## List op Planned Associated Projects in the NORDKEM Project 5/89 on Medical Need for Quality Specifications in Laboratory Medicine

<u>No.</u>	<u>Participants</u>	<u>Title</u>
1.	Hørder M,Olivarius N, Jørgensen PJ	, Diabetes in Primary
	Bihlet I, Hyltoft Petersen P	Health Care
2.	Fraser C, Hyltoft Petersen P	Quality Specifications
	Antonsen S	for Detection Limit
3.	Grodum E, Hyltoft Petersen P,	Reference interval for
	Hangaard J, Bollerslev J et al.	cortisol at CRH-test
4.	Djurhus S, Rohold A, Vadstrup S,	Reference intervals
	Hyltoft Petersen P	for SPotassium
5.	Magid E, Hyltoft Petersen P et al.	Evaluation of clinical
		tests
6.	Heedman P-A, Olsson S, Larsson O	Prevalence of idiop.
	Ritter B, Marsell R, Östergaard H,	hemochromatosis and
	Olander B, Åkerblom Å	hypercholest. 18 y men
7.	Arends J, Hyltoft Petersen P,	Qual. spec. maternal
	Nørgaard-Pedersen B	S-α-Fetoproteinanalys
8.	Uldall A, Blaabjerg O	Planning & implement.
		of Nordic reference S
9.	Tryding N et al.	Analyt. specificity,
		tests in primary care
10.	Nilsson Ehle P	Qual. spec. SCholest
11.	Westgard J O, Hyltoft Petersen P	SCholest.Clin strat,
		anal. qual.,control
		system
12.	Sandberg S	Qual. spec. for tests
		in primary care
13.	Kaikola H-L	_ **
14.	Penttilä I M	Qual. spec. Sα-Feto-
		protein
15.	Lindstedt G	Qual. spec. thyroid
		hormones
16.	Lindstedt G	Qual. spec. prostate
		specific antigen
17.	Lytken-Larsen M, Fraser C,	Qual. spec HbA <sub>lc</sub>
	Hyltoft Petersen P	

18.	Kofstad J, Momsen G, Gade	Qual. spec. BpH,pCO <sub>2</sub> ,
	Christensen N	pO2, Ca <sup>++</sup>
19.	Nilsson Ehle H	Qual. spec. iron de-
		ficiency parame-
		ters
20.	Linnet C	Anal. goals for P
		Bilirubin (unconj)
		determinations
21.	Linnet C	Anal. goals for P
		Creatinine determ.
22.	Dybkaer R, Hyltoft Petersen P,	Nomenclature
	de Verdier C-H et al.	
23.	Stenman U-H	Qual. spec. for
		human choriogonado-
		tropin (hCG)
24.	Brandslund I, Borg Rasmussen J	Conc $\alpha_1$ -antitrypsin
		in workers in dusty
		work areas
25.	Sorto A, Kaihola H-L	Quality goals & orga-
		nization of QC in pri-
		mary care lab.

In the associated projects the quality specification work should be performed for each analytical quantity in each specified situation. Models for estimation of needed quality and for specification of acceptable analytical performance and critical errors have been developed during the past ten years. These models should be applied to the clinical strategy in the project, whether a screening, a diagnostic situation or a time-related pathological process. Each clinical situation should be analyzed concerning the type of model to be used, and - if no relevant model is available - to develop or elaborate new and useful models.

The main project group must be directly involved by providing the associated groups with the available literature and, if needed, to help the groups to find their relevant models or to try to develop new useful models. On the other hand the main project group should try to compile the collected knowledge and make it useful.