

# IMPACT OF EXTRACORPOREAL MEMBRANE OXYGENATION CIRCUIT ON FENTANYL PHARMACOKINETICS IN CRITICALLY ILL PATIENTS



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## Background and importance



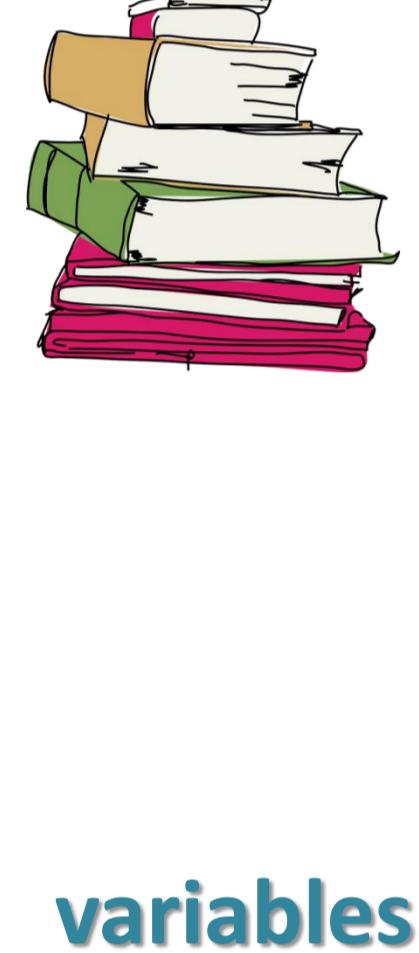
Fentanyl's lipophilicity and protein binding may contribute to a sequestration of the drug in the extracorporeal membrane oxygenation (ECMO) circuit, which could **impact to pharmacokinetic** and **response** in critically ill patient

## Aim and objectives

To assess the **impact of ECMO circuit** on the plasma concentration (**Pk**) of fentanyl in critically ill patients



## Material and methods



Observational, prospective, multidisciplinary, cohort study

variables

### Demographic

Age  
Sex  
BMI

### Therapeutic

Dose and duration  
Sedative concomitant drug  
CYP3A4 inducer-inhibitor

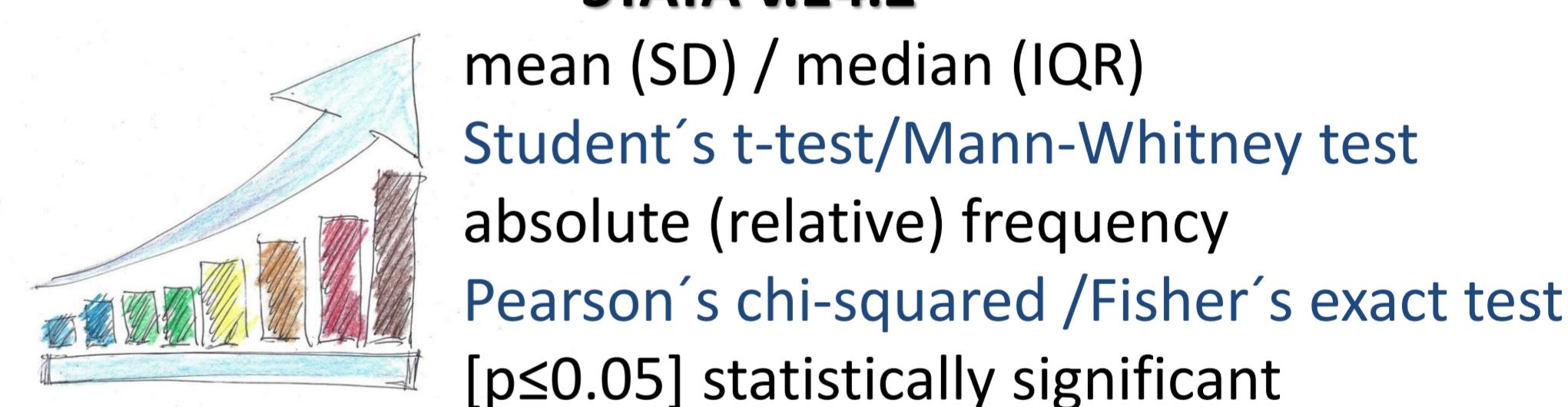
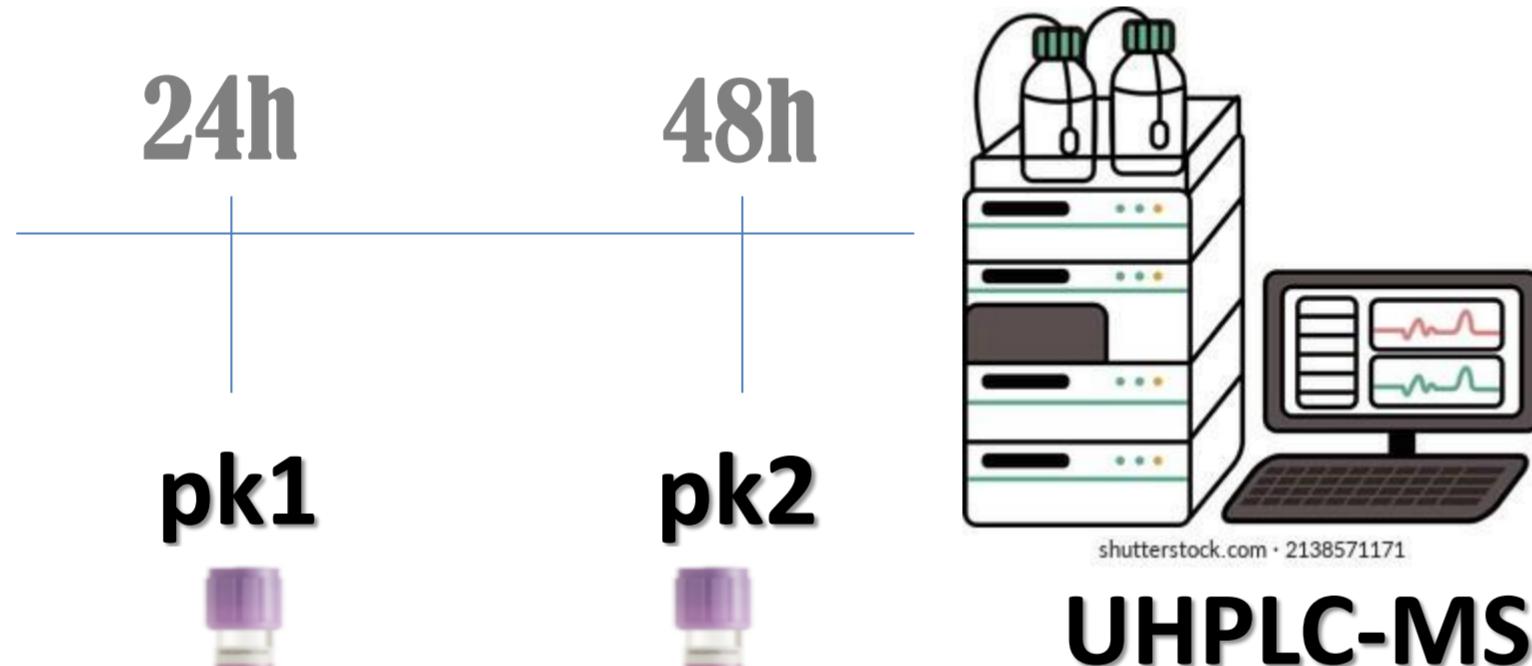
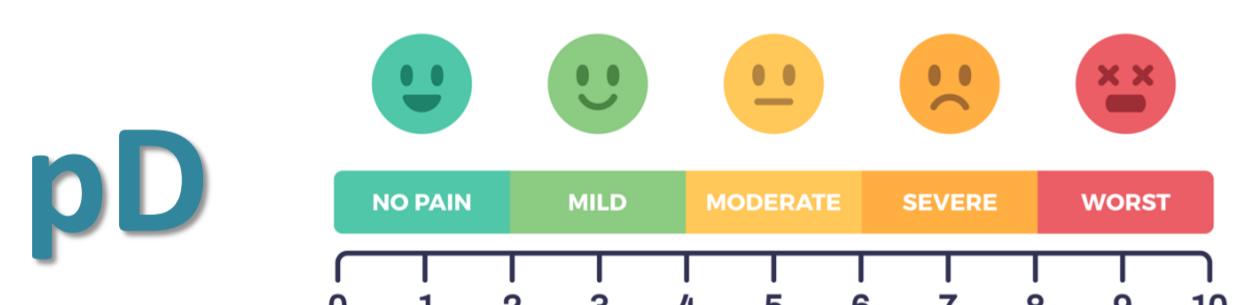
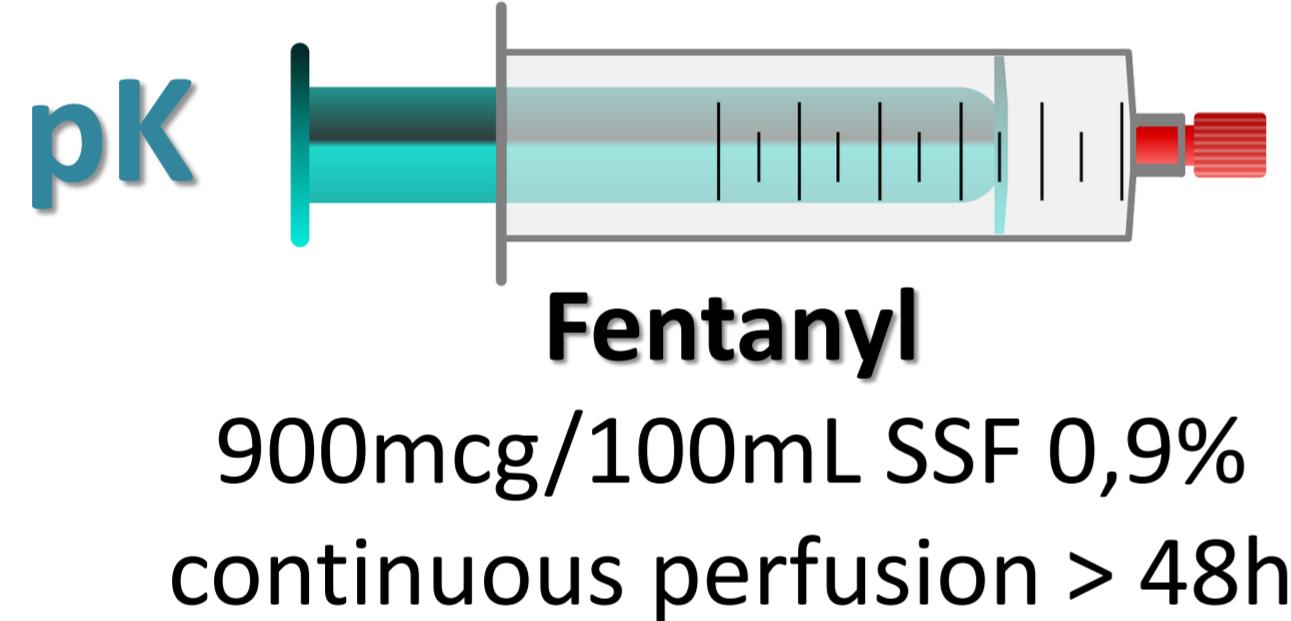
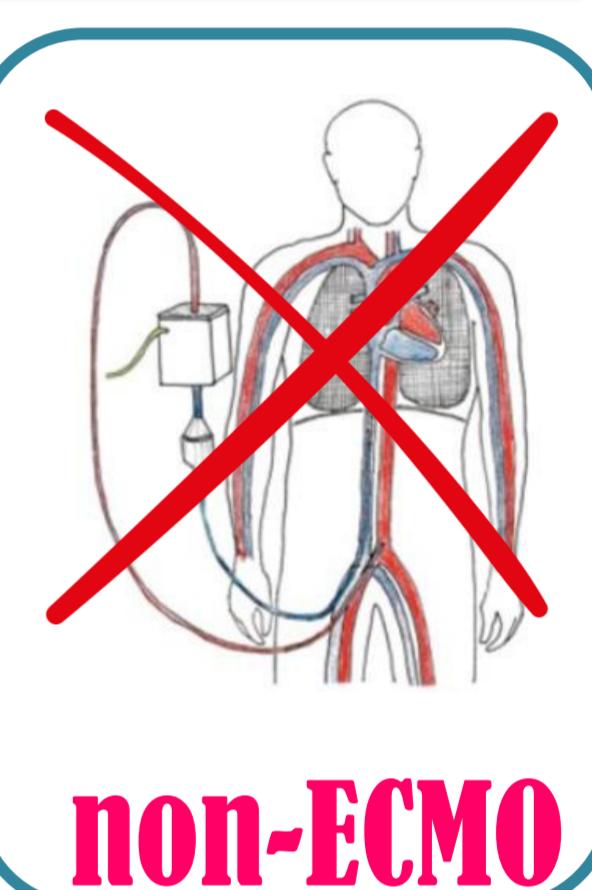
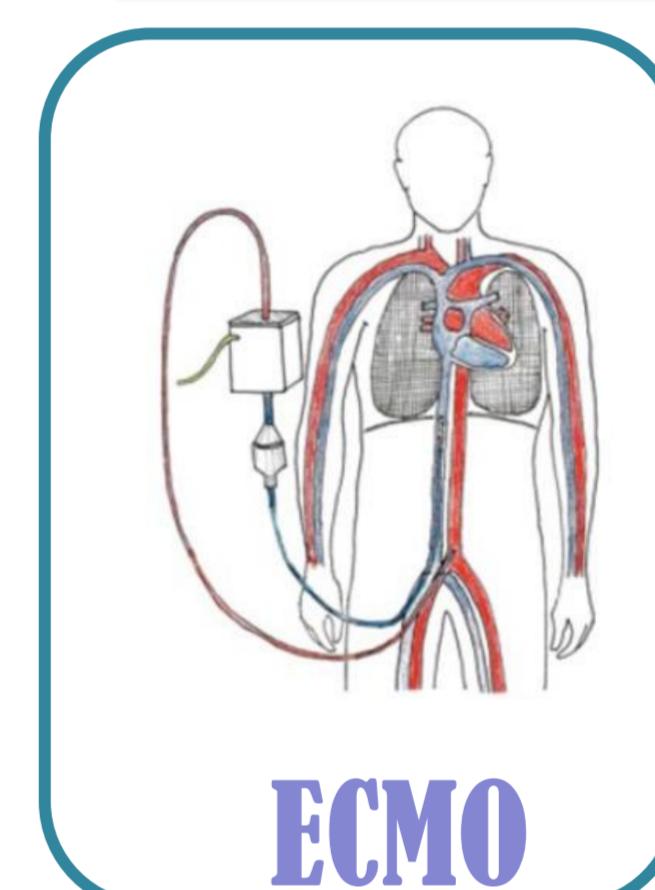
### Clinical

Diagnosis and comorbidities  
SAPS3 score  
CRRT and Mechanical ventilation (MV)  
Clinical parameters (at 0, 24, 48h)  
- BP, HR, RR  
Analytical parameters (at 0, 24, 48h)  
- renal, hepatic, INR, protein, albumin

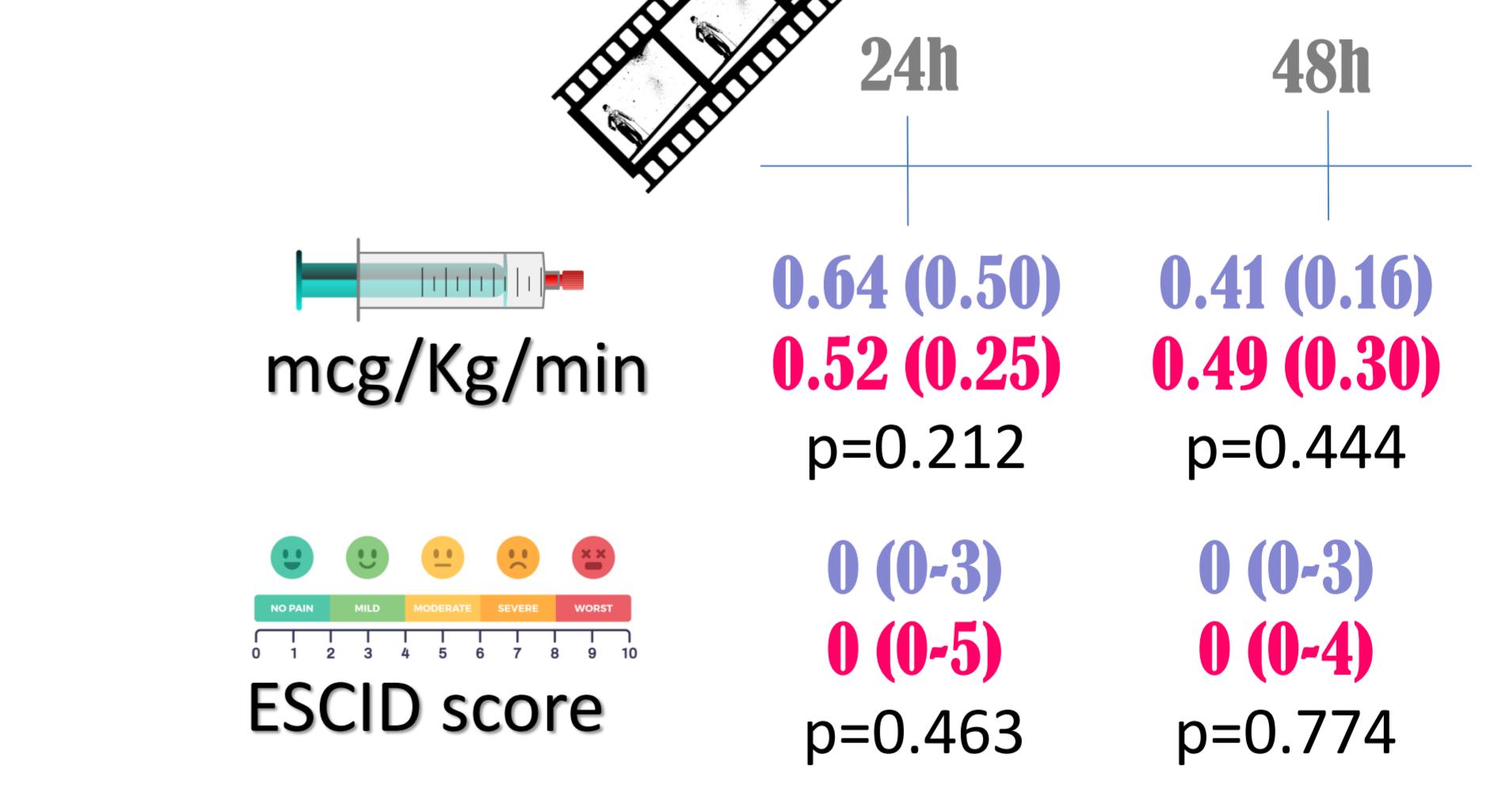
### Adverse events

Respiratory depression  
Bradycardia  
Hypotension  
CNS depression  
Constipation, ileus

### Adult critically ill patients



## Results



No statistically significant differences



Drug interaction  
Concomitant treatment  
Adverse events

## Conclusion and relevance

In our serie the use of **ECMO** had **no significant impact** on fentanyl **Pk** during the **first 48h**. Using a **standard perfusion** in patients on extracorporeal support to dose according to **the response** guided by the **ESCID** scale is **safe** to achieve **adequate pain control**.