



SUMMARY AND CONCLUSION

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Valve prosthesis of different construction may have different traumatic effect in the red blood cells. The hemolytic effect may be changed considerably even by modification of the same valve. The hemolysis may be sufficient to produce clinical hemolytic anemia or may cause a compensated hemolytic state.

Significant hemolytic anemia may be an important determinant of prognosis. In some cases, it may be a sign of leakage around the valve of Ball variance demanding surgical intervention.

The turbulence of blood and direct contact between red cells and solid surfaces are the most important causes.

The purpose of the present work is to evaluate the hemolytic effect of prosthetic materials and designs and to study the influence of a single and double valve replacement on hemolytic process. The study consisted of 39 patients following prosthetic heart valves implantation. They were investigated for:-

Hemoglobin concentration, Hematocrit value, reticulocyte count osmotic fragility test, serum LDH and serum haptoglobin. Patients were divided into three groups:- group I consisted of 14 patients with valvular heart replacement with tilting Disc valves of Bjork - Shiley or omniscience varieties.

Group II consisted of 10 patient with valvular heart replacement by Ball valve of the Starr - Edwards variety. Group III included 15 patients with valvular heart replacement by Ball and tilting Disc valves. Another group (group IV) represented the control group and consisted of 10 normal healthy persons.

The incidence of hemolysis compensated and uncompensated regardless of the type and site of valve replacement was 68%. It is 28% for group I, 16% for group II and 24% for group III.

Hemoglobin level and hematocrit values were below normal in 56% and 68% of cases respectively.

[The mean value of Reticulocyte count in group I, II and III were significantly increased from the control group $P < 0.05$ in all].

In 6 cases (15%) of cases the serum haptoglobin was completely absent, i.e., Zero. There was highly significant decrease of mean haptoglobin level of group I, II and III in comparison to the control group.

Serum LDH was at high level above the normal in all cases. Comparison of mean level of LDH between different groups studied were not significant but, when they were compared with the control group a highly statistically significant increase of LDH level was observed ($P < 0.001$).

In conclusion it is recommended to investigate regularly every patient with prosthetic heart valve for any sign of hemolysis, at least every three months to detect any hemolysis of minimal degree which may be complicated later on by hemolytic anemia or it may predict malfunction of the valve and require valve re-replacement.