

# *Imaging & Physiology Summit*

## ***A New Avenue to The Coronary Microcirculation: Absolute Flow And Resistance Measurement by The Pressure Wire***

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# A NEW AVENUE TO THE CORONARY MICROCIRCULATION

*Presently, there is a growing interest in studying the Microcirculation of the heart, but....*

- Present invasive techniques to assess the microcirculation, are crude, inaccurate, and extremely operator-dependent
- Doppler: - *inaccuracy of measurement  $\geq 20\%$* 
  - *adequate signals in  $\leq 70\%$  of patients*
  - *signal extremely operator-dependent*
- IMR: - *easier to perform*
  - *but also inaccuracy of  $\geq 15\%$*
  - *and also operator-dependent*

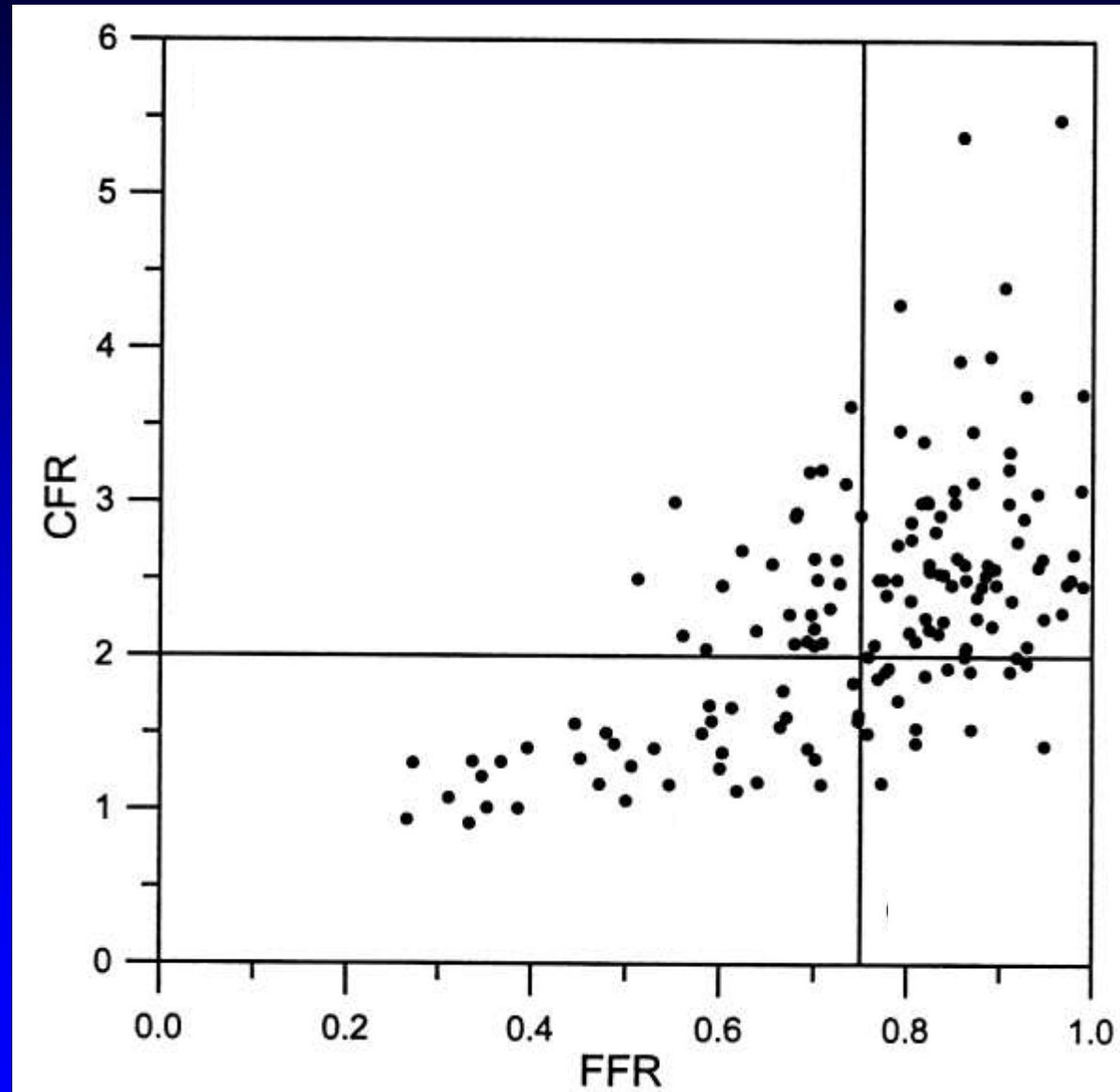
# ***A NEW AVENUE TO THE CORONARY MICROCIRCULATION***

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***Consequently, a lot of our knowledge about the microcirculation is speculative and open for multiple interpretations***

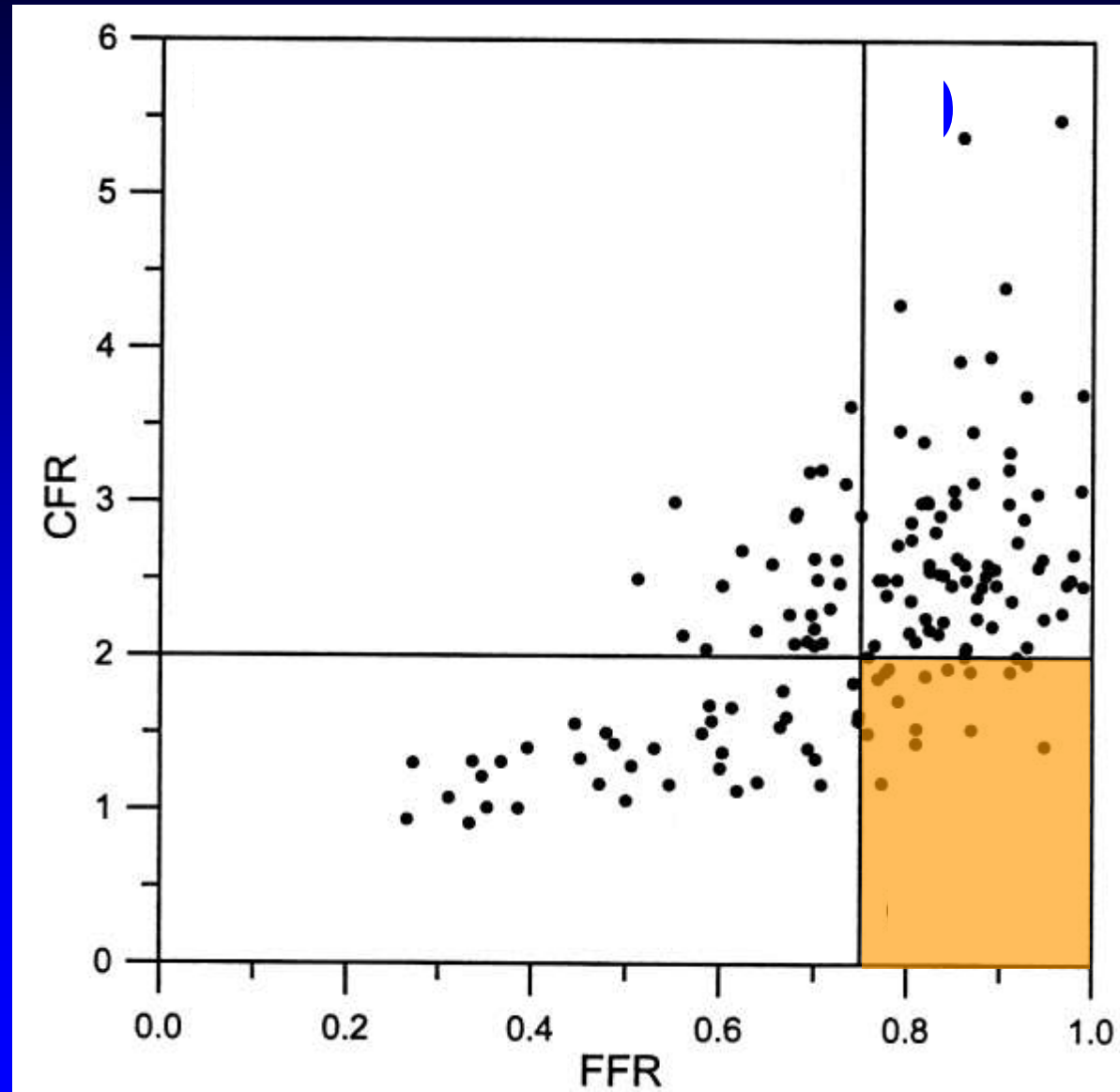
# Use of CFR and FFR to classify microvascular disease

150 patients,  
CFR & FFR  
measurement



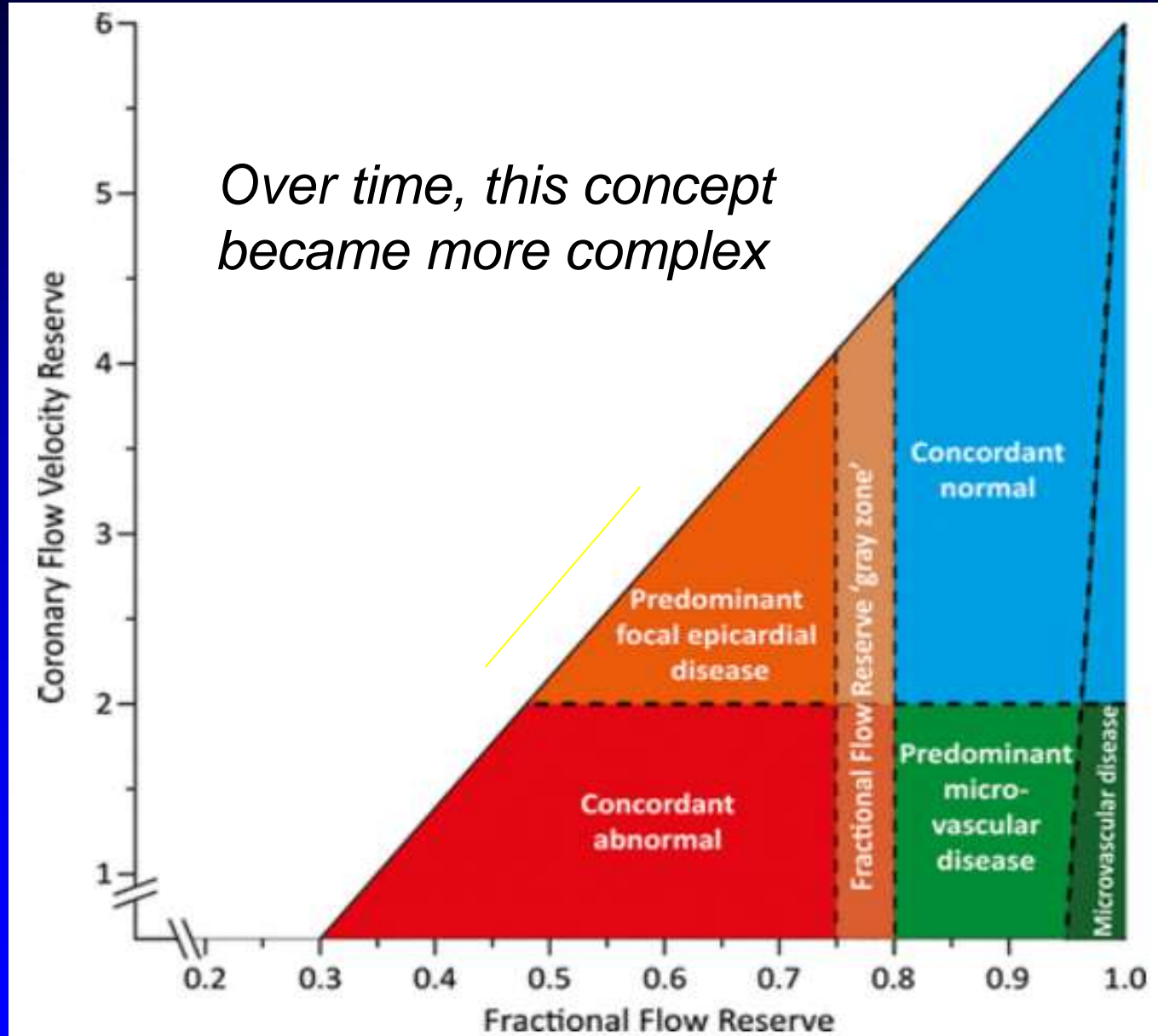
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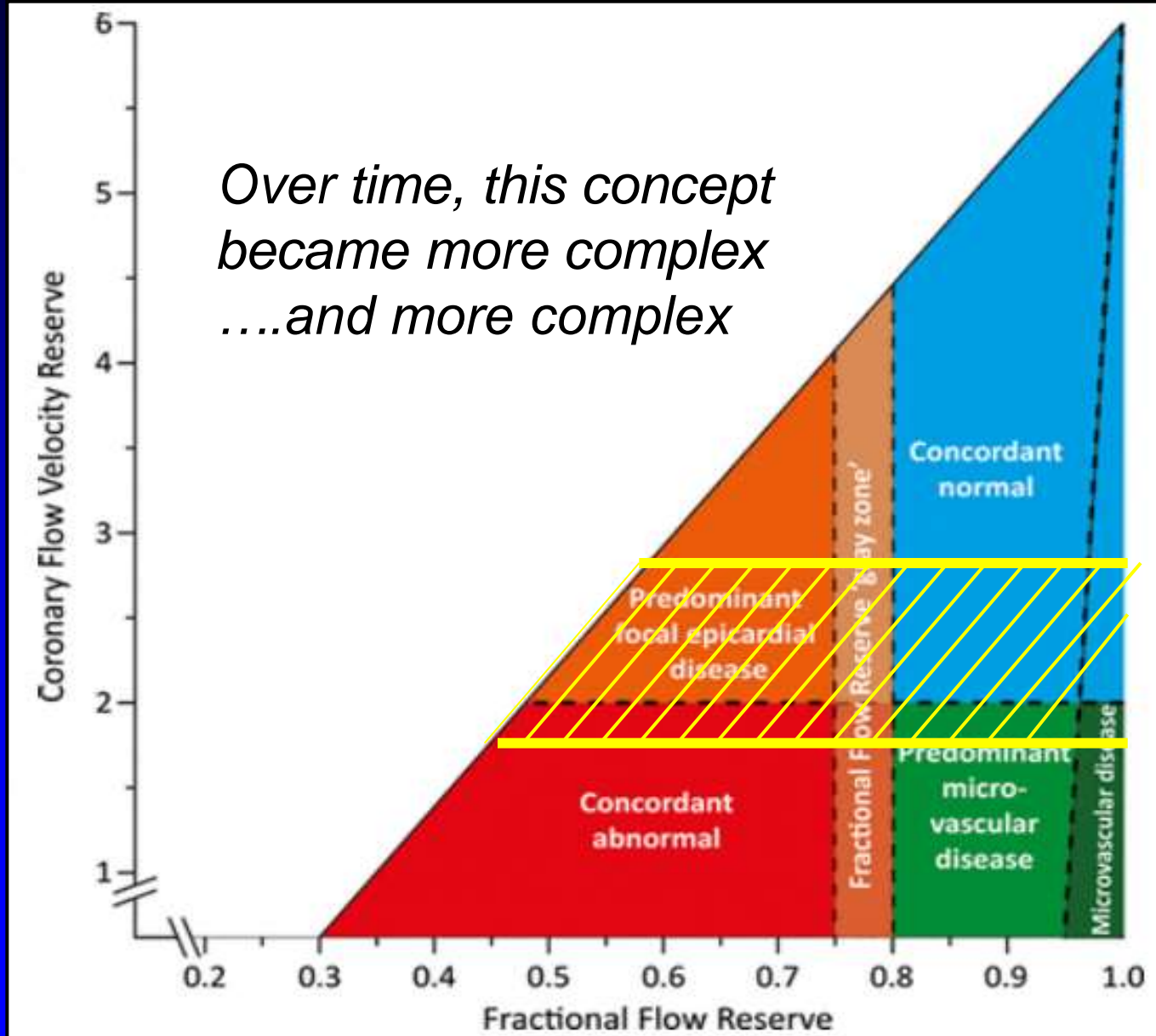
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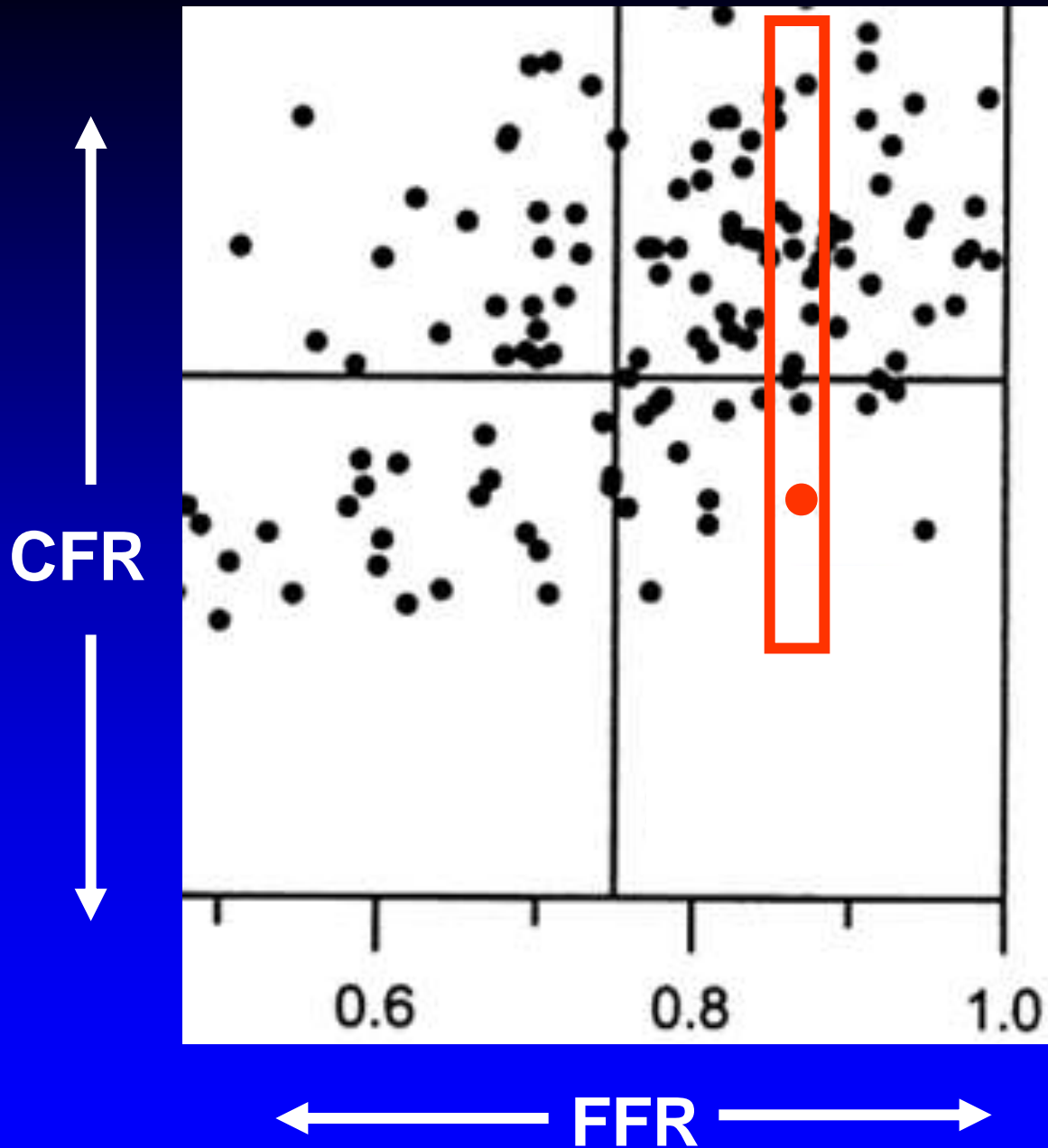
*Over time, this concept became more complex*



# Use of CFR and FFR to classify microvascular disease

*Over time, this concept became more complex  
....and more complex*





Variability in horizontal direction (**FFR**) = **2 %**

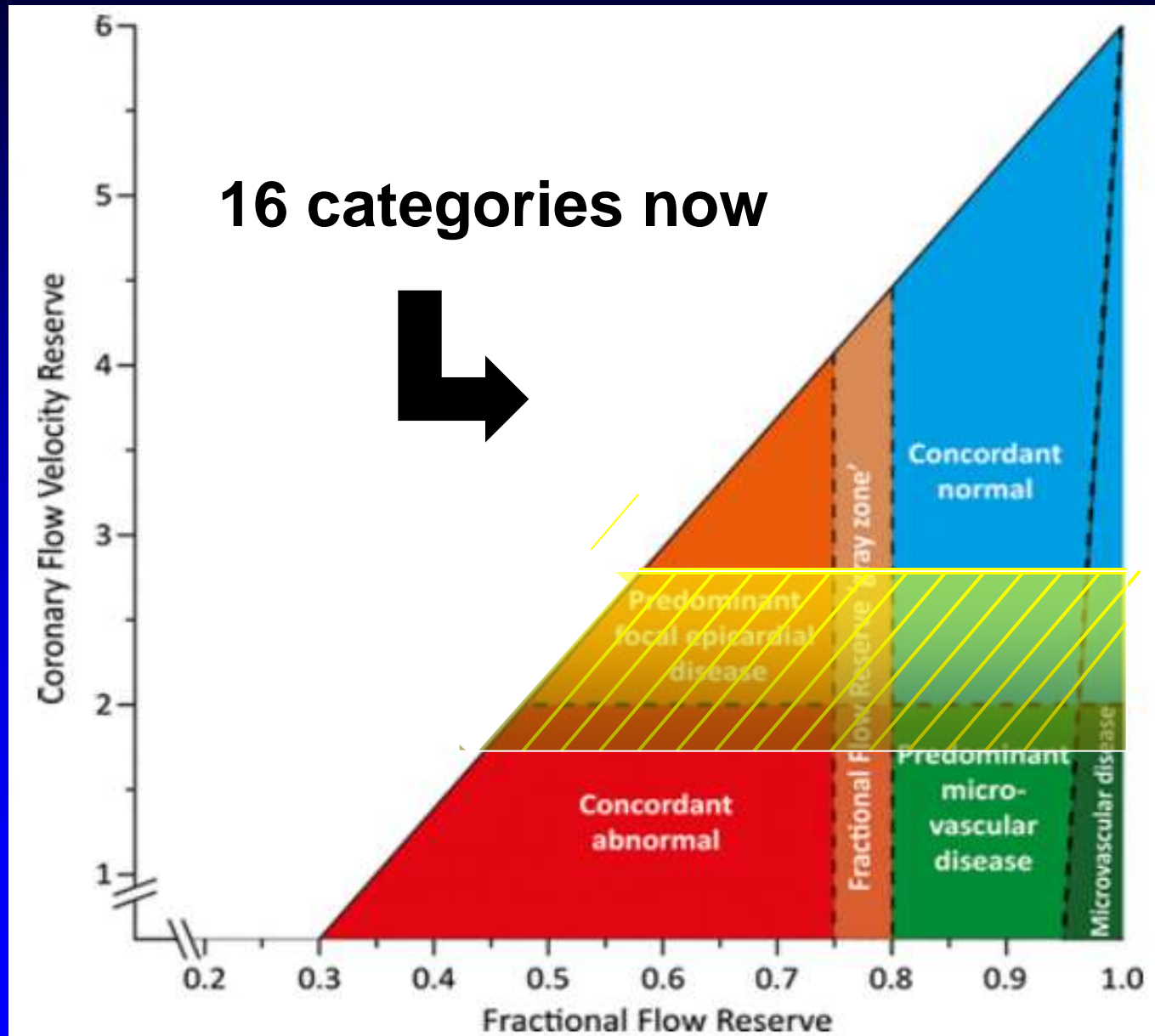
Variability in vertical direction (**CFR**) = **20 %**



*No reliable conclusions about microcirculation possible if flow cannot be assessed more accurately !*



# Use of CFR and FFR to classify microvascular disease



# Use of CFR and FFR to classify microvascular disease

*Next year, it might look like this.....!*



# ***A NEW AVENUE TO THE CORONARY MICROCIRCULATION***

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***So it might be clear:***

***No reliable conclusions about microcirculation are possible if flow cannot be assessed more accurately !***

# ***A NEW AVENUE TO THE CORONARY MICROCIRCULATION***

*The ideal technique to assess the microcirculation, should be:*

- easy to perform with standard PCI equipment
- accurate and reproducible
- operator-independent

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**→** *Measurement of absolute flow and resistance by thermodilution and continuous infusion of saline*

PRINT

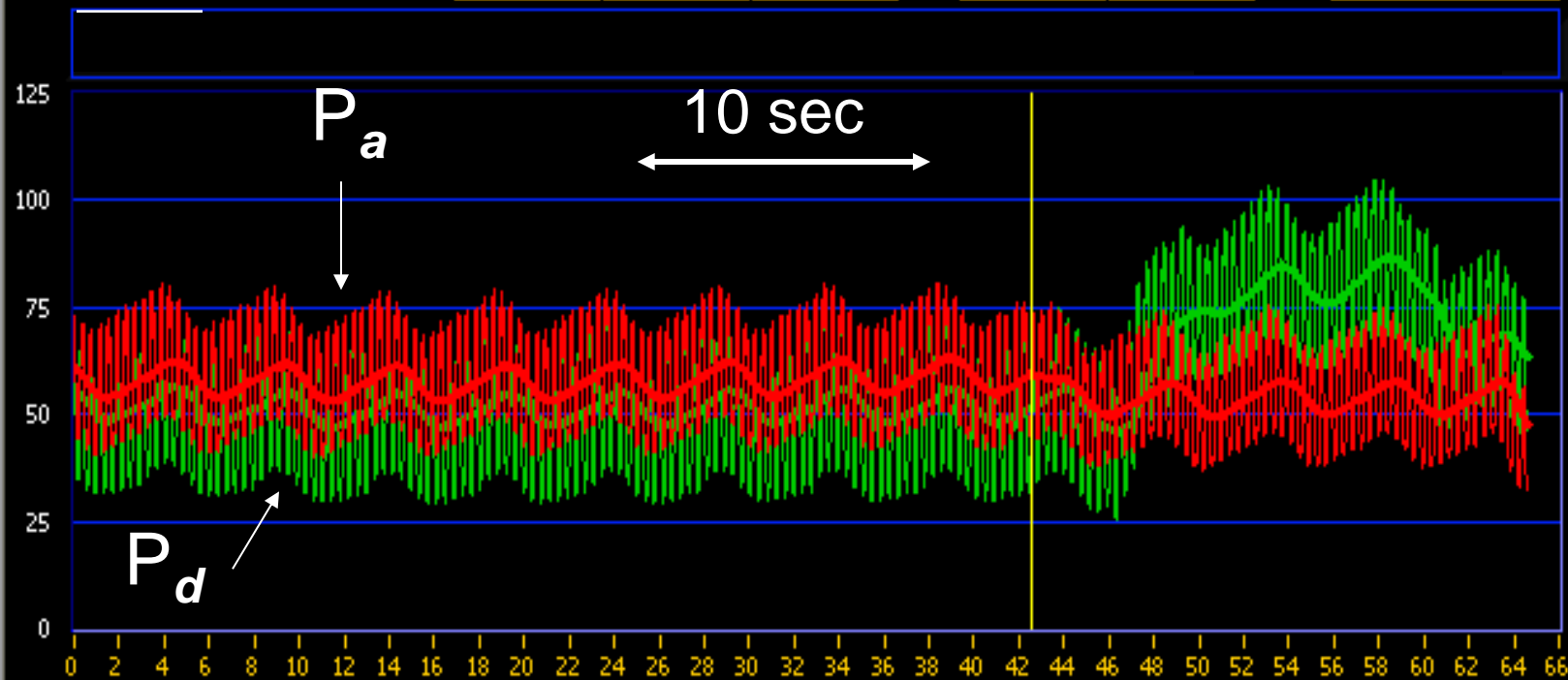
EDIT

RENAME

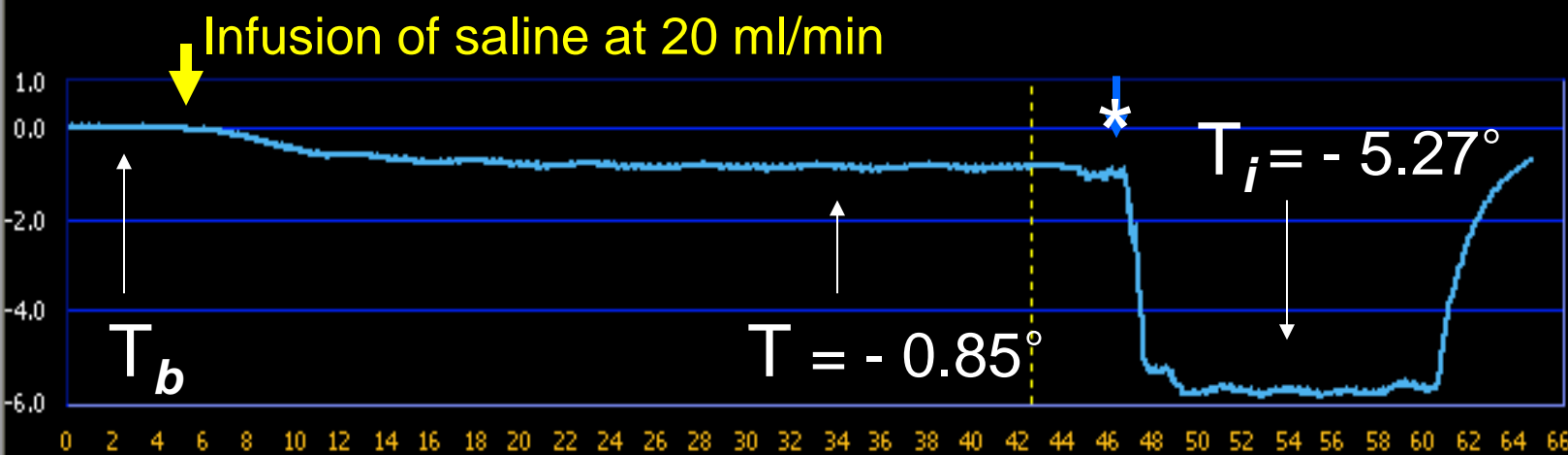
EXPORT

ERASE

SETUP

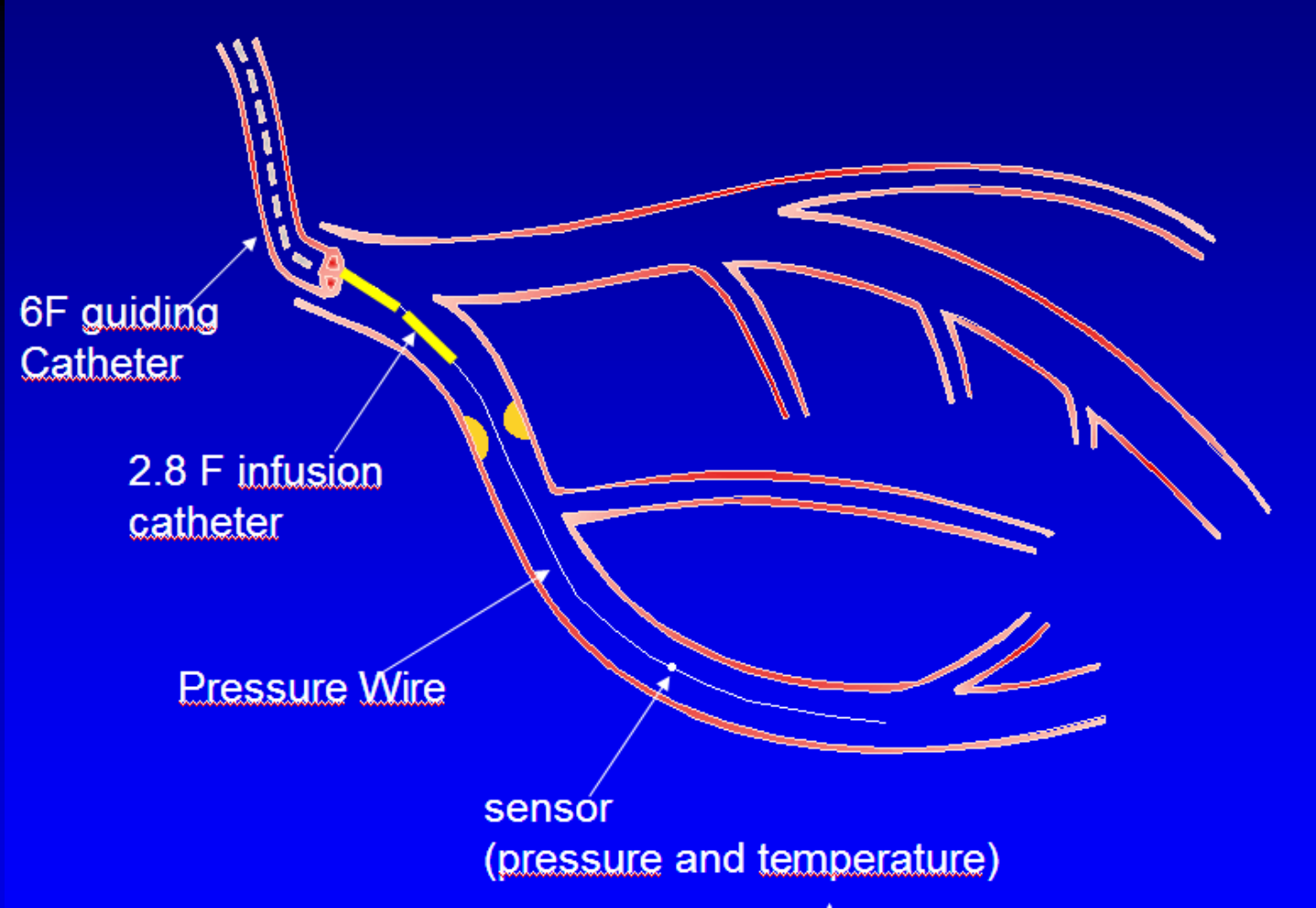
59  
Pa mean51  
Pd mean0.86  
FFR-0.85  
dT42.5  
CURSOR

RESET



$$Qb = 20 \times (-5.27 / -0.85) \times 1.08 = 134 \text{ ml/min}$$





saline infused at 20 ml/min

temperature of saline is 5° below blood temperature

after mixing, temperature of mixtate is 1° below blood temp

→ blood flow must be 5 x infusion flow of saline

Absolute  
max coronary  
blood flow



$$Q_b = Q_i \cdot \frac{T_i}{T} \cdot 1.08$$

infusion flow



temp of infusion at tip  
of guiding catheter

temp of blood at  
sensor position

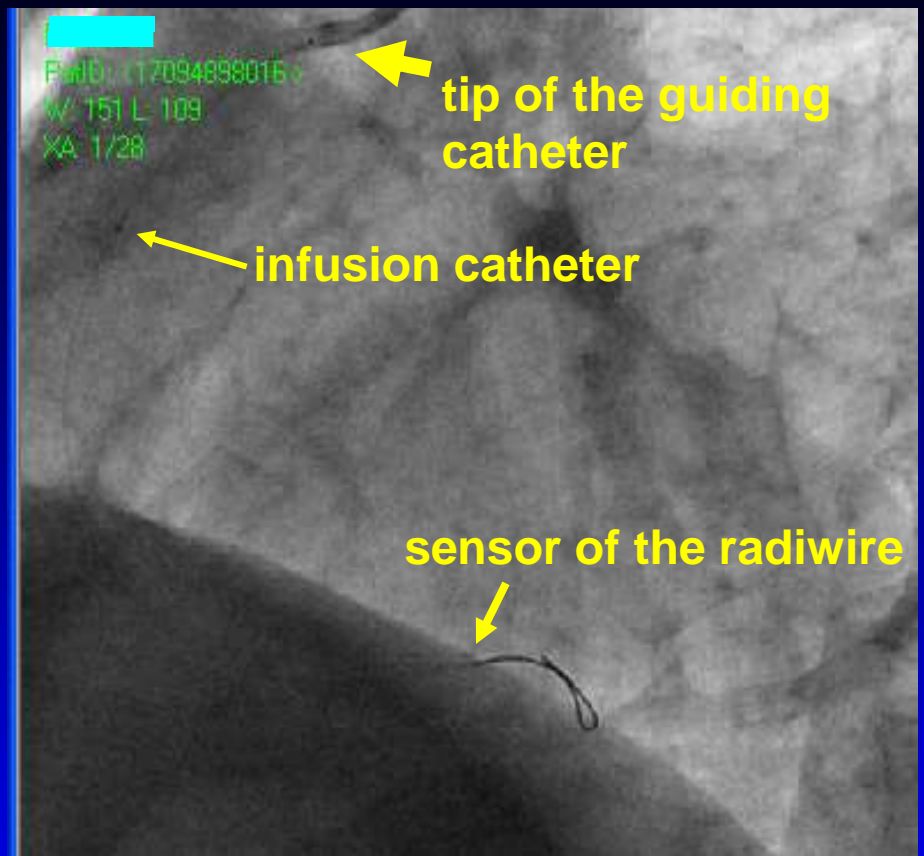
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PRINT

EDIT

RENAME

EXPORT

ERASE

SETUP

59

Pa mean

51

Pd mean

0.86

FFR

-0.85

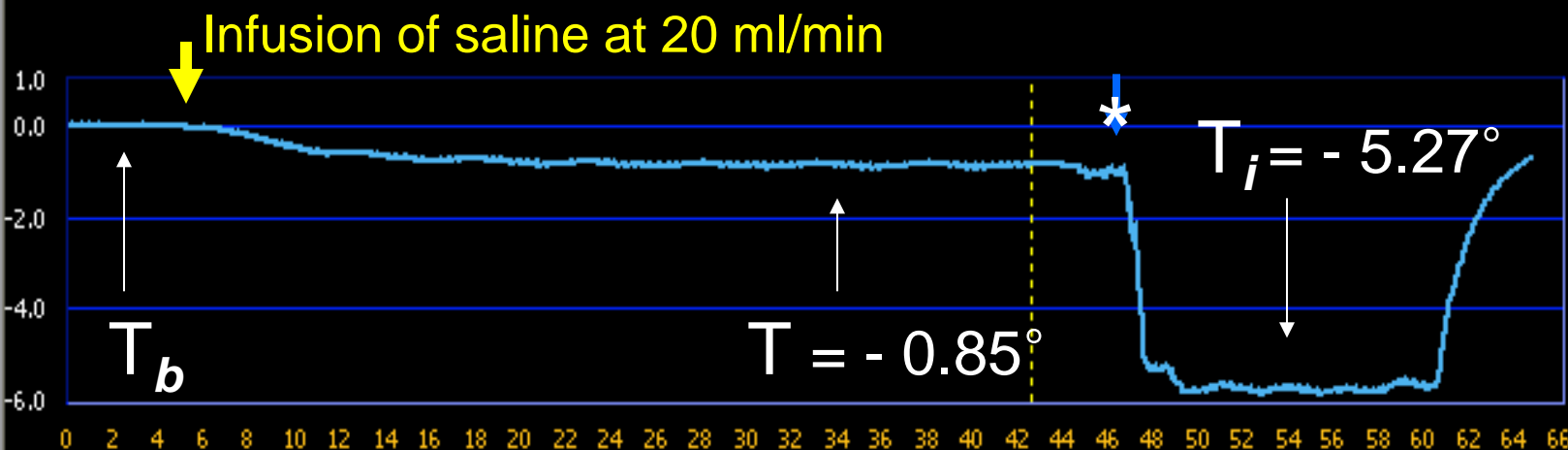
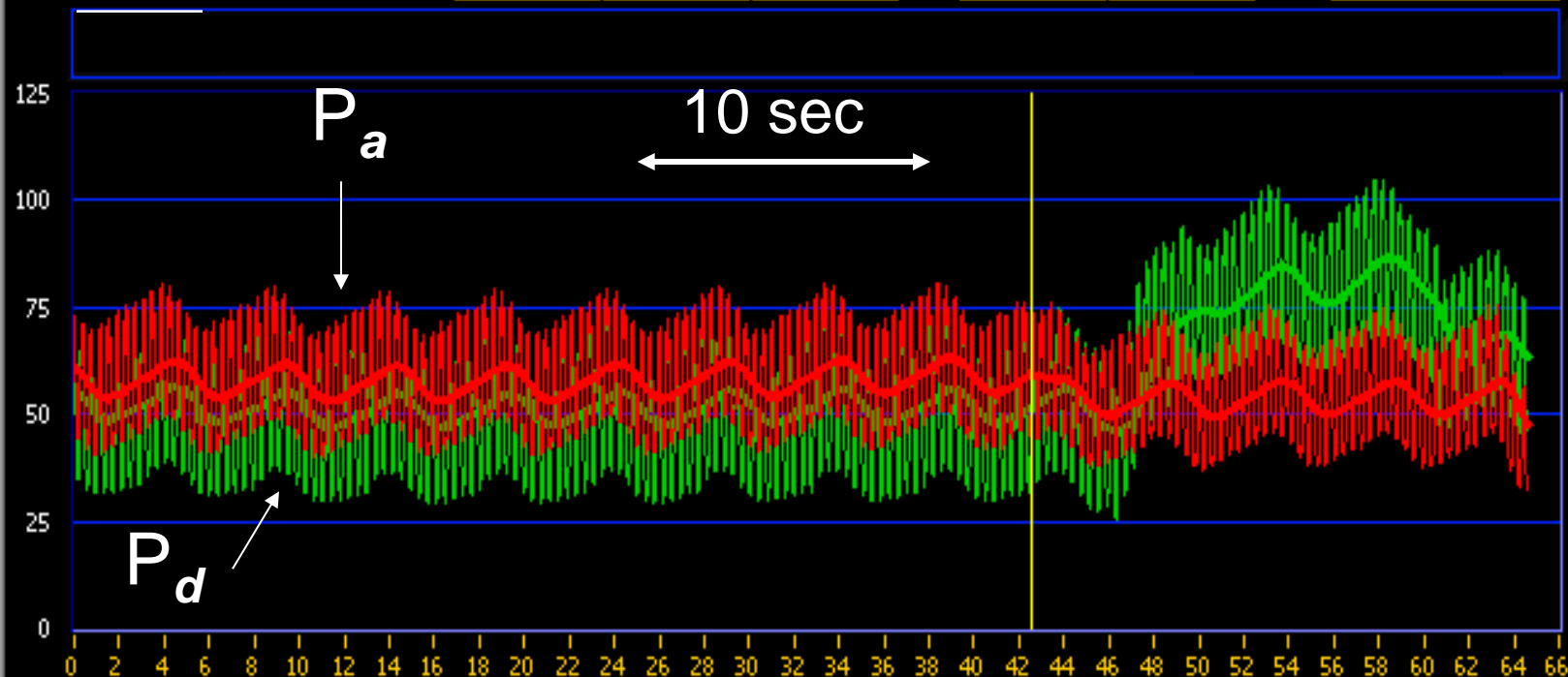
dT

42.5

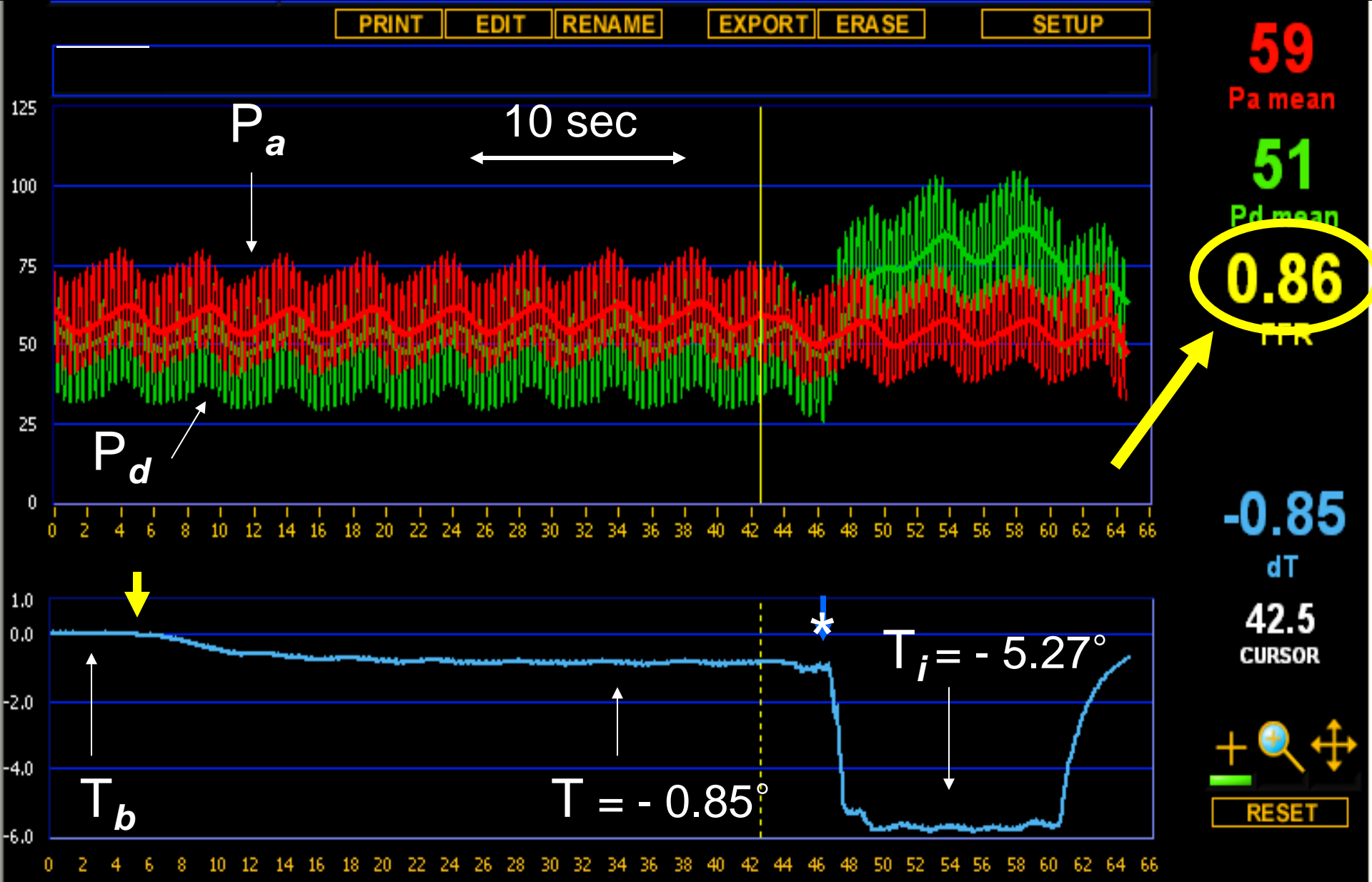
CURSOR



RESET



$$Qb = 20 \times (-5.27 / -0.85) \times 1.08 = 134 \text{ ml/min}$$



$Q_b = 134 \text{ ml/min} \longrightarrow \text{normal max flow} = 100/86 \times 134 = 156 \text{ ml/min}$

	before PTCA	at occlusion	after PTCA
FFR <sub>myo</sub>	0.50	0.18	0.97
FFR <sub>cor</sub>	0.39	-	0.96
Q <sub>c</sub> /Q <sup>N</sup>	0.11	0.18	0.01

note: the values in this matrix are independent of pressure or other hemodynamic variables. Such a matrix completely describes the distribution of flow in that part of the coronary circulation related to the respective artery.

The concept of FFR enables calculation of the separate contribution of coronary and collateral flow to myocardial blood flow

*In the present case:*

$Q_{myo}$  "normal" ( i.e. without stenosis) = 156 ml/min

$Q_{myo} = 146$  ml/min

$Q_{cor} = 134$  ml/min

$Q_{coll} = 12$  ml/min

$R_{micro} = ( P_d ) / Q_{myo}$

$R_{cor} = ( P_a - P_d ) / Q_{cor}$

$R_{collat} = ( P_{w -} - P_d ) / Q_{coll}$

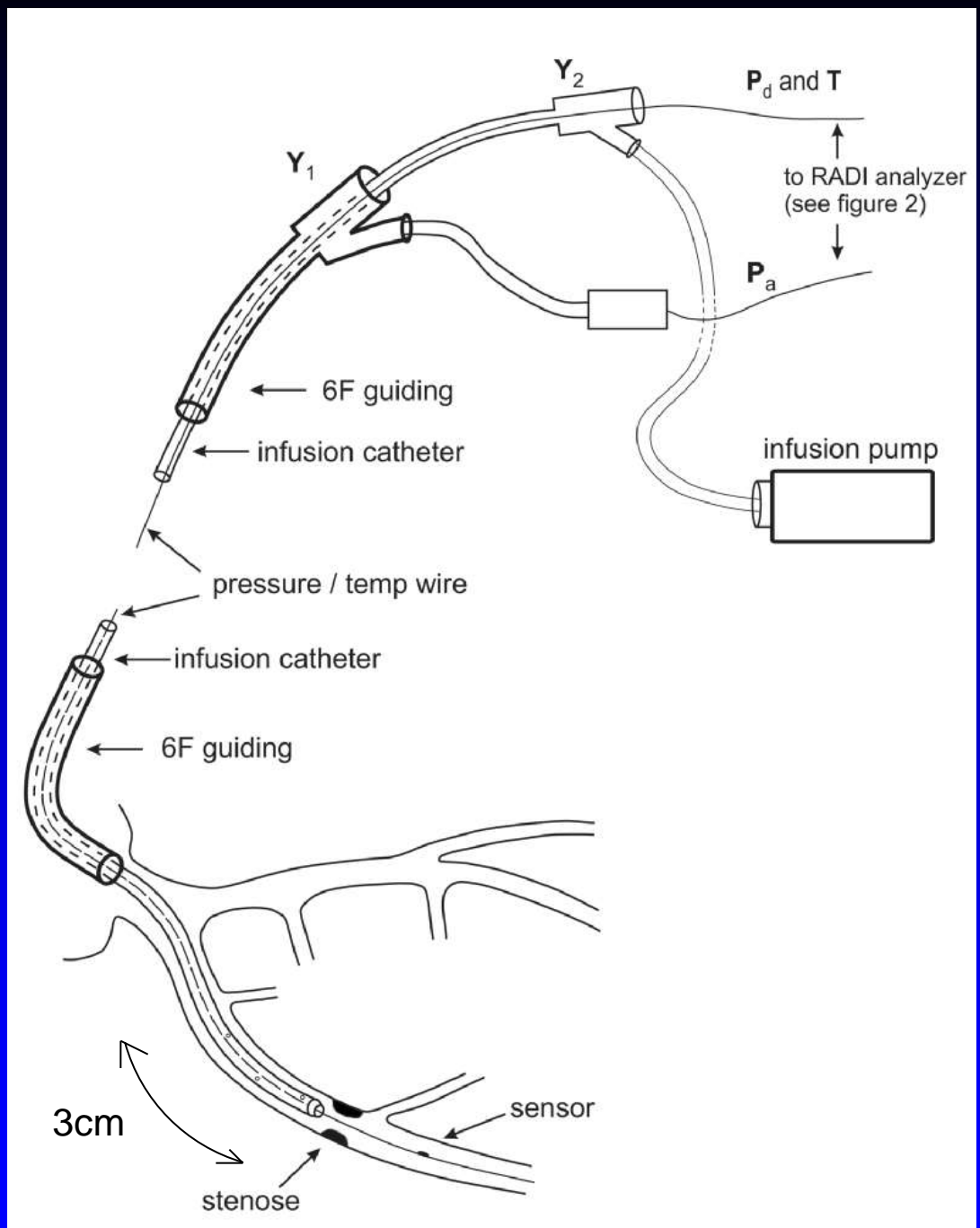
So, you calculate both the actual coronary, myocardial  
And collateral blood flow, **AND** the normal values  
of blood flow ( *in ml/min*)...

*...and because pressure is measured simultaneously,  
you also calculate **absolute microvascular resistance**  
In  $\text{dyn}/\text{sec}/\text{cm}^{-5}$  (as well as its normal value)*

## **Instrumentation up to recently:**

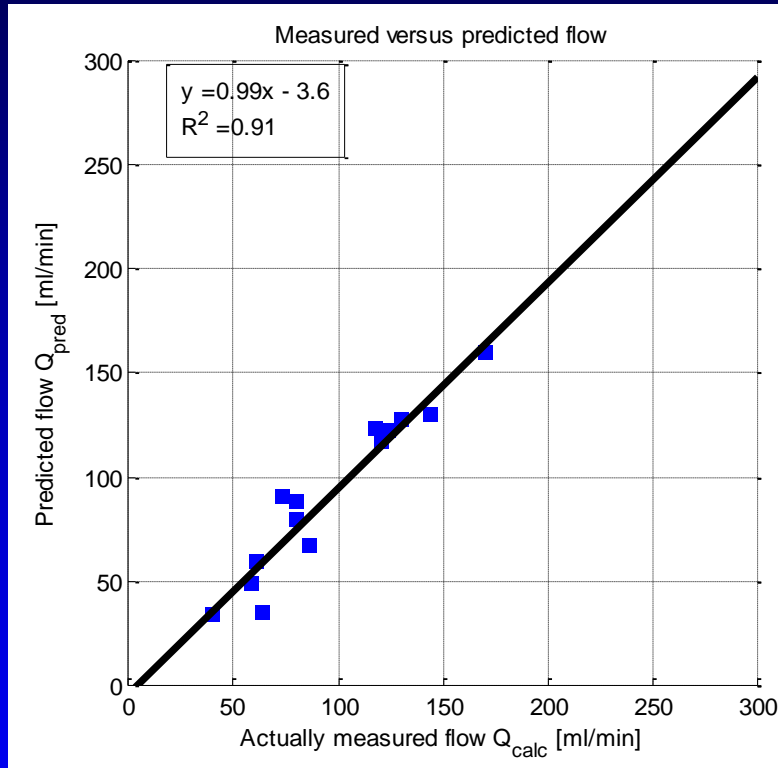
- OTW infusion catheter for infusion of saline
- 2 Y-connectors
- sustained iv infusion of adenosine for maximum hyperemia

*(20-30 minutes and some specific skill of operator)*



# Validation in humans: 36 selected patients

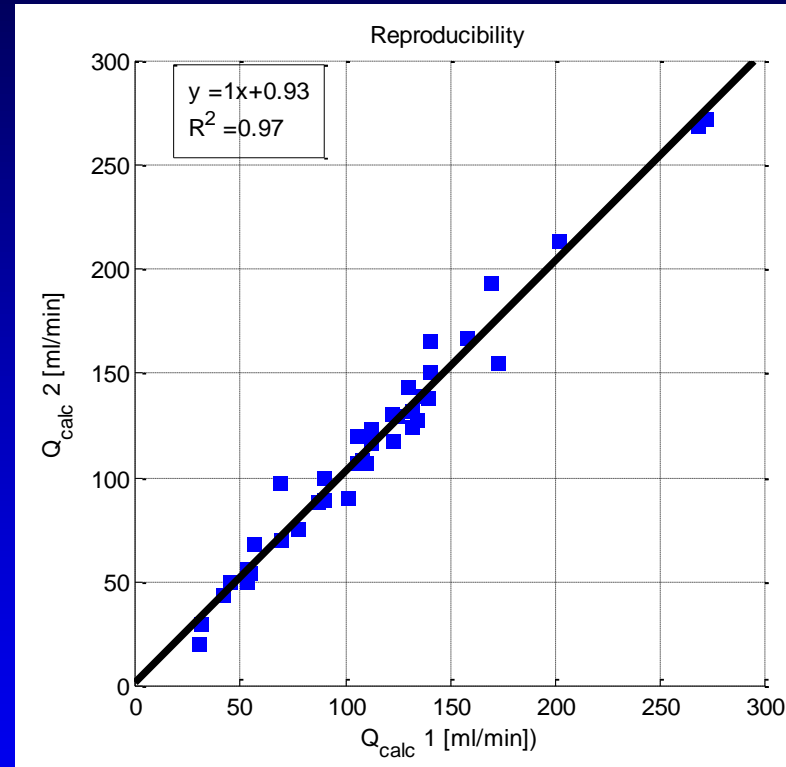
predicted flow



measured flow

N = 14

Measurement # 2



Measurement # 1

N = 22

*Aarnoudse et al, JACC 2007; 50: 2294-2304*  
*Otterspoor et al, Cather Cardiovasc Interv 2015, on-line*

# ***A NEW AVENUE TO THE CORONARY MICROCIRCULATION***

- *recently, 2 major breakthroughs have simplified the technique considerably:*
- *a new **monorail infusion catheter** and*
  - ***saline induced maximum hyperemia***



# What do you need?

1. Thermistor-wire and adequate software

→ *regular St Jude Medical Pressure wire*



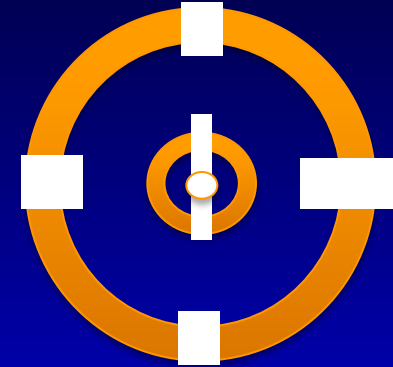
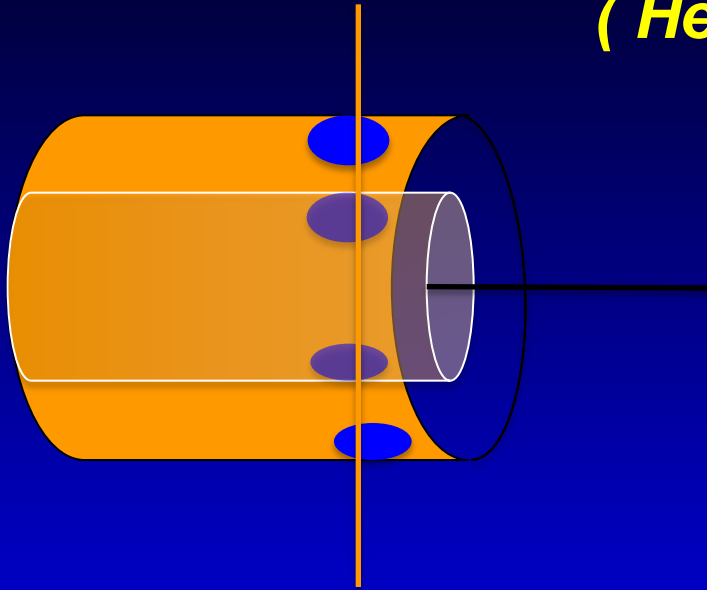
2. Adequate infusion catheter for complete mixing  
→ *Hexacath monorail catheter*



3. Sustained maximum hyperemia  
(= minimal microvascular resistance)

→ *saline infusion itself !*

# Absolute Flow Measurements by Thermodilution First Data with a Novel Catheter ( *HexaFlow*® )



- Monorail infusion catheter with double lumen (Hexacath, Paris)
- Inner lumen to measure the infusion temperature
- Outer lumen to infuse saline via side holes



# **Infusion Catheter For Thermodilution (HexaFlow®)**

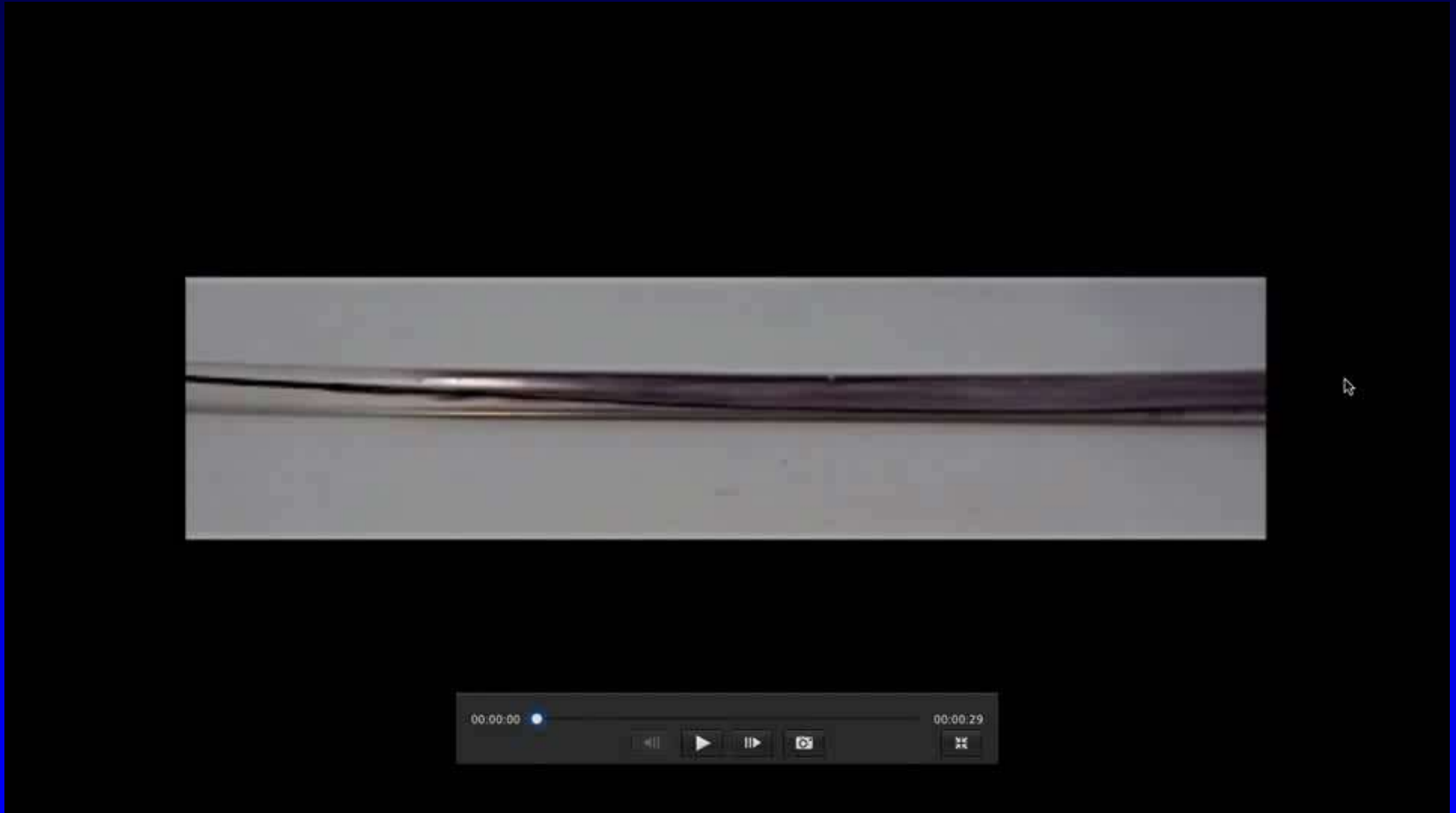
(complete mixing of blood and saline)



without guidewire



with guidewire



Sideholes are needed to obtain complete mixing of the indicator

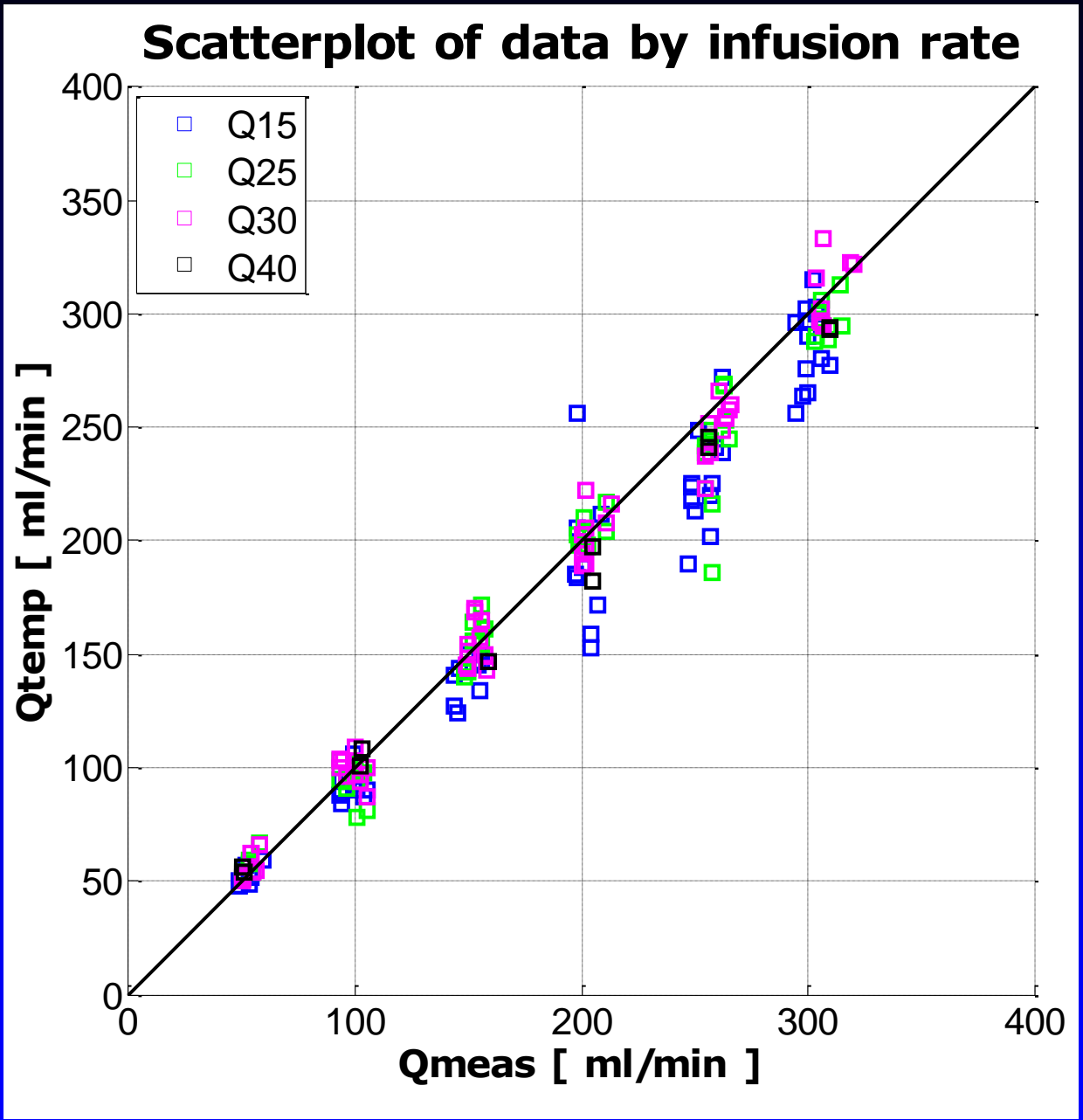


Regular infusion catheter



Special monorail infusion catheter (hexaflow®)

New monorail  
Infusion  
catheter  
(N=239)



measured flow versus calculated flow

## ***Percentual difference between true flow and calculated flow***

**Infusion rate 15 ml/min:  $-6.4 \pm 9.0$  %**

**Infusion rate 30 ml/min:  $-0.1 \pm 6.6$  %**

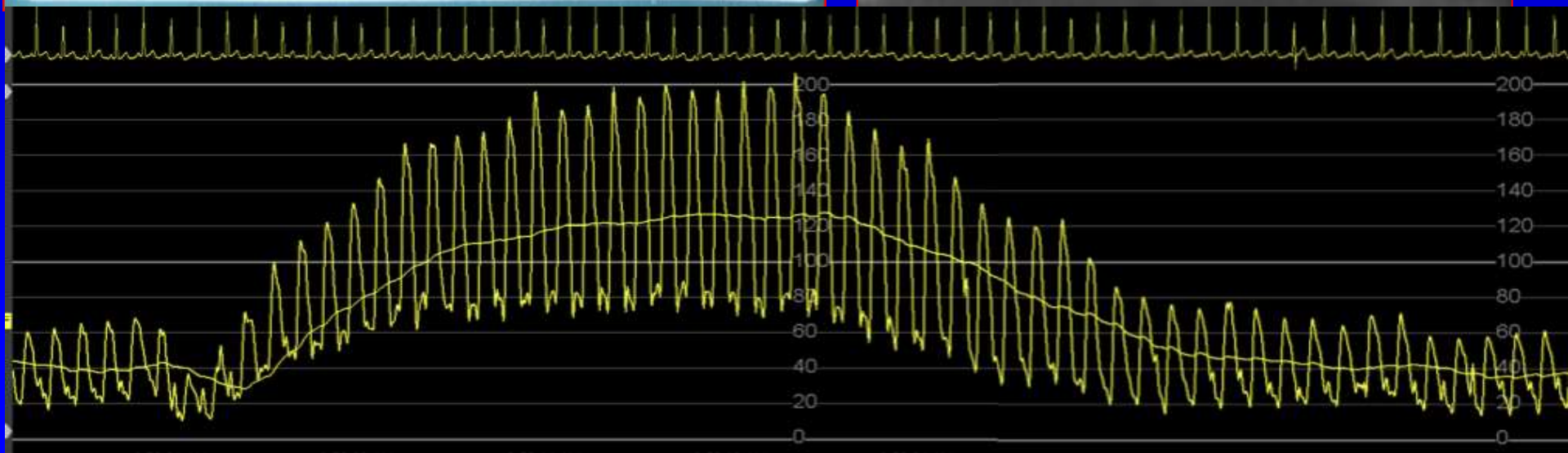
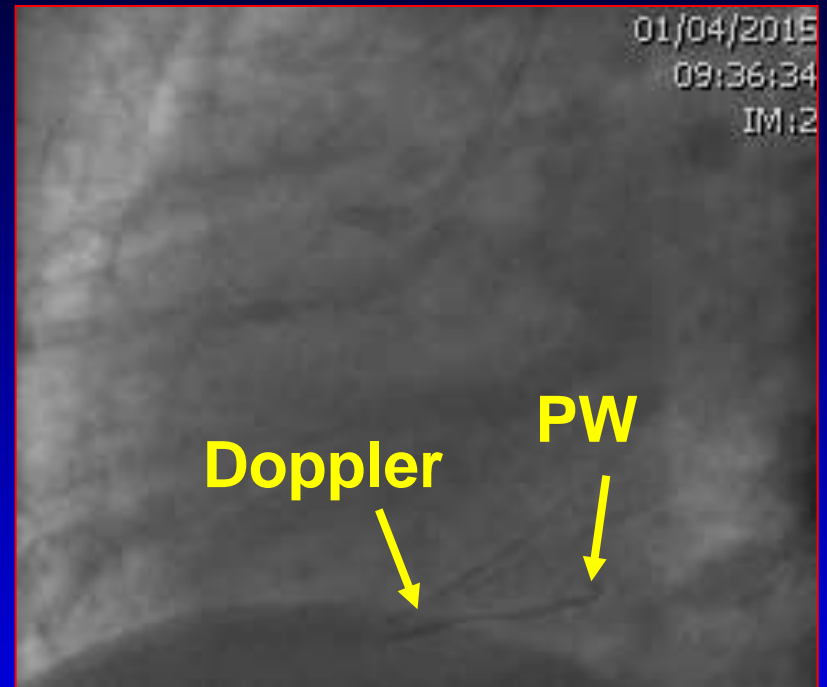
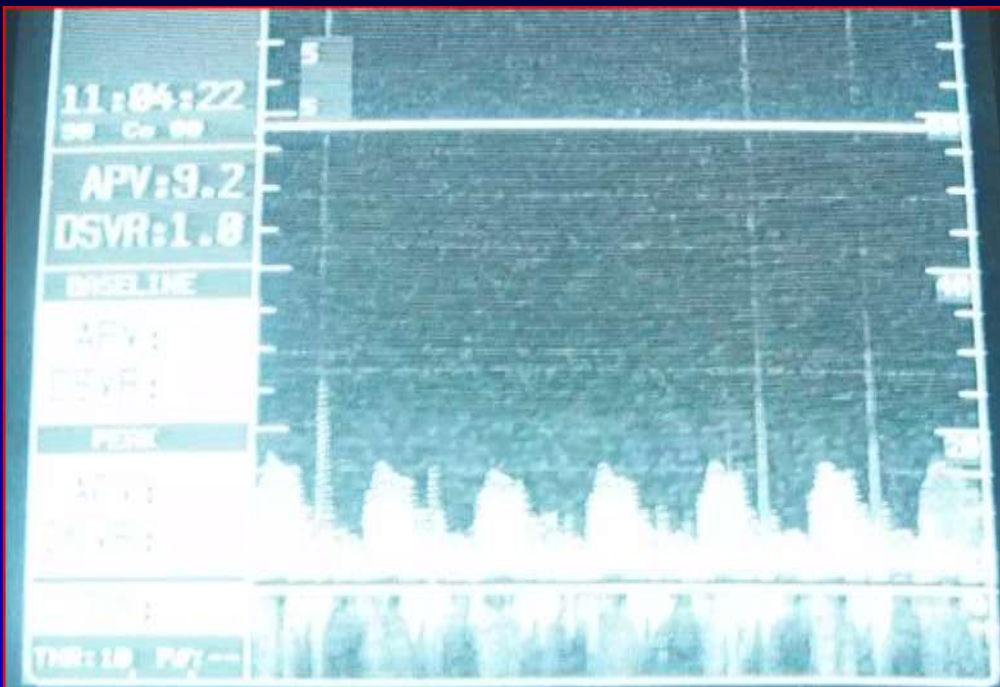
*(most common infusion rate in humans)*

***coronary saline infusion itself induce  
maximum hyperemia !***

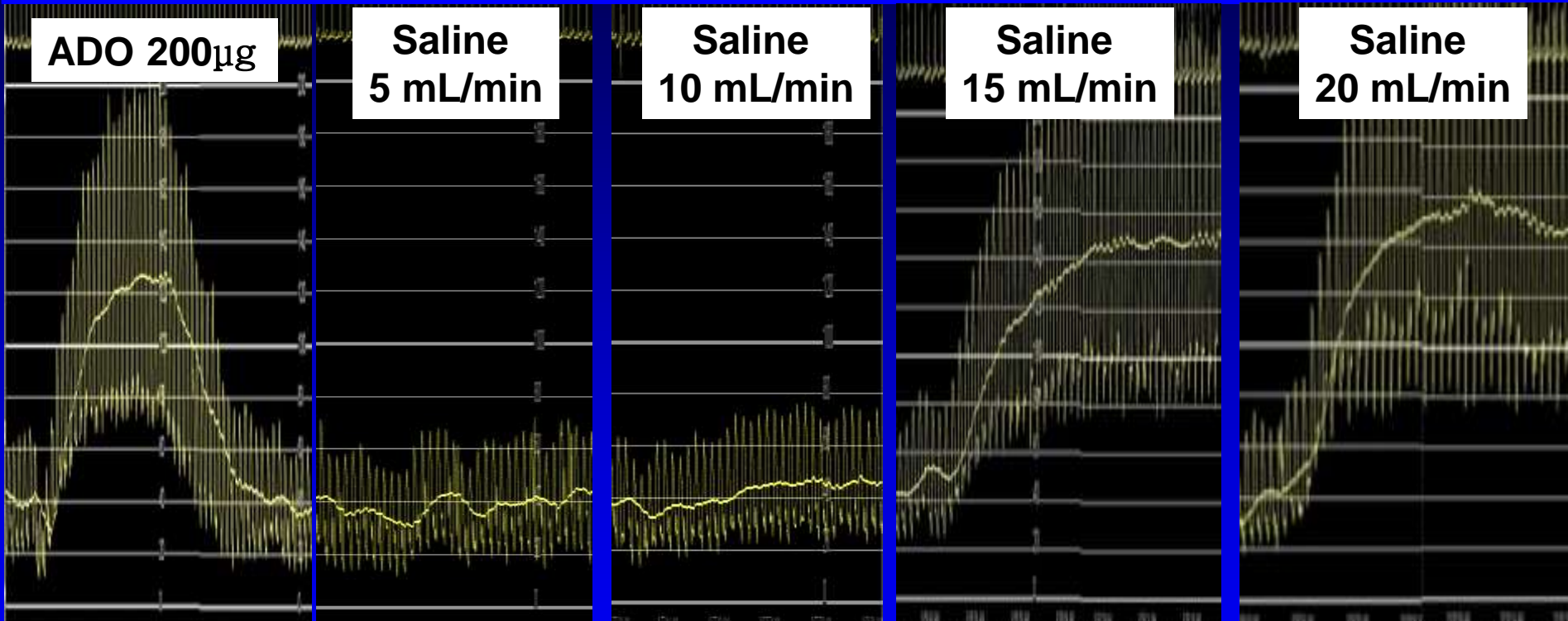
*(thereby avoiding the necessity for separate  
Infusion of adenosine)*



# Flow Velocity Measurements Adenosine IC



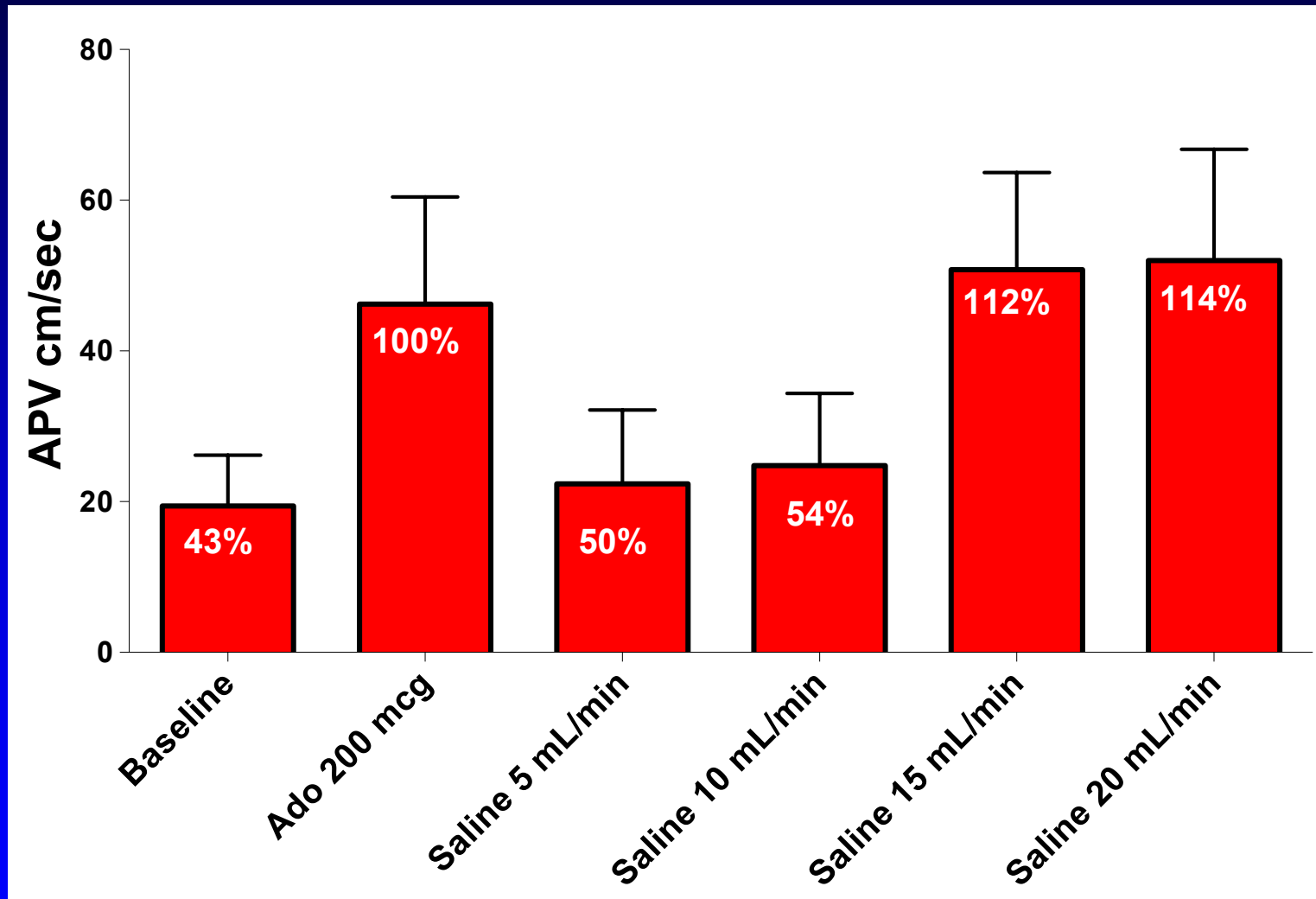
# Effect on Saline Infusion on Coronary Flow



**Continuous coronary infusion of Saline at a rate of 20 ml/min induces maximal hyperemia within 20 seconds**

*N = 18 Courtesy of Julian Adjedh and Bernard De Bruyne*

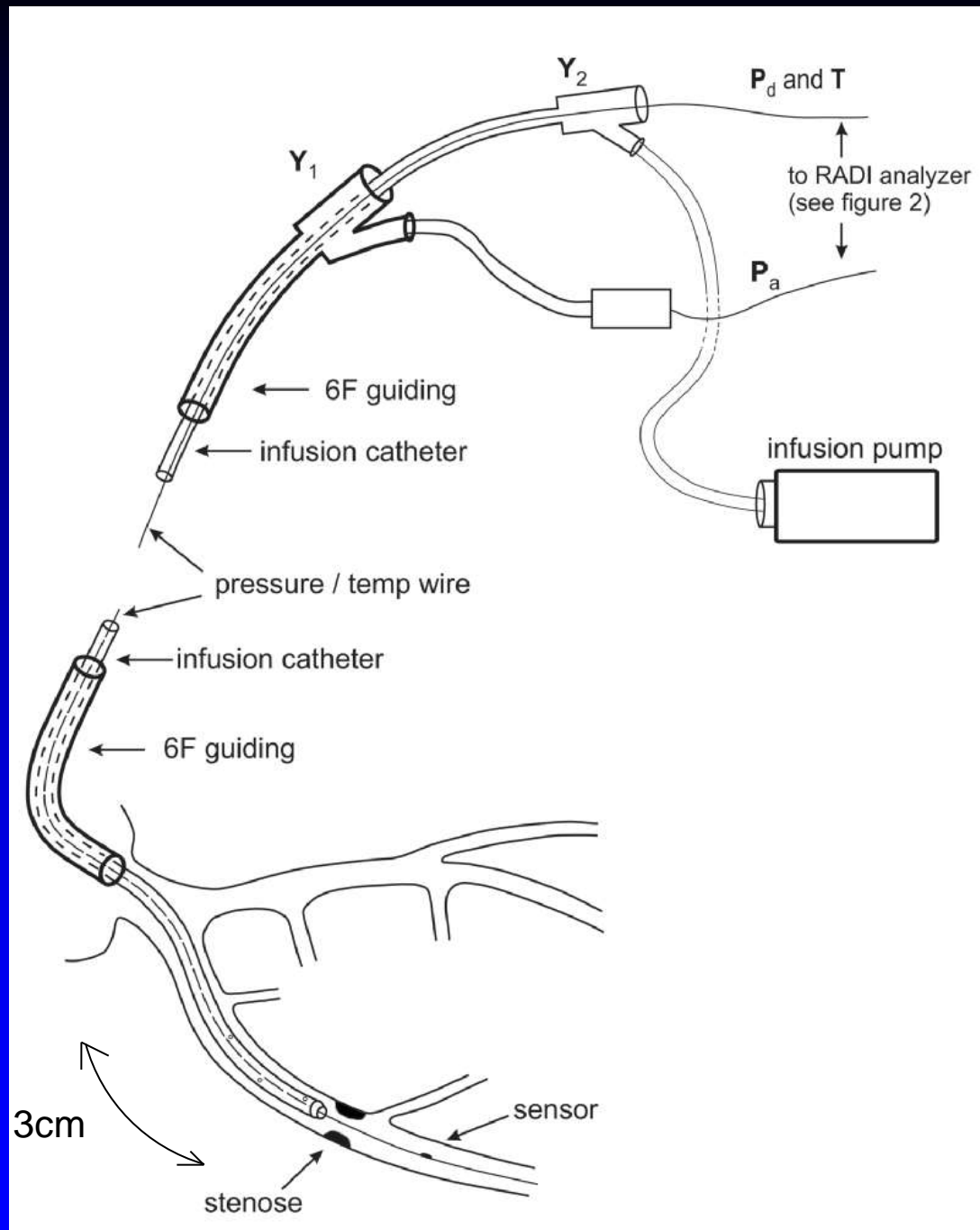
# Coronary saline infusion induces Maximal Hyperemia



*N = 18 Courtesy of Julian Adjedh and Bernard De Bruyne*

## Instrumentation today:

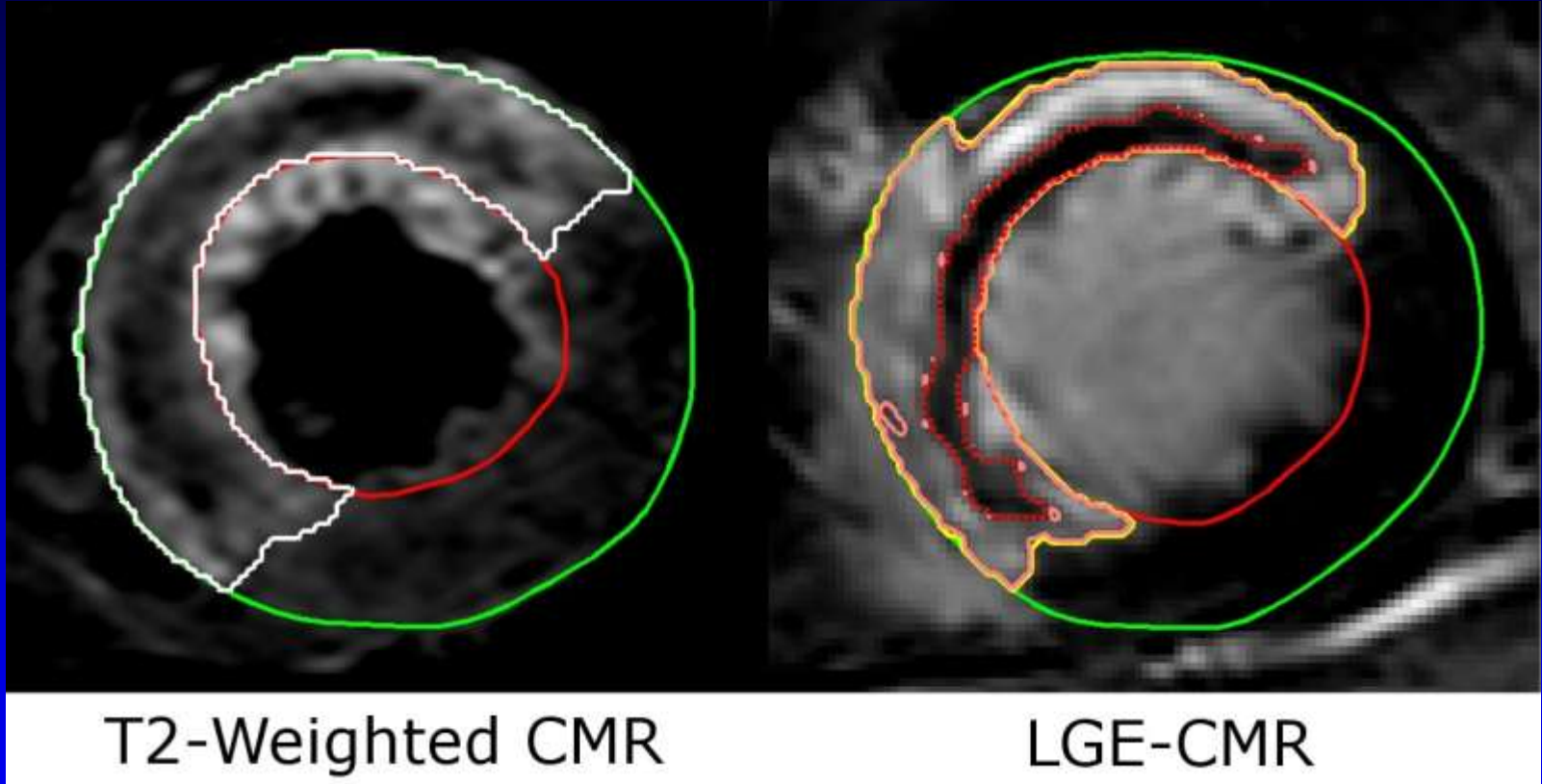
- ~~OTW infusion catheter for infusion of saline~~
- ~~2 Y-connectors~~
- monorail infusion catheter (*FlowCath*®)
- ~~sustained iv infusion of adenosine for maximum hyperemia~~
- Saline infusion itself is hyperemic



Example: Application in acute STEMI:

*Measurement of microvascular resistance  
Immediately after PPCI and after 5 days  
(N = 20)*

→ *evaluation of microcirculatory resistance  
In the first days after STEMI*



Area at risk and infarcted area by MRI



absolute flow  
in the infarct  
area  
(ml/min/g)

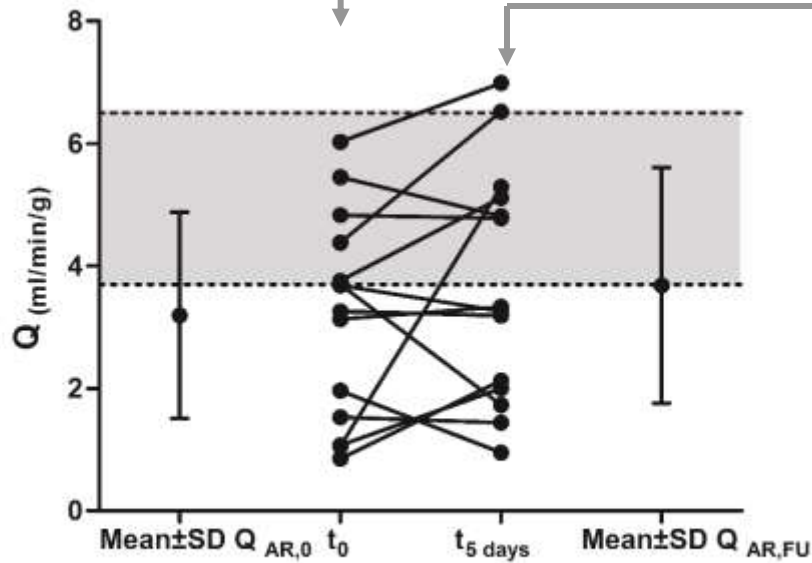


Figure 4a

absolute  
resistance  
in the infarct  
Area  
(dyn.s.cm<sup>-5</sup>)

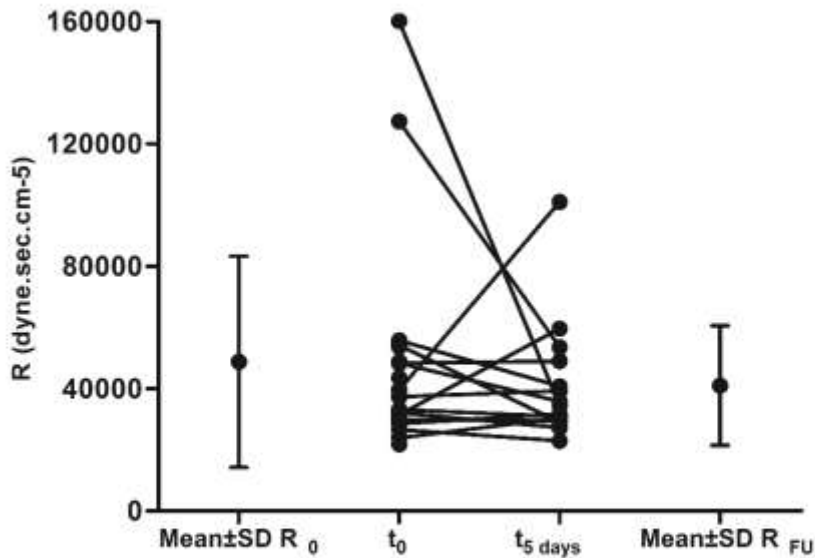
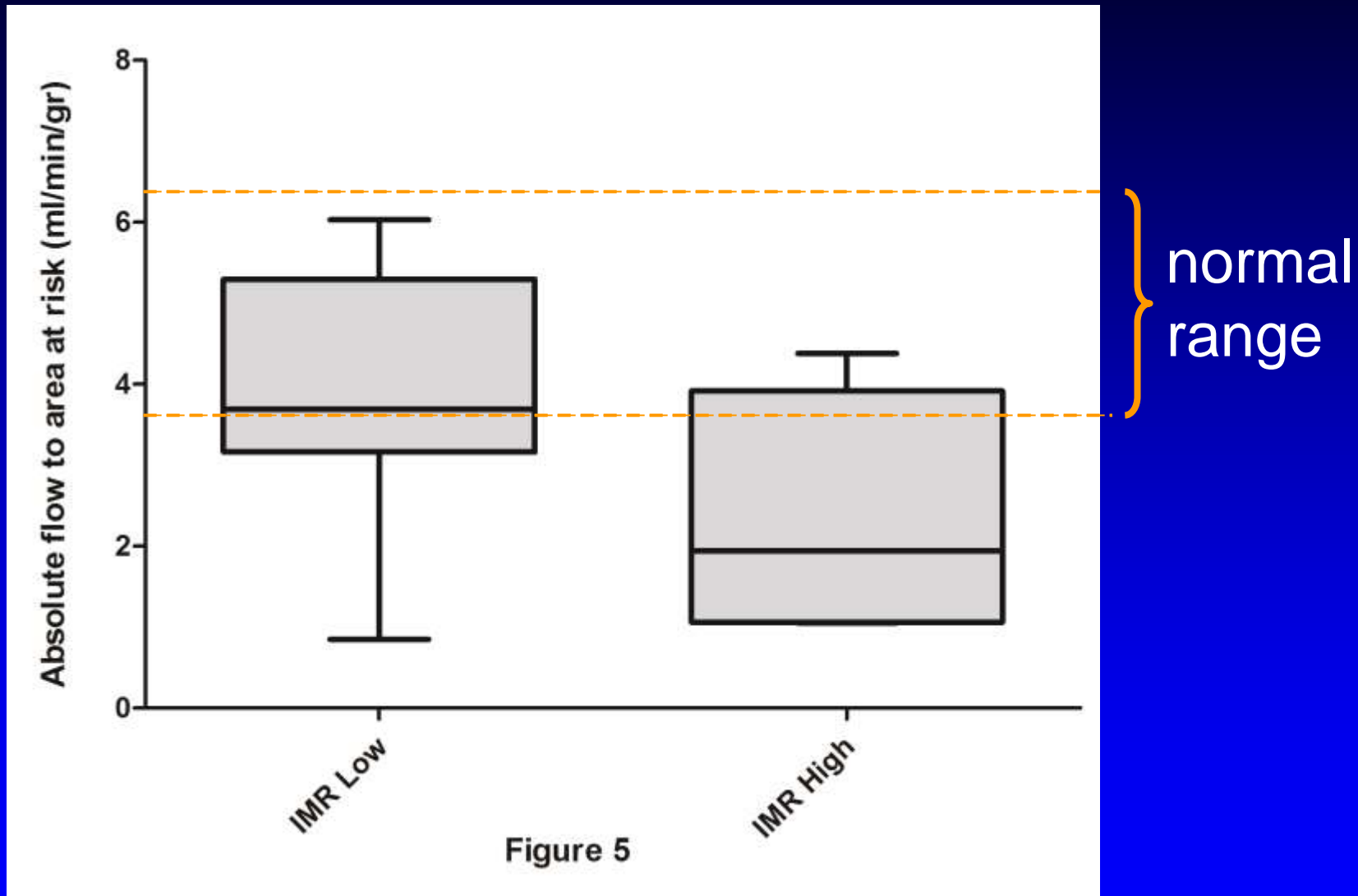


Figure 4b

hyperacute phase  
day 5

# Absolute flow in ml/min/g: correlation with IMR





# ***A NEW AVENUE TO THE CORONARY MICROCIRCULATION***

***continuous infusion of saline and thermodilution enables measurement of absolute coronary flow and microvascular resistance***

- easy to perform with standard PCI equipment
- with the new monorail infusion catheter (*FlowCath*®) it takes only 5 – 10 minutes
- the saline infusion itself guarantees maximum hyperemia; no additional stimulus necessary
- accurate and reproducible
- ***operator-independent***, you can continue the measurement for minutes