

Trapianto del polmone: cosa c'è di nuovo?



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Fondazione IRCCS Ca' Granda
Ospedale Maggiore Policlinico

Sistema Socio Sanitario
 Regione
Lombardia

Agenda

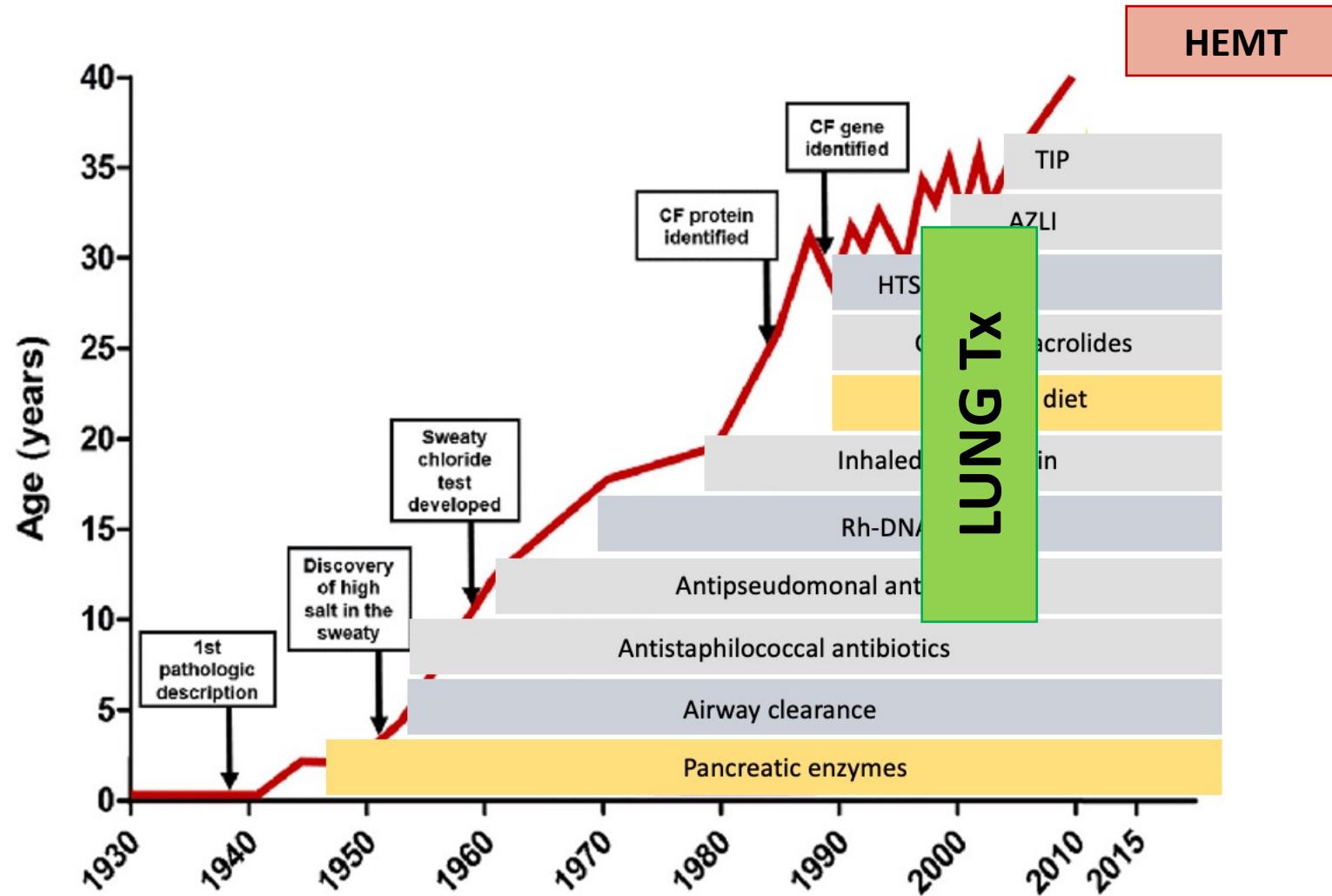
- Introduzione
- Referral & Lista
- Complicanze
- Modulatori CFTR
- La telemedicina

*** Nessun conflitto di interesse per questa presentazione***

LuTx per FC

Introduzione

FC: un paradigma terapeutico in costante evoluzione



Modified from
Elborn S, 2013

Advanced CF lung disease (ACFLD)

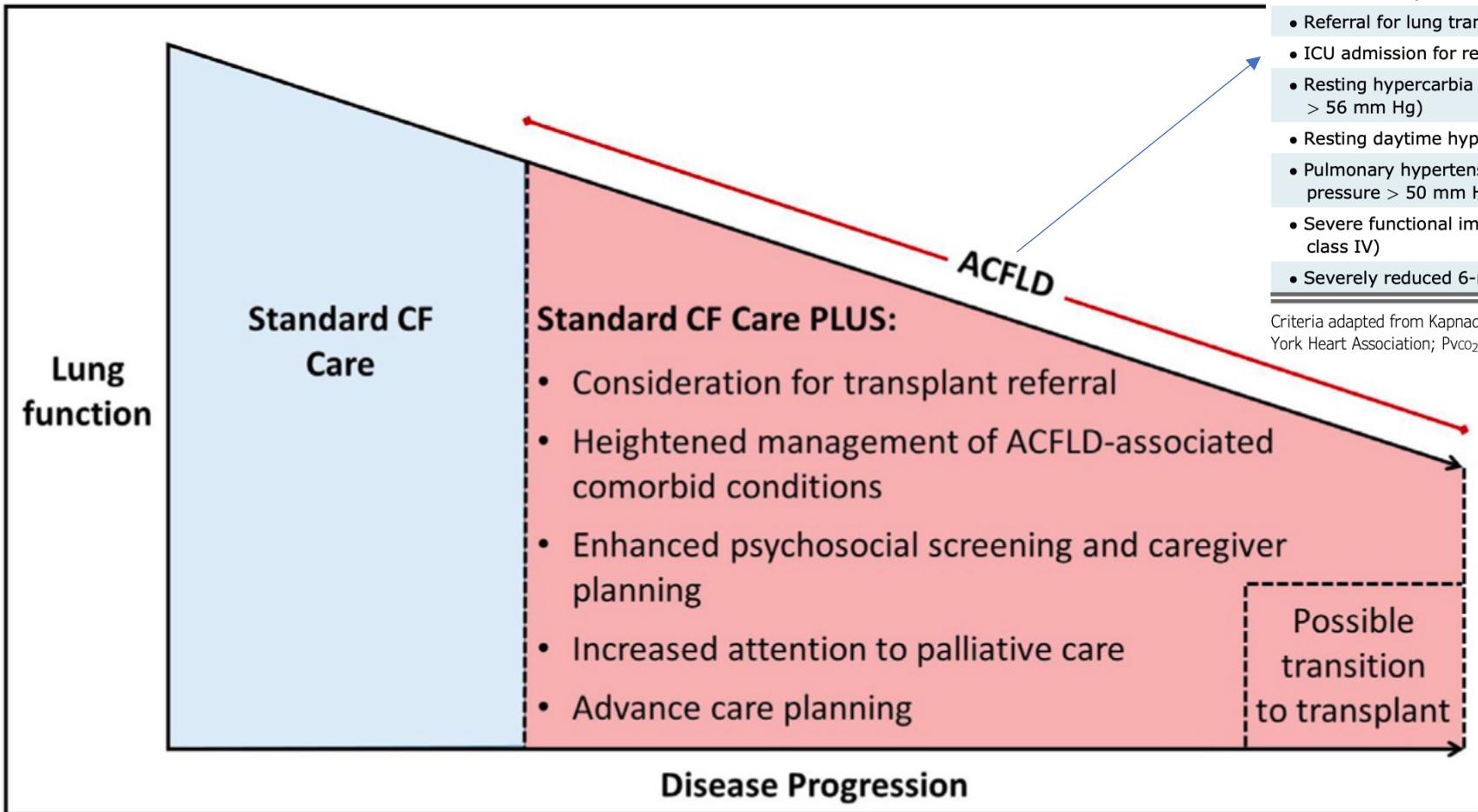
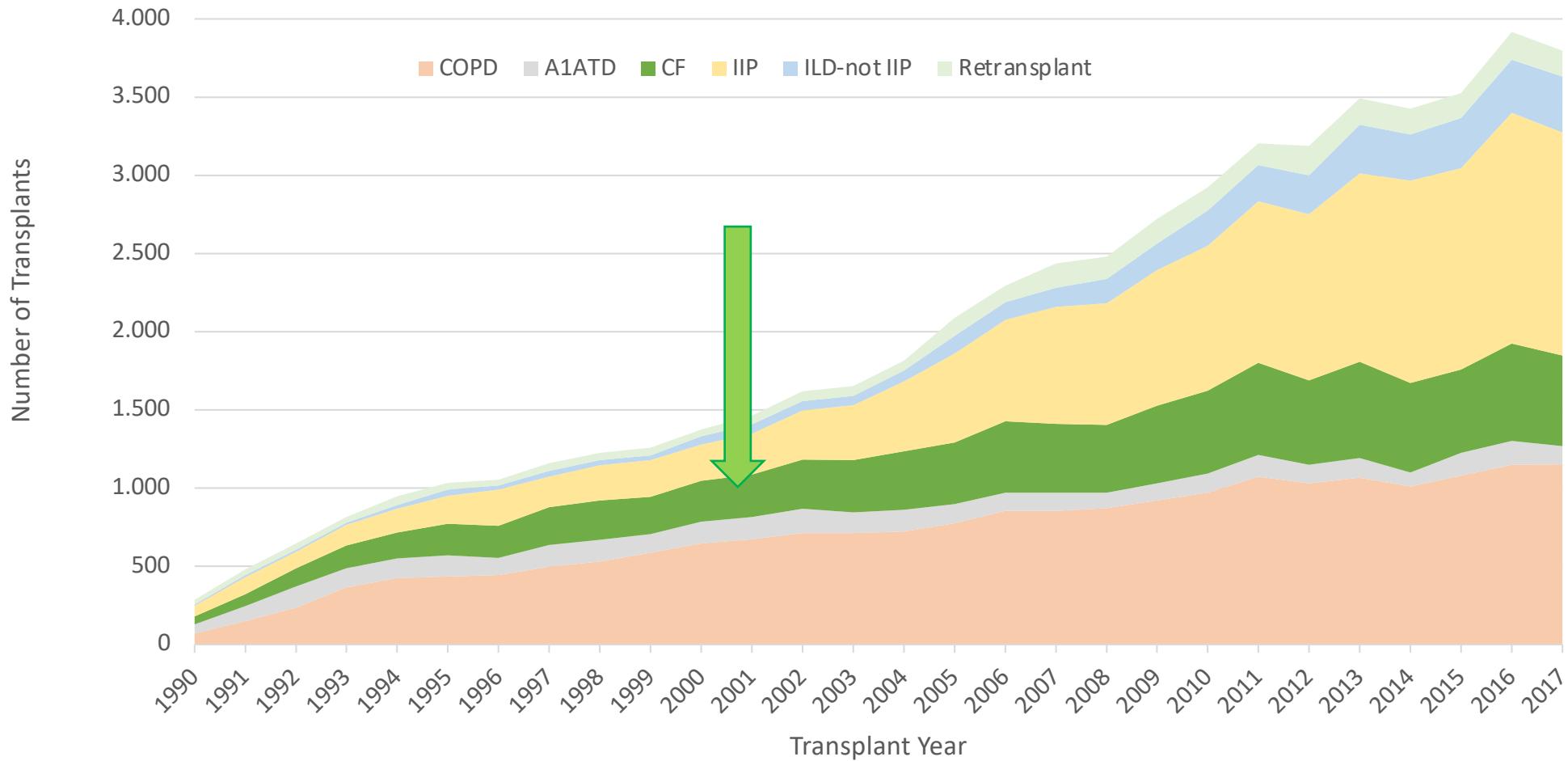


TABLE 2] Defining Advanced CF Lung Disease

Any of the following criteria
• FEV ₁ < 40% predicted
• Referral for lung transplant evaluation
• ICU admission for respiratory failure
• Resting hypercarbia (Paco ₂ > 50 mm Hg or Pvco ₂ > 56 mm Hg)
• Resting daytime hypoxemia
• Pulmonary hypertension (pulmonary arterial systolic pressure > 50 mm Hg on echocardiogram)
• Severe functional impairment (NYHA functional class IV)
• Severely reduced 6-min walk distance (< 400 m)

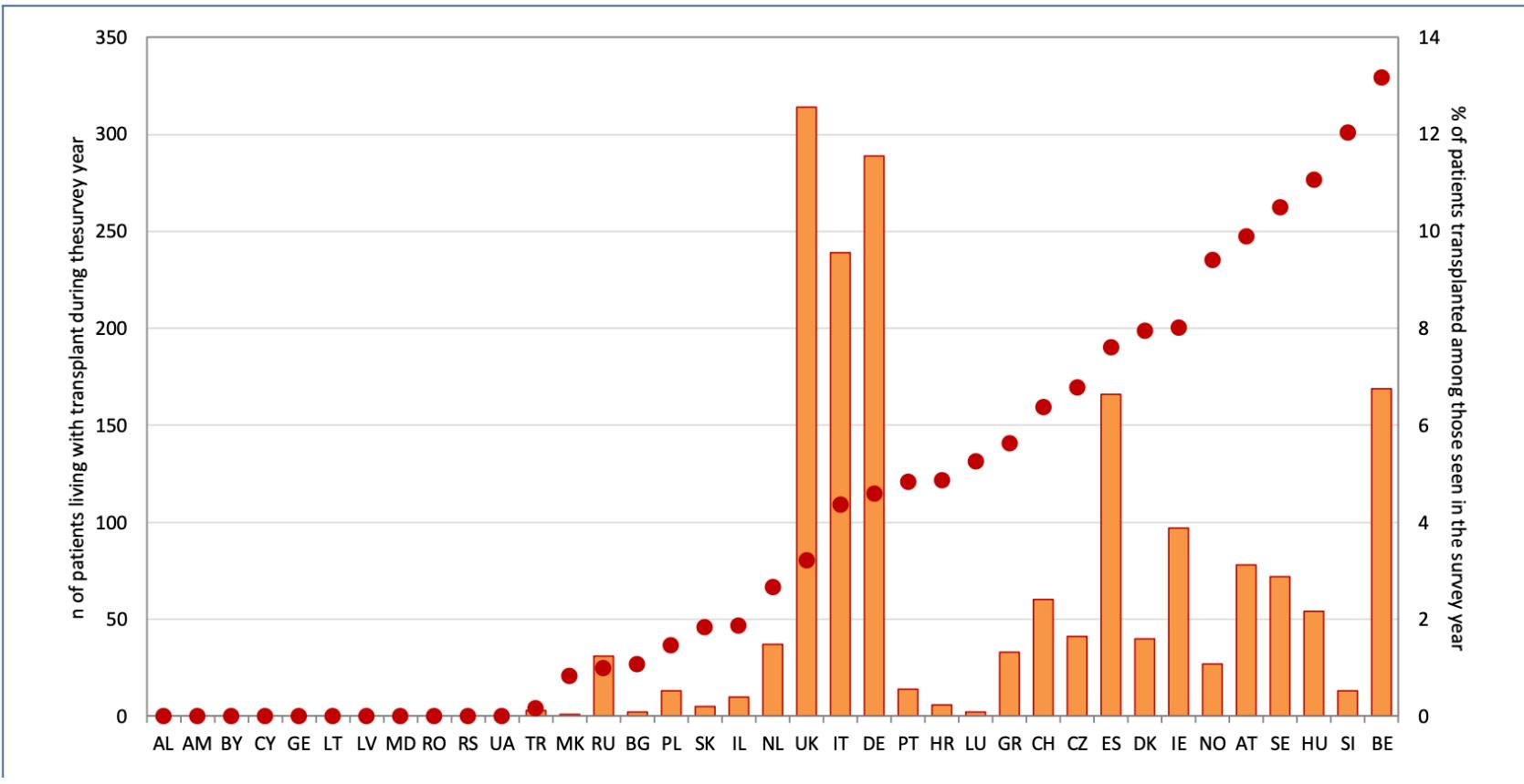
Criteria adapted from Kapnidak et al.³⁰ CF = cystic fibrosis; NYHA = New York Heart Association; Pvco₂ = partial pressure of co₂ in venous blood.

LuTx (adulti) - indicazioni



LuTx (adulti) per FC – dati per ECFSPR

Figure 8.1 Number of patients living in 2018 with transplanted lungs, by country.

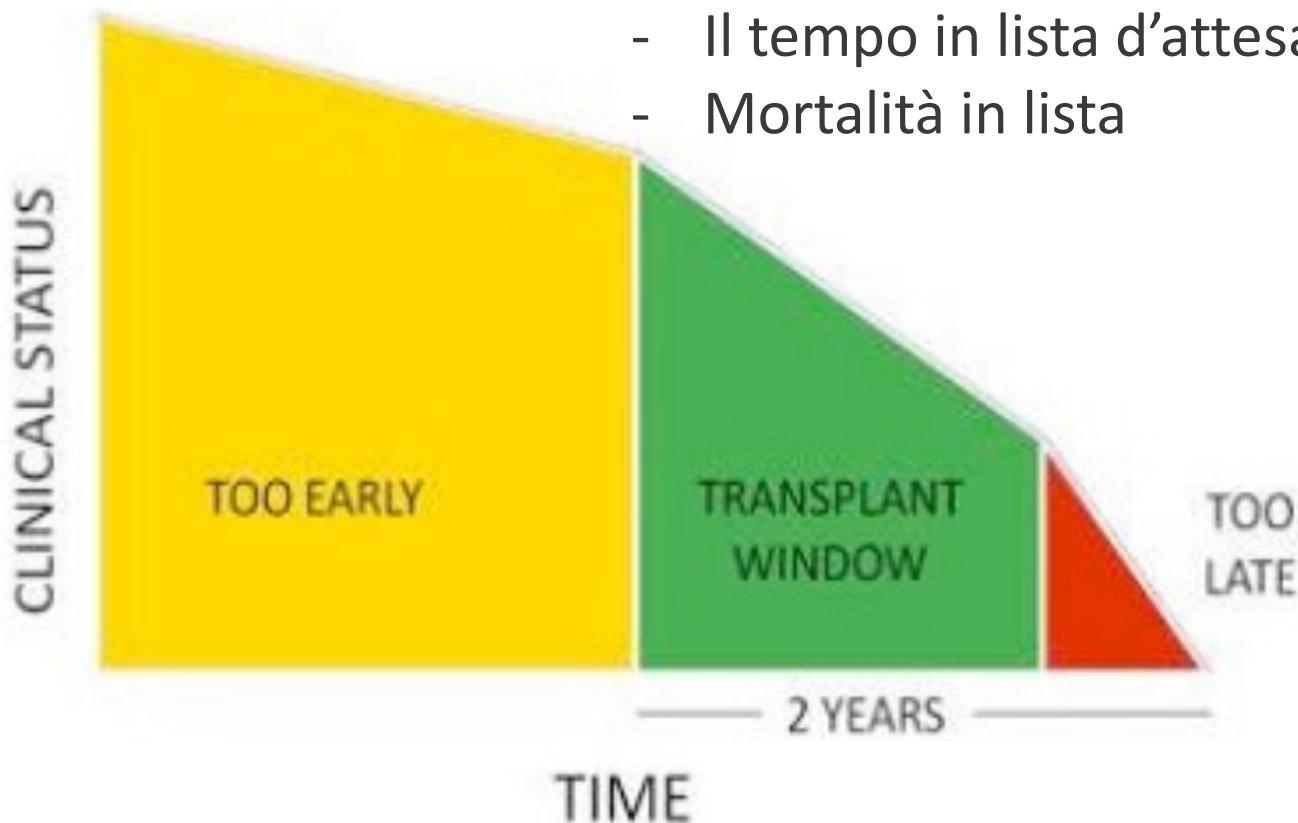


Referral & Listing

Cos'è cambiato negli ultimi anni; il ruolo dei modulatori

La finestra trapiantologica

- Il listing richiede tempo
- Il tempo in lista d'attesa può essere (molto) lungo
- Mortalità in lista



Candidati al LuTx

ARTICLE IN PRESS

Indicazioni generali - Adulti con end stage lung disease con i seguenti criteri:

- Alto rischio di mortalità (> 50%) nei successivi 2 anni
- Alta probabilità (>80%) di sopravvivenza a 5 anni da un eventuale trapianto in caso di adeguata funzione del graft polmonare.

Table 2 Risk factors for poor post-transplant outcomes

Risk factors can change over time and may not be a contraindication for referral, but when present at the time of listing or while listed for lung transplantation may increase risk for poor transplant outcomes. There was 100% consensus (24 committee members) for the content of the entirety of Table 2.

ABSOLUTE CONTRAINDICATIONS:

- Candidates with these conditions are considered too high risk to achieve successful outcomes post lung transplantation.
 - Factor or condition that significantly increases the risk of an adverse outcome post-transplant and /or would make transplant most likely harmful for a recipient.
 - Most lung transplant programs should not transplant patients with these risk factors except under very exceptional or extenuating circumstances.
- [...]
1. Lack of patient willingness or acceptance of transplant
 2. Malignancy with high risk of recurrence or death related to cancer
 3. Glomerular filtration rate < 40 mL/min/1.73m² unless being considered for multi-organ transplant
 4. Acute coronary syndrome or myocardial infarction within 30 days (excluding demand ischemia)
 5. Stroke within 30 days
 6. Liver cirrhosis with portal hypertension or synthetic dysfunction unless being considered for multi-organ transplant

Candidati al LuTx per FC



Consensus document for the selection of lung transplant candidates: An update from the International Society for Heart and Lung Transplantation

**The Journal of
Heart and Lung
Transplantation**

<http://www.jhltonline.org>

Lorriana E. Leard, MD,^a Are M. Holm, MD, PhD,^b Maryam Valapour, MD, MPP,^c

Referral for lung transplantation should occur for an individual with CF meeting any of the following criteria despite optimal medical management including a trial of ELX/TEZ/IVA if eligible:

- FEV1 < 30% pred (o < 40% in children)
- FEV1 < 40% pred (o < 50% in children) and any of the following:
 - 6MWT distance < 400 mt
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 - Hyooxemia
 - Pulmonary hypertension (PAPs > 50 mmHg on echocardiogram or evidence of right ventricular dysfunction)
 - Worsening nutritional status despite supplementation
 - 2 exacerbations per year requiring IV antibiotics
 - Massive hemoptysis (>240 mL) requiring bronchial artery embolization
 - Pneumothorax
- FEV1 < 50% pred + rapidly declining based on PFT or progressive symptoms
- Any exacerbation requiring positive pressure ventilation

Referral: un flusso di informazioni



- Stato funzionale (ultime PFR disponibili e andamento nel tempo)
- Accertamenti per valutazione scambi gassosi
 - EGA
 - 6MWT e SN
- Dati microbiologici:
 - Batteri
 - Miceti
 - NTM
- Storia di riacutizzazioni
- Precedenti embolizzazioni e/o PNX
- Ecocardio
- TC torace con immagini
- Comorbilità e stato nutrizionale

Referral & Listing: il ruolo del fisioterapista

- Passaggio di consegne dal centro di provenienza
- Disostruzione
- Fabbisogno di O₂tp sotto sforzo
- Ottimizzazione ventiloterapia non invasiva
- ALLENAMENTO!

Transplantation. 103(9):e220–e238, SEPTEMBER 2019
DOI: 10.1097/TP.0000000000002806, PMID: 31461743
Issn Print: 0041-1337
Publication Date: September 2019



Exercise for Solid Organ Transplant Candidates and Recipients: A Joint Position Statement of the Canadian Society of Transplantation and CAN-RESTORE

Tania Janaudis-Ferreira;Sunita Mathur;Robin Deliva;Nancy Howes;Catherine Patterson;Agnès Räkel;Stephanie So;Lisa Wickerson;Michel White;Yaron Avitzur;Olwyn Johnston;Norine Heywood;Sunita Singh;Sandra Holdsworth;

Transplantation. 101(9):1967–1968, SEPTEMBER 2017
DOI: 10.1097/TP.0000000000001784, PMID: 29633979
Issn Print: 0041-1337
Publication Date: September 2017



The Emerging Importance of Skeletal Muscle Function in Assessing Candidates for Transplantation

James Walsh;Daniel Chambers;Peter Hopkins;



COMMENTARY

Sarcopenia in lung transplant candidates: A novel biomarker to estimate skeletal muscle mass

James R. Walsh, Peter M. A. Hopkins

First published: 03 March 2018 | <https://doi.org/10.1111/ctr.13200>

RESPIRATORY CARE

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Research Article | Original Research

The 1-Minute Sit-to-Stand Test in Lung Transplant Candidates: An Easy-to-Use Alternative to the 6-Minute Walk Test?

Dario Kohlbrenner, Christian Benden and Thomas Radtke
Respiratory Care October 2019, respcare.07124; DOI: <https://doi.org/10.4187/respcare.07124>

Lista d'attesa – allocazione degli organi

Ethical Principles for the Allocation of Donor Lungs

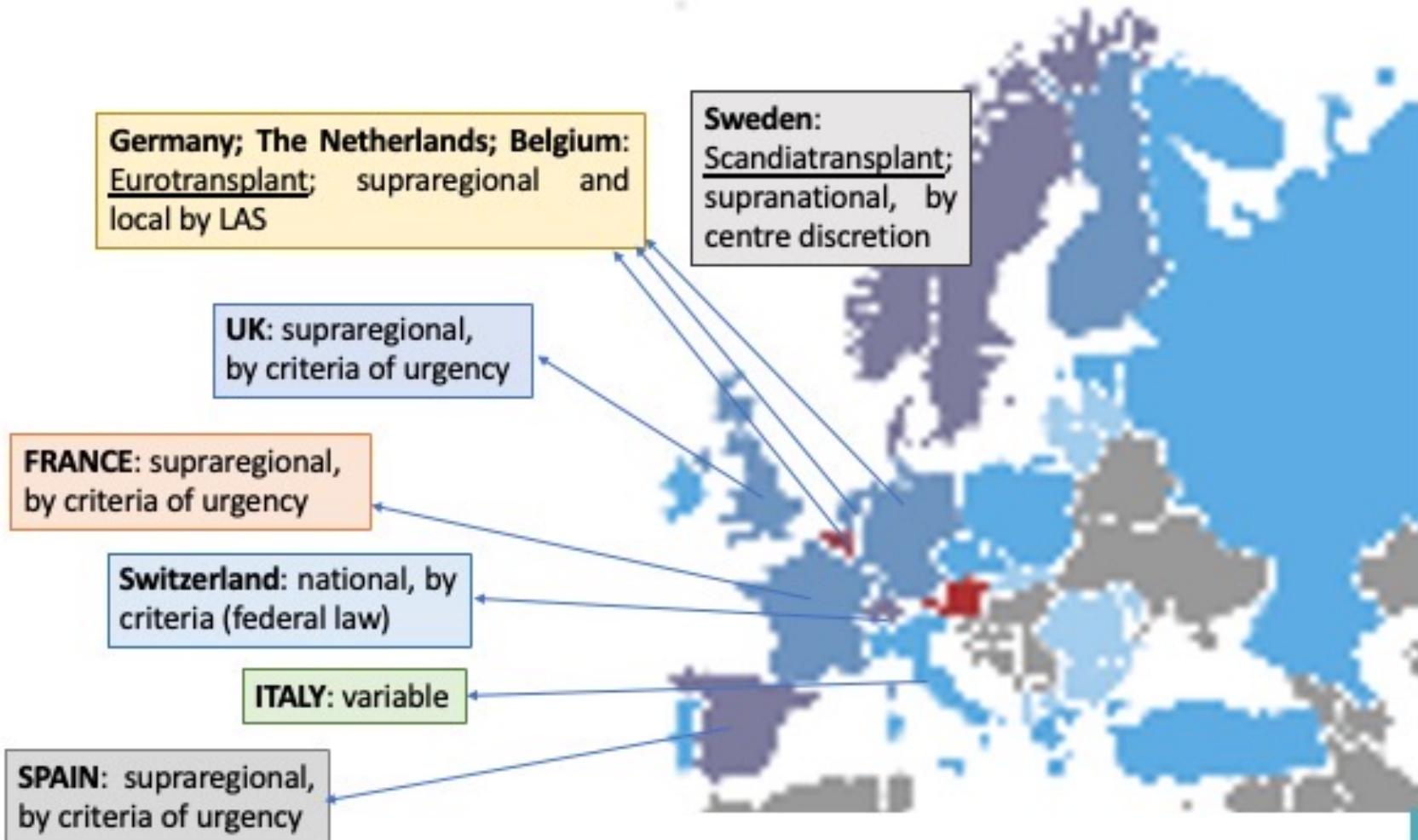
<i>Principle</i>	<i>Application to organ allocation</i>
UTILITY	To maximize net benefit (e.g., using years of survival gained to prioritize allocation)
JUSTICE	To distribute the benefits and burdens of organ allocation system in a fair way (e.g., using medical urgency to prioritize allocation, allowing special consideration for candidates for whom it is difficult to find a suitable organ)
RESPECT FOR PERSONS	To treat persons as autonomous with the right for self-determination (e.g., the right to give or withhold informed consent for a lung transplant)

Modified from:

Leard LE et al., JHLT 2021;

<https://optn.transplant.hrsa.gov/resources/ethics/ethical-principles-in-the-allocation-of-human-organs/>

Eterogenità nell'allocazione degli organi



Lista d'attesa – LAS (Lung Allocation Score)

Goal: net benefit of the transplant to the candidate as well as clinical urgency, and it is calculated using a series of pretransplant clinical diagnostic data that analyses revealed to be predictive of both pre- and posttransplant outcomes.

Range 0-100, the higher, the better (or worse?) >> the higher the more urgent to be transplanted/the better (expected) after LuTx.

- >> decrease in WL mortality
- >> decrease in time on WL
- >> improved survival after LuTx

Table 1. Lung Allocation Score Components^a

Waiting list urgency parameters

Age	O ₂ requirement at rest
Body mass index	Diabetes mellitus
Diagnosis	Six-minute walk distance <150 ft
Functional status (no assistance, some assistance, total assistance)	Continuous mechanical ventilation
FVC (% predicted)	Partial pressure of CO ₂
Pulmonary artery systolic pressure	Increased Pco ₂ of 15% for 6 months

Posttransplant survival variables

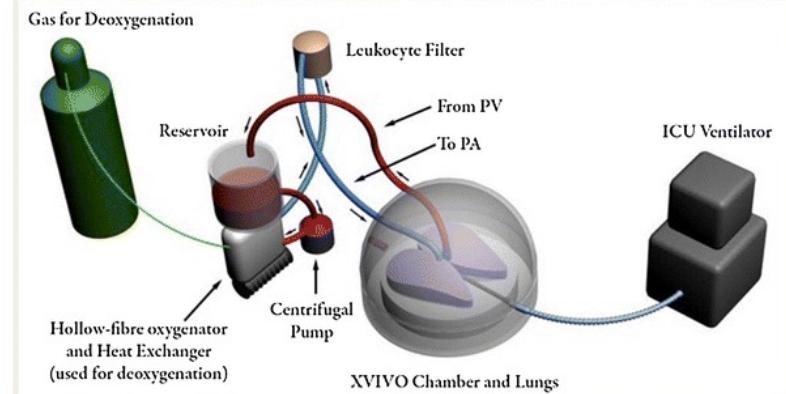
Age	Continuous mechanical ventilation
Functional status	Diagnosis
FVC (% predicted)	Pulmonary capillary wedge pressure

^a Lung allocation score (LAS) calculated from difference between expected 1-year posttransplant survival and waiting list urgency. Score is standardized with range of 0–100 [5].

FVC = forced vital capacity; Pco₂ = partial pressure of carbon dioxide.

Strategie per aumentare il pool di donatori

Medium percentage of solid organ use for transplantation	
	% utilized donors
Kidney	74%
Liver	78%
Heart	36%
Lung	18,6%



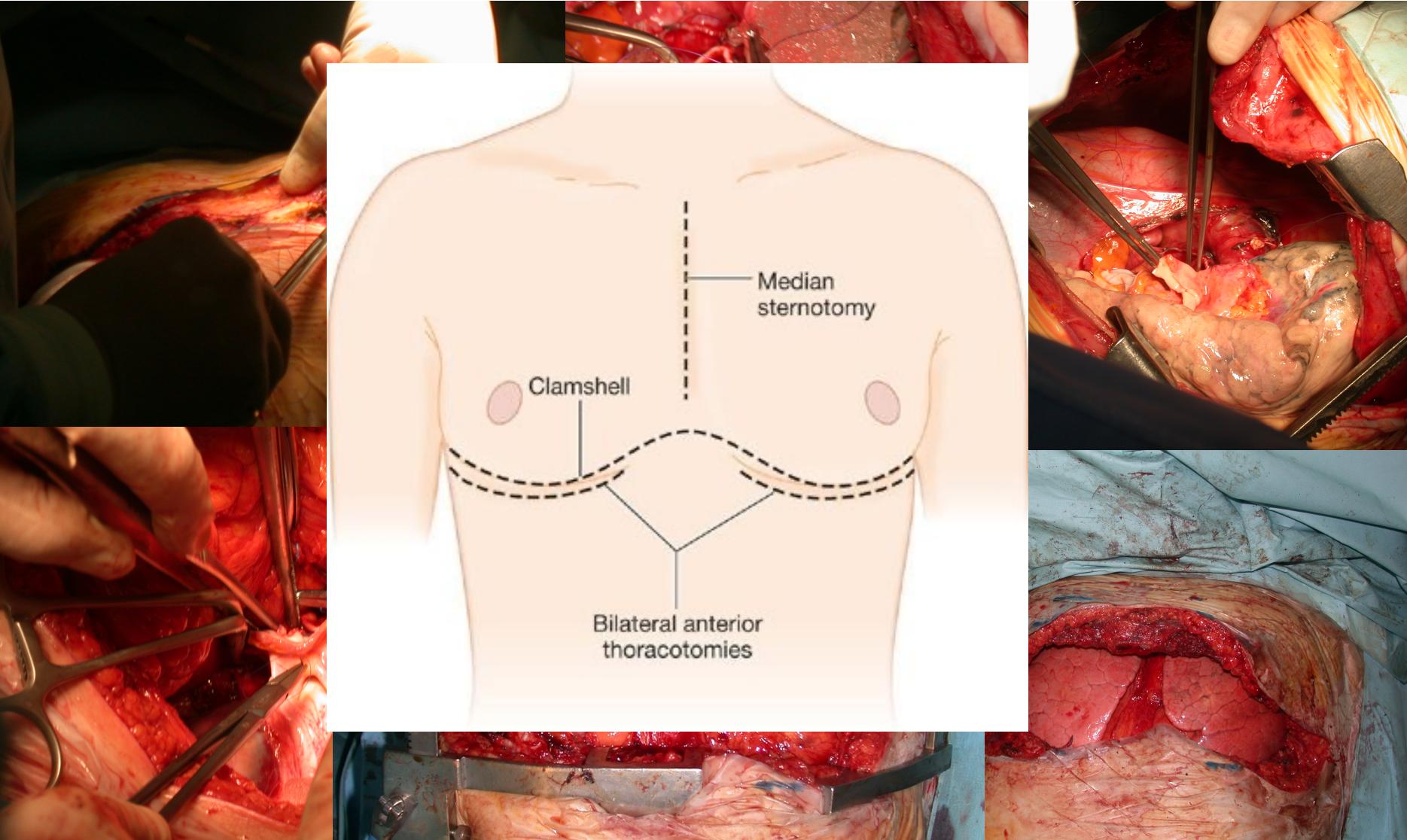
Strategies to expand donor pool:

- Extended criteria – «marginal» donor
- Living lobar
- Ex Vivo Lung Perfusion (EVLP)
- Organ Care System (OCS)
- Donation after cardiocirculatory death (DCD)

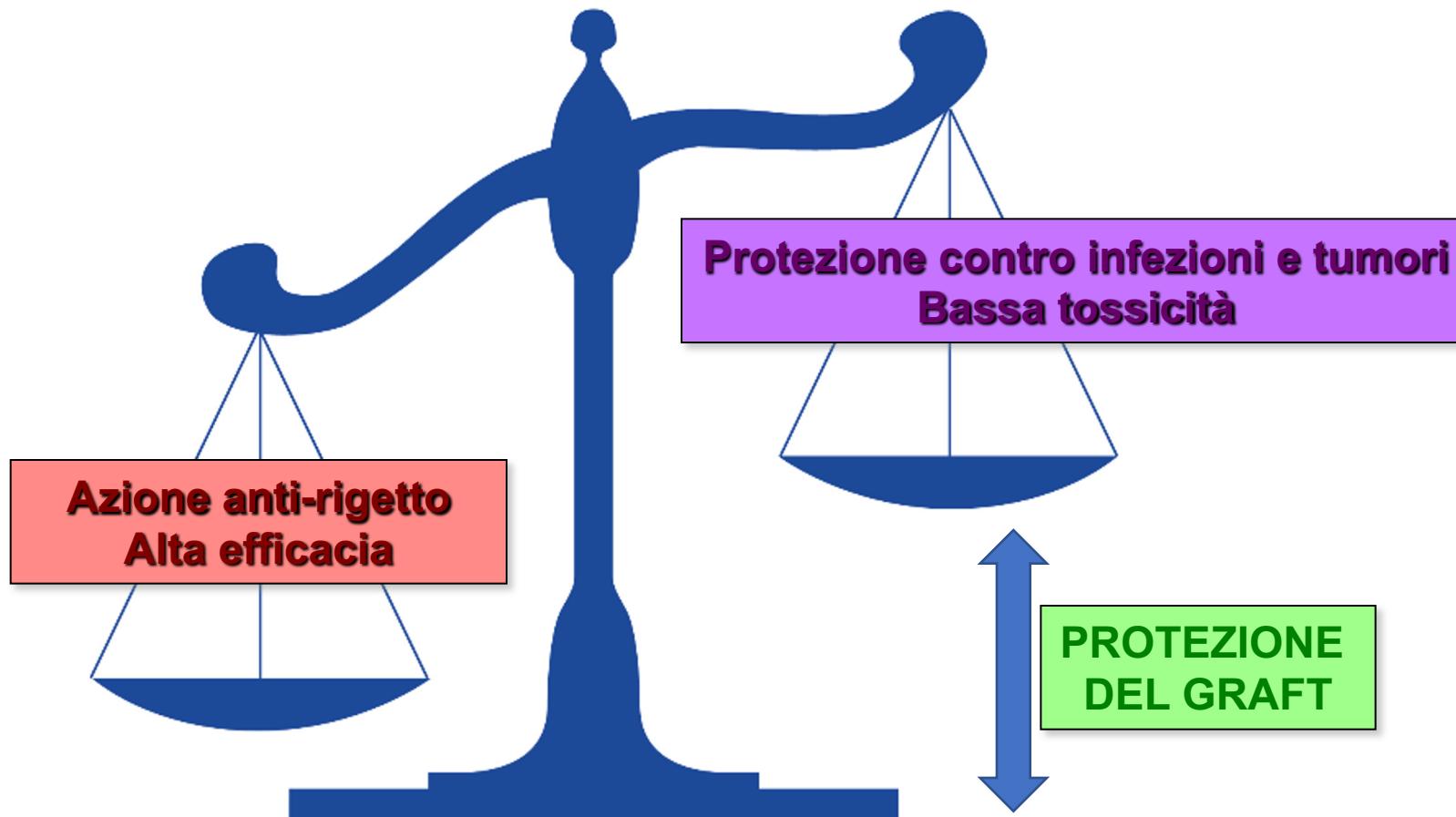
Il trapianto

Sorveglianza, terapie e complicanze

#Pneumi ❤ chirurghi



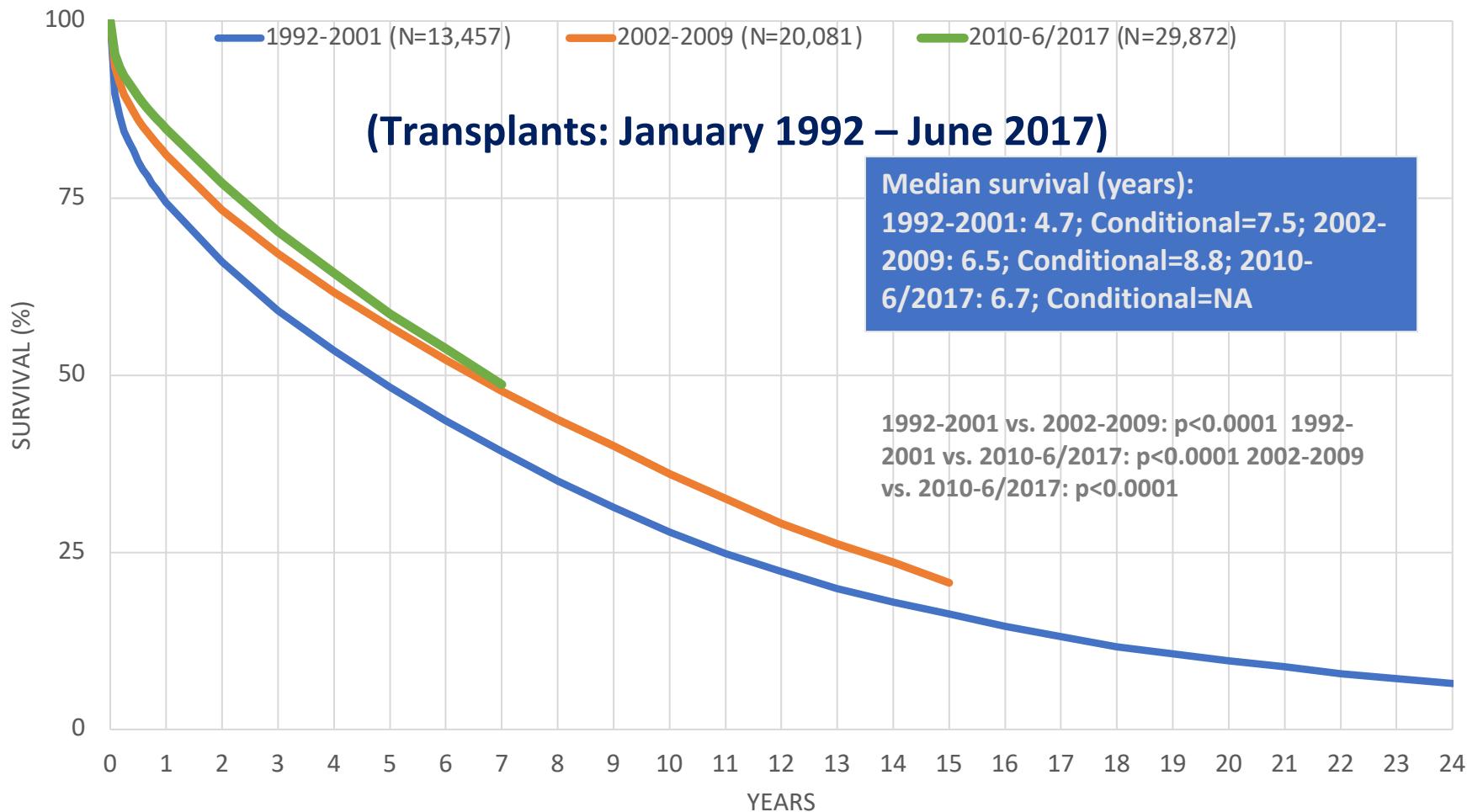
Terapia immunosoppressiva - RAZIONALE



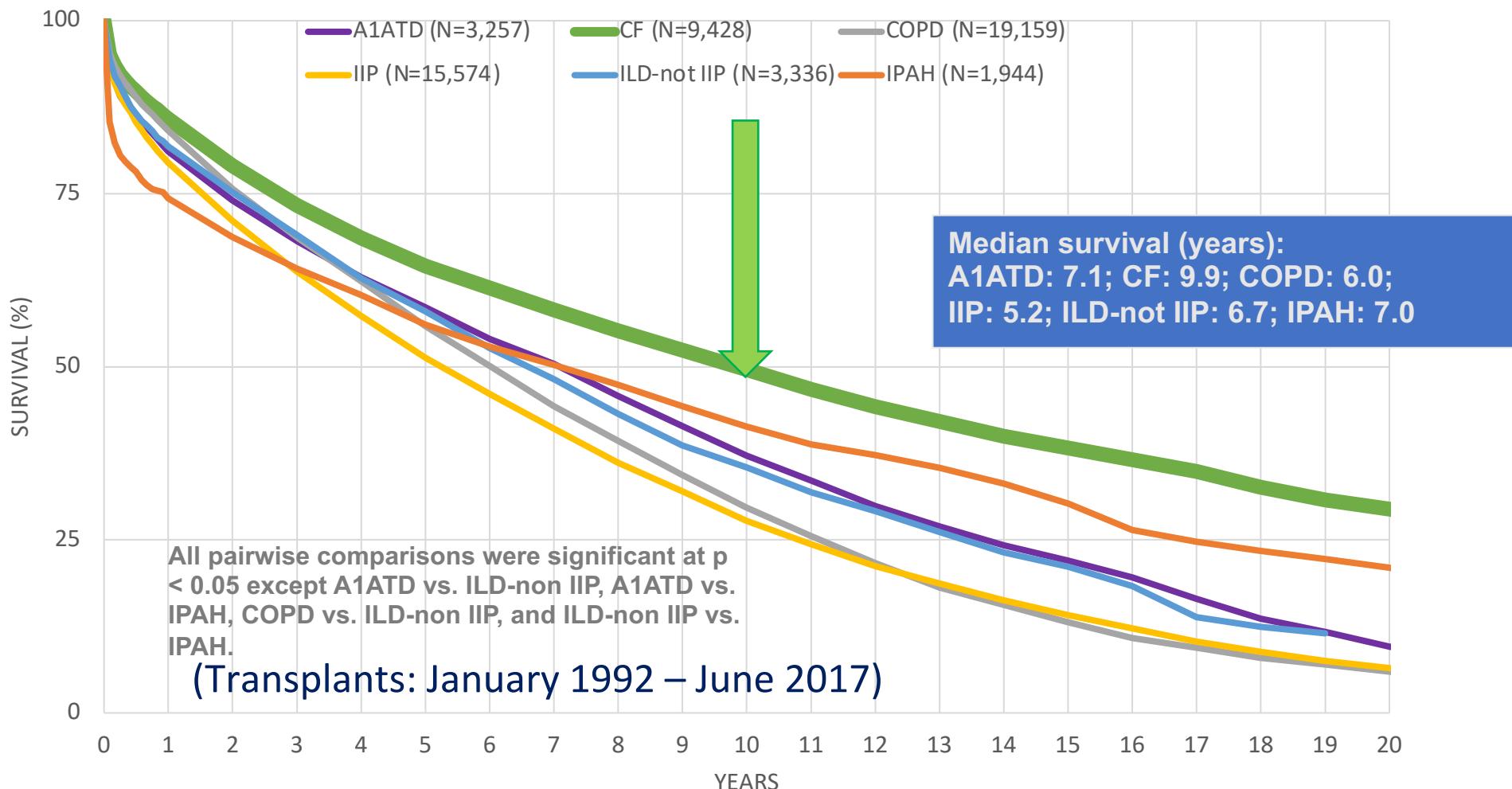
Successful outcomes following lung transplantation depend on our ability to pharmacologically manipulate the immune system to prevent rejection of the lung allograft.

Heeger PS, Dinavahi R. *Transplant immunology for non-immunologist*. Mount Sinai J Med. 2012;79(3):376–87.

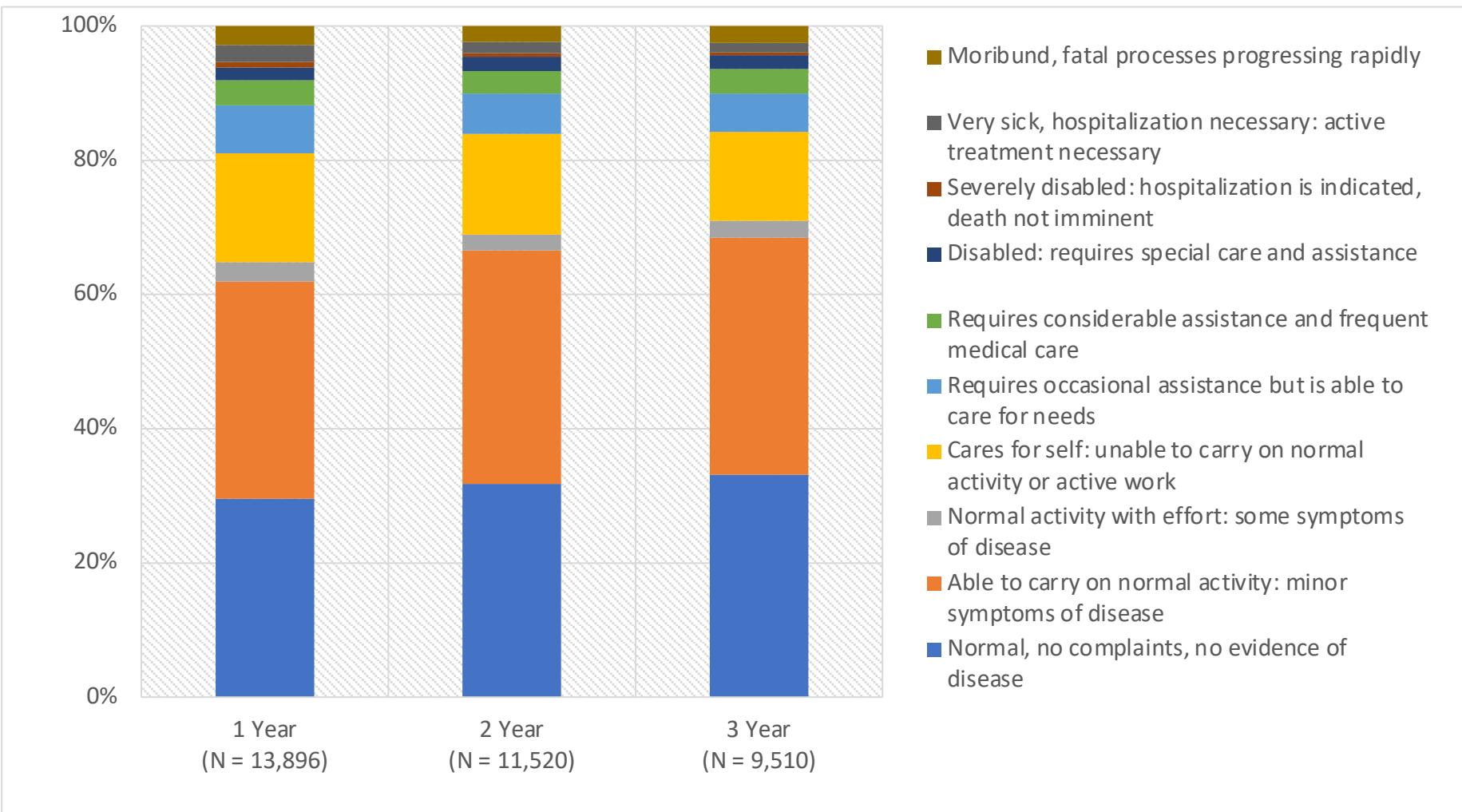
LuTx (adulti) – sopravvivenza per era



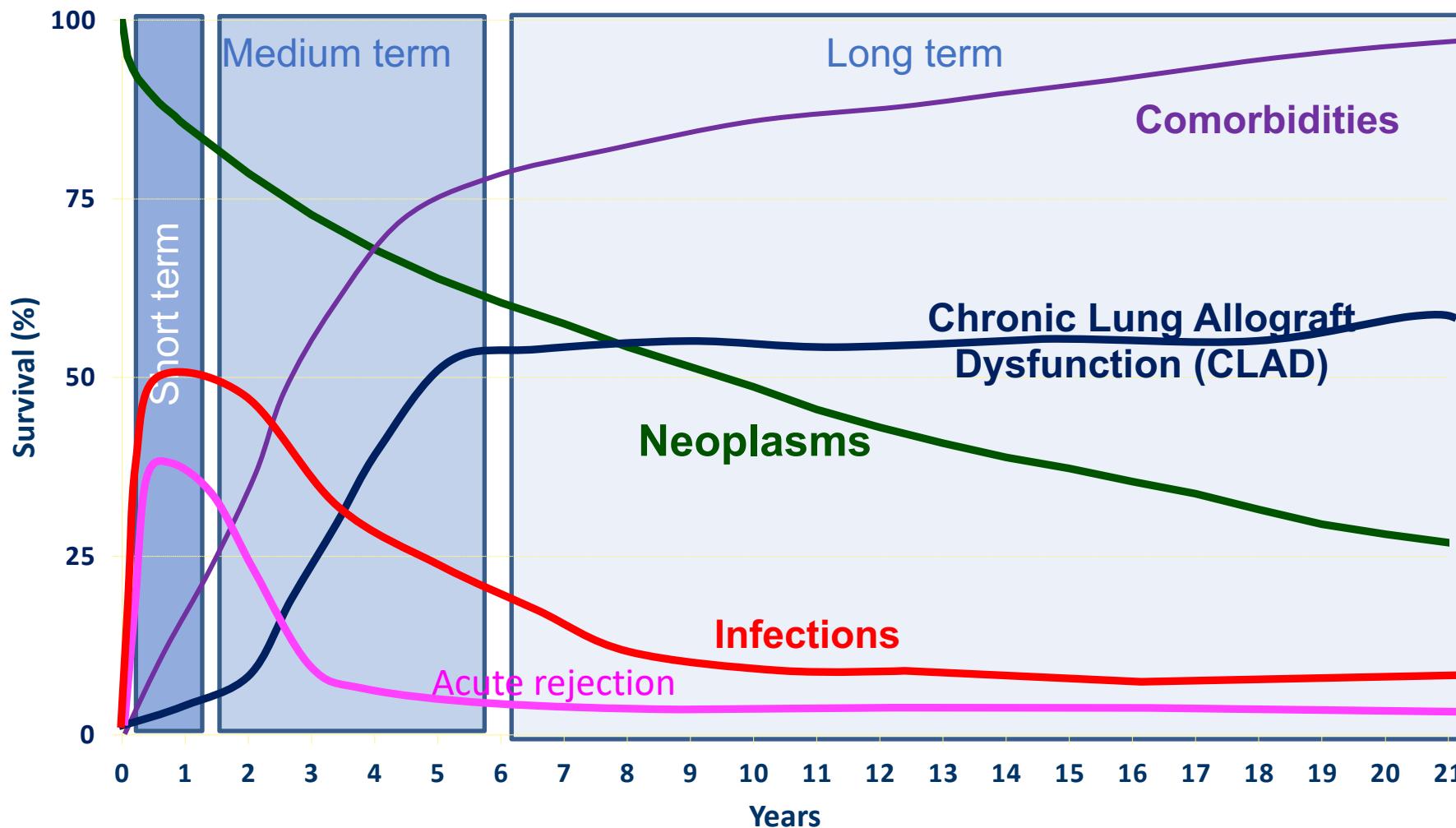
LuTx (adulti) – sopravvivenza per indicazioni



LuTx (adulti) – come si vive dopo?



LuTx - complicanze

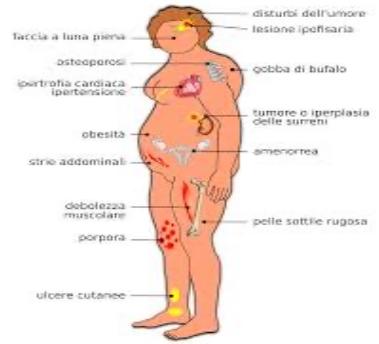


Sorveglianza del paziente e del graft

- Visite di routine in ambulatorio, comprensive di esami del sangue, PFR, RX torace, EGA arterioso e 6MWT
- TC torace e TBB/FBS almeno una o due volte l'anno
- Screening per DSA de-novo (anticorpi specifici del donatore)
- Riferimento ad altri specialisti per la gestione delle comorbilità e delle complicanze extrarespiratorie

Effetti collaterali delle terapie post LuTx

- Ipertensione arteriosa
- Obesità
- Dislipidemia
- Diabete
- Osteoporosi
- Cataratta
- Ritardata guarigione delle ferite
- Distimie
- **Nefrotossicità**
- **Epatotossicità**
- **Fototossicità**
- **Neurotossicità**
- **Mielotossicità**



Outcome	Within <u>1 Year</u>	Within <u>5 Years</u>	Within <u>10 Years</u>
Hypertension	51.7%	80.7%	
Renal Dysfunction	22.5%	53.3%	71.9%
<i>Abnormal Creatinine $\leq 2.5 \text{ mg/dl}$</i>	<i>15.7%</i>	<i>35.3%</i>	<i>41.4%</i>
<i>Creatinine $> 2.5 \text{ mg/dl}$</i>	<i>5.0%</i>	<i>14.3%</i>	<i>18.7%</i>
<i>Chronic Dialysis</i>	<i>1.7%</i>	<i>3.0%</i>	<i>7.3%</i>
<i>Renal Transplant</i>	<i>0.1%</i>	<i>0.8%</i>	<i>4.6%</i>
Hyperlipidemia	26.2%	57.9%	
Diabetes	23.0%	39.5%	
Bronchiolitis Obliterans Syndrome	9.3%	41.1%	64.6%

Terapia immunosoppressiva in FC

- Molto farmaci sono epatotossici → **epatopatia** preesistente FC relata
- Difficoltà a mantenere triplice tp IS nei pazienti con difficile controllo dello **stato infettivo** (germi multiR)
- Difficile controllo glicemico in paziente già affetti da **CF-RD**
- Difficoltà a mantenere TIS nel corretto range in caso di **malassorbimento**
- Leucopenia: AZA/MMF + Bactrim/valganciclovir
- FDR per PTLD

Il rischio neoplastico post LuTx

Cumulative Post Transplant Malignancy Rates in Survivors

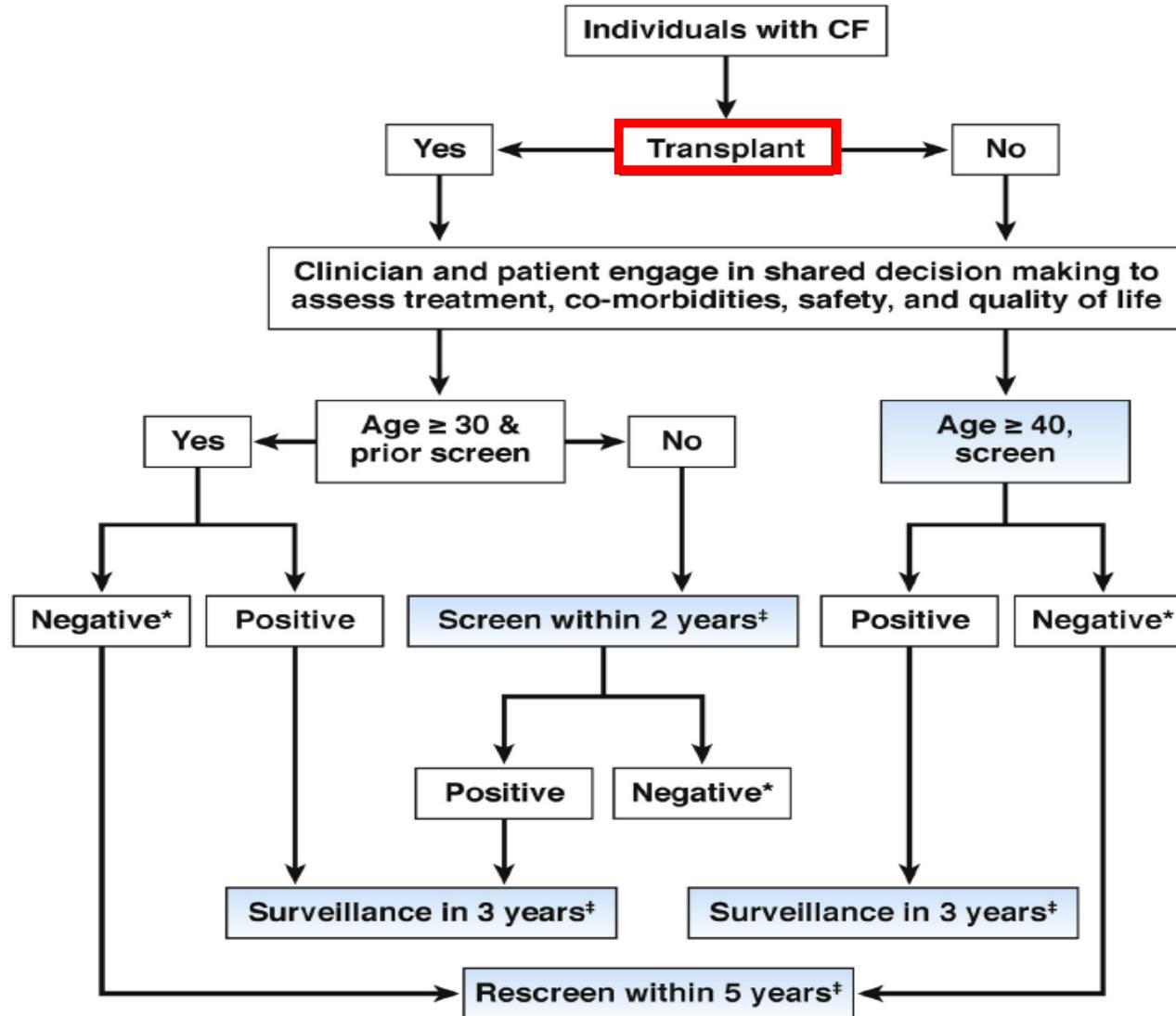
Transplants: July 2004 – June 2019

Malignancy/Type	1-Year Survivors	5-Year Survivors	10-Year Survivors
No Malignancy	23,654 (94.7%)	8,756 (80.4%)	2,297 (68.3%)
Malignancy (all types combined)	1,311 (5.3%)	2,134 (19.6%)	1,066 (31.7%)
Malignancy Type*			
Skin	456	1508	809
Lymphoma	250	163	76
Other	575	583	286
Type Not Reported	30	13	1

Other malignancies reported include: adenocarcinoma (2; 2; 1), bladder (2; 2; 1), lung (2; 2; 0), breast (1; 7; 3); prostate (0; 5; 2), cervical (1; 1; 0); and colon (0; 1; 0). Numbers in parentheses represent the number of reported cases within each time period.

* Recipients may have experienced more than one type of malignancy; therefore, the sum of individual malignancy types may be greater than the total number with malignancy.

Il rischio neoplastico post LuTx



INFEZIONI

– 4 possibili scenari ad alto rischio

- Infezioni del ricevente da parte di microorganismi acquisiti in comunità o in ospedale
- Colonizzazione delle vie aeree e/o di altri distretti (specialmente nei pazienti affetti da fibrosi cistica)
- Infezioni donor-derived
- Infezioni da germi opportunisti

Review Article

Infections after lung transplantation

Mario Nosotti¹, Paolo Tarsia^{2,3}, Letizia Corinna Morlacchi^{2,3}

Vaccinarsi è importante ... prima e dopo!



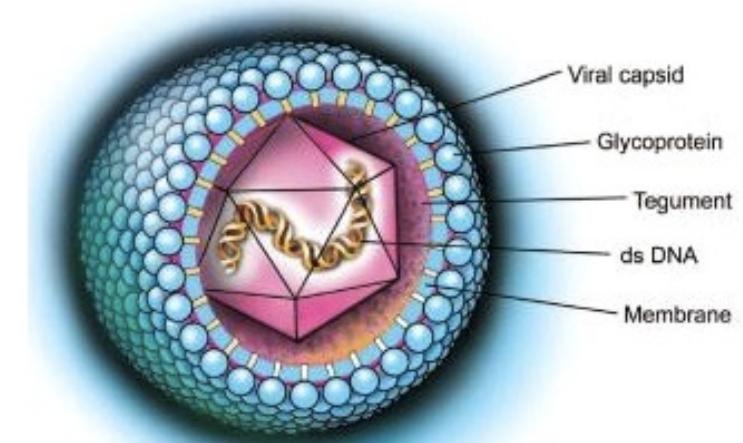
Vaccination: It works

Hey guys - I don't even
feel any rain. Why are
we doing this again? Just
put down the stupid
umbrellas - they're bad for
your arms anyway.

Infezioni - CMV

Dopo il LuTx, l'infezione da CMV può verificarsi con differenti modalità:

- Trasmissione dal graft di un donor CMV sieropositivo
- Riattivazione di un'infezione latente in un ricevente CMV sieropositivo dall'epoca pre-Tx
- Contatto con un individuo infetto da CMV (infezione attiva >> contagio)
- Trasfusione di emocomponenti provenienti da un donatore di sangue CMV sieropositivo



HCMV Human Cytomegalovirus

Infezioni - CMV

Immunity to CMV during post-transplantation prophylaxis

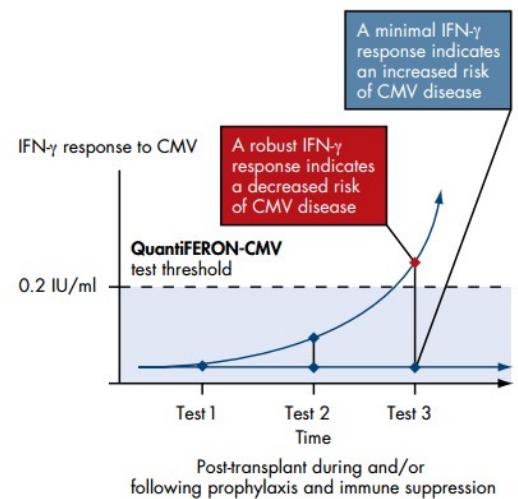
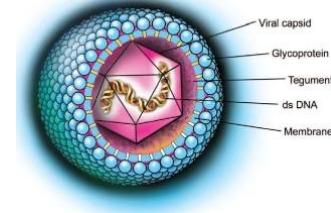


Figure 1. A theoretical model of QuantiFERON-CMV responses in a post-transplant setting during and/or following prophylaxis and immune suppression.

Review



lement of

Transplantation

Treatment and prevention of cytomegalovirus infection in heart and lung transplantation: an update

Luciano Potena , Paolo Solidoro, Filippo Patrucco & Laura Borgese

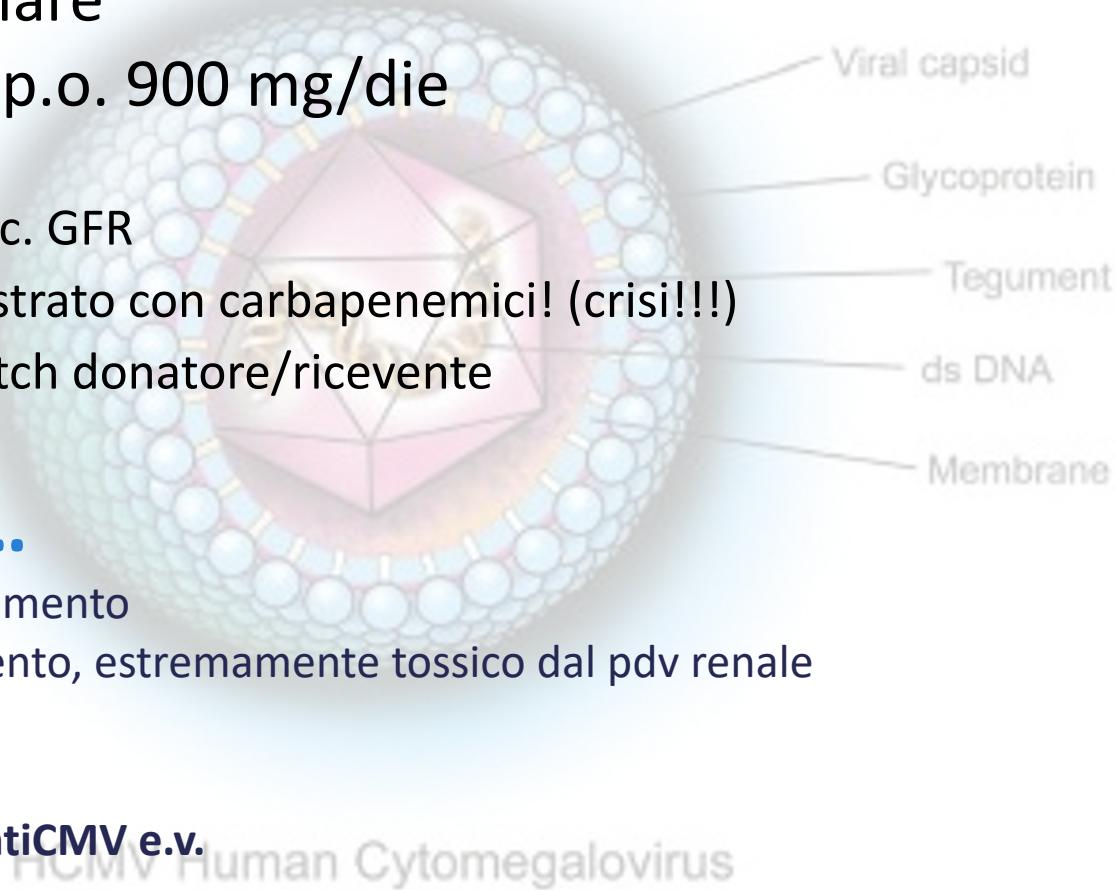
Pages 1611-1622 | Received 04 Mar 2016, Accepted 06 Jun 2016, Accepted author version posted online: 24 Jun 2016, Published online: 30 Jun 2016

 Download citation  <https://doi.org/10.1080/14656566.2016.1199684>

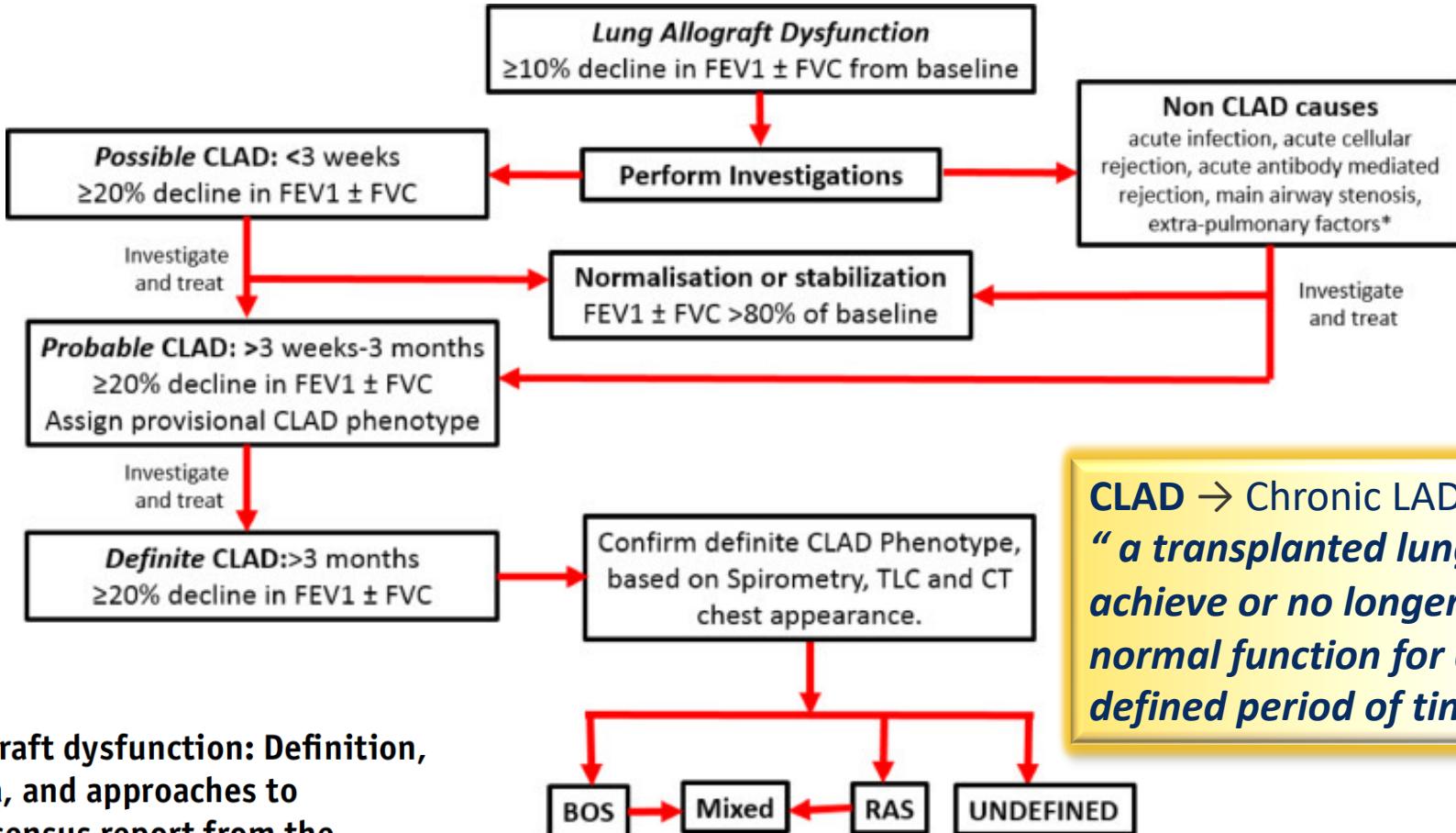
 Check for updates

Infezioni - CMV

- GANCICLOVIR e.v. 5 mg/kg ogni 12 ore; a seguire ogni 24 ore – impianto intraoculare
- VALGANCICLOVIR p.o. 900 mg/die
 - Aggiustamento dose sec. GFR
 - Attenzione se somministrato con carbapenemici! (crisi!!!)
 - Durata secondo mismatch donatore/ricevente
- **ALTERNATIVE...**
 - Foscavir → per trattamento
 - Cidofovir → trattamento, estremamente tossico dal pdv renale
 - Letermovir ???
 - Vaccino antiCMV...
 - **Immunoglobuline antiCMV e.v.**



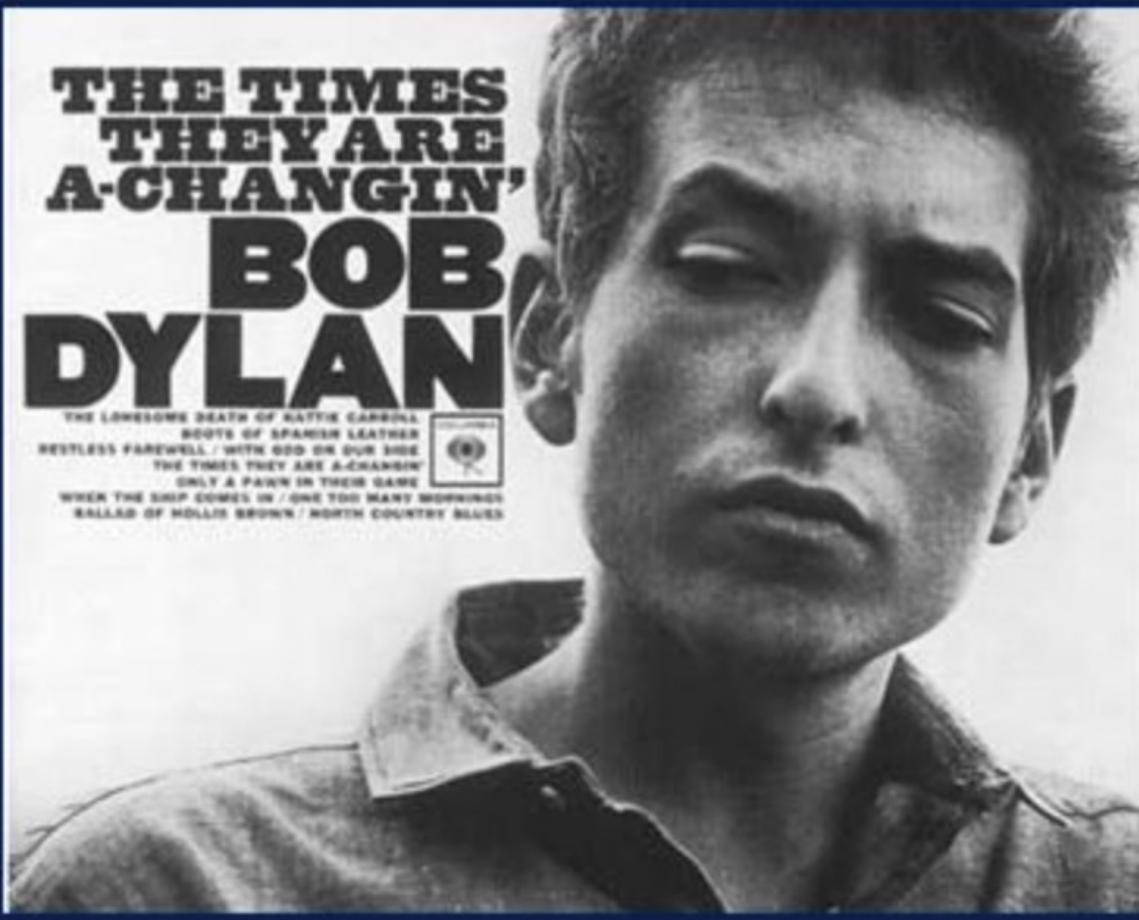
CLAD – il nuovo nome del rigetto cronico



Chronic lung allograft dysfunction: Definition, diagnostic criteria, and approaches to treatment—A consensus report from the Pulmonary Council of the ISHLT

L'arrivo dei modulatori CFTR

Prima. E anche dopo?



**THE TIMES
THEY ARE
A-CHANGIN'**
**BOB
DYLAN**

THE LOWDOWN DEATH OF KATTIE CARROLL
ROOTS OF SPANISH LEATHER
RESTLESS FAREWELL / WITH GOD OR OURSELVES
THE TIMES THEY ARE A-CHANGIN'
ONLY A PAWN IN THEIR GAME
WHEN THE SHIP COMES IN / ONE TOO MANY MORNINGSES
BALLAD OF HOLLIS BROWN / NORTH COUNTRY BLUES

The line it is drawn
The curse it is cast
The slow one now
Will later be fast
As the present now
Will later be past
The order is
Rapidly fadin'
And the first one now
Will later be last

For the times they are a-changin'.

The times they are a-changing ...



Impact of CFTR modulator use on outcomes in people with severe cystic fibrosis lung disease

Michal Shteinberg  ^{1,2} and Jennifer L. Taylor-Cousar³

The CFTR modulators IVA, LUM/IVA, TEZ/IVA and ELX/TEZ/IVA have a beneficial effect not only in individuals with mild to moderate CF, but also in individuals with advanced pulmonary disease, including candidates for lung transplantation. This beneficial impact was clearly demonstrated by randomised

Ivacaftor & LuTx

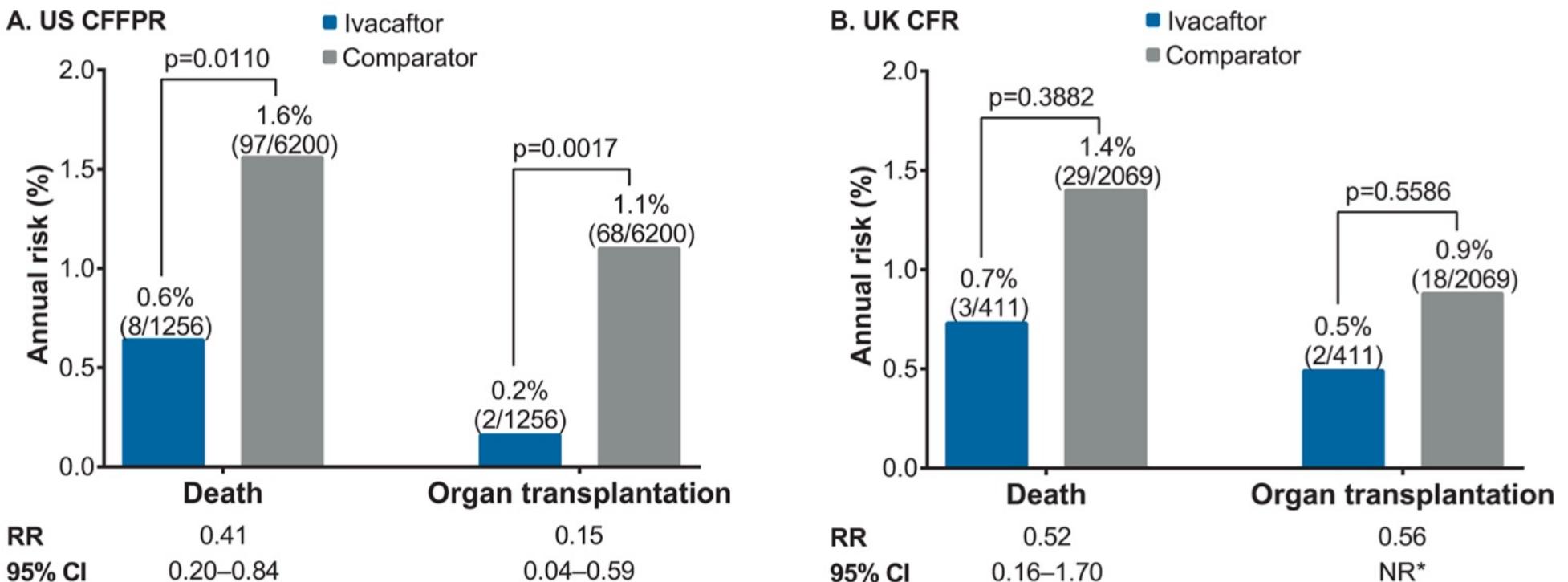


Figure 3 Death and organ transplantation, 2014 ivacaftor and comparator cohorts, (A) US CFFPR and (B) UK CFR. *Fisher's exact p values are shown when the expected value is <5 in at least one cell of the contingency table. CFFPR, Cystic Fibrosis Foundation Patient Registry; CFR, Cystic Fibrosis Registry; NR, not reported; RR, relative risk.

ELX/TEZ/IVA & LuTx

American Journal of Respiratory and Critical Care Medicine



ATS Journals

Rapid Improvement After Starting Elexacaftor-tezacaftor-ivacaftor in Patients with Cystic Fibrosis and Advanced Pulmonary Disease

Pierre-Régis Burgel ; Isabelle Durieu , Raphaël Chiron , Sophie Ramel , Isabelle Danner-Boucher , Anne Prevotat ,
Dominique Grenet , Christophe Marguet , Martine Reynaud-Gaubert , Julie Macey , Laurent Mely , Annlyse Fanton
, Sébastien Quetant , Lydie Lemonnier , Jean-Louis Paillasseur , Jennifer Da Silva , Clémence Martin ; , French
Cystic Fibrosis Reference Network study group... [Show less](#)

Burgel PR, et al. Blue J Feb 2021

ELX/TEZ/IVA & LuTx

Table 4. Numbers of lung transplantation in France by underlying disease comparing 2020 to 2018-2019.

	2018	2019	2020	Variation 2020 vs 2018/2019
Cystic fibrosis	72	80	33	-56.5 %
Pulmonary hypertension	29	22	24	-5.8 %
Pulmonary fibrosis	97	89	82	-11.8 %
COPD	127	136	90	-31.5 %
Alpha 1 antitrypsin deficiency	6	2	3	-25 %
Other diseases	27	36	33	+4.7%
All indications	358	365	265	-26.4%

COPD: chronic obstructive pulmonary disease

Candidati al LuTx per FC



Consensus document for the selection of lung transplant candidates: An update from the International Society for Heart and Lung Transplantation

**The Journal of
Heart and Lung
Transplantation**

<http://www.jhltonline.org>

Lorriana E. Leard, MD,^a Are M. Holm, MD, PhD,^b Maryam Valapour, MD, MPP,^c

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 - Worsening nutritional status despite supplementation
 - 2 exacerbations per year requiring IV antibiotics
 - Massive hemoptysis (>240 mL) requiring bronchial artery embolization
 - Pneumothorax
- FEV1 < 50% pred + rapidly declining based on PFT or progressive symptoms
- Any exacerbation requiring positive pressure ventilation

I candidati al LuTx stanno cambiando



As a consequence, patients might only develop end-stage lung disease later in life, leading to a more elderly CF population being referred for lung transplantation. A change in referral demographics in patients with additional confounding factors, secondary to a different ageing-related profile, may have an adverse impact on transplantation outcomes.

Rang C et al.; ERJ 2020; 55: 1902443

L'ultima frontiera: modulatori post SOT?



Review

Challenges in the use of highly effective modulator treatment for cystic fibrosis

Kathleen J. Ramos ^{a,*}, Joseph M. Pilewski ^b, Jennifer L. Taylor-Cousar ^c



OLT:

- PROs: improved respiratory conditions
- CONs: hepatotoxicity and therapeutic drug monitoring



LUTX

- Possible role to improve:
 - Sinus disease
 - Malnutrition
 - GERD
 - Possible prevention of GIT malignancy and PTLD (CFTR protein has possible plays a tumour suppressor role)
- CONs: hepatotoxicity and therapeutic drug monitoring

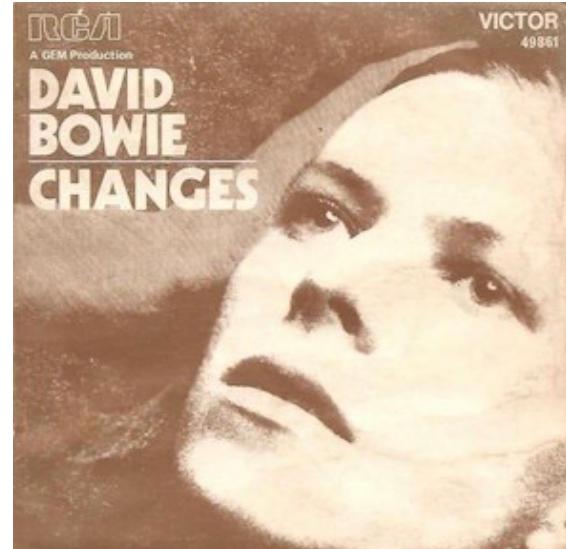
La telemedicina

Un effetto collaterale della pandemia COVID19?

L'impatto della pandemia sulla gestione dei pazienti

- Interruzione del regolare follow up c/o i Centri
- Annullamento o ritardo nell'esecuzione di procedure diagnostiche non essenziali
- Difficoltà nel raggiungere i centri di riferimento
- Impegno dei medici specializzati nelle unità dedicate al COVID19
- Paura dei pazienti nel frequentare gli ospedali

*Ch-ch-ch-ch-Changes
(Turn and face the stranger)
Ch-ch-Changes
Just gonna have to be a different one*



Nuove strategie per la cura e il follow up

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British Journal of Nursing, VOL. 30, NO. 16 | Professional

Providing post-lung transplant care during the time of COVID-19

Sara Winward, Iain Lawrie, Susan Talbot Towell, Nina Sheridan, Patricia Ging

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Tools Share

Abstract

The COVID-19 pandemic is a public health emergency of international concern. Solid organ transplant recipients have been identified as being at high risk of acquiring the virus SARS-CoV-2 and having a more severe COVID-19 disease. This article describes the experience of the National Lung Transplant Centre in Ireland in changing established care pathways for lung transplant recipients during the pandemic. The innovations which were put in place to protect this clinically vulnerable group are discussed. With the advancement of technology and remote monitoring systems available, patient-focused strategies and community-based interventions were implemented. Additional strategies have been implemented so that the new model of care can be safely maintained.

OUTLOOK · 29 JULY 2020

The coronavirus pandemic has forced rapid changes in care protocols for cystic fibrosis

 Lockdowns have accelerated the drive towards telemedicine for people with cystic fibrosis – but the system needs critical appraisal.

Jane Davies



RESEARCH Open Access

New strategies of physical activity assessment in cystic fibrosis: a pilot study

Daniela Savi^{1,2*}, Luigi Graziano¹, Barbara Giordani³, Stefano Schiavetto¹, Corrado De Vito¹, Giuseppe Migliara¹, Nicholas J. Simmonds², Paolo Palange¹ and J. Stuart Elborn⁴



RESEARCH Open Access

Validation of the portable Bluetooth® Air Next spirometer in patients with different respiratory diseases

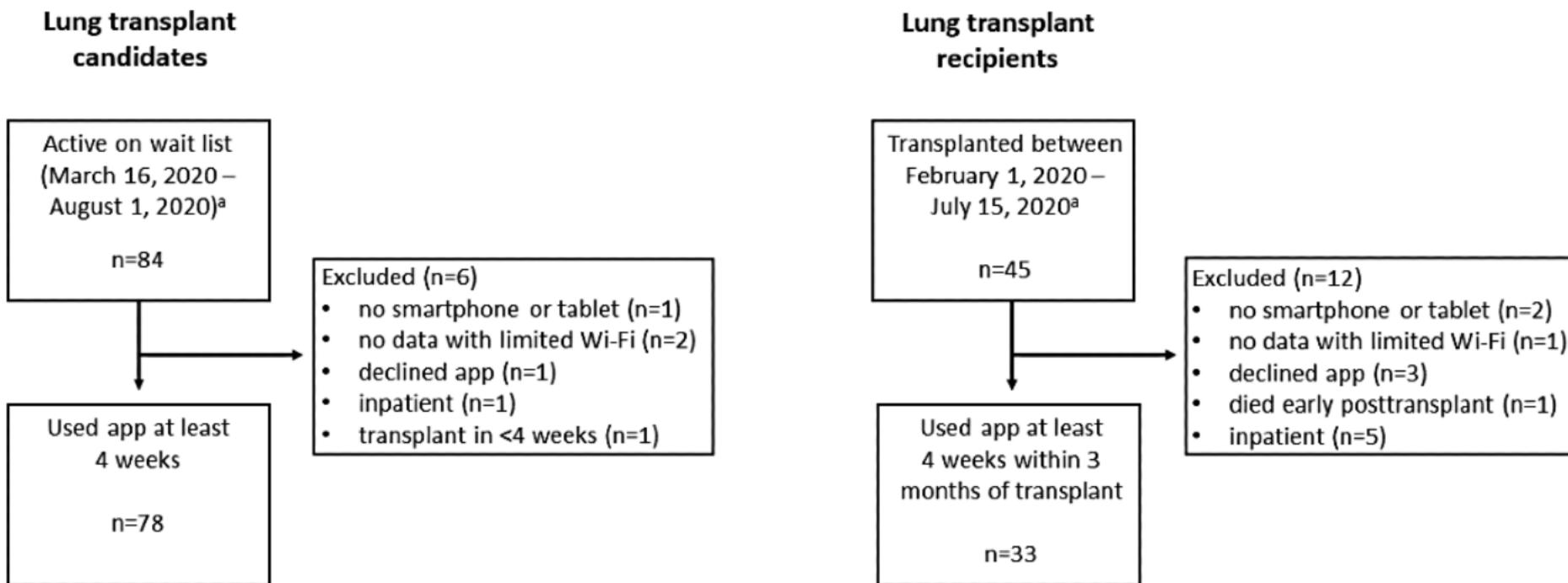
Konstantinos P. Exarchos*, Athena Gogali, Agni Sioutkou, Christos Chronis, Sofia Peristeri and Konstantinos Kostikas



Nuove strategie per la cura e il follow up

Toronto & la teleriabilitazione

Figure 1. Flow and attrition of lung transplant candidates and recipients. ^aThis time frame would permit at least 4 weeks of rehabilitation data to be entered into the app between March 16, 2020, and September 1, 2020, accounting for 2 weeks of hospitalization posttransplant.



Nuove strategie per la cura e il follow up

MHH & la telemedicina

Table 1. Patient Assessment During Video Consultation

PATIENT QUESTIONNAIRE	VITAL SIGNS
Quality of life (visual analog scale 0–10)	FEV1 home spirometry
General state of health: improved/stable/worsened	Respiratory rate
Coughing/sputum, yes/no, onset	Pulse, regular/irregular
Myalgia, malaise, yes/no, onset	Blood pressure
Runny or stuffy nose, yes/no, onset	Oxygen saturation
Sore throat, hoarseness, yes/no, onset	Body temperature
Fatigue, yes/no, onset	Body weight
Change in medication, yes/no, detail	
Physical fitness (flights of stairs without rest)	
Local laboratory report if available	
Current dose of calcineurin inhibitor	

Summary of medical history and vital signs gathered systematically during video consultation.

FEV1, forced expiratory volume in one second.

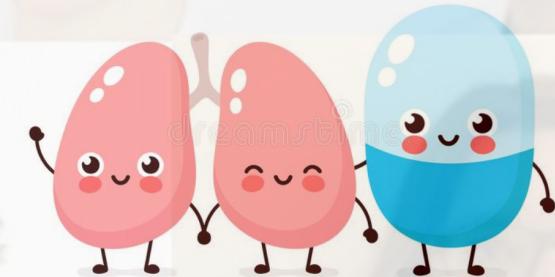
Table 3. Summary of Video Consultations

COMPLETED VCS (n=75)	
Patient device, n (%)	
Laptop/computer	59 (79)
Tablet	3 (4)
Smartphone	13 (17)
Duration of contact, minutes	
Median (25, 75 quartile)	27 (19, 34)
Reason for consultation, n (%)	
Routine surveillance	24 (32)
Follow-up	27 (36)
Clinical indication	24 (32)
Availability of diagnostic parameters, n (%)	
Pulse oximetry	60 (80)
Respiratory rate	67 (89)
FEV1	66 (88)
Body temperature	57 (76)
Pulse	59 (79)
Patient diary	33 (44)
Outcome of consultation, n (%)	
Medication changed	27 (36)
Hospitalization	2 (3)
OSV	3 (4)
Visit local practitioner	14 (19)
No specific measures	28 (38)
Reported technical problems during consultation, n (%)	
e.g., poor sound and frozen picture	11 (15)
FEV1, forced expiratory one second capacity.	

“Every accomplishment starts with a decision to try.” JFK



Q&A



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