



INTRODUCTION

Combined liver-lung transplantation (cLiLuTx) is a life-saving procedure for patients with dual organ failure. **The classic sequence dictates LuTx priority over LiTx, due to the tolerable ischemic time, which is considered shorter for the lung than for the liver.** However, recent reports describe successful LuTx following longer cold ischemic times as well as safe extension of the cross clamp time with ex-vivo lung perfusion (EVLP). Therefore, an **inversed sequence -liver-first- may have several benefits.**

AIM

To create a *theoretical framework* to outweigh the potential benefits of a liver-first sequence

PATIENTS

15 CLiThTx performed between 1/2000 and 12/2017

**N = 4 'liver-first'**

DEMOGRAPHICS / INDICATIONS

Demographics

Age (median-range)	43 y (17-63)
Gender (M/F)	3/1

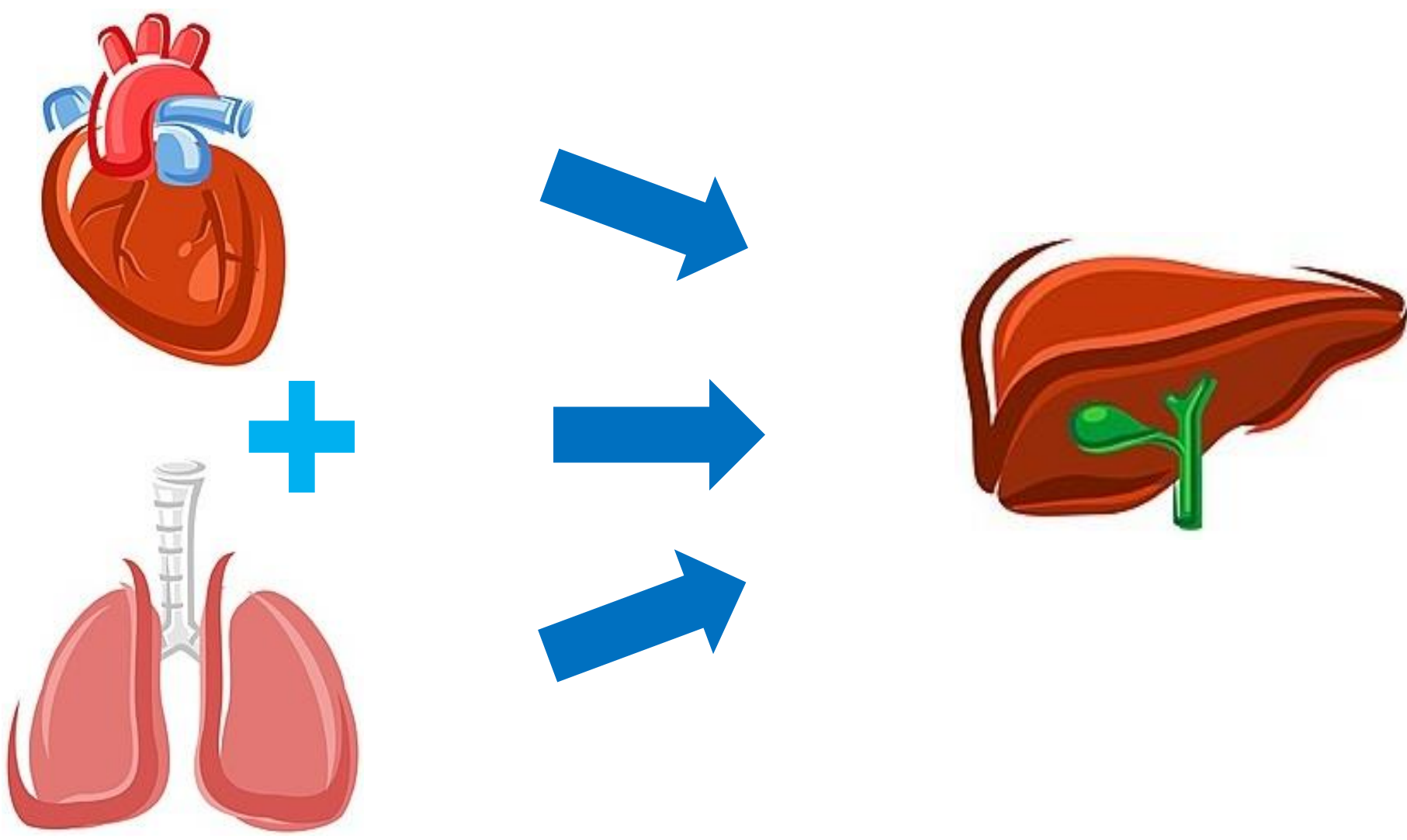
Indications

Liver indication	Pulmonary indication
Cystic fibrosis induced cirrhosis with portal hypertension (2)	Cystic fibrosis (2)
Epithelioid hemangioepithelioma	Epithelioid hemangioepithelioma
Tuberculostatics (Isoniazid) induced acute liver failure	COPD GOLD IV

**Surgical sequence**

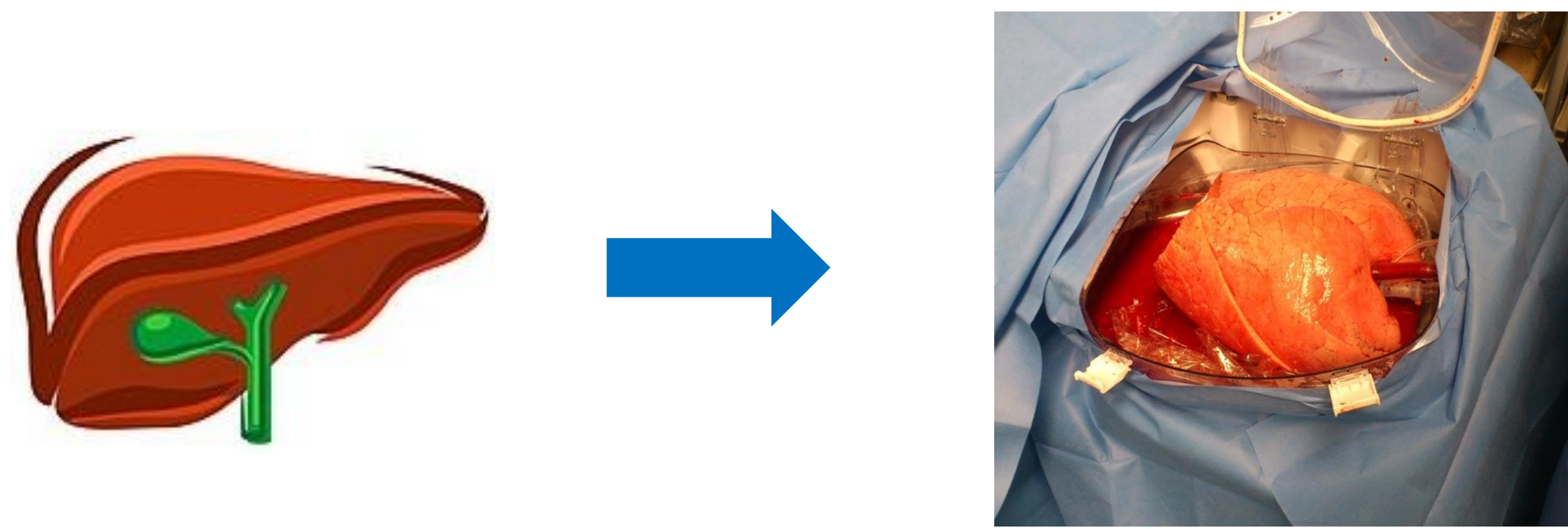
**Classic**  
HEART/LUNG FIRST  
Liver Second

STATIC, COLD  
preservation



**NEW principle**  
LIVER FIRST  
Lung (heart) second

Liver: COLD  
Lung: EVLP\*  
\*OCS™, Andover, USA



First case: drug-induced liver failure + COPD IV (AJT 2014 Oct)

CLINICAL OUTCOME (m Follow-up: 3.2 year)

Primary Graft Dysfunction at 72 hours:  
Case 1: 2; Case 2: 1; Case 3: 0; Case 4: 0

Rejection	Liver	Lung
Early (<3 mo)	0	0
Late (>3 mo)	0	1* B2
Chronic	0	0
Graft loss	0	0

B2: moderate bronchial rejection, treated with steroids

**Leuven 'LIVER-FIRST' principle**

*If the native lungs can withstand the primary LiTx*

- 1/ **Immunological benefit:** transplanted liver neutralizes donor specific HLA antibodies => ↓ rejection
- 2/ **Liver surgery** harms the native lungs instead of lung allografts => ↓ lung edema
- 3/ **Liver IRI** captured by native lungs instead of lung allografts => ↓ lung edema
- 4/ Restoration of **coagulation status** prior to LuTx => ↓ tranfusion during LuTx
- 5/ Avoids **anhepatic phase** and ↓ coagulation factors after LuTx => ↓ risk of thoracic bleeding
- 6/ **Shorter liver cold ischemia time** => ↓ biliary strictures

CONCLUSION

*In our 4 cLiLuTx the sequence could safely be inversed -liver-first- with successful outcome. For every cLiLuTx the liver-first versus lung-first principle should be discussed in team, assessing the organ-specific disease severity.*