

Seroprevalence Of Parvo Virus B19 Igg Antibodies Among Voluntary Blood Donors In Blood Bank At Tertiary Centre

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ABSTRACT

Background: Parvovirus B19 is a human pathogen classified under the genus Erythroparvovirus of the Parvoviridae family 1,2. It is a small, non-enveloped virus possessing a single-stranded DNA genome. Among the known Parvoviruses, B19 is the only strain pathogenic to humans (1). The virus primarily targets erythroid precursor cells, particularly in the bone marrow and foetal liver, through its affinity for the P blood group antigen located on red blood cell membranes (2).

Aim: To determine the seroprevalence of specific IgG antibodies among voluntary blood donors in tertiary care blood bank.

Materials And Methods: This prospective study was carried out in the blood bank of the pathology department of Sree Balaji Medical College in Chromepet, Chennai, over the course of two years, from December 2022 to December 2024. A total of 91 voluntary donors of blood were chosen who gave their consent to participate for the study.

Results: 33% of the donors were A positive, 22% were B positive, 32% were O positive, and 13% were AB positive. Based on the titre values, when the IgG value is less than 0.91(negative), it shows a lack of previous exposure to Parvovirus B19. When the values are between 0.91 and 0.99 (grey zone), re-testing the sample is necessary, and when the titre values are higher than 1.0 (positive), means the donor has had a history of Parvovirus B19 exposure.

Conclusion: This research emphasizes the need for heightened awareness and vigilant monitoring of Parvovirus B19 in transfusion medicine. It advocates from large-scale, multicentric serological and molecular studies to further elucidate the epidemiology and transfusion risk, which can guide the development of evidence-based national blood safety policies in India and other similar setting.

KEYWORDS: *Parvovirus, anaemia, hydrops fetalis, TTI.*

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1. INTRODUCTION

Parvovirus B19 is a human pathogen classified under the genus Erythroparvovirus of the Parvoviridae family 1,2. It is a small, non-enveloped virus possessing a single-stranded DNA genome. Among the known Parvoviruses, B19 is the only strain pathogenic to humans (1). The virus primarily targets erythroid precursor cells, particularly in the bone marrow and foetal liver, through its affinity for the P blood group antigen located on red blood cell membranes (2). Epidemiological studies indicate that seroprevalence rates of Parvovirus B19 increase with age, ranging from 5–10% in children to nearly 50% in adults, suggesting widespread exposure throughout life (2). Although respiratory droplets remain the predominant route of transmission, exposure to infected blood or blood products constitutes a significant transmission risk (1,2). This becomes particularly significant in settings involving transfusion-dependent individuals, pregnant women, and

immunocompromised patients, in whom Parvovirus B19 infection can cause complications such as anaemia or hydrops fetalis (1). Given its ability to persist in donor blood and its resistance to standard inactivation methods, Parvovirus B19 presents a notable transfusion-transmissible risk (6). Consequently, understanding the prevalence of anti-B19 antibodies among voluntary blood donors becomes essential to guide screening practices and minimize the chances of transfusion-related transmission (5).

AIMS AND OBJECTIVES

AIM:

To determine the seroprevalence of specific IgG antibodies among voluntary blood donors in tertiary care blood bank.

OBJECTIVES:

1. To assess the presence of specific Parvo B19- IgG antibodies among healthy voluntary blood donors, indicating past exposure or immunity.
2. To analyse the distribution of seropositivity across different age groups, genders, and blood groups.
3. To highlight the potential risk of transfusion-transmitted B19V infection in asymptomatic donors.
4. To emphasize the importance of screening for Parvovirus B19 in blood donation, practices, especially in settings.

2. MATERIALS AND METHODS

This prospective study was carried out in the blood bank of the pathology department of Sree Balaji Medical College in Chromepet, Chennai, over the course of two years, from December 2022 to December 2024. A total of 91 voluntary donors of blood were chosen who gave their consent to participate for the study. The Sree Balaji Medical College ethical committee in Chromepet, Chennai, gave its approval for the study. Donors who experienced a deferment. In a sterile-capped tube, five millilitres of blood were extracted from each donor collecting bag. The plasma was then separated and kept until needed, after which it was centrifuged.

3. RESULTS

A total of 91 donors were selected between November 2022 and April 2022 to evaluate the seroprevalence of Parvo Virus B19 based on gender, age, and blood group.

CHART 1: Study group's gender distribution

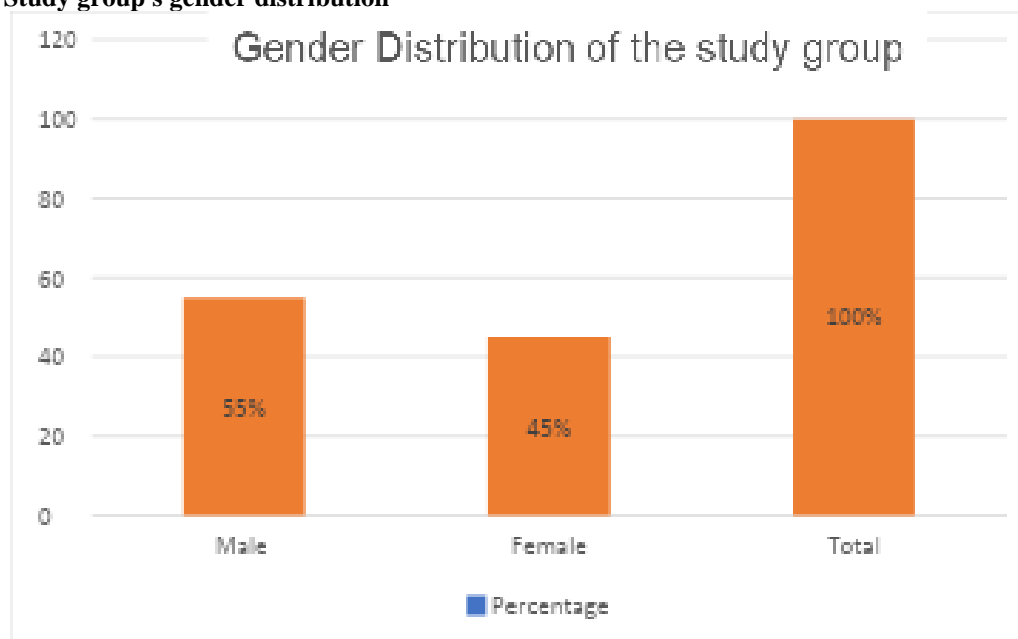


Chart 1 shows 50 (54.9%) of the 91 donors were men, and 41 (45.1%) were women.

Table 1: The study group's age distribution

| Age group in yrs | No. of donors | Percentage % |
|------------------|---------------|--------------|
| >20 and <25 | 30 | 33% |

| | | |
|--------------|----|------|
| >26 and <30 | 24 | 26% |
| >31 and <35 | 18 | 20% |
| >36 and <39 | 08 | 9% |
| More than 40 | 11 | 11% |
| | 91 | 100% |

The blood donors' ages were distributed as follows: 11% in >40 years, 9% in thirty-six to thirty-nine years, 20% in thirty-one to thirty-five years, 26% in twenty-six to thirty years, 33% in twenty to twenty-five years and 34.1% in eighteen to twenty years.

Table 2: The study group's blood group's distribution

| Blood group | No. of. Patients | Percentage |
|--------------------|-------------------------|-------------------|
| A+ | 28 | 33% |
| B+ | 18 | 22% |
| O+ | 34 | 32% |
| AB+ | 11 | 13% |
| Total | 91 | 100% |

33% of the donors were A positive, 22% were B positive, 32% were O positive, and 13% were AB positive.

Table 3: Status of transfusion-transmitted infections of the study group

| TTI status | Donor's(total) | Percentage% |
|-------------------|-----------------------|--------------------|
| Reactive | 0(91) | 0 |
| Non-reactive | 91(91) | 100% |

None of the 91 donors tested positive for the hepatitis B surface antigen (HbsAg).

Table 4: Analyzation of results based on titre value (Parvovirus B19 IgG by EIA)

| IgG Titre Value | Interpretation | No. Of Donors |
|------------------------|-----------------------|----------------------|
| IgG<0.91 | Negative | 69 |
| IgG<0.91-0.99 | Grey Zone | 0 |
| IgG>1.0 | Positive | 22 |

Based on the above titre values, when the IgG value is less than 0.91(negative), it shows a lack of previous exposure to Parvovirus B19. When the values are between 0.91 and 0.99 (grey zone), re-testing the sample is necessary, and when the titre values are higher than 1.0 (positive), means the donor has had a history of Parvovirus B19 exposure.

Table 5: Anti-Parvovirus B19 EIA-based antibody screening

| Anti- Parvovirus B19 Ab | Positive (Total) | Negative |
|--------------------------------|-------------------------|-----------------|
| Immunoglobulin G | 69(91) | 22 |

69 people tested positive and 22 tested negatives for IgG during the EIA screening process for anti-Parvovirus B19 antibodies, yielding a 64.6% total Parvovirus B19 prevalence rate.

Table 6: The study group's prevalence of IgG seropositive by age

| Age range expressed in years | Donors who test positive for IgG (total donors) | Donors who tested negative | Positive Percentage % |
|-------------------------------------|--|-----------------------------------|------------------------------|
| 21-25 | 24(30) | 6 | 80% |
| 26-30 | 18(24) | 6 | 75% |
| 31-35 | 12(18) | 6 | 66% |
| 36-40 | 6(8) | 2 | 75% |
| >40 | 9(11) | 2 | 81% |
| Total | 69(91) | 22 | 76% |

P-<0.05

An increased prevalence of IgG positivity among individuals aged twenty-six and above.

Table 7: The study group's prevalence of IgG seropositive by gender

| Sex | Donors who test positive for IgG (total donors) | Donors who tested negative for IgG | Positive Percentage % |
|--------|---|------------------------------------|-----------------------|
| Male | 38(50) | 12 | 76% |
| Female | 32(41) | 09 | 78% |
| Total | 70(91) | 21 | 77% |

P-value <0.05

Among the IgG serology-positive individuals, the number of males who showed positivity was 38 and females were 32.

Table 8: The study group's prevalence of IgG seropositive by blood group

| Blood group | IgG positive donors (Total) | IgG negative donors | Percentage% |
|-------------|-----------------------------|---------------------|-------------|
| A+ | 19(28) | 9 | 68% |
| B+ | 7(18) | 11 | 64% |
| O+ | 28(34) | 6 | 82% |
| AB+ | 9(11) | 2 | 82% |
| Total | 63(91) | 28 | 100% |

P<0.05

The immunoglobulin G positivity was 19 (A positive), 7 were (B positive), 28 were (O positive), and 9 were AB positive.

4. DISCUSSION

Parvovirus B19 is a common viral infection, particularly prevalent among children. In immunocompetent individuals, the infection follows a self-limiting and mild course. However, in certain high-risk groups, B19 infections can result in serious complications. The high-risk populations include pregnant women, patients with underlying haematological problems, and immunodeficient patients who constantly receiving multiple transfusions. Routine Screening is not mandatory for all blood donations. However current blood safety practice, use multiple sensitive screening assays, to minimize the risk of transfusion-transmitted infections. These protocols primarily aim to detect and eliminate blood units contaminated with known infectious agents. Other than primary respiratory route of transmission, infection is also acquired through: Nosocomial (hospital-acquired) transmission, Intra-familial spread, Transmission between patients or from patients to healthcare workers. Of potential concern for blood safety is increasing evidence of long term B19 persistence in the circulation and tissues of not only immunocompromised but also immunocompetent individuals (7). Majority of the problems are due to prevalence of asymptomatic carriers in the society, as well as blood donations during the window period of infections. Transfusion transmissible infections (TTI) is a major challenge to the blood transfusion service all over the world. The problem of TTI is directly proportional to SSION: the prevalence of infection in the blood donor's community. Patients requiring blood transfusion are more prone to acquire HBV, HIV and HCV (7). B19V infection has a biphasic clinical course. There is a phase of high viremia during the first few days initially, which gradually diminishes to undetectable levels in the later phase by the eighth day as IgM production begins. The antibody levels will peak which may coincide with a decline in viral load. By 10–12 days post-infection, seroconversion from IgM to IgG antibodies occurs. In patient with defined indication if we detect B19V IgG in two separate blood samples taken 6 months apart, it is considered & B19V-safe for patients with defined indications. Blood donors who test positive for B19V IgM should be tested for B19V DNA by NAT; qPCR is the gold-standard method for most viral nucleic acid-based clinical diagnostics (8).

Age wise distribution of voluntary blood donors in the present study.

| Age group (years) | Number of donors(n) | Percentage (%) | Findings in present study | Comparison with literature |
|-------------------|---------------------|----------------|------------------------------------|--|
| 21-25 | 30 | 33.0% | Most common age group among donors | Kumari S et al.: similar donor age trend (8) |
| 26-30 | 24 | 26.4% | Second most common age group | Jaiswal R et al.: High IgG positivity in younger adults. |
| 31-35 | 18 | 19.8% | Moderate representation. | Limited literature details for this specific age subgroup |
| 36-39 | 8 | 8.8% | Less frequently represented. | Older age groups underrepresented in most donor studies. |
| More than 40 | 11 | 12.0% | Least represented age group. | Often grouped as more than 35 or more than 40 without specific analysis. |

Gender wise distribution of voluntary blood donors in the present study.

| Gender | Number of donors (n) | Percentage (%) | Findings in present study | Comparison with literature |
|--------|----------------------|----------------|--|--|
| Male | 50 | 55% | Higher participation among male donors | Kumar S et al.: Reported male predominance in donor population |
| Female | 41 | 45% | Lower proportion compared to males | Memory Chirambo et al.: Found higher parvovirus IgG in males. |

Blood Group: In our study, 28 donors were A positive, 18 donors were B positive, 34 donors were O positive, and 11 donors were AB positive.

Parvovirus B19 Positivity Among Our Voluntary Blood Donors Based on EIA (IgG/IgM) Titre Values

Interpretation criteria for IgG titre values in Parvovirus B19 serology

| IgG Titre range | Result | Interpretation |
|----------------------|----------------|---|
| Less than 0.91 | Negative | No previous exposure to parvovirus B19 |
| 0.91 -0.99 | Equivocal zone | Borderline, repeat testing advised. |
| More or equal to 1.0 | Positive | Indicates past infection and immune response. |

Serological Results of Donors for Parvovirus B19 Antibodies

| Antibody type | Total tested(n) | Positive (n) | Negative(n) | Interpretation |
|---------------|-----------------|--------------|-------------|---|
| IgG | 91 | 69 | 22 | Majority showed past exposure to Parvovirus B19 |

Age-wise and Gender-wise Distribution of IgG Anti-Parvovirus B19 Seropositivity

| Age Group (in years) | Total Donors (n) | IgG Positive (n) | IgG Positivity (%) | Interpretation |
|----------------------|------------------|------------------|--------------------|---|
| 21–25 | 30 | 24 | 80.0% | Most IgG reactivity observed in this age group |
| 26-30 | 24 | 18 | 75.0% | High seropositivity in second most common group |
| 31–35 | 18 | 12 | 66.7% | Moderate IgG positivity |
| 36-40 | 8 | 6 | 75.0% | Comparable to younger groups |
| >40 | 11 | 9 | 81.8% | Slightly higher, suggesting age-linked exposure |

Overall age wise differences were statically significant ($P < 0.05$).

IgG Anti-Parvovirus B19 Positivity by ABO Blood Group

| Blood group | IgG Positive(n) | Observation |
|-------------|-----------------|---|
| A positive | 19 | Moderate number of seropositive cases. |
| B positive | 7 | Lowest IgG positive among blood groups |
| O positive | 28 | Highest IgG seropositivity; aligns with previous studies (Jaiswal R et al.) |
| AB positive | 9 | High relative to its small population. |

Other TT-Infection: Screening Our present study group's screening for required tests for other transfusion transmitted infections like human immunodeficiency virus, Hep B, Hep C, syphilis, and malaria was low because all the blood donors included in the study gave their consent voluntarily. Out of 91 blood donors in our study, none had a HbsAg positive result. According to a study by Kumari S et al., none of them shows HbsAg positivity.

5. CONCLUSION

The study evaluated the seroprevalence of Parvovirus B19 among voluntary blood donors at a tertiary care centre to assess its potential as a transfusion-transmitted infection (TTI). A high proportion of donors demonstrated IgG Seropositivity, indicating past exposure. These findings imply that although Parvovirus B19 exposure is common in the donor population, the immediate risk of transmission via blood transfusion is low (9). However, the ability of B19V to persist in circulation, its resistance to standard viral inactivation procedures and its potential to cause serious complication in immunocompromised and transfusion-dependent patients under its clinical relevance. Given these concerns, routine universal screening for B19V may not be cost-effective in low prevalence regions. Nevertheless, the results of this study

support the study support the implementation of targeted screening or nucleic acid testing (NAT) in high-risk recipient populations, such as paediatric oncology patients, pregnant women, or individuals with chronic haemolytic anaemia (9). This research emphasizes the need for heightened awareness and vigilant monitoring of Parvovirus B19 in transfusion medicine. It advocates from large-scale, multicentric serological and molecular studies to further elucidate the epidemiology and transfusion risk, which can guide the development of evidence-based national blood safety policies in India and other similar setting.

REFERENCES

- [1] Reno ML, Cox CR and Powell EA. Parvovirus B19: Clinical and Diagnostic Review, Clinical Microbiology Newsletter, Vol 44(12),2022.107-114.
- [2] Qiu J, Soderlund-Venermo M, Young NS. Human parvoviruses. Clin. Microbial. 2017. Rev. 30, 43–113. <https://doi.org/10.1128/CMR.00040-16>.
- [3] Agbandje M, Kajigaya S, McKenna R, Young NS, Rossmann MG. The structure of human parvovirus B19 at 8 Å resolution. Virology 1994; 203, 106–115. <https://doi.org/10.1006/viro.1994.1460>.
- [4] Servant A, Laperche S, Lallemand F, Marinho V, Maur DSG, Meritet JF, Garbarg-Chenon A. Genetic diversity within human retroviruses: identification of three genotypes. J. Virol. 2002; 76, 9124–9134. <https://doi.org/10.1128/jvi.76.18.9124-9134.2002>.
- [5] Gallinella G, Venturoli S, Manaresi E, Musiani M, Zerbini M. B19 virus genome diversity: epidemiological and clinical correlations. J. Clin. Virol.;2003; 28, 1–13. [https://doi.org/10.1016/s1386-6532\(03\)00120-3](https://doi.org/10.1016/s1386-6532(03)00120-3).
- [6] Norja P, Hokynar K, Aaltonen LM, Chen R, Ranki A, Partio EK, et al., Bio portfolio: lifelong persistence of variant and prototypic erythrovirus DNA genomes in human tissue. Proc. Natl. Acad. Sci. U. S. A. 2006; 103, 7450–7453. <https://doi.org/10.1073/pnas.0602259103>.
- [7] Kumar S, Gupta RM, Sen S, Sarkar RS, Philip J, Kotwal A, Sumathi SH. Seroprevalence of human parvovirus B19 in healthy blood donors. Medical journal armed forces India. 2013 Jul 1;69(3):268-72.
- [8] Kumari S, Kuruvilla Thomas R, Sruthi S, Barani R, Sangvi S, Krishnamoorthy R, Srikanth P. Increased parvovirus B19 seropositivity in healthy blood donors in India. Scientific Reports. 2024 Sep 3;14(1):20497.
- [9] Crabol Y, Terrier B, Rozenberg F, Pestre V, Legendre C, Hermine O, et al. Intravenous immunoglobulin therapy for pure red cell aplasia related to human parvovirus b19 infection: a retrospective study of 10 patients and review of the literature. Clin. Infect. Dis. 2013; 56, 968–977. <https://doi.org/10.1093/cid/cis1046>.
- [10] Pyoria L, Toppinen M, Mantyla E, Hedman L, Aaltonen LM, Vihinen-Ranta, et al. Extinct type of human parvovirus B19 persists in tonsillar B cells. Nat. Commun. 8, 14930. <https://doi.org/10.1038/ncomms14930>.
- [11] Arvia R, Margheri F, Stincarelli MA, Laurenzana A, Fibbi G, Gallinella G. 2020. Parvovirus B19 activates in vitro normal human dermal fibroblasts: a possible implication in skin fibrosis and systemic sclerosis. Rheumatology 59, 3526–3532. <https://doi.org/10.1093/rheumatology/keaa230>.
- [12] Bua G, Conti I, Manaresi E, Sethna P., Foster, S., Bonvicini, F., Gallinella, G., 2019. Antiviral activity of brincidofovir on parvovirus B19. Antivir. Res. 162, 22–29. <https://doi.org/10.1016/j.antiviral.2018.12.003>.
- [13] Guan, W., Wong, S., Zhi, N., Qiu, J. The genome of human parvovirus B19 can replicate in nonpermissive cells with the help of adenovirus genes and produces infectious virus. J. Virol. 2009;83, 9541–9553. <https://doi.org/10.1128/JVI.00702-09>.
- [14] Pozzuto T, Kietzell VK, Bock T, Schmidt-Lucke C, Poller W, Zobel T, et al. 2011. Transactivation of human parvovirus B19 gene expression in endothelial cells by adenoviral helper functions. Virology 411, 50–64.
- [15] Bock CT, Duchtig A, Uta F, Brunner E, Sy BT, Klingel K, et al.. Molecular phenotypes of human parvovirus B19 in patients with myocarditis. World J. Cardiol. 2014. Vol 6, 183–195. <https://doi.org/10.4330/wjc.v6.i4.183>.
- [16] Lindblom A, Heyman M, Gustafsson I, Norbeck O, Kaldensjo T, Vernby, A et al.. Parvovirus B19 infection in children with acute lymphoblastic leukemia is associated with cytopenia resulting in prolonged interruptions of chemotherapy. Clin. Infect. Dis. 2008; vol 46, 528–536. <https://doi.org/10.1086/526522>.
- [17] Eid AJ, Chen SF, Astidco P. Human parvovirus B19 in solid organ transplantation. Am. J. Transplant. 2013. 13 (Suppl. 4), 201–205. <https://doi.org/10.1111/ajt.12111>.
- [18] Koduri, P.R., Parvovirus B19-related anemia in HIV-infected patients. AIDS Patient Care STDS 2000;14, 7–11. <https://doi.org/10.1089/108729100318082>.
- [19] Adamson-Small LA, Ignatovich IV, Laemmerhirt MG, and Hobbs JA.. Persistent parvovirus B19 infection in non-erythroid tissues: possible role in the inflammatory and disease process. Virus Res. 2014; 190, 8–16. <https://doi.org/10.1016/j.virusres.2014.06.017>.
- [20] Thammasri K, Rauhamaki S, Wang L, Filippou A, Kivovich V, Marjomaki V, et al. Human parvovirus B19 induced apoptotic bodies contain altered self-antigens that are phagocytosed by antigen presenting cells. 2013; PLoS One 8, e67179. <https://doi.org/10.1371/journal.pone.0067179>.