

## CHAPTER VI

## STUDY ON HUMAN CONTROL SERA AND SLE SERA

Human control sera and SLE sera specimens were collected from the patients at the Institute of Dermatology of Thailand, as indicated below:-

	healthy human sera	20 cases			
Control sera	Other autoimmune disease (Alopecia,	20 cases			
	Renal disease , Pemphigus, RA,				
	PSS, Bullous pemphigoid)				

		patients with SLE followed ARA	oriteria	
Sample sera	- Active	20	cases	
	- Inactive	20	cases	
		patients with SLE not followed	20	cases
		ARA criteria		

All specimens were studied for Anti-DNA antibodies by the proposed method in comparison with Farr technique (Radio immuno assay), and antinuclear antibody by immunofluorescence technic (ANF or FANA)

The principle of the proposed method used and the other methods were shown in table 8.

TABLE 8 COMPARISON OF THE PRINCIPLE OF THE ELISA PROPOSED

METHOD AND OTHERS

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	Enzyme-linked immunosorbent assay (proposed method)	Immuno- fluorescence (FANA)	Farr ammonium sulfate precipitation (Radio immuno assay)
Layer 1.	Polystyrene	Rat Liver cells	test serum (Anti-DNA)
Layer 2.	DNA (calf thymus)	test serum	isotopically labeled
		(antinuclear antibody)	Ag (C <sup>14</sup> DNA)
Layer 3.	test serum	Fluorescence	sat. (NH <sub>4</sub> ) <sub>2</sub> SO <sub>4</sub>
	(Anti DNA)	anti-human IgG conjugate	
Layer 4.	anti-human IgG	***	
	labeled with peroxid	ase	
	Color yield	fluorescence	radioactive counting
System		microscopy	
Quanti-	serial dilution	Serial dilution	radioactivity
tation	enzyme reaction		proportional to Ab
	/	\ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \	binding capacity