cat faeces

Aim

- ▶ Identify antigenic T-cell and B-cell epitopes
- ▶ Design a potentially cost-efficient peptide-based vaccine by exploring the *T. gondii* antigens through immunoinformatics techniques.

MATERIAL & METHODS



RESULTS AND DISCUSSION

- IDENTIFIED:
 - 1 CD8+ and 1 CD4+ T cell epitopes
 - 10 B cell epitopes
- Multiepitope vaccine construct design via attachment of linkers(GPGPG, and KK) and Adjuvant (Cholera toxin B subunit)-3Dpro
- VALIDATION
- ProSa: Z score = -2.93

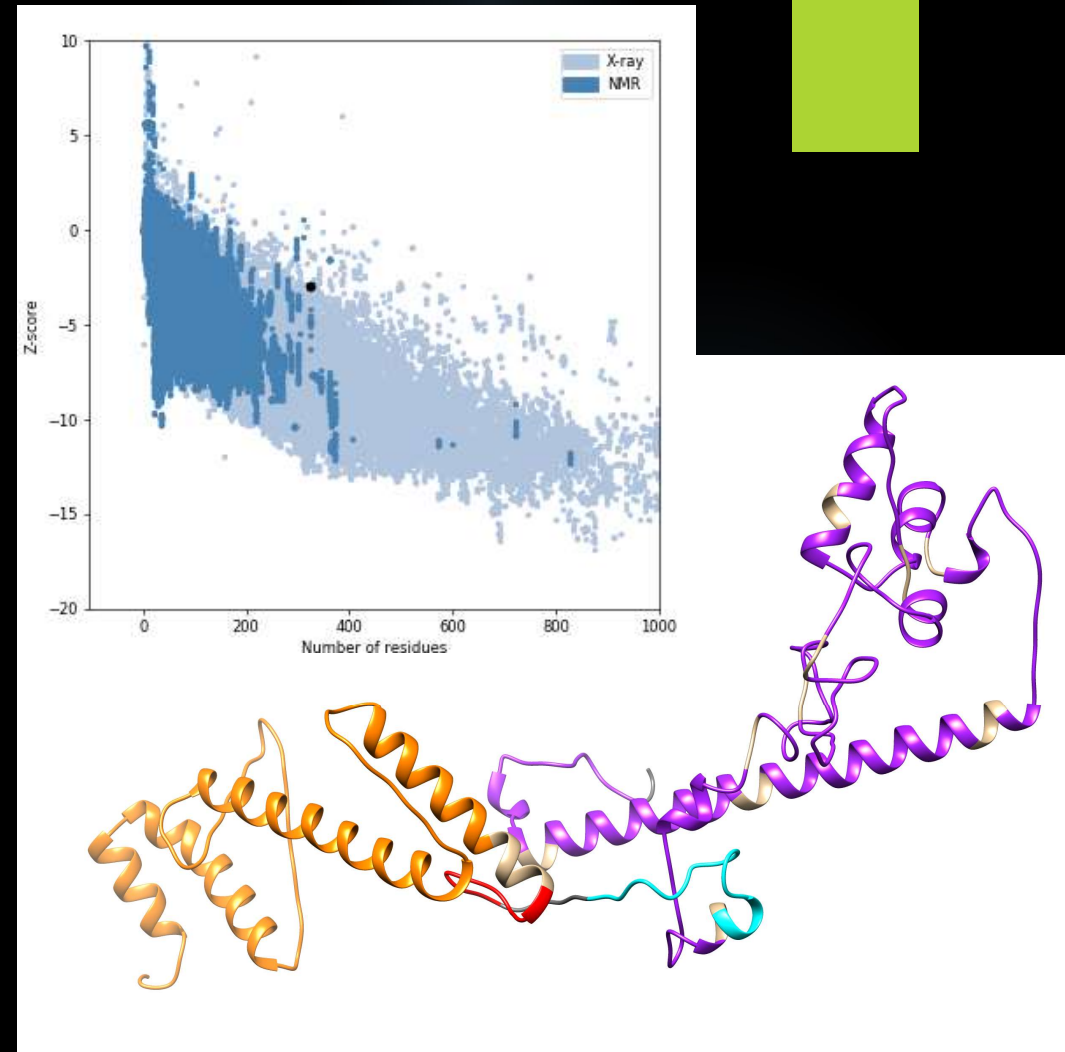
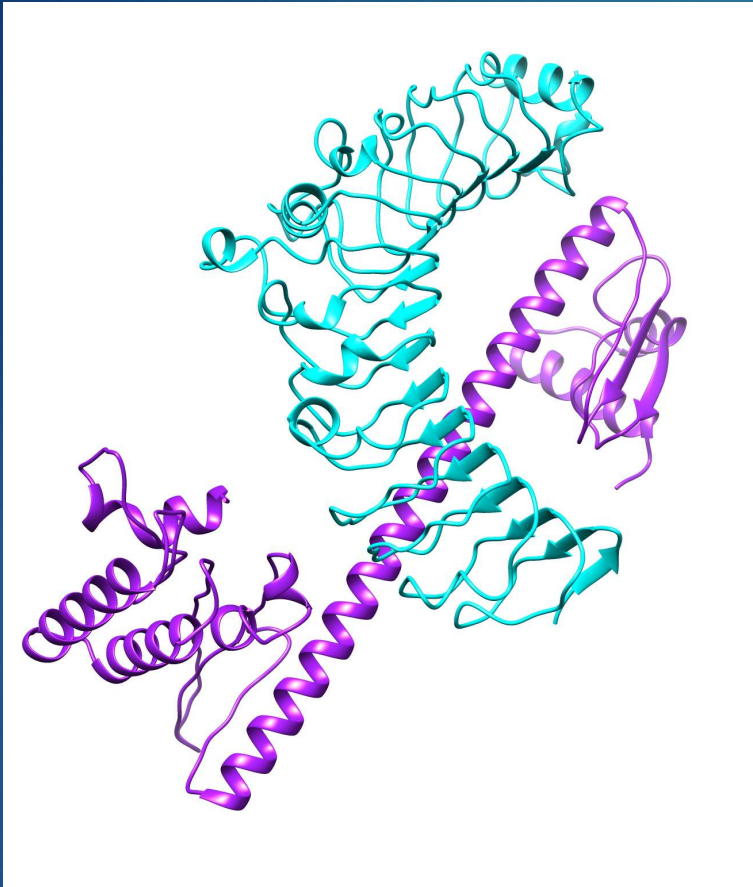


Fig. 1: Tertiary structure of final refine multiepitope designed vaccine and validation ProSA plot



- DOCKING: Toll-like receptor-4
 - ✓ Binding affinity: -151.645 kcal/mol
- Physicochemical properties of vaccine construct
 - ✓ Antigenicity: 0.9135
 - ✓ Solubility: 0.9485
 - ✓ pI: 9.81
 - ✓ Molecular weight: 34.08 kDa
 - ✓ Instability Index: 29.36
 - ✓ Aliphatic Index: 75.62
 - ✓ GRAVY: -0.425

Fig. 2 Tertiary structure of designed vaccine docked against Toll like receptor 4 (TLR2)

CONCLUSION

- ▶ Current study designed a promising multiepitope vaccine from a cocktail of *T. gondii* antigens.
- ▶ Designed vaccine was highly antigenic and non-allergenic, appropriate for production.
- ▶ Great promise in conferring complete host protection as it effectively elicited host immune response
- ▶ Valuable for future studies focusing on vaccine development against *T. gondii*.