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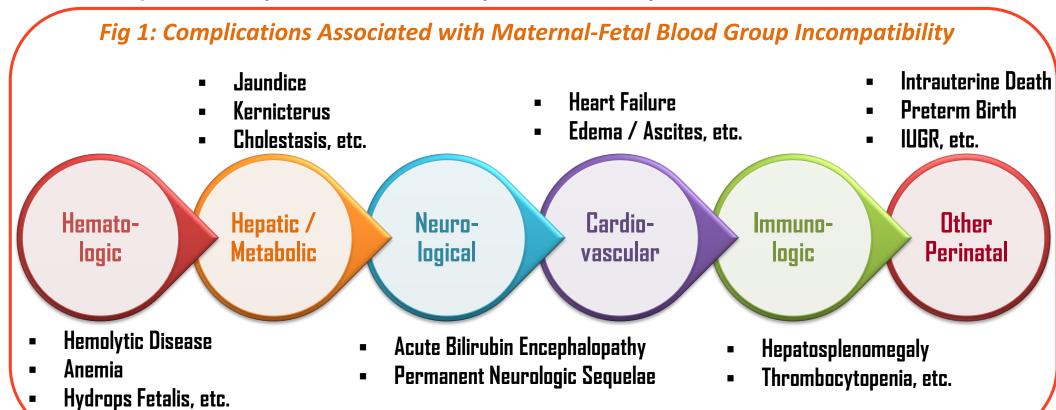
Maternal-Child Blood Group Discordance & Congenital Heart Disease Severity: An Immunological Risk Perspective from a Tertiary Hospital-Based Study

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INTRODUCTION

- ✓ Congenital heart disease (CHD) affects ~1 % of all live births, with outcomes influenced by both anatomical complexity and clinical severity.^[1]
- ✓ While genetic and environmental factors are well-recognized, the role of maternal-foetal immuno-haematological interactions—particularly ABO and Rh incompatibility—remains underexplored, despite their known impact in haemolytic disease of the newborn. [2]



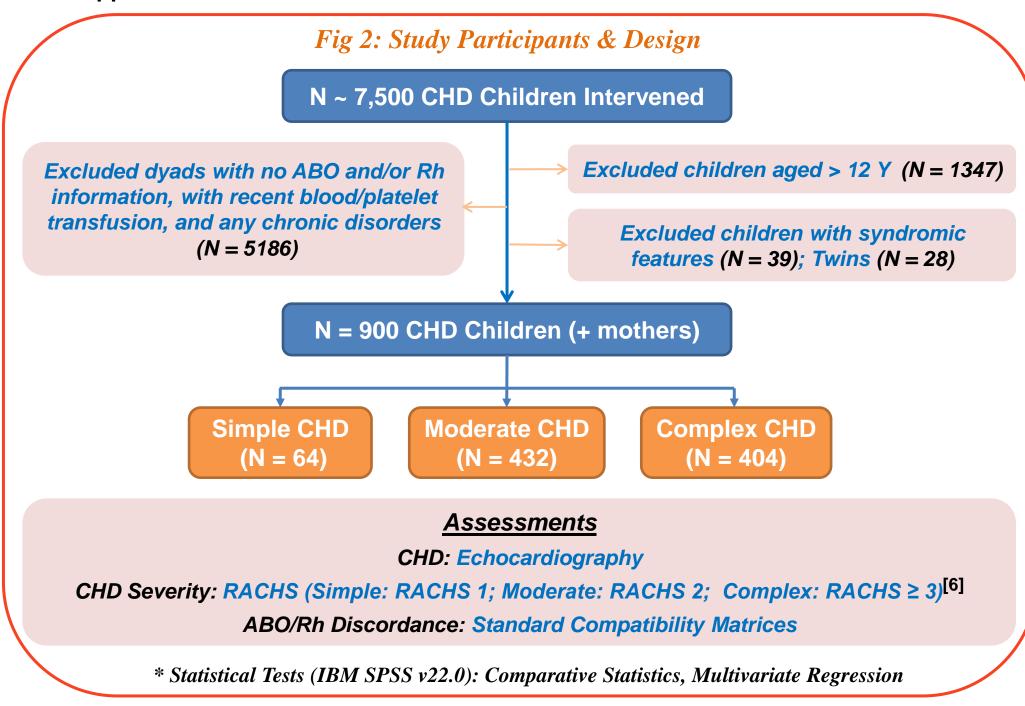
Prior studies suggest a link between ABO blood groups and CHD risk.^[3] Still, evidence on ABO and Rh discordance is limited and inconclusive, with <u>no research yet examining their association with CHD severity</u>.^[4,5]

OBJECTIVE

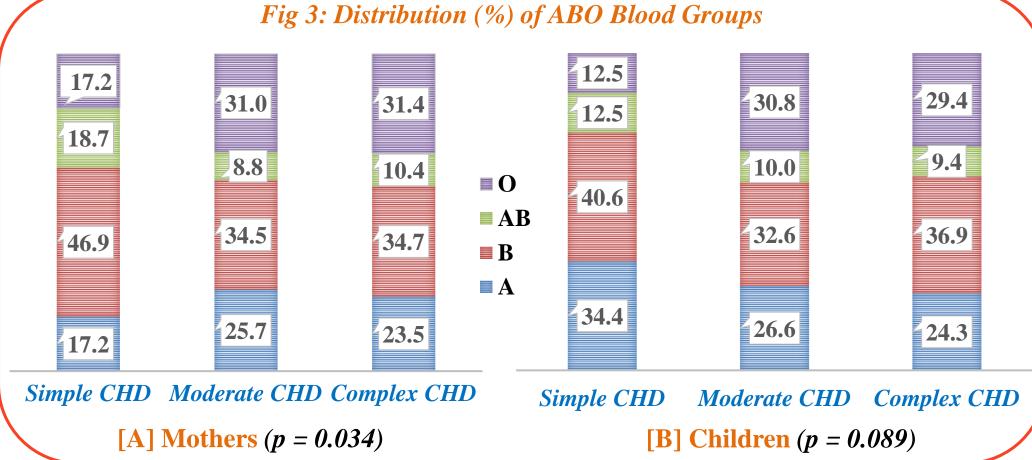
To investigate the association between maternal-child blood group compatibility patterns (ABO and Rh), and the severity of CHD in affected children

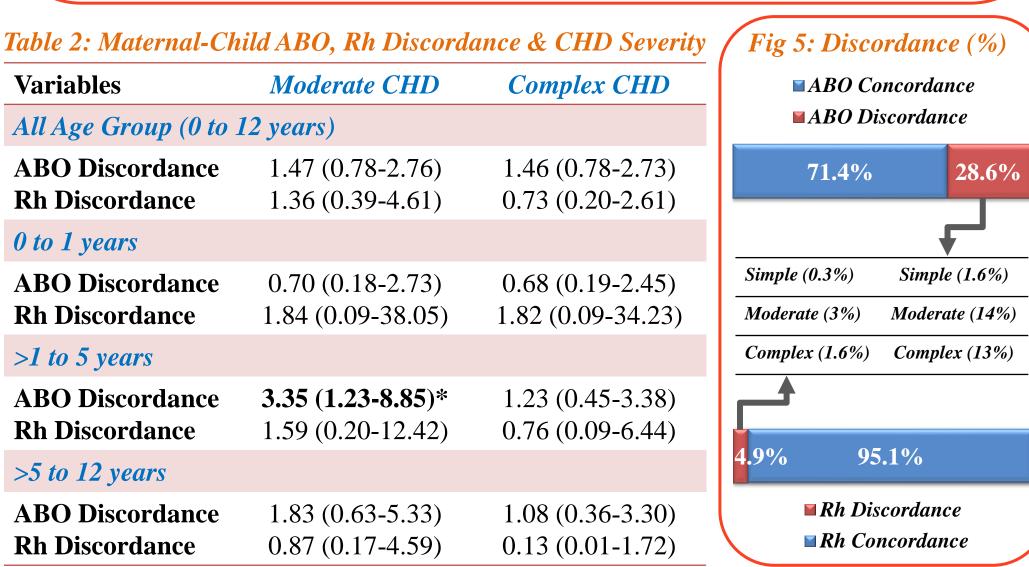
METHODOLOGY

- ✓ Design: Retrospective Cross-Sectional Observational Study
- **✓** Participants: Indian mother-child dyads underwent surgical correction for CHD
- **✓** Duration: *March 2022 to July 2025*
- Site: Sri Sathya Sai Sanjeevani Hospital- A free-of-cost tertiary cardiac centre
- **✓ IEC Approved with Informed Consent**



RESULTS & DISCUSSION





<u>Strengths</u>: Large, clinically well-characterized cohort, standardized RACHS severity classification and robust statistical adjustment

<u>Limitations</u>: Single-centre design, potential referral bias and lack of genetic / mechanistic biomarkers to validate causality

Table 1: Association Between ABO, Rh Factors & CHD Severity Fig 4: Rh (-) Distribution Adjusted Odds Ratio (95% CI) **Variables** (p = 0.224)Moderate CHD Complex CHD (p = 0.566)Maternal ABO Blood Groups (w.r.t. O) 0.75 (0.31-1.79) 0.83 (0.35-1.98) 5.2 0.47 (0.29-0.81)* 0.38 (0.15-0.85)* 0.25 (0.11-0.70)** 0.30 (0.10-0.73)** AB 3.9 Maternal Rh Antigen (w.r.t. Rh +) 0.95 (0.27-3.32) **Rh Negative** 1.57 (0.47-5.29) Child ABO Blood Groups (w.r.t. O) 6.3 0.47 (0.29-0.97)** 0.31 (0.15-0.79)** 0.31 (0.16-0.77)** 0.38 (0.16-0.90)* 0.39 (0.29-0.87)* 0.36 (0.19-0.85)* AB Children **Mothers** Child Rh Antigen (w.r.t. Rh +) **■ Complex CHD Rh Negative** 0.62 (0.20-1.89) 0.82(0.27-2.48)**■ Moderate CHD ■ Simple CHD** * $p \le 0.05$; ** $p \le 0.01$

CONCLUSION

- ✓ <u>First large-scale evidence</u> from an Indian cohort showing that maternal-child ABO discordance may modulate CHD severity, especially in early childhood.
- ✓ Highlights the importance of ABO incompatibility profiling for early risk stratification frameworks, particularly in resource-limited clinical settings with high CHD burden.
- ✓ Calls for multicentric studies integrating immunology, genomics, and placental biology to uncover underlying mechanisms.

REFERENCES

- [1] Meng X, et al. MedComm. **2024**; 5(7): e631. doi: 10.1002/mco2.631
- [2] Hall V, et al. StatPearls. 2025. https://www.ncbi.nlm.nih.gov/books/NBK557423/
- [3] Zu B, et al. Sci Rep. 2017; 7: 42804. doi: 10.1038/srep42804
- [4] Lubs ML, et al. Lancet. **1972**; 2(7781): 825-826. 10.1016/s0140-6736(72)92187-3
- [5] Kandur Y, et al. J Pediatr Genet. **2023**; 13(4): 272-276. doi: 10.1055/s-0043-1774292
- [6] Jenkins, et al. J Thorac Cardiovas Surg. 2002; 123(1): 110-118. doi: 10.1067/mtc.2002.119064