

Blood Impedance Characterization from Pulsatile Measurements

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Introduction

Bioimpedance Measurements have been widely applied because of

- low cost
- easy application
- non-invasiveness
- on-line monitoring
- etc

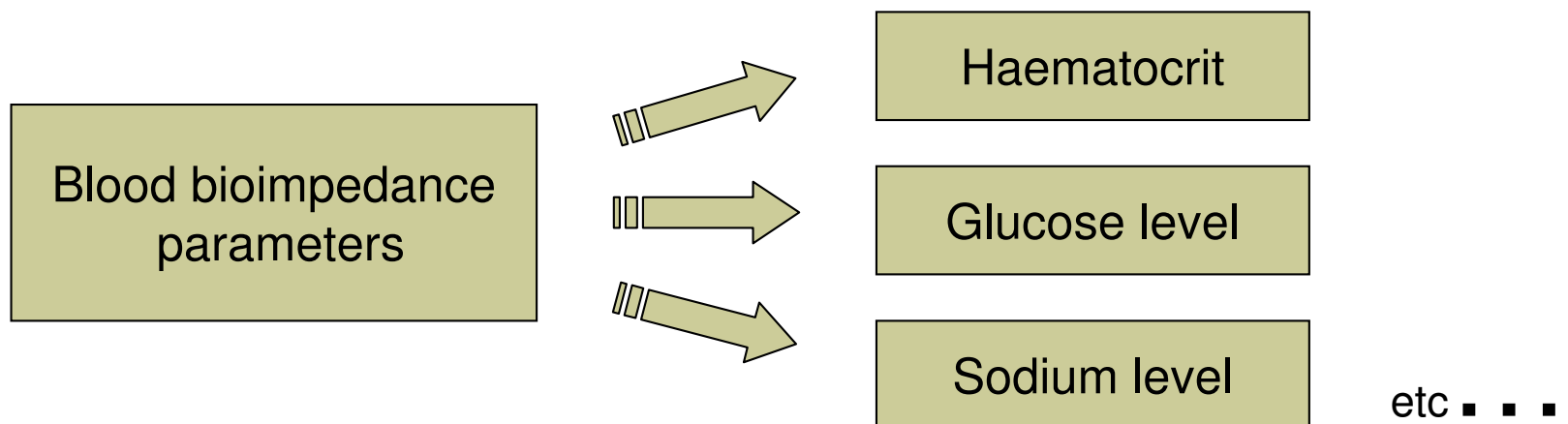
Introduction

- Bioelectrical impedance analysis (BIA)
- Bioelectrical impedance spectroscopy (BIS)
 - Body fluid measurement
e.g. ECF, ICF, TBW.
 - Tissue volume change
e.g. cardiac stroke volume
 - Tissue characterization
e.g. ischemic organ identification.

Introduction

Why do we characterize blood by bioimpedance method ?

-- to find correspondence between clinical indices and blood bioimpedance parameters.

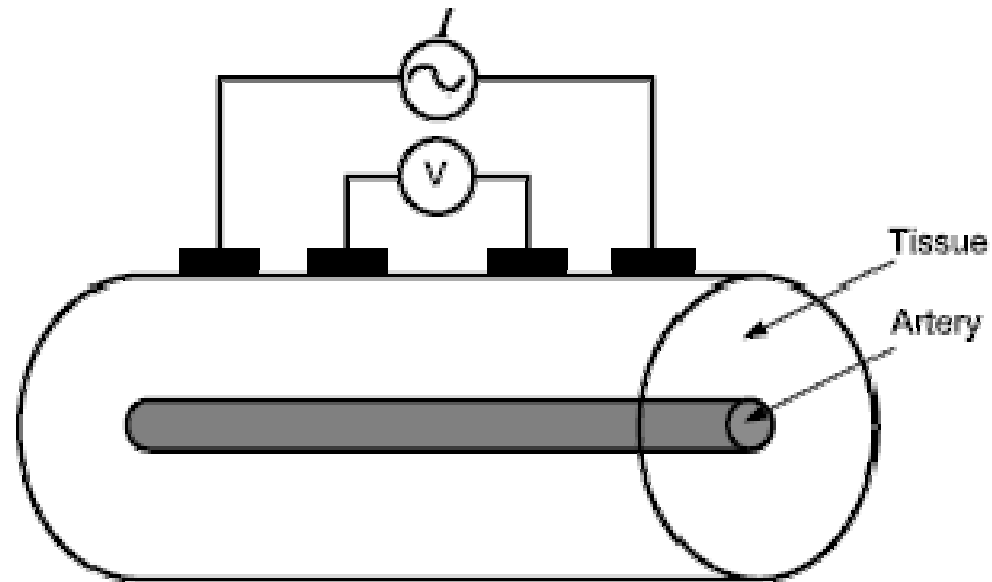


Introduction

Difficulties to measure blood impedance ***in vivo***

- *Physiological noises*
- *Body disturbances*
- *Heterogeneity*

Modeling



A tetrapolar BIS sensor is applied on a finger segment (cylindrical model)

Modeling

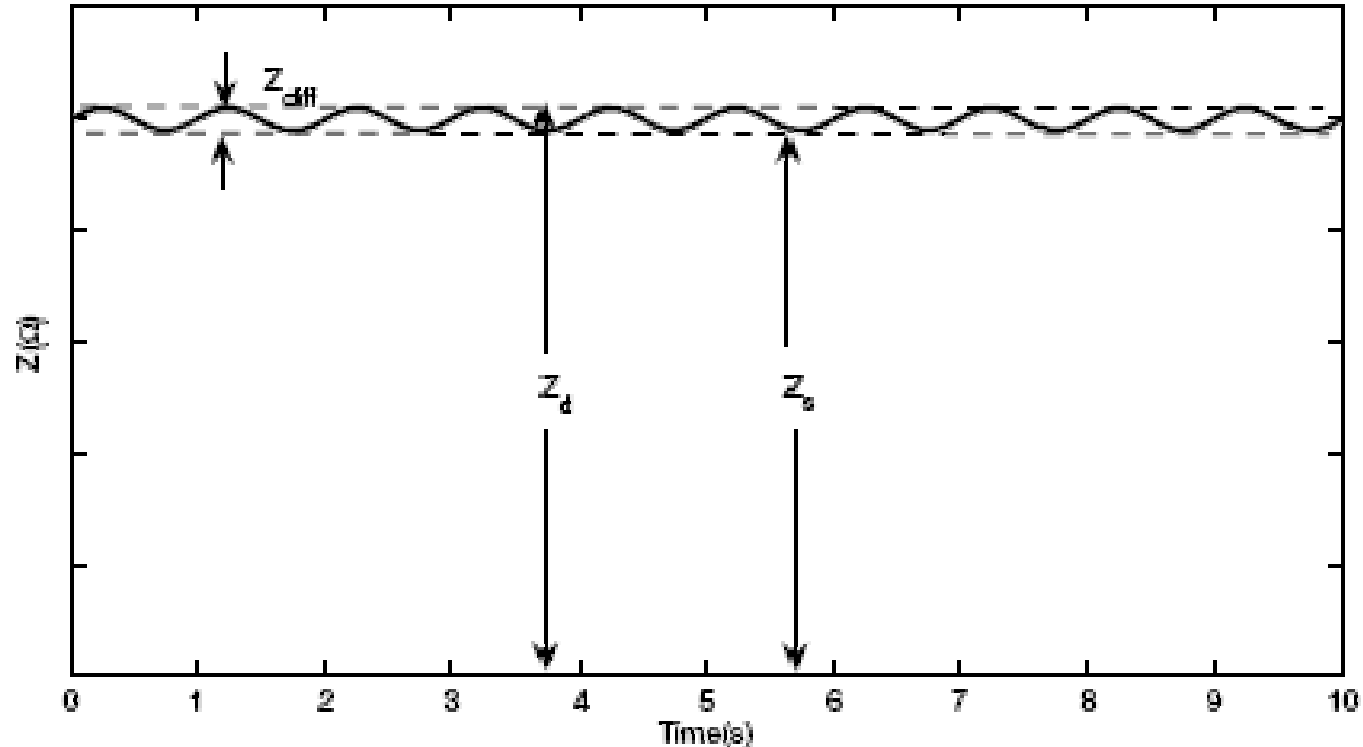


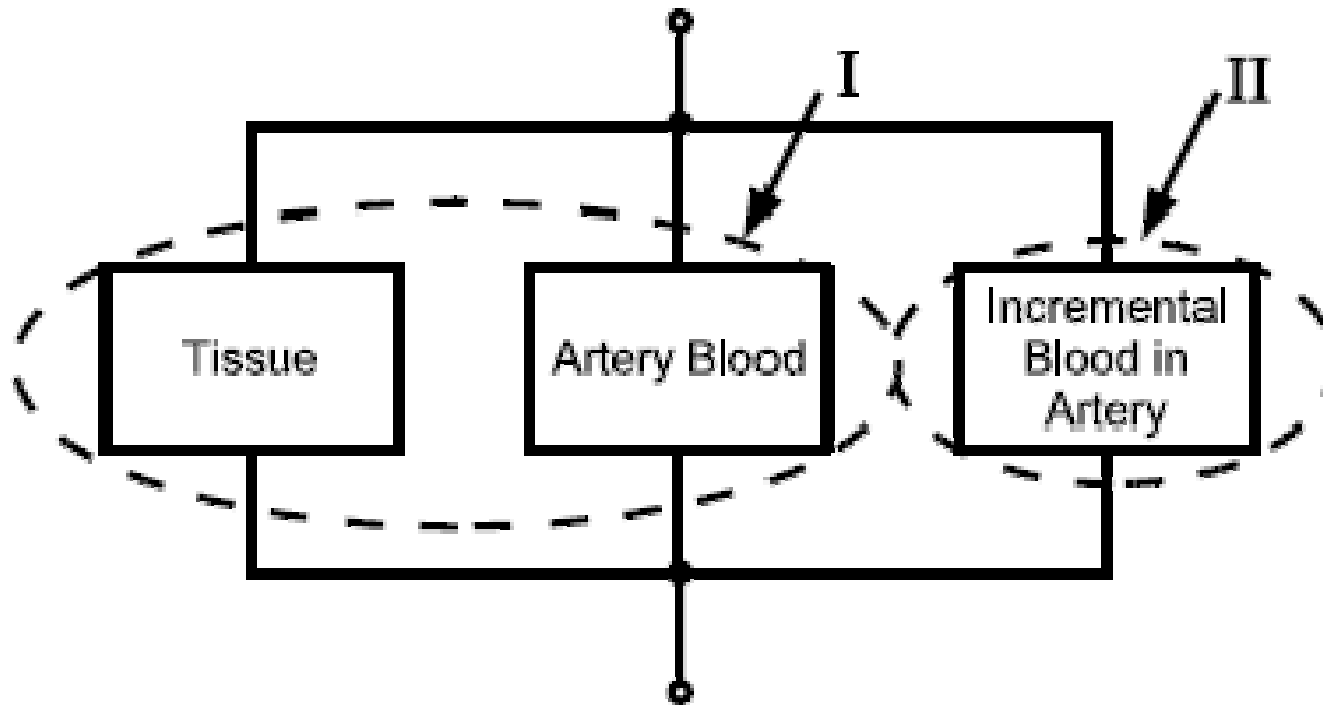
Illustration of the pulsatile impedance wave.

Z_d : impedance corresponding to the heart diastole;

Z_s : impedance corresponding to the heart systole;

Z_{diff} : The difference between Z_d and Z_s

Modeling



The bioimpedance model of pulsatile wave is composed of three parallel impedances.

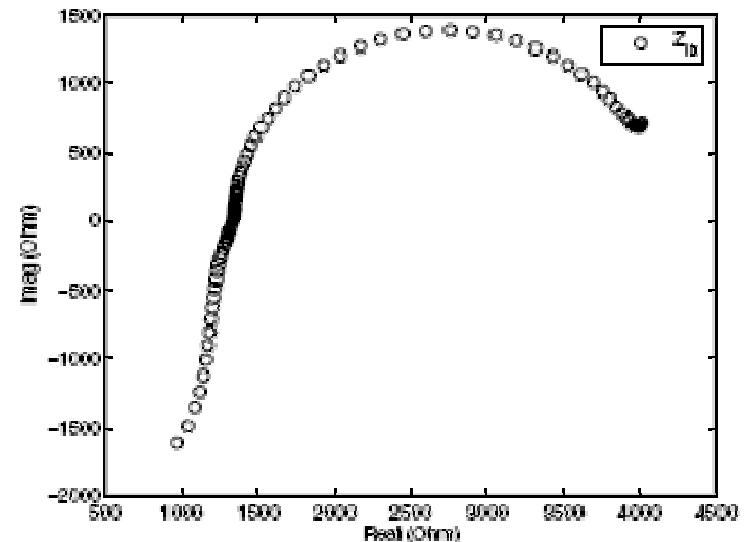
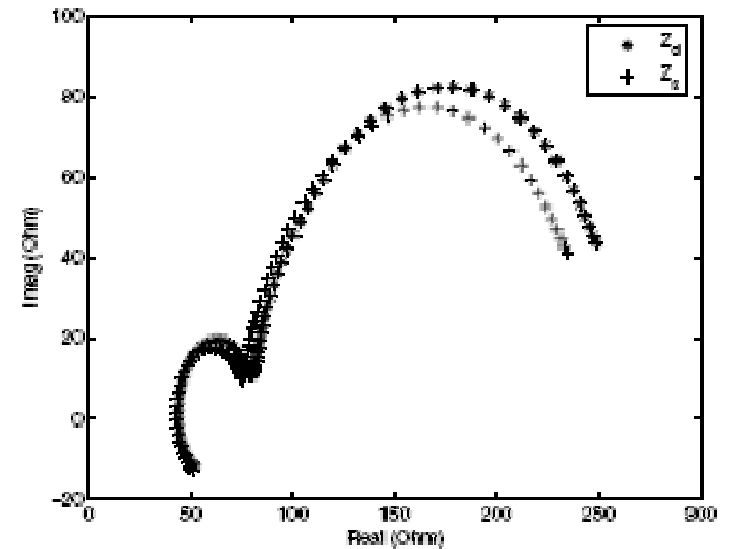
During diastole, $Z_d = Z_I$
systole, $Z_s = Z_I \parallel Z_{II}$

A impedance spectrum from blood

$$Z_d(f) // Z_{ib}(f) = Z_s(f)$$

⇓

$$Z_{ib}(f) = \frac{Z_d(f)Z_s(f)}{Z_d(f) - Z_s(f)}$$



Cole-Cole Model

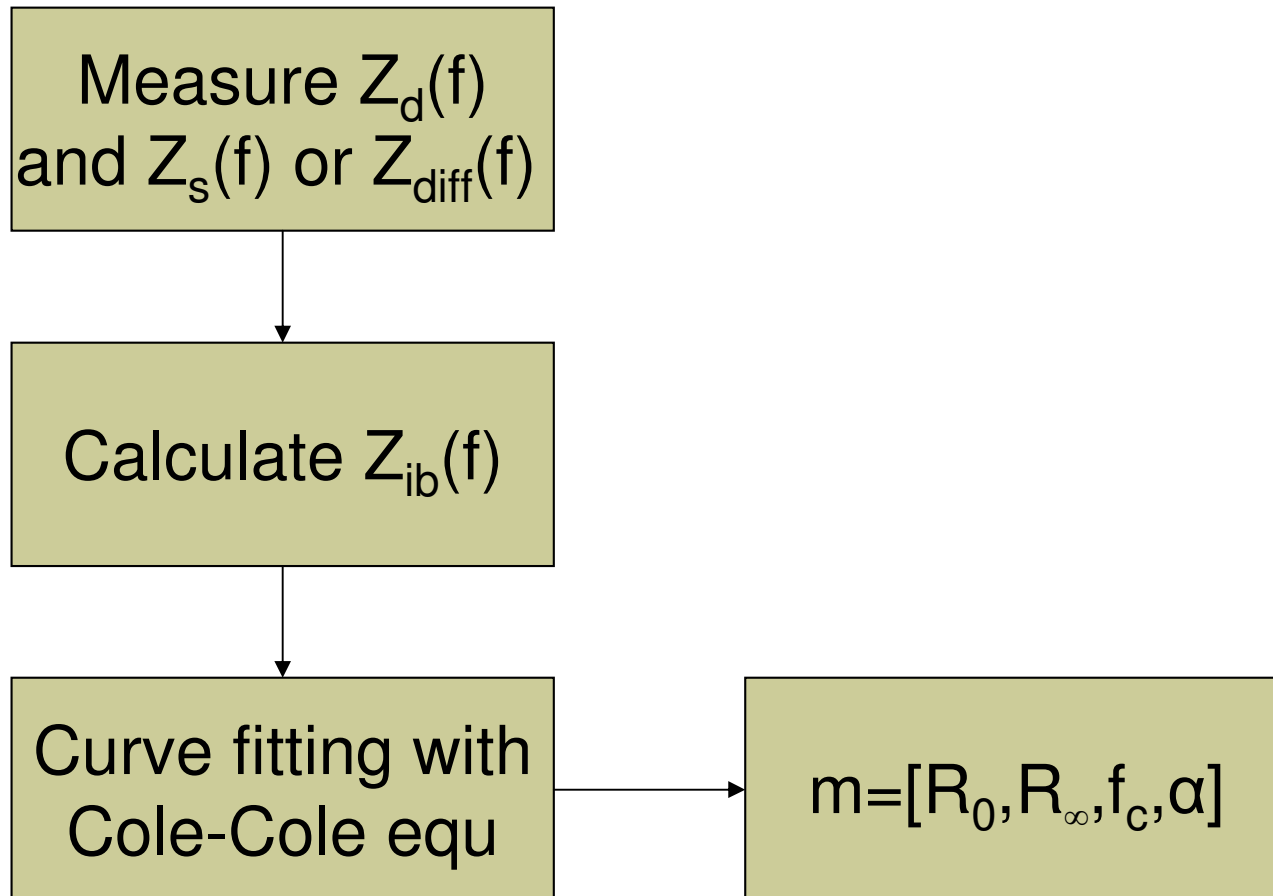
Cole-Cole equation

$$Z(f) = R_{\infty} + \frac{R_0 - R_{\infty}}{1 + j(f / f_c)^{1-\alpha}}$$

Cole-Cole parameters

$$m = [R_0 \ R_{\infty} \ f_c \ \alpha]$$

Method Description



Simulations

A Pspice model is designed to simulate

- physiology structure of finger
- skin-electrode contact interface

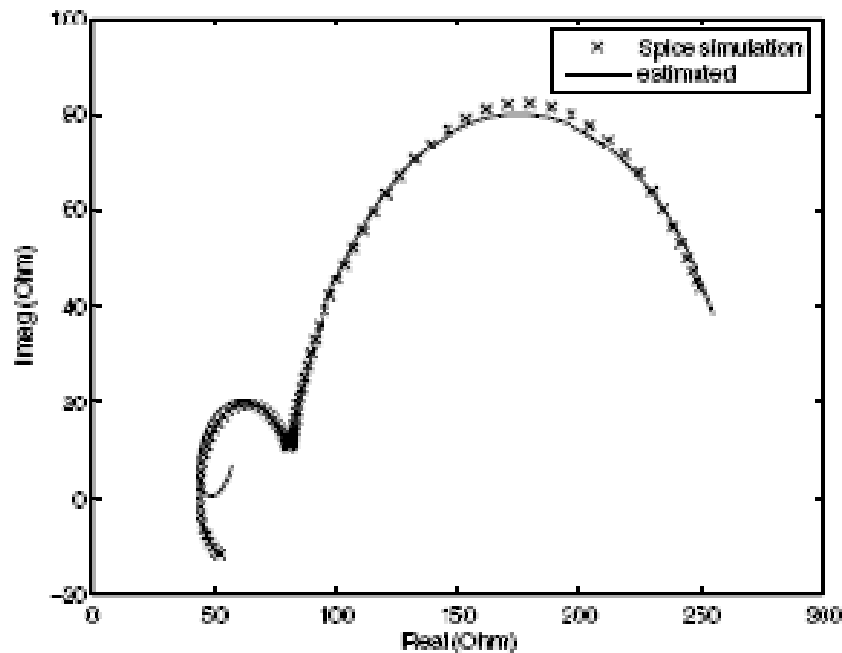
Frequency range: 5k~100MHz

Sweeping frequencies: 88 points, logarithmly distributed

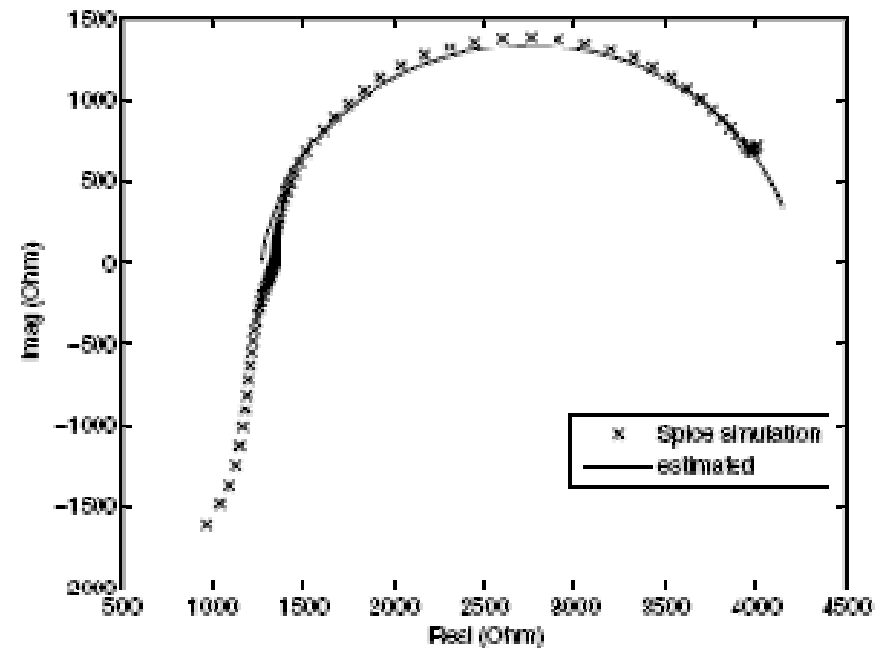
Artery volume change: 10%

Nonlinear curve fitting: LM (Levenberg-Marquardt) Method

Simulations



$Z_d(f)$ is fitted in an extended Cole-Cole model



the pulse curve $Z_{ib}(f)$ is fitted in a 1st order Cole-Cole model.

Simulations

NMSE-Normalized Mean Square Error: (1000 trials)

| SNR(dB) | 0 | -5 | -10 | -20 |
|--------------------|--------|--------|--------|-------|
| Err R_0 (%) | 0.1911 | 0.3240 | 0.6594 | 1.867 |
| Err R_∞ (%) | 0.5772 | 0.9659 | 1.511 | 5.791 |
| Err f_c (%) | 0.884 | 1.614 | 2.453 | 8.302 |
| Err α (%) | 5.588 | 11.76 | 17.70 | 62.37 |

*Measurement error and corresponding Cole-Cole parameter error, as a function of SNR.
(Sampling rate is 1 kHz)*

Conclusions

■ Advantages:

- Parameters estimated are of a homogeneous medium--blood;
- Inductive effect is alleviated;
- No need to use multiple Cole-Cole model

■ Disadvantage:

- Due to poor SNR in real measurements, it is hard to estimate pulsatile impedance wave amplitude.