



# Institutional Repository - Research Portal

## Dépôt Institutionnel - Portail de la Recherche

researchportal.unamur.be

## RESEARCH OUTPUTS / RÉSULTATS DE RECHERCHE

### Is Thrombin Time Useful To Guide Peri-Procedural Management For Patients On Dabigatran Etxilate?: An In Vitro Validation Study

Mullier, François; Douxfils, Jonathan; Baes, Hélène; Baudar, Justine; Lessire, Sarah; Gourdin, Maximilien; Classen, Jean-François; Chatelain, Christian; Chatelain, Bernard; Dogne, Jean-Michel

*Publication date:*  
2013

*Document Version*  
Early version, also known as pre-print

[Link to publication](#)

*Citation for published version (HARVARD):*

Mullier, F, Douxfils, J, Baes, H, Baudar, J, Lessire, S, Gourdin, M, Classen, J-F, Chatelain, C, Chatelain, B & Dogne, J-M 2013, 'Is Thrombin Time Useful To Guide Peri-Procedural Management For Patients On Dabigatran Etxilate?: An In Vitro Validation Study', New Orleans, United States, 7/12/13 - 10/12/13, .

#### General rights

Copyright and moral rights for the publications made accessible in the public portal are retained by the authors and/or other copyright owners and it is a condition of accessing publications that users recognise and abide by the legal requirements associated with these rights.

- Users may download and print one copy of any publication from the public portal for the purpose of private study or research.
- You may not further distribute the material or use it for any profit-making activity or commercial gain
- You may freely distribute the URL identifying the publication in the public portal ?

#### Take down policy

If you believe that this document breaches copyright please contact us providing details, and we will remove access to the work immediately and investigate your claim.

# Is thrombin time useful to guide Peri-procedural Management for patients on dabigatran?:

## An *in vitro* validation study

François Mullier<sup>1</sup>, Jonathan Douxfils<sup>2</sup>, Hélène Baes<sup>1</sup>, Justine Baudar<sup>1</sup>, Sarah Lessire<sup>3</sup>, Maximilien Gourdin<sup>3</sup>, Jean-François Classen<sup>1</sup>, Bernard Chatelain<sup>1</sup>, Jean-Michel Dogné<sup>2</sup>

<sup>1</sup>Hematology laboratory Namur Thrombosis and Hemostasis Center (NTHC), Namur Research Institute for Life Sciences (NARILIS), CHU Dinant-Godinne UCL Namur, Belgium; <sup>2</sup>Department of Pharmacy, Namur Thrombosis and Hemostasis Center (NTHC), Namur Research Institute for Life Sciences (NARILIS) University of Namur, Namur, Belgium; <sup>3</sup>Department of Anesthesiology, Namur Thrombosis and Hemostasis Center (NTHC), Namur Research Institute for Life Sciences (NARILIS), CHU Dinant-Godinne UCL Namur, Belgium

### Background

Possibilities to monitor the intensity of dabigatran etexilate (DE) treatment may be valuable before urgent intervention. The Working Group on Perioperative Haemostasis proposed that the drug plasma concentration (I) should be less or equal to 30ng/mL. However, plasma levels where it is safe to carry out an invasive procedure or surgery have not been confirmed prospectively. In addition, no biological test has been correlated with bleeding risk. Literature showed heterogeneous data regarding the arrest of DE depending on the renal function and the haemorrhagic risk. In addition, physicians request guidance for patients excluded from the clinical trials. Consequently, there is still a need for a rapid and widely available biological test.

Some authors make proposals based on activated Partial Thromboplastin Time (aPTT), Hemoclot Thrombin Inhibitor® (HTI) or Thrombin Time (TT).<sup>1,2</sup> TT displayed several advantages over aPTT. However, TT is affected by numerous analytic variables.<sup>1</sup>

### Objectives

1. To determine the optimal [thrombin] with a variety of instruments and reagents
2. To assess the repeatability of TT at optimized conditions
3. To compare the sensitivity and linearity of TT at residual [dabigatran] (DA) with those of aPTT and HTI
4. To validate the fibrinogen reagent (with heparin inhibitor) for TT experiments

### Methods

Dabigatran was spiked at increasing [ ] in pooled citrated normal human platelet-poor plasma (NPP). The following [DA] were prepared: 0, 5, 10, 20, 30, 40 and 50ng/ml.

#### Optimal [thrombin] determination.

- The optimal [thrombin] was defined with 2 reagents: bovine thrombin (HemosIL® TT) and human thrombin (STA®-Thrombin) on 4 instruments: STA-R Evolution®, ACL-TOP®, CS2000i® and KC10®.
- TT higher than Tmax is not informative for the physician. Thus, the optimized [thrombin] was defined as the maximum [thrombin] giving, in a reproducible way, a TT at 50ng/ml lower than Tmax and a TT at 0ng/ml higher than the minimum TT. The Tmax was set arbitrarily at 120 sec.
- The repeatability of optimized TT was assessed by running aliquots of NPP spiked with all [DA] on 10 consecutive days
- TT (1.5 NIH/ml, STA®-Thrombin), aPTT with SynthasIL® and STA®-C.K.Prest, and HTI were also determined within 1 hour on STA-R® on replicates of all NPP solutions.
- STA®- Fibrinogen-5 was diluted to the optimized [thrombin] (1.5 NIH/ml) on STA-R® and used as a thrombin reagent.

### Conclusions

TT may be more informative than aPTT and HTI to provide with guidance to carry out an urgent procedure or surgery for patients receiving DE. However, TT is affected by a lot of analytic variables that should be understood by laboratories. Each laboratory should optimize its TT procedure according to its combination coagulometer-reagent.

### Results and Discussion

The thrombin origin is a more important variable in comparison to the type of coagulometer. At [DA] of 30ng/ml and [thrombin] of 1.5 NIH/ml, the STA®-TT ranges from 56 sec to 74 sec according to the instrument, whereas on a same instrument, the TT varied of minimum 43 sec, depending on the thrombin origin.

#### The optimized [thrombin] are the following :

- a) STA-R®, STA®-Thrombin : 1.5 NIH/ml,
- b) STA-R®, HemosIL® TT : 3.8 NIH/ml,
- c) ACLTOP®, STA®-Thrombin 3.8 NIH/ml,
- d) ACLTOP®, HemosIL® TT 5.0 NIH/ml,
- e) KC10®, STA®-Thrombin 1.5 NIH/ml,
- f) KC10®, HemosIL® TT 5.0 NIH/ml,
- g) CS2000i®, STA®-Thrombin 1.4 NIH/ml,
- h) CS2000i®, HemosIL® TT 3.8 NIH/ml.

Except for STA®-Thrombin on STA-R®, the [optimized] is not the one recommended by the manufacturer. Figure 1 shows the results of optimized TT according to thrombin origin and coagulometer.

Repeatability experiments showed that variability increased with the [DA] (i.e. STA®-Thrombin (STA-R®): coefficient of variation: 8.0% and 28.9% for 0ng/ml and 50ng/ml, respectively) and that the variability depends on the coagulometer and the reagent.

Comparison of sensitivity and linearity of TT, aPTT and HTI are presented in Table 1. APTT is not sensitive enough in low [DA], whatever the reagent, whereas HTI is not suitable in [DA] lower than 50ng/ml.

TT performed with fibrinogen reagent shows longer times in comparison to thrombin reagent (>120 sec vs 109 sec at 30ng/ml DA).

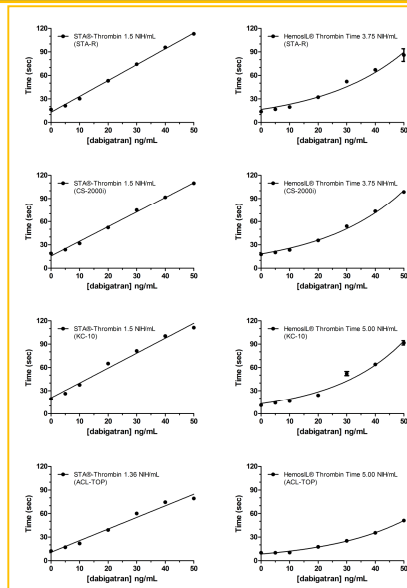


Figure 1: Comparison of the clotting time obtained with the different platform-reagent combinations

Table 1: Clotting time of optimized thrombin time, aPTT with different reagents and comparison with HTI

[Dabigatran] (ng/ml)	TT (sec)	aPTT (sec, SynthasIL®)	aPTT (sec, STA®-C.K.Prest)	HTI (ng/ml)
0	17.9	30.3	31.2	0.0
5	22.0	31.2	33.1	0.0
10	31.5	32.4	34.5	0.0
20	59.3	34.9	38.6	7.2
30	82.9	36.0	39.6	10.3
40	100.9	38.0	42.8	21.3
50	120.0	39.3	44.2	27.4

### Disclosure

The authors have no relevant conflicts of interest to disclose.

### References

1. Douxfils J, Mullier F, Robert S, et al. Thromb Haemost 2012 May;107(5):985-97.
2. Douxfils J, Dogné J-M, Mullier F, Chatelain B, Ronquist-Nil Y, Malmström R.E, Hjemdahl P. Thromb Haemost. 2013; 110(3): 543-9

### Contacts

J. Douxfils : jonathan.douxfils@unamur.be,  
J.M. Dogné : jean-michel.dogne@unamur.be,  
F. Mullier : mullierfrancois@gmail.com