Comparative Thromboelastography on Peripheral vs Uterine Blood as an Early Marker of Postpartum Coagulation Changes

Tyler Guidugli DO, Allison A. Mootz MD, Noor Raheel MBChB, Sebastian Seifert MD, John J. Kowalczyk MD, Michaela K. Farber MD, MS

Introduction

Postpartum hemorrhage (PPH): Leading cause of maternal mortality worldwide¹ Early prediction of systemic coagulopathy may facilitate prompt treatment



Viscoelastic Testing (TEG & ROTEM): Rapid detection of coagulopathy, targeted therapy² Prior case series of ROTEM during early PPH \rightarrow coagulopathy in vaginal vs peripheral blood³



AIM: Compare peripheral and uterine blood using TEG in routine cesarean delivery (CD)

1. Say et al, 2014. 2. Collins et al, 2022. 3. Ma'ayeh et al, 2021





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Methods



A priori power analysis for paired comparison: n=7 to detect a 30% reduction in TEG maximum amplitude between peripheral and uterine samples





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Results





Peripheral TEG samples \implies n=10, normal coagulation for pregnant patients

Uterine cavity TEG samples \implies n=5, rapid clotting, unable to perform TEG \implies n=5, clot in citrated tube, TEG showed profound coagulopathy



No association between PPH & uterine TEG coagulopathy (p=0.524) Unable to compare traditional labs – all uterine samples rejected due to clot







Discussion



Ma'ayeh *et al.* demonstrated profound **coagulopathy of vaginal blood** compared to **normal peripheral coagulation** in 3 patients with PPH.



Confounders: Amniotic fluid, cervical mucosa, vaginal fluid, vaginal flora, and clot

Some similarity with our TEG results ...

Figure 1: Peripheral blood TEG normal coagulation.



Figure 2: Uterine blood TEG profound coagulopathy.



References: 1. Ma'ayeh et al, 2021





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Discussion



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Formation of **clot despite citrate**. Likely unable to inhibit since clotting cascade already initiated.

Limitations:

- Applies only to uterine samples
- TEG 6s uses microfluidic channels to transport sample

Figure 3: Citrated vacutainer of uterine blood sample.



Figure 4: Clotted TEG cartridge from uterine blood sample.





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Uterine vs peripheral blood coagulation at time of delivery, prior to PPH, demonstrates **immediate clotting** of the **uterine** samples.



No correlation between PPH and ability to run TEG on uterine blood samples.



Uterine blood appears to be a **non-viable** method of assessing **early coagulopathy** at the time of hysterotomy closure.



Figure 5: Fresh uterine sample, immediate clot formation.





