CHAPTER V

CONCLUSION

- 1. The HFE polymorphisms (H63D and C282Y) was not found in the studied subjects at Maharaj Nakorn Chiang Mai Hospital.
- Iron overload was obvious in the patients with β-thalassemia/HbE disease and homozygous β-thalassemia. ZPP levels were higher in β-thalassemia patients, indicating that although iron overload complicates the clinical symptom in these patients, the utilization of iron are impaired.

