

<p>What does BIA measure? And what does it estimate?</p>	<ul style="list-style-type: none"> • BIA measures the electrical response of the body to an electric current applied by single- or multifrequency devices • It estimates TBW, FFM, and FM or other parameters using SF-BIA and MF-BIA predictive equations
<p>How to choose the right device?</p>	<ul style="list-style-type: none"> • Choose a device that also provides raw bioimpedance measurements as outputs (e.g., resistance, reactance, and impedance) in addition to the body composition parameters
<p>How do you select an equation to estimate body composition?</p>	<ul style="list-style-type: none"> • Ideally, equations should match: <ul style="list-style-type: none"> - the characteristics of the population being evaluated (e.g., age, sex, sexual maturation, ethnicity, health status, obesity degree) - the device being used (e.g., brand, model/version, frequency, whole-body or segmental, supine/standing)
<p>If available equations are not population-specific or device-specific, how to proceed?</p>	<ul style="list-style-type: none"> • Perform a cross-validation study • Choose a reference standard measuring body composition at the same level (e.g., multicompartiment, DXA, dilution method) • Use agreement analysis to evaluate the validity of the selected equations. A guide is provided in Earthman (4) • If agreement analysis is not satisfactory, develop a new equation and test its external validity using an external sample or the bootstrapping method
<p>What alternatives to body composition assessment exist when using BIA?</p>	<ul style="list-style-type: none"> • Raw BIA measurements can be used, such as resistance, reactance, and impedance • These measurements can be adjusted by height or used to compute BIA parameters, including phase angle, impedance ratio, and BIVA
<p>What protocol to follow?</p>	<ul style="list-style-type: none"> • We advise following the guidance provided by Lyons-Reid et al (11) and Brantlov et al. (12) until a standard protocol for the pediatric population is established; or, if available, the study protocol by the device's manufacturer • Use the same protocol for all subjects and during all follow-up visits • When deviations from the recommendations are necessary, record modifications and report them in future publications