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Letters to the Editor

at any current frequency [4]. Linearity supports validity of monitoring with only pre- and postdialysis measurements.

> ANTONIO PICCOLI and MARTA CODOGNOTTO Padova, Italy

Correspondence to Prof. Antonio Piccoli, Dept. Scienze Mediche e Chirurgiche, Policlinico IV piano, Via Giustiniani, 2, I-35128 Padova, Italy. E-mail: apiccoli@unipd.it

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## **Reply from the Authors**

We appreciate Piccoli's interest in our article [1], and thank him for further analysis of the data we presented. We fully agree that a cyclical variation in BIA variables was apparent in hemodialysis patients during both dialysis and interdialysis periods according to the concept that dialysis causes a reduction of total body water, and especially (or only) of extracellular body water, and that total body water progressively increases between dialysis sessions due to water retention.

Indeed, change in resistance, reactance, and phase angle during a hemodialysis session can be affected by other factors, such as an increase in hematocrit, variations in electrolytes concentration, a rapid shift from intracellular to extracellular water, and others [2]. As a matter of fact, acute changes in body water induced by dialysis are not predicted well by data derived from BIA [3], and when BIA was applied to estimate the fluid loss during hemodialysis, overestimation usually occurred. As a consequence, changes in BIA variables during either dialysis or interdialysis periods are expected to reflect variation in total body water and its extracellular/intracellular distribution, but also to be affected by other factors. Further studies seem necessary to us to understand to which extent these data can be compared with those obtained in a healthy population or in predialysis patients. In addition, our results indicate that the measurement timing with respect to dialysis session is a crucial aspect in assessing BIA in such patients. Furthermore, since phase angle is considered an independent marker of survival [4-6], the

point of time for performing the analysis may influence the prognostic significance of such a parameter.

> VINCENZO BELLIZZI, LUCA SCALFI, and BIAGIO R. DI IORIO Avellino, Italy

Correspondence to Dr. Vincenzo Bellizzi, Unità Operativa Complessa di Nefrologia e Dialisi, Ospedale "A. Landolfi," Via Melito, 83029 Solofra (AV), Italy. E-mail: vincenzo.bellizzi@tin.it

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## End-stage renal disease increases plasma transcobalamin and neutralizes influence of TCN 776C>G polymorphism

To the Editor: A lack of influence of TCN 776C>G has been recently reported on transcobalamin and homocysteine plasma levels in two series of patients with kidney transplant and end-stage renal disease (ESRD), contrary to what was previously observed in a healthy population [1–3]. In the ESRD series, which included 66 hemodialysis patients, influences of *MTHFR 677TT* (P=0.024) and TCN 776CC (P=0.036) on homocysteine disappeared in a multivariate model that included a combination of 677TTx776CC genotypes, a confounder of each polymorphism [2]. The lack of influence of *MTHFR* agreed with some of the previous data [4]. We performed a similar study in 55 nonsupplemented hemodialysis patients. We confirmed the lack of influence of either TCN