

## Reply to the letter to the editor regarding the article “Pressure wire versus microcatheter for FFR measurement: a head-to-head comparison”



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Emerging technologies to measure fractional flow reserve (FFR) are increasing. Novel technologies suggest potential advantages to promote the use of new devices or software.

As interventional cardiologists we have to focus on the results of these new technologies and on their reproducibility among different studies. As mentioned by Demir et al in their letter, FFR measured with a microcatheter (FFR<sub>MC</sub>) compared with a pressure wire (FFR<sub>w</sub>) has been evaluated by our team<sup>1</sup>, and also by Menon et al<sup>2</sup>, Wijntjens et al<sup>3</sup>, Fearon et al<sup>4</sup> and Ali et al<sup>5</sup>. These studies found a difference between FFR<sub>w</sub> and FFR<sub>MC</sub> of 0.83±0.07 vs. 0.80±0.10 in 77 patients, 0.81±0.11 vs. 0.79±0.12 in 58 patients, 0.86±0.06 vs. 0.82±0.07 in 28 patients, 0.81 vs. 0.83 in 169 patients and 0.83±0.09 vs. 0.78±0.11 in 74 patients, respectively. All previous studies, including 406 patients, agreed that FFR<sub>MC</sub> overestimates FFR compared to FFR<sub>w</sub>. We believe that the reproducibility of the results of five studies with a similar design, in which each lesion was measured twice with both devices, is reliable enough to support our conclusion. We strictly

included all consecutive patients with FFR measurement indications and reference diameter above 2.5 mm<sup>1</sup>. Therefore, without selection bias we had lower crossability with FFR<sub>MC</sub> compared to pressure wire in most calcified and tortuous vessels. We disagree with the message from Demir et al which is trying to get to the following simplest conclusion: if the mean difference is 0.03 therefore it should be negligible because misclassification could mostly concern patients in the “diagnostic grey zone”. It is important to note two points from these five studies. First, standard deviations are between 0.06 and 0.12 and, second, the underlying mechanism is probably due to the larger size of the FFR<sub>MC</sub> device compared to FFR<sub>w</sub>. Similar to our study<sup>1</sup>, Fearon et al<sup>4</sup> and Ali et al<sup>5</sup> suggest that reference diameter is an independent predictor of FFR<sub>MC</sub> overestimation. Therefore, we believe that increasing the FFR<sub>MC</sub> cut-off value or limiting the problem of inaccuracy of FFR<sub>MC</sub> to patients within the “grey zone” for clinical decision making could not be adopted from currently available data. However, we believe that FFR<sub>MC</sub> in a large vessel could be similar and

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accurate (difference close to zero with  $FFR_w$ ), while in a smaller vessel it might have a higher degree of inaccuracy (difference of 0.10 and more). It is therefore unfortunate, from our point of view, that the main potential advantage of FFRMC, which is easy and accurate investigation of lesions that a pressure wire cannot cross, has become its weakest point.

### Conflict of interest statement

The author has no conflicts of interest to declare.

### References

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