

# OXYGEN EXTRACTION RATIO (OER): A POSSIBLE PARAMETER TO MEASURE CARDIOVASCULAR (CV) STRESS IN HAEMODIALYSIS (HD).

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## Introduction

- Hemodialysis treatment may cause peripheral tissue hypoxia not routinely evaluated
- Intra-HD hypoxia (Arterial Oxygen saturation, SaO<sub>2</sub>, <90%), automatically detected with dedicated sensors, has been reported to associate with increased morbidity and mortality.
- The Oxygen extraction ratio (OER, n.v 20%-25%), obtained with the measurement of SaO<sub>2</sub> and Central Venous Oxygen Saturation (SvcO<sub>2</sub>), is routinely employed in ICU to quantify tissue hypoxic stress. Theoretically more sensitive than SaO<sub>2</sub>, OER increments may indicate parenchymal hypoperfusion and inadequate adaptation to hypoxia (Figure 1).

## Aims

We aimed to evaluate if OER:

- increased significantly during HD sessions;
- was different between long (HD<sub>Long</sub>) and short (HD<sub>Short</sub>) HD intervals;
- was differently affected by Isolated UF (iUF) or Isolated Diffusion (iD).

## Methods

We enrolled 20 clinically stable patients on HD since >6 months, with Central Venous Catheter.

We contemporarily sampled arterial SO<sub>2</sub> (by capillary oxymeter) and ScvO<sub>2</sub> (by blood gas analysis) to calculate the OER basally, 15', 30', 60', 120' and end of HD<sub>Long</sub>.

In 10 of them (group A), OER was re-measured in the following two HD<sub>Short</sub> sessions.

In the other 10 (group B), OER was re-measured during the first hour of the first and second HD<sub>Short</sub> performed by applying iUF or iD respectively.

During each HD, UF rate was kept at <10 ml/kg/h and symptoms were recorded.

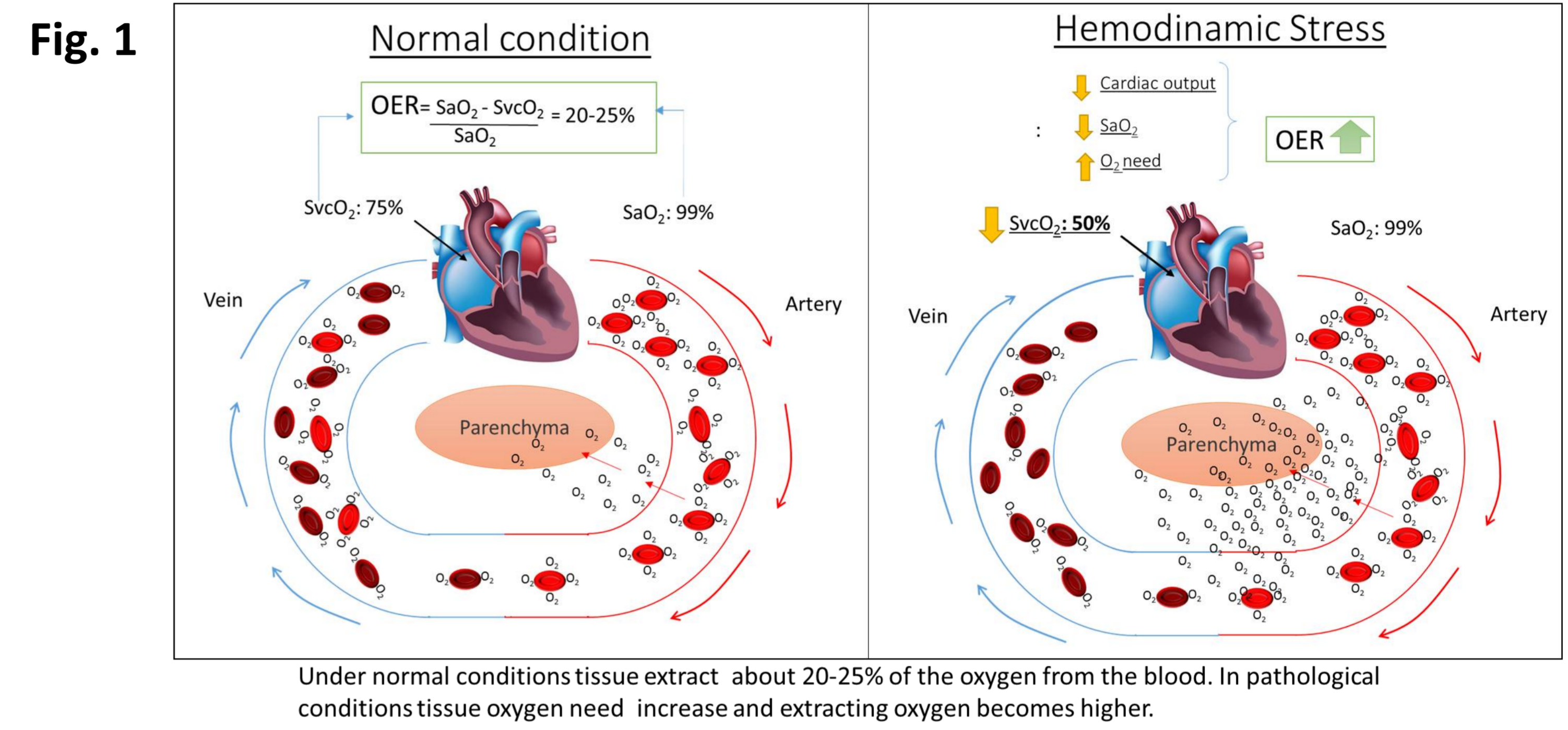
## Results

Clinical and biochemical features are reported in Tab.1.

In the HD<sub>Long</sub> session, OER increased within 30' (post hoc test p<.05) and then progressively up to the end of HD, by 38%. Mean basal OER of HD<sub>Long</sub> (34,4 ± 7), HD<sub>Short1</sub> (33,8 ± 7) and HD<sub>Short2</sub> (34,2 ± 7) were not different (Tab.2).

In the two HD<sub>Short</sub> sessions, OER changes overlapped (no statistical difference) with those in HD<sub>Long</sub> (Fig.2).

During the first hour of the HD<sub>Short1</sub> session with iUF, the increment of OER from basal was not significant, at variance with the significant increase recorded during the HD<sub>Short2</sub> with iD (Tab. 2). Importantly, all HD sessions were asymptomatic with no change (ANOVA = n.s.) in blood pressure (either systolic or diastolic) nor of heart rate. During sessions, no significant change was evident for capillary SO<sub>2</sub> (average 98±1 %), while ScvO<sub>2</sub> progressively decreased, thus allowing to refer the recorded O<sub>2</sub> consumption to peripheral extraction and not to other possible causes of systemic SO<sub>2</sub> reduction.



Characteristic	Value
N.	20
Men/Females; N (%)	8 (40)/12 (60)
Age, yr	75±13
HD Vintage, yr	3,7±3,3
BMI, Kg/m <sup>2</sup>	25,1±5,6
Comorbidity; N (%)	12 (60)
ACS, N (%)	2 (10)
Peripheral ischemia, N (%)	3 (15)
Cerebral ischemia, N (%)	0
Diabetes, N (%)	7 (35)
Hb, g/dl	10,3±1,2
Calcium, mg/dl	8,5±0,4
Phosphate, mg/dl	5,2±1,2
PTH, pg/ml	338±154
ALP, UI/L	84±26

HD, type	Paz. (n)	Basale	15'	30'	60'	120'	End HD	Anova
HD <sub>Long</sub>	20	34,4 ± 7	39,0 ± 7	40,0 ± 8*	40,7 ± 6#	42,5 ± 8°	46,9 ± 6^	.0001
HD <sub>Short1</sub>	10	33,8 ± 7	40,2 ± 8*	41,3 ± 6#	42,4 ± 5#	42,0 ± 7#	45,9 ± 8°	.004
HD <sub>Short2</sub>	10	34,2 ± 7	40,6 ± 9	42,2 ± 7*	42,0 ± 8*	42,9 ± 10*	45,9 ± 7°	.003
HD <sub>Short</sub> iUF	10	33,0 ± 6	35,9 ± 7	36,6 ± 6	37,5 ± 7			n.s
HD <sub>Short</sub> iD	20	32,4 ± 6	37,6 ± 6	38,8 ± 5*	38,4 ± 7			.04

Tab. 2, Mean values of OER. iUF = Isolated Ultrafiltration; iD: isolated diffusion; Bonferroni Post-hoc test vs Basale: \*p<.05; # p<.005; ° p<.001; ^ p<.0001

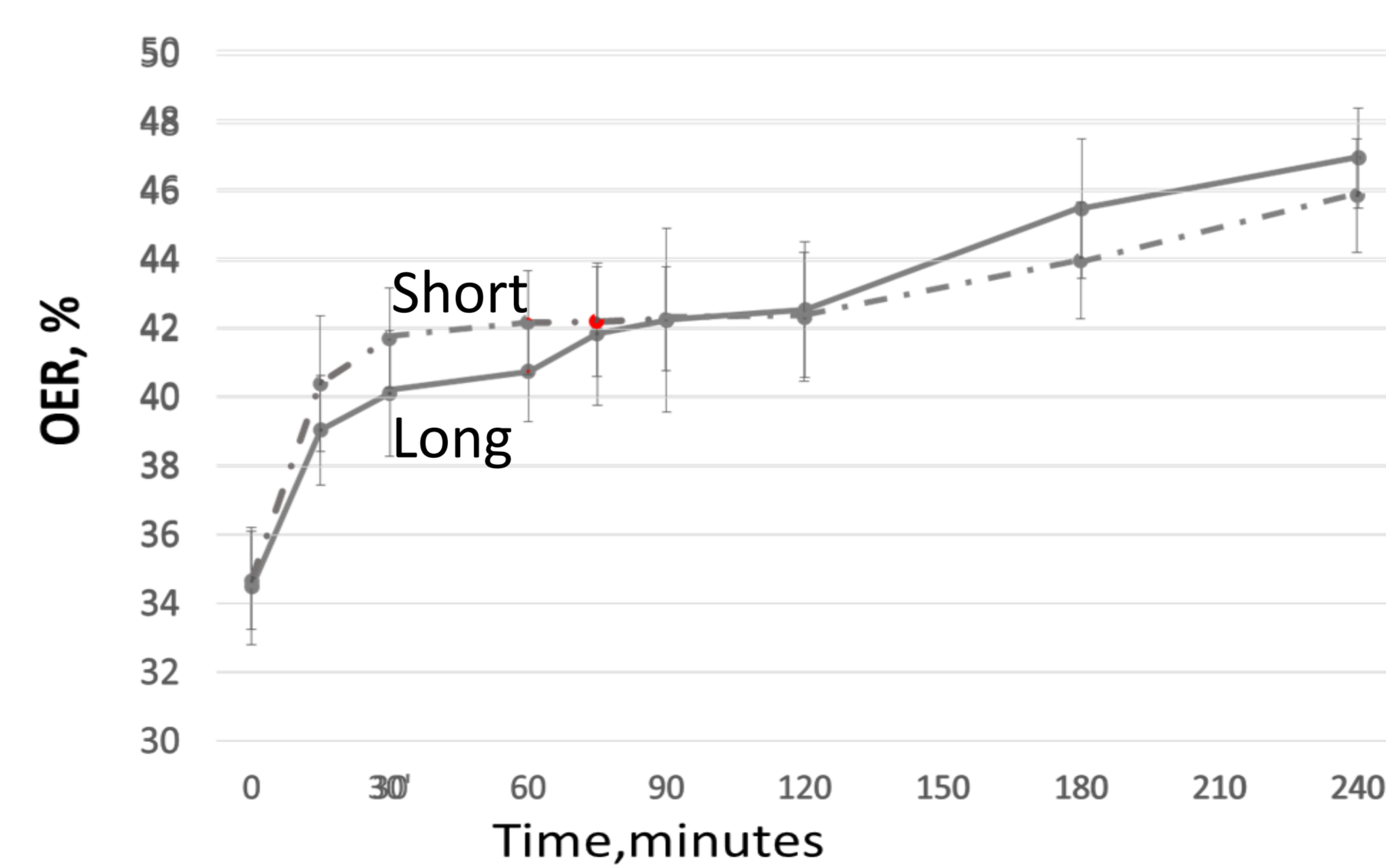


Fig. 2 OER trend during HD Short (median) vs Long interval

## Discussion and Conclusions

a. Even asymptomatic HD sessions increase O<sub>2</sub> consumption

in amounts that are measurable by OER;

b. HD intervals marginally modify the adaptive process to hypoxia;

c. iD affects this adaptive response possibly more than iUF.

Our results indicate that OER is a possible marker of HD stress, potentially useful in fragile patients.