Home sampling as alternative to venipuncture: results, patient experience and implementation

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Venipuncture

Benefits

- Scalable in volume
- Short logistics
- Processed same day

Challenges

- Trained personnel
- Collection at hospital
- Extra visit to hospital
- Geographically limited



Finger prick

Benefits

- sample at home
- no extra visit to hospital/outpatient clinic
- easily scalable studies
- Testing is done centrally in laboratory

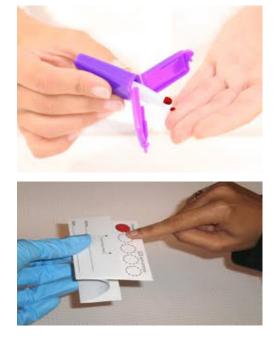
Challenges

- Small volumes
- Sample stability
- Self sampling
- Logistics



Fingerprick collection devices

- Sample type and processing
- Determine volume Hct
- Labour involved



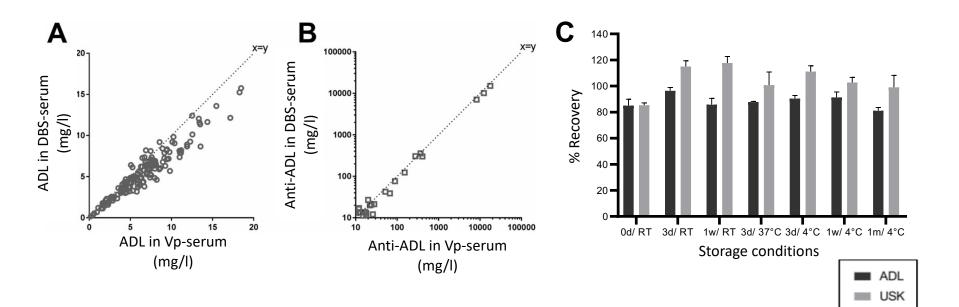


Berends SE et al. Ther Drug Monit 2020; Berends SE et al. Br J Clin Pharmacol 2019; Bloem K et al. Bioanalysis 2018; Kneepkens EL et al. Br J Clin Pharmacol.2017



Tuesday, 21 December 2021

Finger prick comparability and stability in DBS





Our finger prick set





Our finger prick set















> 90% succesful finger prick

Table 2. Evaluation of the fingerprick								
		fingerprick 2454)	Failed fingerprick (n=223)					
	Patients (n=1648)	Controls (n=806)	Patients (n=171)	Controls (n=52)				
Failed fingerprick – no. (%)								
Succeeded to collect blood in the tube								
$Yes \ge 3 drops$	1280/1361 (94)	656/680 (96)	57/110 (52)	24/37 (65)				
Yes, but < 3 drops	80 (6)	23 (3)	36 (33)	12 (32)				
No	1 (0.1)	1 (0.1)	17 (15)	1 (3)				
Reason for little to no collection*								
Not enough blood to form a whole drop	N.A.	N.A.	29 (26)	11 (30)				
Drop formation but it did not fall into the tube	N.A.	N.A.	26 (24)	7 (19)				
Enough blood formation, but unable to collect it in the tube	N.A.	N.A.	22 (20)	6 (16)				

Participant experience is positive

Table 2. Evaluation of the fingerprick									
	Successful fingerprick (n=2454)			Failed fingerprick (n=223)					
	Patients (n=1648)			Controls (n=806)		Patients (n=171)		Controls (n=52)	
Experience fingerprick - no. (%)									
Experience of the prick									
Less painful than expected	172/875	(20)	70/410	(17)	11/94	(12)	2/22	(9)	
As painful as expected	616	(70)	316	(77)	71	(76)	20	(91)	
More painful than expected	87	(10)	24	(6)	12	(13)	0		
Experience of the complete process									
Positive	492/875	(56)	238/410	(58)	25/94	(27)	6/22	(27)	
Neutral	321	(37)	160	(39)	39	(42)	13	(59)	
Negative	62	(7)	12	(3)	30	(32)	3	(14)	

Finger prick is prefered in research and study setting

Table 2. Evaluation of the fingerprick								
	Successful fingerprick (n=2454)			Failed fingerprick (n=223)				
	Patients (n=1648)			trols 806)	Patients (n=171)		Controls (n=52)	
Preferences for blood sampling meth	od – no. (%)							
Preference for scientific research								
Fingerprick at home	476/1167	(41)	312/437	(71)	27/96	(28)	11/22	(50)
Venepuncture at healthcare	336	(29)	39	(9)	57	(59)	6	(27)
institute								
No preference	355	(30)	86	(20)	12	(13)	5	(23)
Preference for routine care								
Fingerprick at home	222	(19)	168	(38)	12	(13)	6	(27)
Venepuncture at healthcare	503	(43)	83	(19)	58	(61)	6	(27)
institute								
No preference	441	(38)	186	(43)	25	(26)	10	(46)

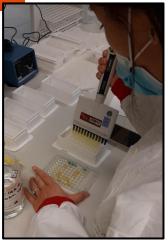


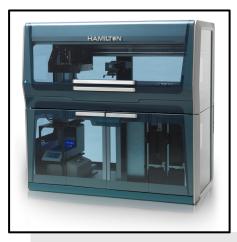
Sample process under control

As little as 5-10 μl serum

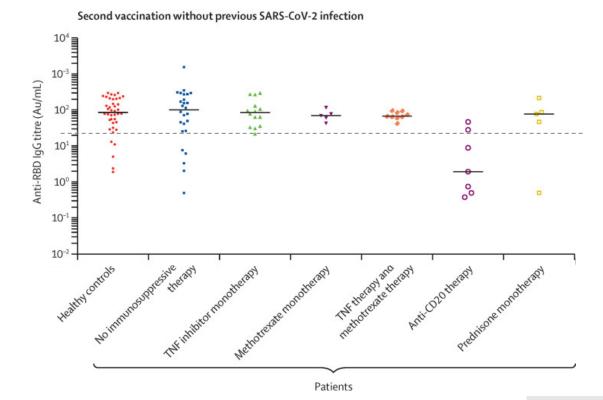








Finger prick used in SARS-CoV2 study



Brakenhoff et al. Trials (2021) 22:412 https://doi.org/10.1186/s13063-021-05241-5

The RECOVAC IR study: the immune response and safety of the mRNA-1273 COVID-19 vaccine in patients with chronic kidney disease, on dialysis or living with a kidney transplant

Marcia M.L. Kho¹, Marlies E.J. Reinders¹, Carla C. Baan¹, Debbie van Baarle^{2,3}, Frederike J. Bemelman⁴, Dimitri A. Diavatopoulos⁵, Ron T. Gansevoort⁶, Fiona R.M. van der Klis³, Marion P.G. Koopmans⁷, A. Lianne Messchendorp⁶, Renate G. van der Molen⁸, Ester B.M. Remmerswaal⁹, Nynke Rots³, Priya Vart^{6,10,11}, Rory D. de Vries⁷, Luuk B. Hilbrands ⁽¹⁾ ¹² and Jan-Stephan F. Sanders⁶ and RECOVAC Collaborators^{*}

LETTER

A prospective, randomized, single-blinded, crossover trial to investigate the effect of a wearable device in addition to a daily symptom diary for the remote early detection of SARS-CoV-2 infections (COVID-RED): a structured summary of a study protocol for a randomized controlled trial

Timo B. Brakenhoff¹⁺¹, Billy Franks^{1†}, Brianna Mae Goodale², Janneke van de Wijgert³, Santiago Montes⁴, Duco Veen^{1,5,6}, Eskild K. Fredslund⁷, Theo Rispens⁸, Lorenz Risch^{9,10,11}, Ariel V. Dowling¹², Amos A. Folarin^{13,14,15}, Patricia Bruijning³, Richard Dobson¹⁴, Tessa Heikamp¹, Paul Klaver¹, Maureen Cronin^{2†}, and Diederick E. Grobbee^{15†} On behalf of the COVID-RED Consortium¹⁶

Antibody development after COVID-19 vaccination in patients with autoimmune diseases in the Netherlands: a substudy of data from two prospective cohort studies

Laura Boekel, Maurice Steenhuis, Femke Hooijberg, Yaëlle R Besten, Zoé L E van Kempen, Laura Y Kummer, Koos P J van Dam, Eileen W Stalman, Erik H Vogelzang, Olvi Cristianawati, Sofie Keijzer, Gestur Vidarsson, Alexandre E Voskuyl, Luuk Wieske, Filip Eftimov, Ronald van Vollenhoven, Taco W Kuijpers, S Marieke van Ham, Sander W Tas, Joep Killestein, Maarten Boers, Michael T Nurmohamed, Theo Rispens, Gertjan Wolbink

Summary

Background Data are scarce on immunogenicity of COVID-19 vaccines in patients with autoimmune diseases, who are Lancet Rheumatol 2021



Trials











Summary

We can accurately measure antibodies in small volumes









Acknowledgements

Sanquin

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Study partnerships Participants