

CURRICULUM VITAE

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Education: 1971: B.Sc. (Medical Science)
Chiangmai University, Thailand
1973: M.D.
Chiangmai University, Thailand
1975-6: Intern,
Westsuburban Hospital, Oak Park, Illinois, USA.
1977: Diploma in Thai Board of Internal Medicine
1983: Diploma in Thai Board of Hematology
Positions: 1979-1980: Instructor
1981-1983: Assistant Professor
1983-1989: Associate Professor
1990: Professor
Professional Societies: 1982: The Society of Hematology of Thailand
1982: The Genetic Society of Thailand
1986: The Royal College of Physicians of Thailand
1988: The Science Society of Thailand
1989: The New York Academy of Sciences
1992: The American Society of Advances in Sciences
1997: Thai Academy of Science and Technology Foundation

1998: American Association for Clinical Chemistry
1999: American society of Hematology
2001: IMBN, Asia-Pacific Division

Member of Committees and Boards-University, National, International

1986-1990: Scientific Committee, 12th-16th Conference on Science and Technology of Thailand
1988-1994: Scientific committee, Mahidol Research Grant, Mahidol University
1989: Member of the Executive Committee and Chairman of The Academic Affair, Thalassemia Foundation of Thailand
1992-8: Member of the Research Committee, Faculty of Medicine Siriraj Hospital, Mahidol University
1992: Member of the Committee, Mahidol Award, Mahidol University
1994: Member of the Editorial Board, HEMOGLOBIN
1996: Member of the Editorial Board, Journal of the Medical Association of Thailand

Scholarship, Fellowship, Research Grant:

1984-1987: NIH Research Grant (co-investigator)
1985-1986: EEC (CEC) Research Grant (STD1)
1986: JSPS-NRCT Research Fellow, Kyoto University (Professor Takashi Murachi)
1987-1990: Monbusho Research Grant
1988-1993: JSPS-NRCT Research Grant
1989-1992: NIH Research Grant (co-investigator)
1989-1992: EEC (CEC) Research Grant (STD2)
1993-1995: EEC (CEC) Research Grant (STD3)
1994-1998: Prajadhipok-Rambhai Barni Foundation
1997-2000: IAEA
1997-2000: Senior Research Fellow Grant, National Science and Technology Development Agency, Thailand
2002: Outstanding Scientist Awards, Foundation for the Promotion of Science and Technology Under the Patronage of His Majesty The King of Thailand
2002-2005: Senior Research Scholar, Thailand Research Fund, Thailand
2006: Golden Silk Award, Guangxi Medical University
2002-2006: Genomewide search for disease modifier genes in thalassemia (NIH Research Grant)

- 2004-2006: Molecular mechanism of anemia in thalassemia mice and new therapeutic intervention (Mahidol University Research Grant)
- 2005-2008: Apoptosis (Mahidol University Research Grant)
- 2005-2008: Pathophysiologic changes and therapeutic intervention in thalassemia (Senior Research Scholar, Thailand Research Fund, Thailand)
- 2005-2010: Genetic modifier genes in beta thalassemia (BIOTEC, National Science and Technology Development Agency, Thailand)

Publications (selected publications from more than 350 articles)

1. Fucharoen S, Winichagoon P, Thonglairuam V, Siriboon W, Siritanaratkul N, Kanokpongsakdi S, Vantanasiri C. Prenatal diagnosis of thalassemia and hemoglobinopathies in Thailand: experience from 100 pregnancies. *Southeast Asian J Trop Med Public Health* 1991; 22: 16-29.
2. Winichagoon P, Fucharoen S, Wilairat P, Chihara K, Fukumaki Y, Wasi P. Identification of five rare mutations including a novel frameshift mutation causing β^0 -thalassemia in Thai patients with β^0 -thalassemia/hemoglobin E disease. *Biochim Biophys Acta* 1992; 1139: 280-6.
3. Fucharoen S, Winichagoon P. Thalassemia in Southeast Asia: problems and strategy for prevention and control. *Southeast Asian J Trop Med Public Health*. 1992; 23: 647-55.
4. Winichagoon P, Fucharoen S, Kanokpongsakdi S, Fukumaki Y. Detection of α -thalassemia-1 (Southeast Asian type) and its application for prenatal diagnosis. *Clin Genet* 1995; 47: 318-20.
5. Fucharoen S, Siritanaratkul N, Winichagoon P, Chowthaworn J, Siriboon W, Muangsup W, Chaicharoen S, Poolsub N, Chindavijak B, Pootrakul P, Piankijagum A, Schechter AN, and Rodgers GP. Hydroxyurea increases hemoglobin F levels and improves the effectiveness of erythropoiesis in β -thalassemia/hemoglobin E disease. *Blood* 1996; 87: 887-92.
6. Fucharoen S, Winichagoon P, Wisedpanichkij R, Sae-Ngow B, Sriphanich R, Oncoung W, Muangsapaya W, Chowthaworn J, Kanokpongsakdi S, Bunyaratvej A, Piankijagum A, Dewaele C. Prenatal and postnatal diagnoses of thalassemias and hemoglobinopathies by HPLC. *Clinical Chemistry* 1998; 44: 740-8.
7. Fucharoen S, Winichagoon P, Siritanaratkul N, Chowthaworn J, Pootrakul P. Alpha and beta-thalassemia in Thailand. *Ann NY Acad Sci* 1998; 850: 412-4.
8. Winichagoon P, Saechan V, Sriphanich R, Nopparatana C, Kanokpongsakdi S, Maggio A, Fucharoen S. Prenatal diagnosis of β -thalassaemia by reverse dot-blot hybridization. *Prenat Diagn* 1999; 19: 428-35.
- *9. Fucharoen S, Winichagoon P. Clinical and hematologic aspects of hemoglobin E β -thalassemia. *Current Opinion in Hematology* 2000; 7: 106-112.
10. Lacerra G, Sierakowska H, Carestia C, Fucharoen S, Summerton J, Weller D, Kole R. Restoration of hemoglobin A synthesis in erythroid cells from peripheral blood of thalassemic patients. *PNAS* 2000; 97: 9591-6.

11. Fucharoen S, Ketvichit P, Pootrakul P, Siritanaratkul N, Piankijagum A, Wasi P. Clinical manifestation of β -thalassemia/hemoglobin E disease. *J Pediatric Hematology/Oncology* 2000; 6: 552-7.
12. Winichagoon P, Fucharoen S, Chen P, Wasi P. Genetic factors affecting clinical severity in beta-thalassemia syndromes. *J Pediatr Hematol Oncol* 2000; 22: 573-80.
13. Fucharoen S, Winichagoon P. Molecular epidemiology of α -thalassemia. *J Clin Genetics and Tribal Research* 2000; 5: 1-15.
14. Imai K, Tientadakul P, Opartkiattikul N, Luenee P, Winichagoon P, Svasti J, Fucharoen S. Detection of haemoglobin variants and inference of their functional properties using complete oxygen dissociation curve measurements. *Br J Haematol* 2001; 112: 483-7.
15. Tang D.C., Fucharoen S, Ding I, Rodgers G.p. Rapid differentiation of five common α -thalassemia genotypes by polymerase chain reaction. *J Lab Clin Med* 2001; 137: 290-5.
16. Aulia D, Timan I.S., Tatsumi N, Funahara Y, Asada Y, Bunyaratvej A, Fucharoen S, kondo T, Kondo H. Field evaluation of WHO hemoglobin color scale in West Java in 2000. Reprinted from *Journal of Analytical Bio-Science* Vol. 24, No. 2, 2001; 159; 24 : 159-61.
17. Old J, Khan S, Verma I, Fucharoen S, Kleanthous M, Ioannou P, Kotea N, Fisher C, Riazuddin S, Saxena R, Wichagoon P, Kyriacou K, A1-Quobaili F, Khan B. A multi-center study in order to further define the molecular basis of β -thalassemia in Thailand, Pakistan, Sri Lanka, Mauritius, Syria, and India and to develop a simple molecular diagnostic strategy by amplification refractory mutation system-polymerase chain reaction. *Hemoglobin* 2001; 25: 397-407.
18. Svasti S, Yodsowan B, Sriphanich R, Winichagoon P, Boonkhan P, Suwanban T, Sawangaretrakul P, Srisomsap C, Ketudat-Cairns J, Svasti J, Fucharoen S. Association of Hb Hope [β 136(H14)Gly \rightarrow Asp] and Hb H disease. *Hemoglobin* 2001; 25: 429-35.
19. Turbpaiboon C, Svasti S, Sawangareetakul P, Winichagoon P, Srisomsap C, Siritanaratkul N, Fucharoen S, Wilairat P, Svasti J. Hb Siam [α 15(A13) Gly(Arg(α 1) (GGT \rightarrow CGT)] is a typical α chain hemoglobinopathy without an α -thalassemic effect. *Hemoglobin* 2002; 26: 77-81.
20. Sawangaretrakul P, Svasti S, Yodsowan B, Winichagoon P, Srisomsap C, Svasti J, Fucharoen S. Double Heterozygosity for Hb Pyrgos [β 83(EF7)Gly \rightarrow Asp] and Hb E [β 26(B8)Glu \rightarrow Lys] found in Association with α -Thalassemia. *Hemoglobin* 2002; 26(2): 191-6.
21. Fucharoen S, Winichagoon P. Thalassemia and abnormal hemoglobin. *Int J Hematology (Supplement II)* 2002; 76: 83-89.
22. Suwanmanee T, Sierakowska H, Lacerra G, Svasti S, Kirby S, Walsh CE., Fucharoen S, Kole R. Restoration of human β -globin gene expression in murine and human IVS2 654 thalassemic erythroid cells by free uptake of antisense oligonucleotides. *Mol Pharmacol* 2002; 62: 545-53.
23. Svasti S, Hieu TM, Munkongdee T, Winichagoon P, Be TV, Binh TV, Fucharoen S. Molecular analysis of β -thalassemia in South Vietnam. *Am J Hematol* 2002; 71: 85-8.
24. Winichagoon P, Sriphanich R, Sae-ngow B, Chowthaworn J, Tantisirin P, Kanokpongsakdi S, Fucharoen S, Wasi P. Application of automated HPLC in prenatal diagnosis of thalassemia. *Laboratory Hematology* 2002; 8: 29-35.

25. Suwanmanee T, Sierakowska H, Fucharoen S, Kole R. Repair of a splicing defect in erythroid cells from patients with β -thalassemia/H bE disorder. *Molecular Therapy* 2002; 6: 718-26.
26. Jamsai D, Nefedov M, Narayanan K, Orford M, Fucharoen S, Williamson R, Ioannou P A. Insertion of common mutations into the human β -globin locus using GET Recombination and an EcoRI endonuclease counterselection cassette. *J Biotechnology* 2003; 101: 1-9.
27. Chui D H K, Fucharoen S, Chan V. Hemoglobin H disease: not necessarily a benign disorder. *Blood* 2003; 101: 791-800.
28. Phumala N, Porasuphatana S, Unchern S, Pootrakul P, Fucharoen S, Chantharaksri U. Hemin: a possible cause of oxidative stress in blood circulation of β -thalassemia/Hemoglobin E disease. *Free Radical Research* 2003; 37: 129-35.
29. Jamsai D, Orford M, Fucharoen S, Williamson R, Ioannou P A. Insertion of modifications in the β -globin locus using GET Recombination with single-stranded oligonucleotides and denatured PCR fragments. *Molecular Biotechnology* 2003; 23: 29-36.
30. Jamsai D, Orford M, Nefedov M, Fucharoen S, Williamson R, Ioannou P A. Targeted modification of a human β -globin locus BAC clone using GET recombination and an I-*SceI* counterselection cassette. *Genomics* 2003; 82: 68-77.
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33. Unchern S, Laoharuangpanya N, Phumala N, Sipankapracha P, Pootrakul P, Fucharoen S, Wanachivanawin W, Chantharaksri U. The effects of vitamin E on platelet activity in β -thalassemia patients. *Br J Haematol* 2003; 123: 738-44.
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40. Jamsai D, Zaibak F, Khongnium W, Vadolas J, Voullaire L, Fowler K J, Gazeas S, Fucharoen S, Williamson R, Ioannou P A. A humanized mouse model for a common β^0 -thalassemia mutation. *Genomics* 2005; 85: 453-61.
41. Ngiwsara L, Srisomsap C, Winichagoon P, Fucharoen S, Sae-Ngow B, Svasti J. Hb kurosaki [α 7(A5)Lys \rightarrow Glu (AAG \rightarrow GAG)]: an 2α -globin gene mutation found in Thailand *Hemoglobin* 2005; 29(2): 155-59.
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44. Sripichai O, Whitacre J, Munkongdee T, Kumkhaek C, Makarasara W, Winichagoon P, Abel K, Braun A, Fucharoen S. Genetic analysis of candidate modifier polymorphisms in Hb E- β^0 -thalassemia patients. *Ann NY Acad Sci* 2005; 1054: 433-8.
45. Fucharoen S. Genotypes and phenotypes of thalassemia: A discussion. *Ann NY Acad. Sci.* 2005; 1054: 518-21.
46. Watanapokasin Y, Chuncharunee S, Sanmund D, Kongnium W, Winichagoon P, Rodgers G P, Fucharoen S. In vivo and in vitro studies of fetal hemoglobin induction by hydroxyurea in β -thalassemia/Hb E patients. *Exp Hematol* 2005; 33: 1486-92.
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48. Chumpia W, Peerapittayamongkol C, Angchaisuksiri P, Komanasin N, Muta K, Kuaha K, Iida H, Inoue S, Wada Y, Kurihara M, Hamasaki N, Fucharoen S. Single nucleotide polymorphisms and haplotypes of protein C and protein S genes in the Thai population. *Blood Coagulation and Fibrinolysis* 2006; 17: 13-8.
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50. Peng C, Fucharoen S, Kontoghiorghes G J, Tsai C. Report on the proceedings of the 15th international conference on oral chelation (ICOC) in the treatment of thalassemia and other diseases at Taichung, Taiwan, April 22-26, 2005 *Hemoglobin* 2006; 30: 63-8.
51. Svasti S, Kole R, Fucharoen S. Repair of incorrectly spliced β -globin mRNA by antisense oligonucleotide. *J Med Tech Assoc Thailand* 2006; 34: 1369-77.287.
52. Jamsai D, Zaibak F, Vadolas J, Voullaire L, Fowler K J, Gazeas S, Peters H, Fucharoen S, Williamson R, Ioannou P A. A humanized BAC transgenic/knockout mouse model for HbE/ β -thalassemia. *Genomics* 2006; 88: 309-15.
53. Changsri K, Akkarapathumwong V, Jamsai D, Winichagoon P, Fucharoen S. Molecular mechanism of high Hb F production in Southeast Asian-type HPFH. *Int J Hematol* 2006; 83: 229-37.

54. Srichairatanakool S, Ounjaijean S, Thephinlap C, Khansuwan U, Phisalpong C, Fucharoen S. Iron-chelating and free-radical scavenging activities of microwave-processed green tea in iron overload. *Hemoglobin* 2006; 30: 311-27.
55. Jintaridith P, Srisomsap C, Vichittumaros K, Kalpravidh R W, Winichagoon P, Fucharoen S, Svasti M.R. J, Kasinrerak W. Chicken egg yolk antibodies specific for the γ chain of human hemoglobin for diagnosis of thalassemia. *Int J Hematol.* 2006; 83: 408-14.
56. Luechapudiporn R, Phumala Morales N, Fucharoen S, Chantharaksri U. The reduction of cholesteryl linoleate in lipoproteins: and index of clinical severity in β -thalassemia/Hb E. *Clin Chem Lab Med* 2006; 44: 574-81.
57. Tankanitlert J, Howard T A, Tamsakulphong A, Sirankapracha P, Morales N P, Sanvarind Y, Fucharoen P, Ware R E, Fucharoen S, Chantharaksri U. A pharmacokinetic study of paracetamol in Thai β -thalassemia/Hb E patients. *Eur J Clin Pharmacol* 2006; 62: 743-48.
58. Morales N P, Charlermchoung C, Luechapudiporn R, Yamanont P, Fucharoen S, Chantharaksri U. Lipid fluidity at different regions in LDL and HDL of β -thalassemia/Hb E patients. *Biochemical and Biophysical Research Communications* 2006; 350: 698-703.
59. Svasti S, Paksua S, Nuchprayoon I, Winichagoon P, Fucharoen S. Characterization of a novel deletion causing $(\delta\beta)^0$ -thalassemia in a Thai family. *Am J Hematol* 2006; 82: 155-61.
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64. Sripichai O, Fucharoen S. Genetic polymorphisms and implications for human diseases. *J Med Assoc Thai* 2007; 90: 394-8.
65. Winichagoon P, Svasti S, Winichagoon P, Chitchumroonchokchai C, Fucharoen S. Expression of β^E and γ -globin genes in infants heterozygous for hemoglobin E and double heterozygous for hemoglobin E and α -thalassemia. *Haematologica.* 2007; 92: 702-3.
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74. Limenta Lie <ichael George, Jirasomprasert T, Tankaniltert J, Svasti S, Wilairat P, Chantharaksri U, Fucharoen S, Phumala Morales N. UGT1A6 genotype-related pharmacokinetics of deferiprone (L1) in healthy volunteers. *Br J Clin Pharmacol*; 2008; 65: 908-916.
75. Jearawiriyapaisarn N, M Moulton H, Buckley B, Roberts J, Sazani P, Fucharoen S, L Iversen P, Kole R. Sustained Dystrophin Expression Induced by Peptide-conjugated Morpholino Oligomers in the Muscles of *mdx* Mice. *American Society of Gene Therapy*; 2008; 16: 1624-1629.
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