

Letter to the Editor:

Semi-Quantification Limitations:

FMTVDM[®] Demonstrates Quantified Tumor Response to Treatment
With Both Regional Blood Flow and Metabolic Changes

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True quantification¹⁻⁶ is the actual measurement of material within a tested region. In molecular imaging, the ability to accurately measure isotope accumulation is dependent upon the demonstration that the measuring device, be it SPECT or PET camera, is accurately calibrated, is measuring the correct isotope and can be counted and reproduced serially.

The publication by Humbert⁷ *et al*, is important because it raises the question of whether PET cameras can detect actual changes in disease following treatment. In order to accurately measure changes in regional blood flow and metabolism it is necessary to rely on a truly quantified¹⁻⁶ method and not upon a method that produces only a calculated value. The Humbert⁷ *et al* method makes two flawed presumptions. First, it applies the wrong pharmacologic kinetic model that the isotope absent from the arterial bed traveled only to the site of interest. Second, it uses a matrix setting, which has been demonstrated to produce a loss of signal data, which produces a significant error rate²⁻⁶. This method produces a *semi*-quantified value derived from “first-pass extraction”, not an accurate measurement of the amount of isotope within the tissue of interest.

We have demonstrated that using a true quantification method provides an actual measurement of change in regional blood flow and metabolism, which is useful in assessment of treatment response.

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