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Morimont, Laure; Didembourg, Marie; Haguet, Helene; Modaffari, Elise; Tillier, Maxence; Lebreton, Aurélien; Bouvy, Céline; Dogné, Jean-Michel; Douxfils, Jonathan

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Inter-laboratory variability of the standardized ETP-based APC resistance assay





L. Morimont^{1,2}, M. Didembourg², H. Haguet², E. Modaffari¹, M. Tillier³, C. Bouvy¹, A. Lebreton³, J-M. Dogné², J. Douxfils^{1,2}

¹Qualiblood sa, Namur, Belgium; ²University of Namur, Faculty of Medicine, Department of Pharmacy, Namur Research Institute for Life Sciences (NARILIS), Namur Thrombosis and Hemostasis Center (NTHC), Namur, Belgium; ³ Service d'hématologie biologique, CHU Clermont-Ferrand, Clermont-Ferrand, France.

Laure.morimont@qualiblood.eu

BACKGROUND

- Regulatory bodies recommend assessing the impact of steroid contraceptives on ETP-based APC resistance assay during their development.
- This assay was recently validated and standardized.

Table 1:Intra-run (N=5) and inter-run (N=3) repeatability (N=5) of the commercial reference plasma and the 3 quality controls. Results are expressed as mean inhibition $\% \pm SD$

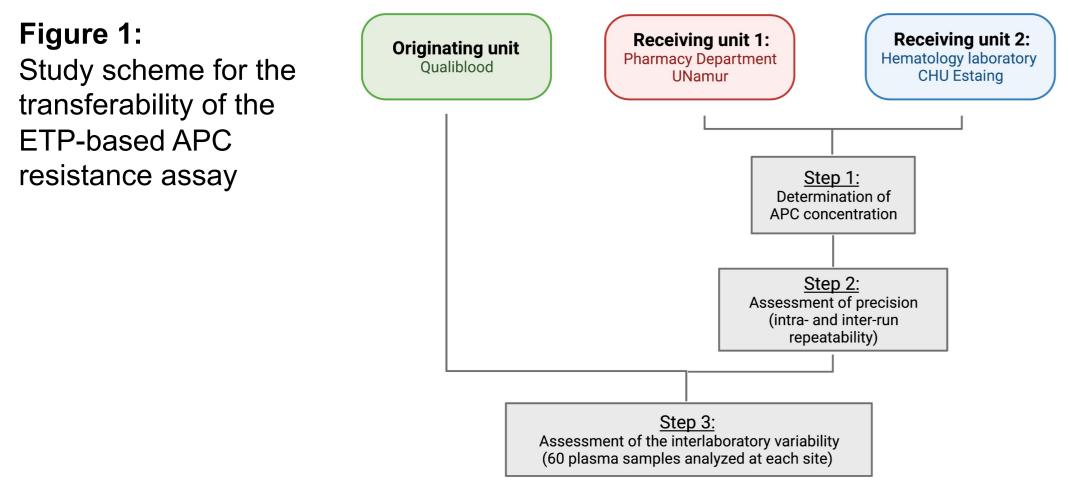
Tested plasma	Intra-run repeatability		Inter-run repeatability	
	Unit 1	Unit 2	Unit 1	Unit 2
Reference plasma	78.8% ±0.4%	75.3% ±1.2%	80.9% ±01.9%	78.5% ±1.2%
QC low	100.0% ± 0.0%	100.0% ±0.0%	100.0% ± 0.0%	97.9% ±1.9%
QC intermediate	40.2% ± 0.7%	37.7% ±0.9%	42.8% ± 2.6%	39.6% ±2.3%
QC high	2.8% ± 0.7%	4.0% ±1.5%	5.0% ± 2.3%	6.0% ±2.1%

AIMS

To assess the inter-laboratory transferability of the ETP-based APC resistance assay.

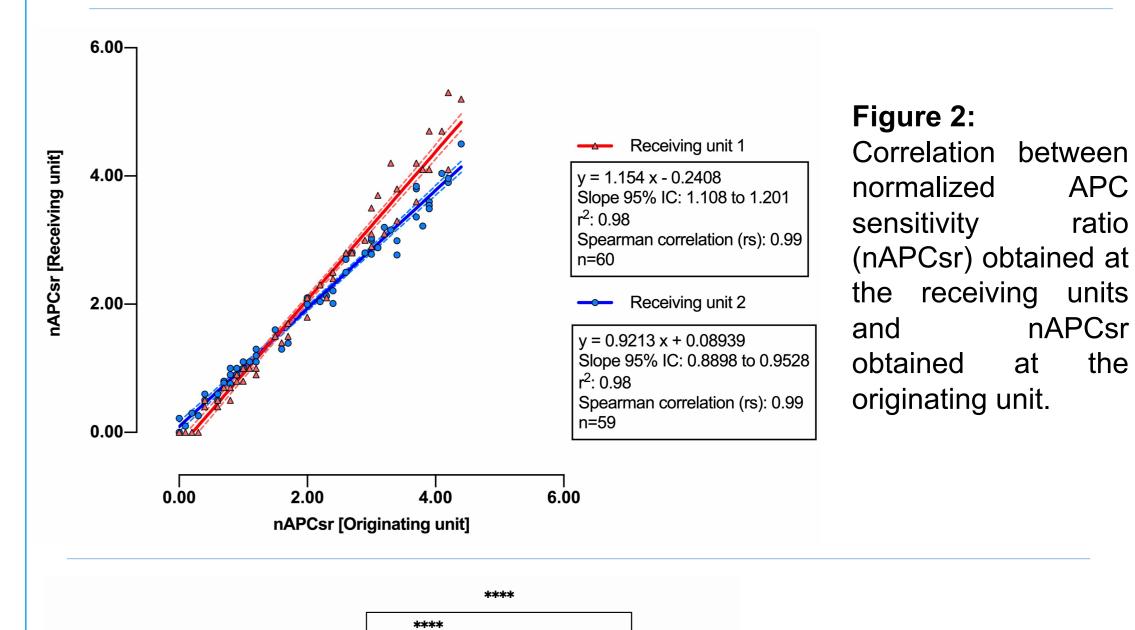
METHODS

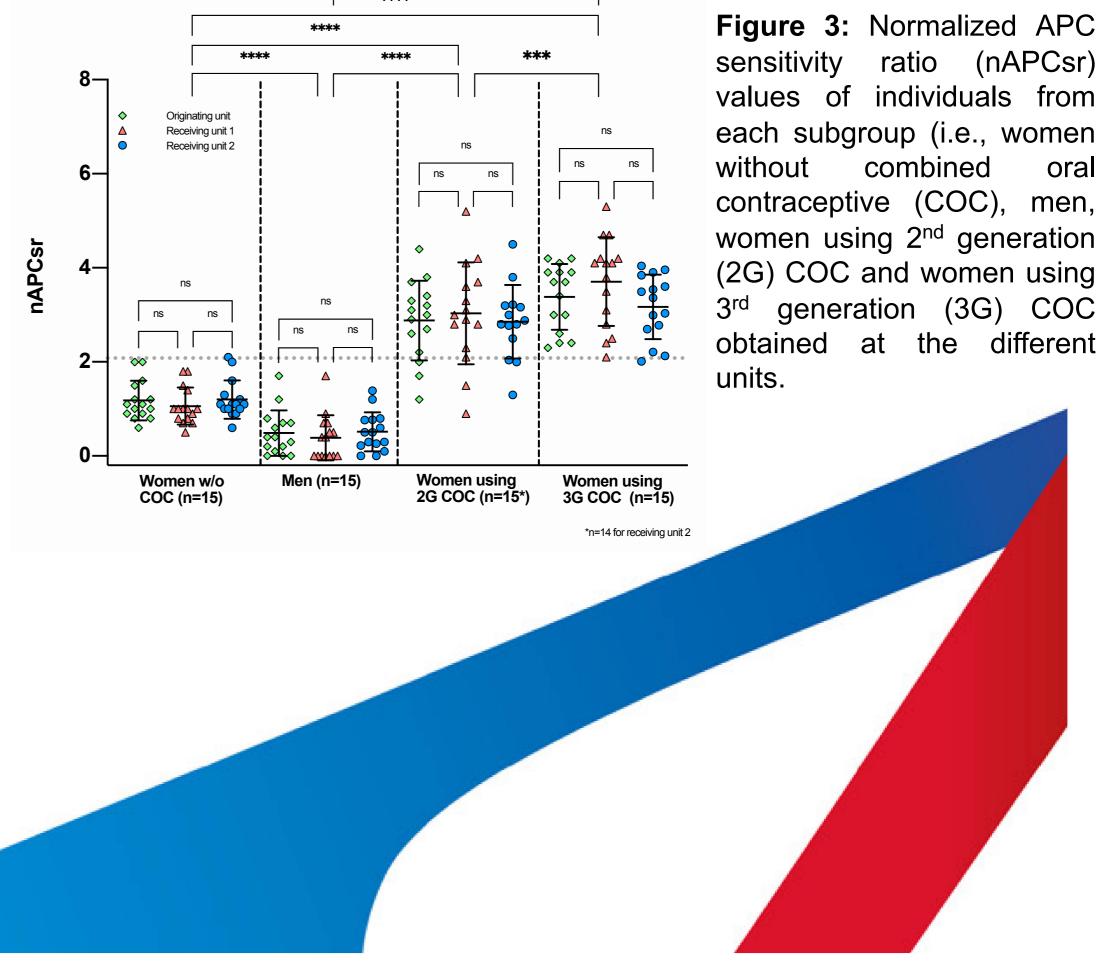
The study scheme is described in figure 1



RESULTS

- APC concentration was defined at 1.21 µg/mL and 1.14 µg/mL in receiving unit 1 and receiving unit 2 respectively.
- Intra- and inter-run (Table 1) repeatability showed SD below 3% in both receiving units.
- Spearman correlation showed effective pairing between the originating and the receiving units (Figure 2).
- The sensitivity of the test was maintained and subgroups analysis still reported significant differences between healthy individuals and women using combined oral contraceptives (Figure 3).





SUMMARY/CONCLUSION

- Excellent intra-laboratory precision and inter-laboratory reproducibility.
- The normalized APC sensitivity ratio obtained with this validated methodology, provides an appropriate sensitivity irrespective of the laboratory in which the analysis is performed.

