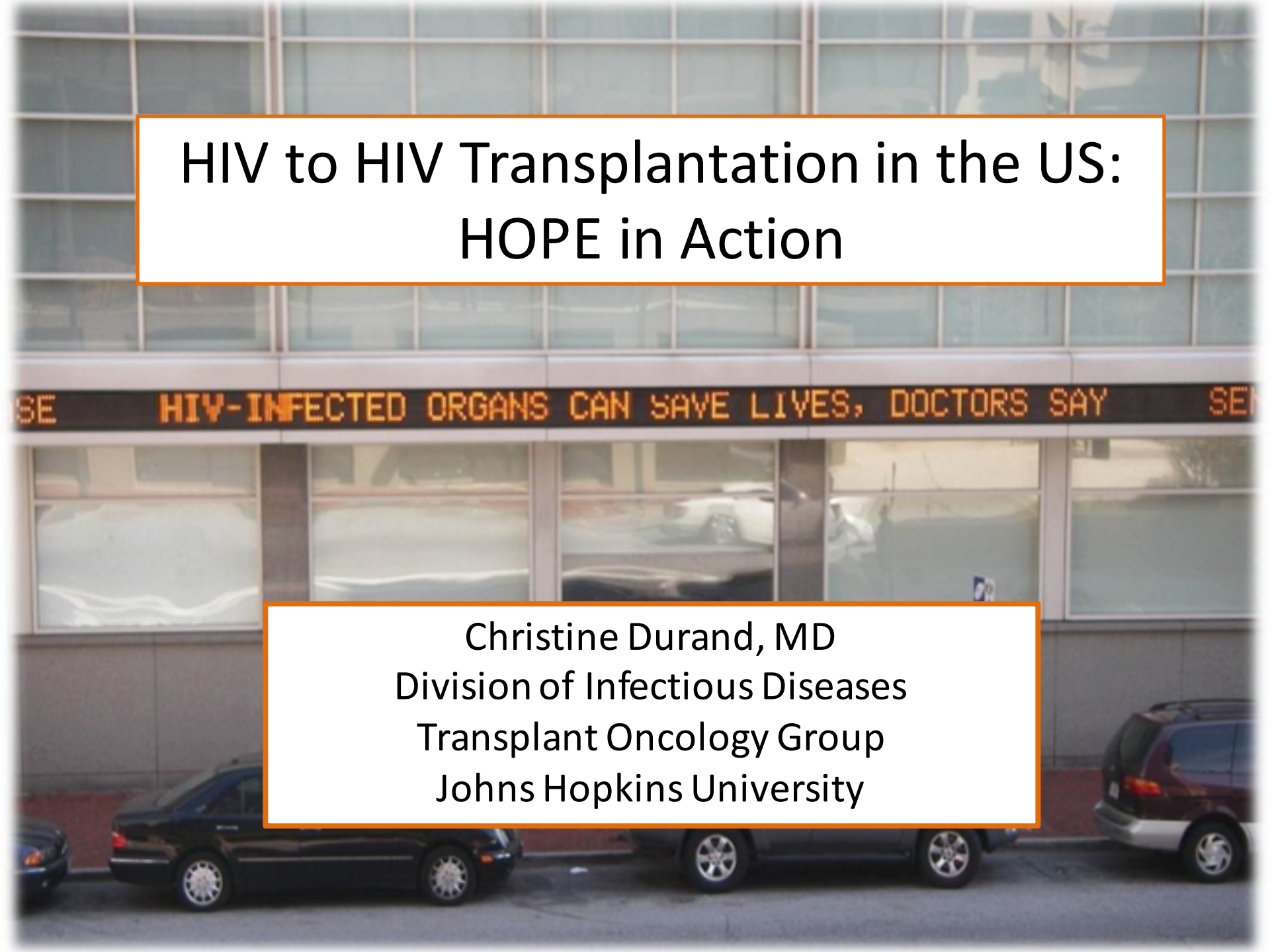


The background image shows a multi-story building with a large window. A digital sign is visible on the building, displaying the text "HIV-INFECTED ORGANS CAN SAVE LIVES, DOCTORS SAY". The sign is orange and black. The building has a modern architectural style with large glass windows. The text "HIV to HIV Transplantation in the US: HOPE in Action" is overlaid on the image in a white box with an orange border.

HIV to HIV Transplantation in the US: HOPE in Action

Christine Durand, MD
Division of Infectious Diseases
Transplant Oncology Group
Johns Hopkins University

Disclosures

- Advisory boards and research grants to JHU from Merck Pharmaceuticals, Gilead Sciences, Bristol Meyers Squibb

Funding:

- Living Legacy Foundation, Maryland
- JHU, Discovery Fund
- JHU Center for AIDS Research 1P30AI094189
- NIH R01-AI120938 (Tobian)
- NIH R34-AI123023 (Durand)
- NIH K23-CA177321 (Durand)

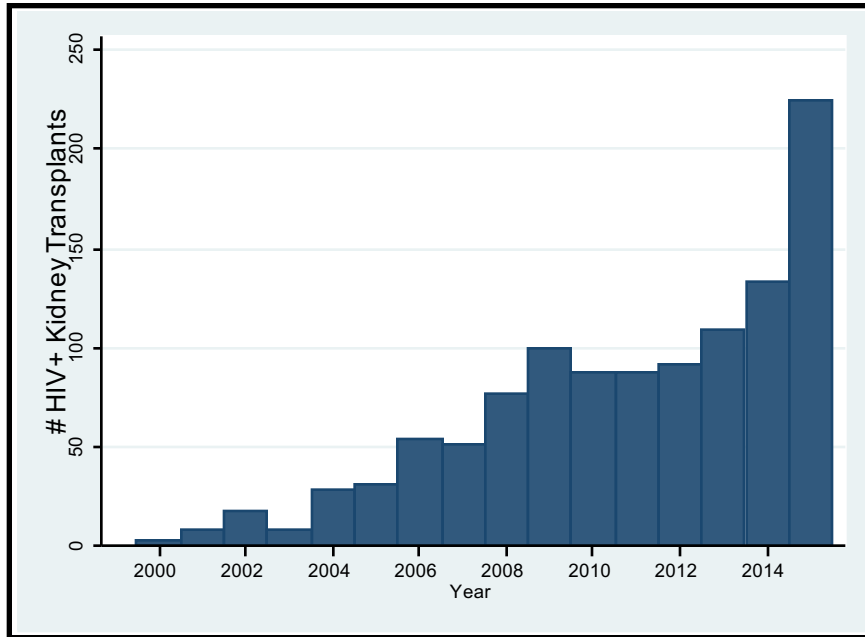
Outline

- Motivation for HIV+ donor organs (D+) for HIV+ recipients (R+) in need of transplant
- Knowledge gaps, risks
- Clinical trials
 - Parent/pilot trial – early outcomes
 - NIH funded multicenter kidney trial

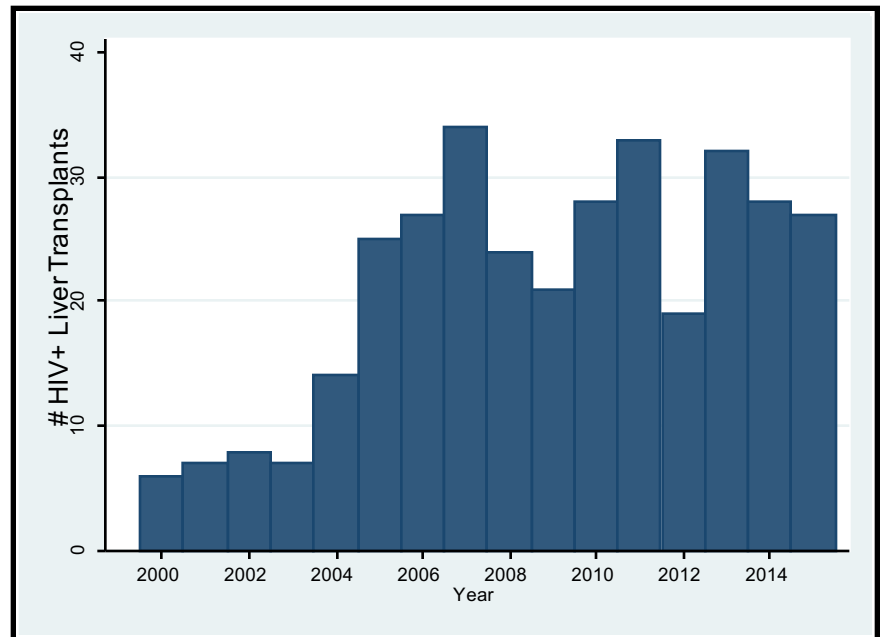
Motivation/Significance

- Prevalence of end stage kidney and liver disease in HIV+ increasing (Lucas CID 2014; Smith Lancet 2014)
- HIV+ candidates have higher waitlist mortality compared to HIV- (Trullas AIDS 2011, Ragni Liver Transpl 2005)
- Excellent outcomes with HIV- organs for HIV recipients (HIV D-/R+) (Roland AIDS 2016, Locke JASN 2015; Locke Transplantation 2016)

US HIV D-/R+ transplant volumes



HIV+ Kidney Transplants



HIV+ Liver Transplants

S Africa: HIV D+/R+ Kidney Transplant

The NEW ENGLAND JOURNAL of MEDICINE

Renal Transplantation between HIV-Positive Donors and Recipients



Table 1. Clinical Characteristics of HIV-Positive Recipients of a Transplant from an HIV-Positive Donor.*

Characteristic	Patient 1	Patient 2	Patient 3	Patient 4
Age (yr)	47	56	37	29
Sex	Male	Male	Male	Female
Before transplantation				
Diagnosis on renal biopsy	HIV-associated nephropathy	HIV-associated nephropathy and hypertensive nephropathy	Malignant hypertension	HIV-associated nephropathy
Creatinine ($\mu\text{mol/liter}$)	678	582	1712	725
CD4 count (cells/mm ³)	288	258	132	147
HIV viral load (copies/ml)	<50	<50	<50	<50
Antiretroviral regimen	Tenofovir, lamivudine, and lopinavir–ritonavir	Stavudine, lamivudine, and efavirenz	Stavudine, lamivudine, and nevirapine	Zidovudine, lamivudine, and nevirapine

Muller et al, NEJM 2010; 362: 2336-7

National Organ Transplant Act, 1984/88

42 U.S.C. 274(b) Sect 372(b):

“requires the OPTN to adopt and use standards for preventing the acquisition of organs from individuals known to be infected with HIV.”

Estimating the Potential Pool of HIV-Infected Deceased Organ Donors in the United States

**B. J. Boyarsky^a, E. C. Hall^{a,b}, A. L. Singer^a,
R. A. Montgomery^a, K. A. Gebo^{c,d,e}
and D. L. Segev^{a,d,*}**

^aDepartment of Surgery, Johns Hopkins School of Medicine, Baltimore, MD

^bDepartment of Surgery, Georgetown University School of Medicine, Washington, DC

^cDepartment of Medicine, Johns Hopkins University School of Medicine, Baltimore, MD

^dDepartment of Epidemiology, Johns Hopkins School of Public Health, Baltimore, MD

^eHIV Research Network, Baltimore, MD

*Corresponding author: Dorry L. Segev, dorry@jhmi.edu

- 2000-2008
- 2 national registries (NIS, HIVRN)
- Excluded those with missing data and medical contraindication
- **500-600 donors per year**

HOPE (HIV Organ Policy Equity) Act

November 21, 2013



HOPE Act Mandates

- Directs the Secretary to revise current regulations (specifically, 42 CFR 121.6)
- June 2015
- Directs Secretary to publish research criteria relating to HIV+ to HIV+ transplant
- November 2015
- Requires the OPTN to revise standards for the acquisition and transportation of donated HIV+ organs
- November 2015

HIV D+/R+: research only for now

The Hope Act states “not later than 4 years after the date of enactment and annually thereafter, the Secretary shall review the results of scientific research in conjunction with the OPTN to determine whether the results warrant revision of the standards of quality.”

- IRB approved protocol
- Open Variance from OPTN



Learn if the use of HIV+ deceased donors is *safe*
and effective in the US

Minireview

Challenges and Clinical Decision-Making in HIV-to-HIV Transplantation: Insights From the HIV Literature

**B. J. Boyarsky¹, C. M. Durand²,
F. J. Palella Jr.³ and D. L. Segev^{1,4,*}**

¹Department of Surgery, Johns Hopkins School of Medicine, Baltimore, MD

²Department of Medicine, Johns Hopkins School of Medicine, Baltimore, MD

³Department of Medicine, Feinberg School of Medicine, Northwestern University Chicago, Chicago, IL

⁴Department of Epidemiology, Johns Hopkins School of Public Health, Baltimore, MD

* Corresponding author: Dorry Segev, dorry@jhmi.edu

Biologic risks

- HIV superinfection
- HIV nephropathy
- Donor derived infections
- Rejection

Key differences: US and S Africa

	South Africa	US
Population	53 million	316 million
Persons living with HIV	5.6 million	1.1 million
HIV+ prevalence	17.8%	0.6%
Predominant subtype	C	B
Annual HIV+ deaths	310,000	17,000
Transmitted drug resistance	< 5%	10-18%
Transplant wait list	4300	123,992
Transplant per year	229	16,896

**Lower risk of HIV superinfection
and drug resistance**

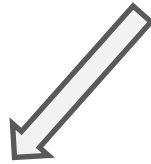
**Higher risk of HIV superinfection
and drug resistance**



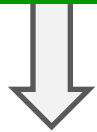
- **First line ART regimen**
- **R5 tropic virus**
- **Potential for PI Ritonavir -sparing regimen**
- **Higher CD4+ T cell count**

- **Second-line ART**
- **X4 tropic virus**
- **History of drug resistance**
- **Requires PI/Ritonavir-based regimen**
- **Lower CD4+ T cell count**

Standard donor criteria
plus positive HIV ELISA
and/or NAT*



Undetectable HIV
(RNA <200 c/mL)



Antiretroviral
treatment

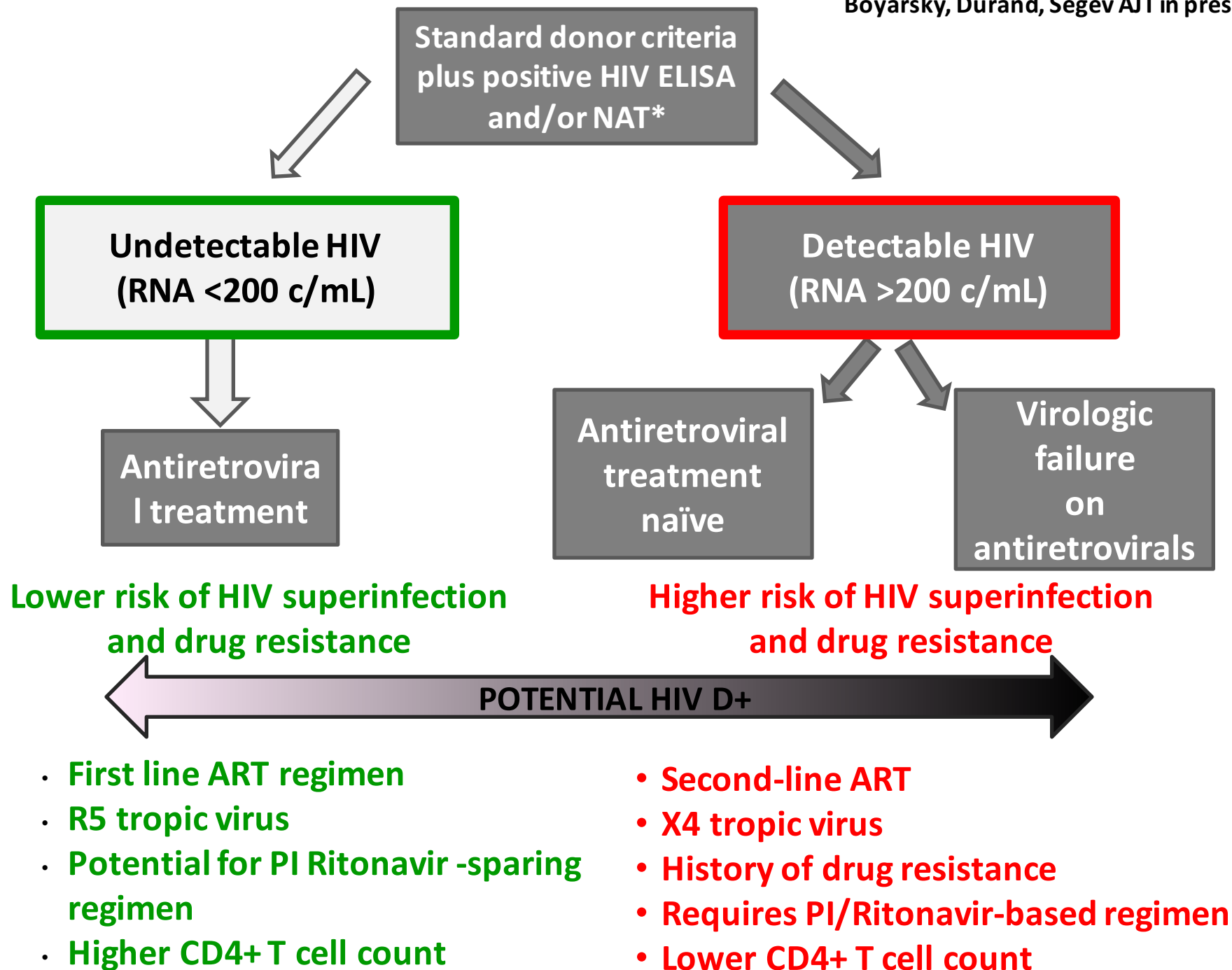
Lower risk of HIV superinfection
and drug resistance

Higher risk of HIV superinfection
and drug resistance



- First line ART regimen
- R5 tropic virus
- Potential for PI Ritonavir -sparing regimen
- Higher CD4+ T cell count

- Second-line ART
- X4 tropic virus
- History of drug resistance
- Requires PI/Ritonavir-based regimen
- Lower CD4+ T cell count



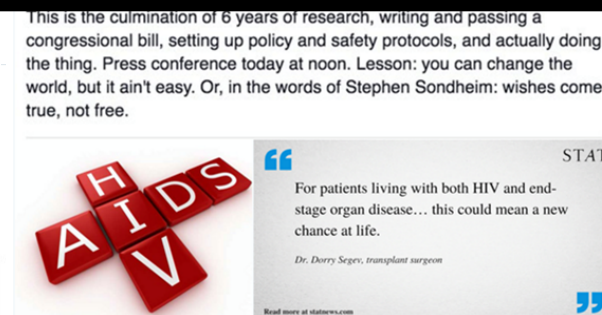
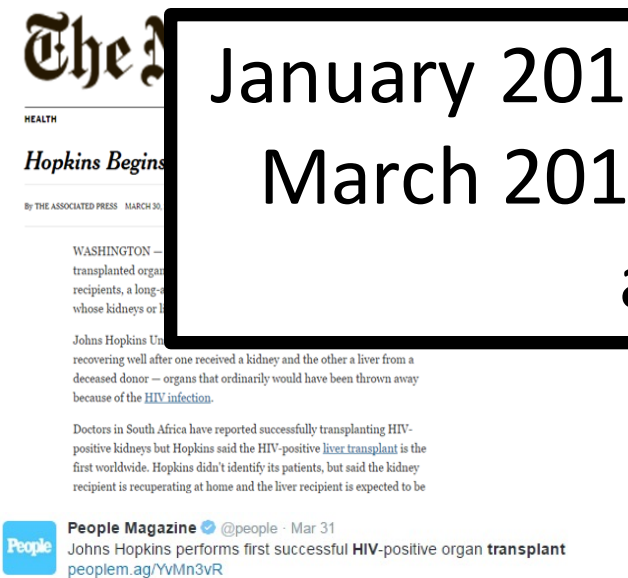


Learn if the use of HIV+ deceased donors in the is
safe and effective in the US

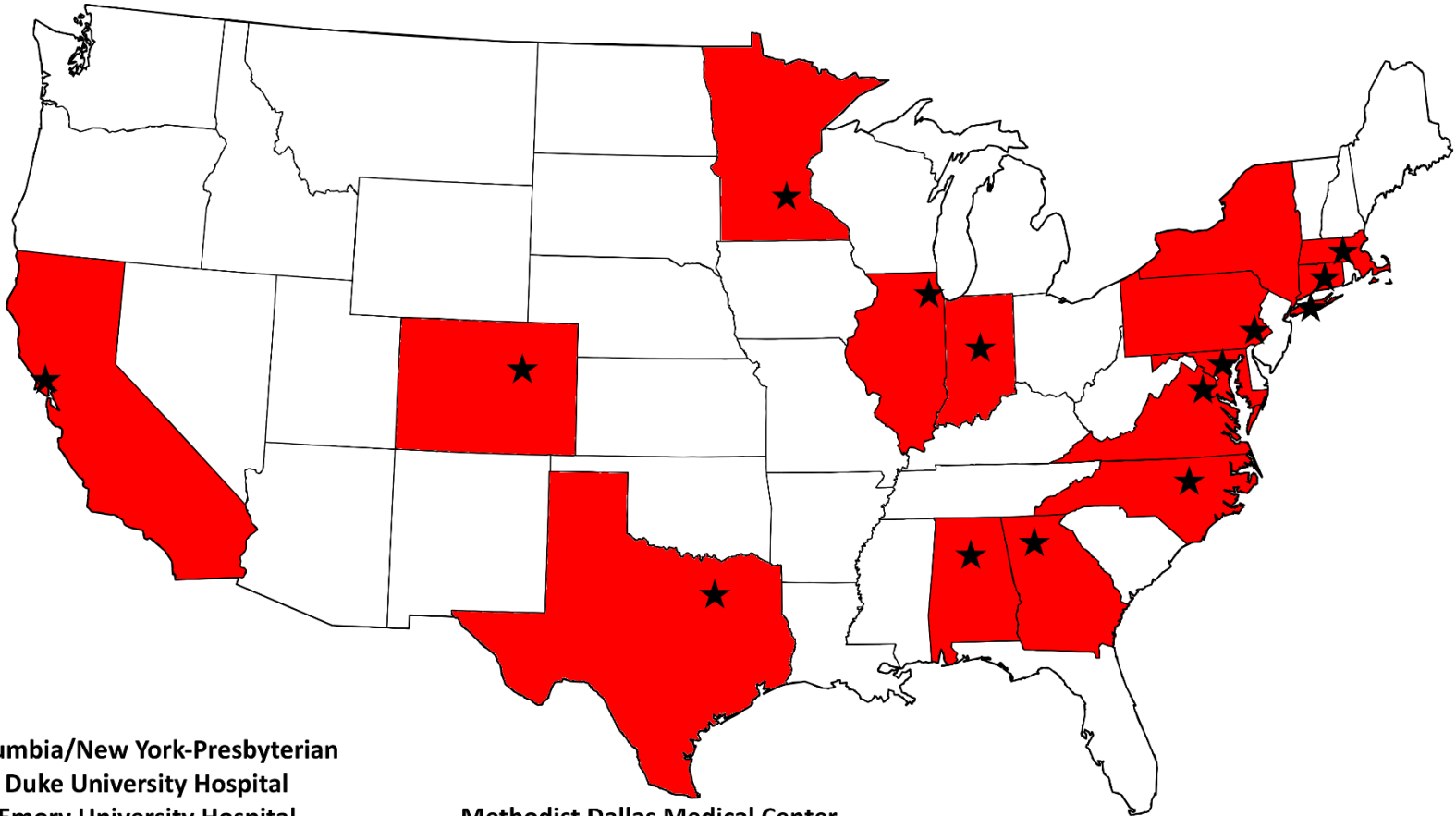
- Parent/pilot transplant trial



January 2016, Pilot protocol (NCT02602262)
March 2016, First in US HIV D+/R+ kidney
and liver transplants



UNOS variance – 19 active centers

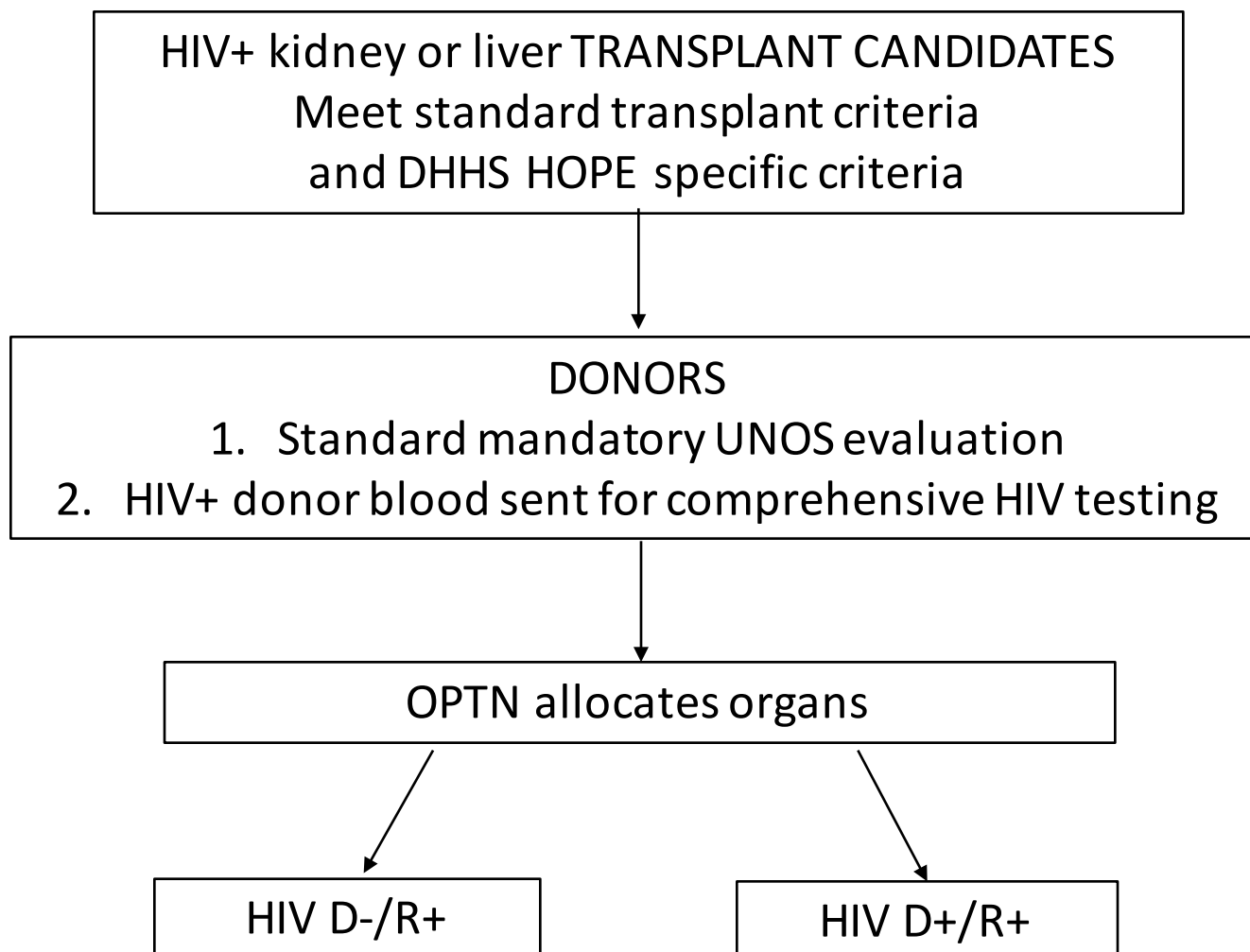


Columbia/New York-Presbyterian
Duke University Hospital
Emory University Hospital
Georgetown University Medical Center
Hahnemann University Hospital
Indiana University Health
The Johns Hopkins Hospital
Massachusetts General Hospital

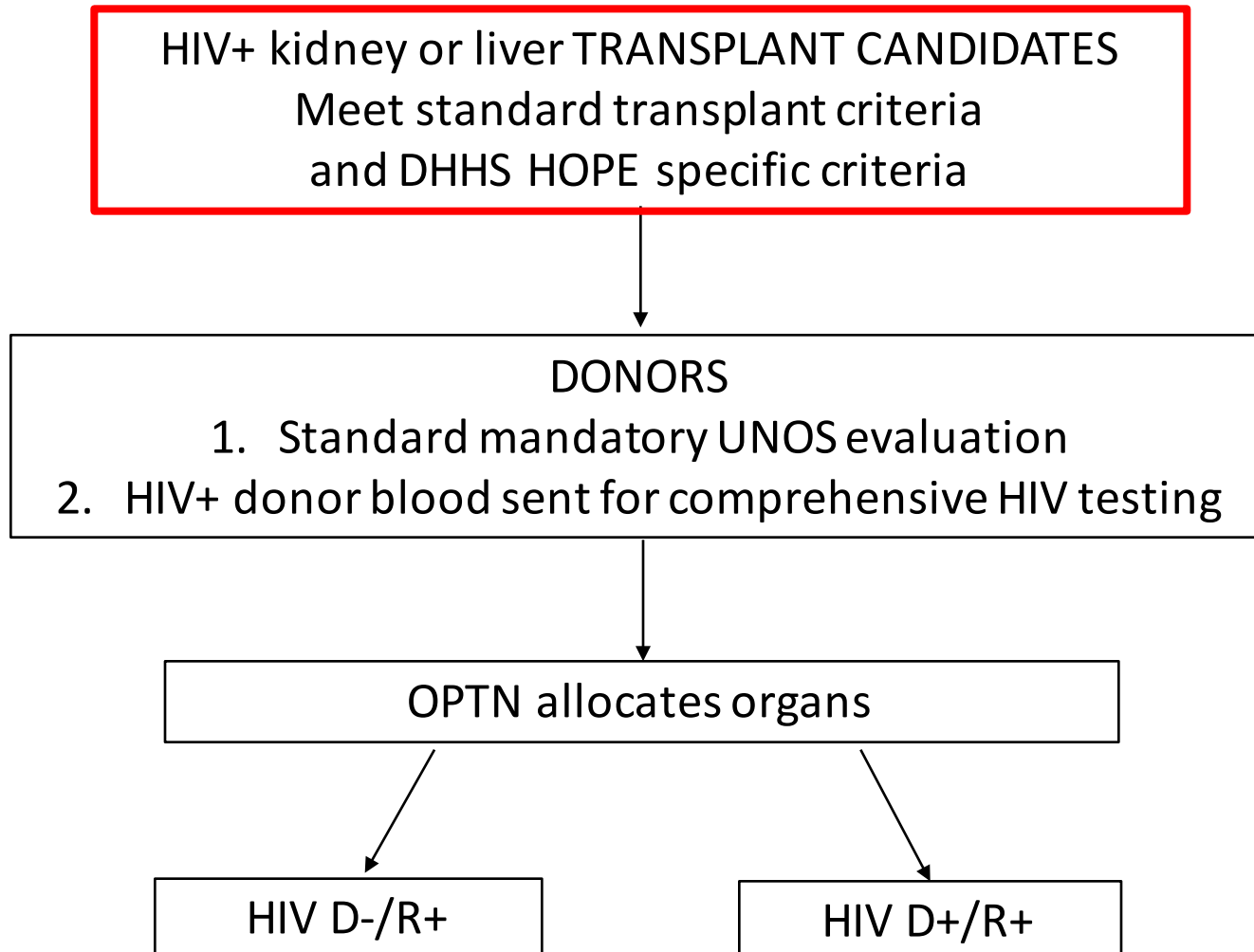
Methodist Dallas Medical Center
Montefiore Medical Center
Mount Sinai Medical Center
Rush University Medical Center
University of Alabama at Birmingham
University of California, San Francisco

University of Colorado, Denver
University of Maryland Medical System
University of Minnesota
Virginia Commonwealth University Medical Center
Yale New Haven Hospital

Methods: Trial Design



Methods: Trial Design

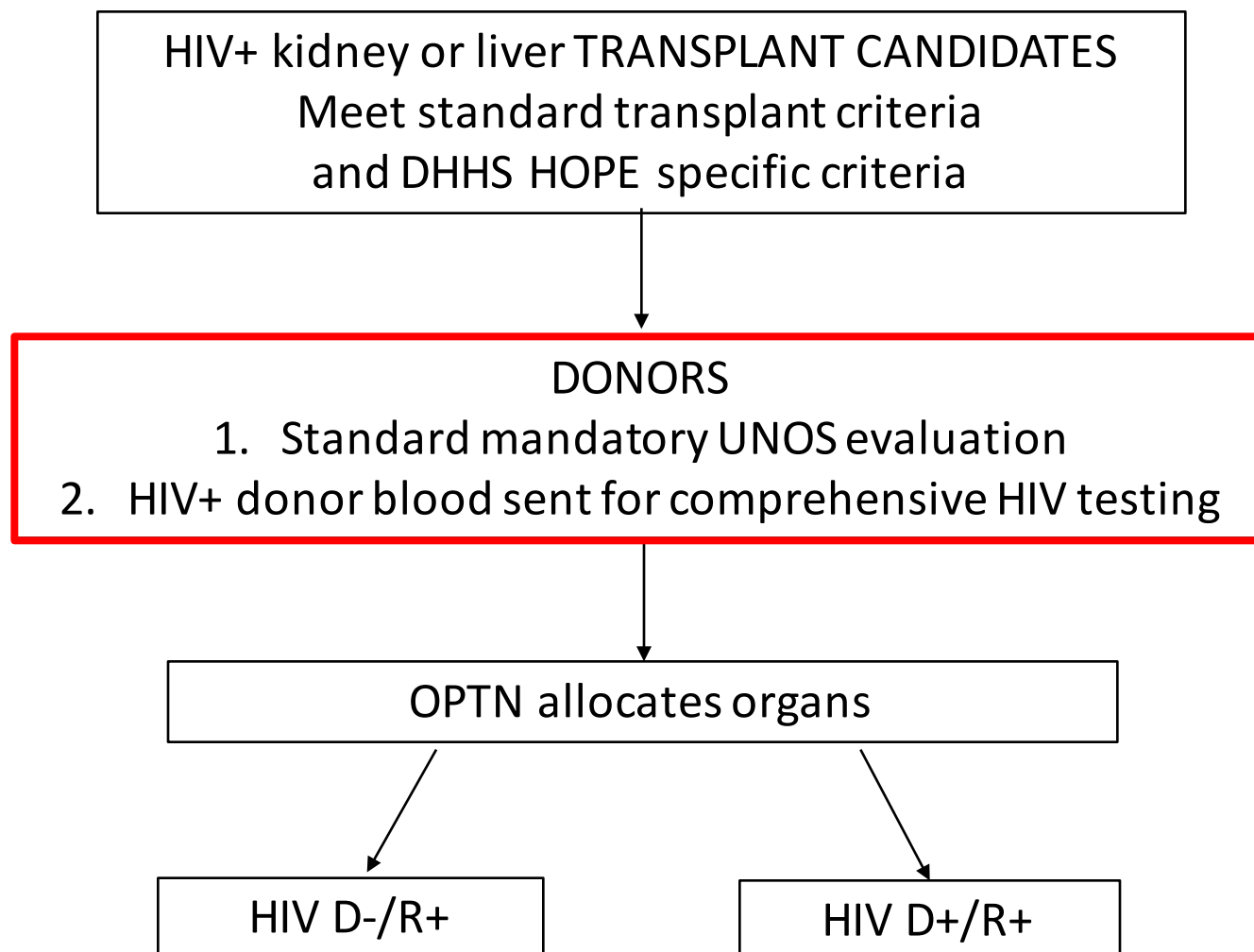


HIV+ Candidate Inclusion Criteria



- No active opportunistic infections or cancer
- Kidney CD4 > 200 cells
 - HIV RNA suppressed on effective ART
- Liver CD4 > 100 cells
 - If unable to tolerate ART, an effective regimen is anticipated post-transplant

Methods: Trial Design



HIV D+ Inclusion Criteria



- No active opportunistic infections or cancer
- Any HIV VL or CD4 count is allowed but study team must describe effective post-transplant antiretroviral regimen for the recipient

HIV D- eligibility

- Per transplant center study team
- Including donors with no history of HIV who test positive during donor screening – suspected false positives
- Historically organs from these donors were not recovered now allowed under HOPE Act

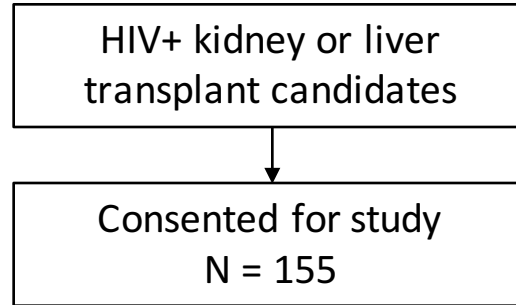
Suspected HIV False Positive Donor: Definition

1. Clinical suspicion of false positive: no history of HIV per chart review and Donor Risk Assessment Interview
2. Discordant HIV Ab and NAT on standard OPTN screening*

*Must fulfill both criteria, discordant Ab and NAT alone does not indicate a false positive

- HIV+ donors on ART can be HIV Ab+/NAT- (treatment suppresses HIV nucleic acid)
- HIV+ donors acutely infected can be HIV Ab- and NAT+ (before seroconversion)

Trial Enrollment



HOPE
IN ACTION →

Consented patients, by site

Characteristic	Johns Hopkins	Mount Sinai	Yale University	Indiana University	Emory University	University of Alabama
	N = 42	N = 94	N = 4	N = 2	N = 9	N = 1
Age, yr, Med (IQR)	52 (42-57)	53 (47-58)	58 (51-61)	52 (45-59)	50 (37-51)	33 (33-33)
Female; N (%)	11 (33%)	26 (31%)	1 (25%)	0 (0%)	1 (11%)	0 (0%)
Race; N (%)						
White	6 (15%)	29 (31%)	1 (25%)	0 (0%)	0 (0%)	0 (0%)
Black/African American	35 (83%)	62 (66%)	3 (75%)	2 (100%)	9 (100%)	1 (100%)
Asian	0 (0%)	2 (2%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)
Native American	0 (0%)	1 (1%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)
Missing	1 (3%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)
Ethnicity; N (%)						
Hispanic	4 (9%)	10 (12%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)
Non-Hispanic	28 (85%)	74 (88%)	4 (100%)	2 (100%)	9 (100%)	1 (100%)
Missing	3 (9%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)
Organ; N (%)						
Kidney only	32 (76%)	84 (87%)	4 (100%)	2 (100%)	9 (100%)	1 (100%)
Liver only	7 (17%)	12 (13%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)
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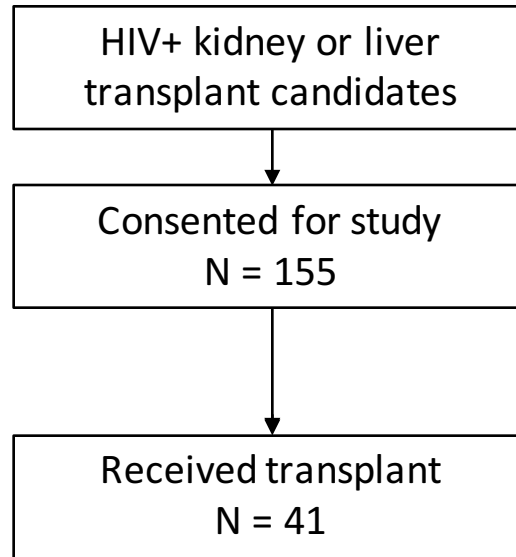
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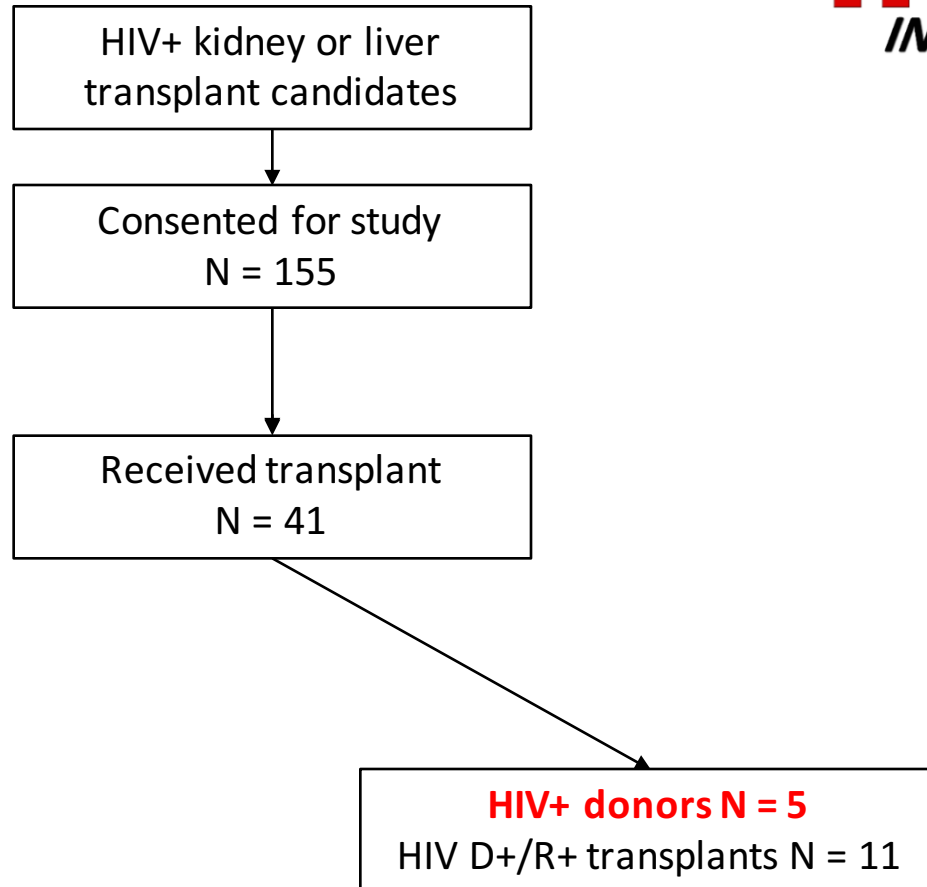
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Trial Enrollment

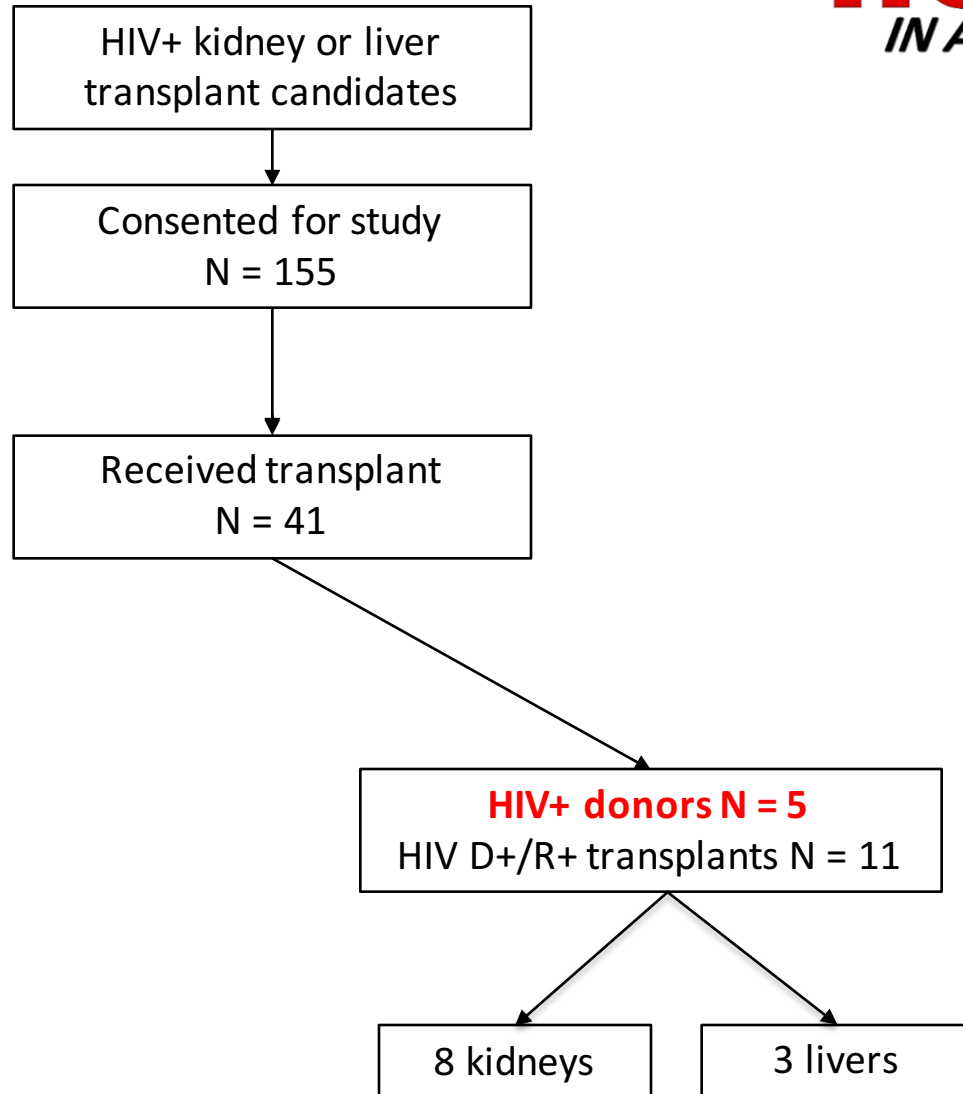


Trial Enrollment

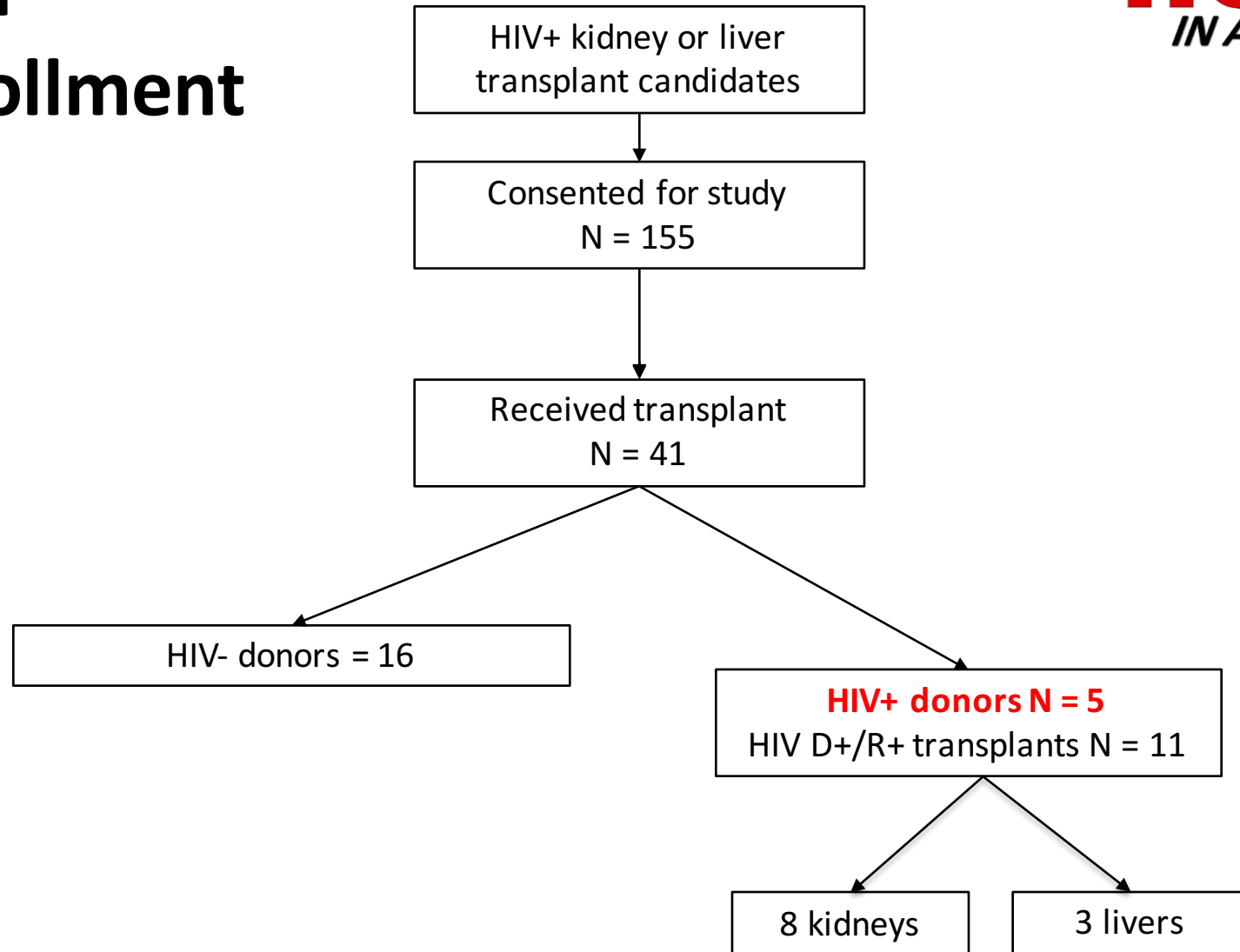


Trial Enrollment

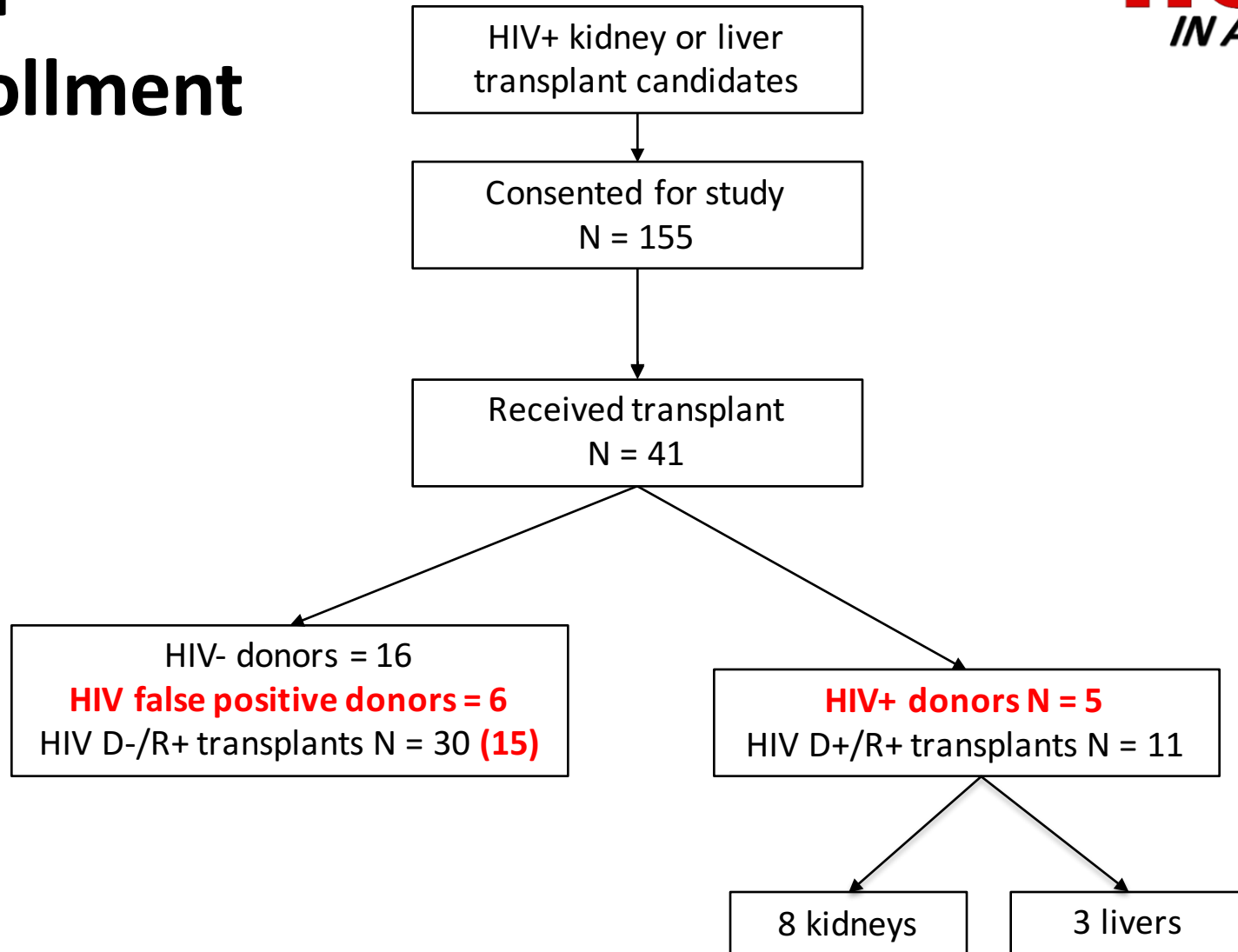
HOPE
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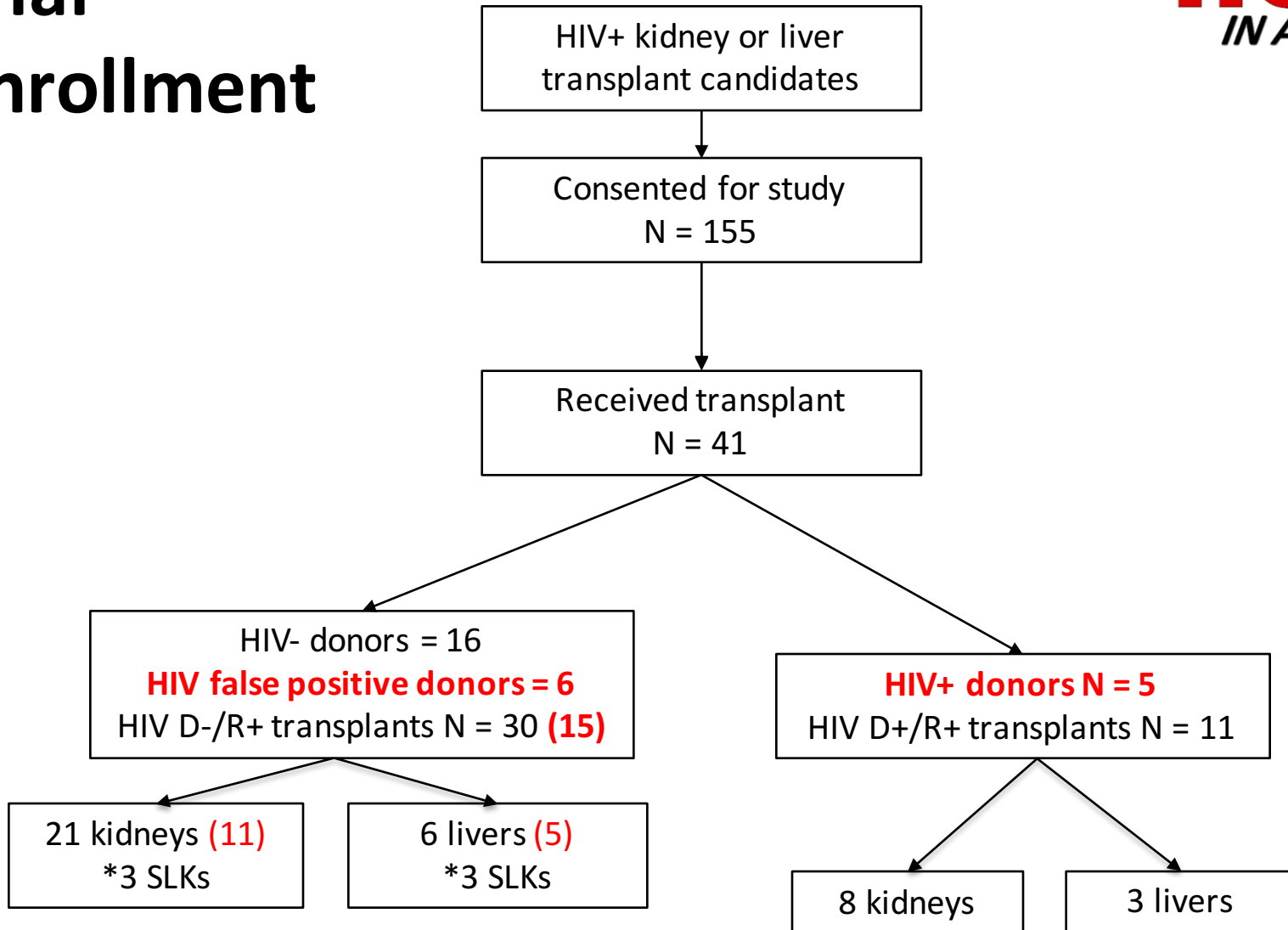
Trial Enrollment



Trial Enrollment



Trial Enrollment



Donor characteristics

Characteristic	HIV- donor (N=22)	HIV+ donor (N=5)
Age, yr, Med (IQR)	31 (25-39)	31 (28-35)
Female; N (%)	7 (32%)	3 (60%)
Race/Ethnicity; N (%)		
White	8 (36%)	1 (20%)
Black/African American	10 (45%)	3 (60%)
Asian	1 (5%)	0 (0%)
Hispanic	3 (14%)	1 (20%)
KDPI; Med (IQR)*	44 (23-54)	30 (28-54)
BMI, kg/m ² ; Med (IQR)	25 (21-27)	23 (23-26)

Donor characteristics

Characteristic	HIV- donor (N=22)	HIV+ donor (N=5)
Age, yr, Med (IQR)	31 (25-39)	31 (28-35)
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Race/Ethnicity; N (%)		
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KDPI; Med (IQR)*	44 (23-54)	30 (28-54)
BMI, kg/m ² ; Med (IQR)	25 (21-27)	23 (23-26)

HIV+ donor characteristics (N=5)

Characteristic	HIV+ donor (N=5)
CD4+ T-cell count, cells/mm ³	980 (476-1533)
HIV plasma RNA, copies/mL; N (%)	
Detectable plasma RNA (140 & 2200 c/mL)	2 (40%)
Undetectable (<20 copies/mL)	3 (60%)
Hepatitis C Antibody Positive; N (%)	0 (0%)
Hepatitis C NAT Positive; N (%)	0 (0%)
Hepatitis B Surface Ag Positive; N (%)	0 (0%)
Hepatitis B Core Antibody Positive; N (%)	0 (0%)
ART Resistance by Clinical Review; N (%)	1 (20%)

HIV+ donor characteristics (N=5)

Characteristic	HIV+ donor (N=5)
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Hepatitis B Surface Ag Positive; N (%)	0 (0%)
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ART Resistance by Clinical Review; N (%)	1 (20%)

HIV D+/R+ ART

Donor	Donor ART	Recipient	Recipient Post-transplant ART*	Organ
1	3TC DTG RTV DRV	1	FTC TAF DTG MVC	Liver
		2	FTC TAF DTG MVC	Kidney
2	ATV RTV TDF FTC	3	RAL ABC 3TC	Kidney
		4	ABC 3TC DTG	Kidney
3	FTC+TAF ATV/c	5	MVC DTG DRV	Kidney
4	EVG/c/TAF/FTC	6	3TC AZT RAL	Kidney
		7	FTC TDF EFV	Kidney
		8	DTG FTC TAF	Liver
5	ATV DTG RTV TDF+FTC	9	FTC RPV DTG TDF	Kidney
		10	DTG FTC/RPV/TAF	Kidney
		11	DRV DTG FTC/TAF	Liver

*FTC = emtricitabine; TAF = tenofovir alafenamide fumarate; DTG = dolutegravir; MVC = maraviroc; RAL = raltegravir; ABC = abacavir; 3TC = lamivudine; DRV = darunavir; AZT = zidovudine; TDF = tenofovir disoproxil fumarate; EFV = efavirenz; ATV/c = atazanavir; EVG = elvitegravir; c = cobicistat

Recipient ART Changes

Donor	Recipient	ART changed	Reason for Change
1	1	Yes	-Stopped ABC due to donor HLA B5701 status associated with hypersensitivity reaction -Changed RAL to DTG for less pill burden -Stopped ETV due to interaction with DTG
	2	No	N/A
2	3	No	N/A
	4	No	N/A
3	5	No	N/A
4	6	No	N/A
	7	No	N/A
	8	No	N/A
5	9	Yes	Clinical history with reported resistance to multiple classes of medications (M41L, V188L, M184V, T215F, K103N)
	10	Yes	
	11	Yes	

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1	1	Yes	-Stopped ABC due to donor HLA B5701 status associated with hypersensitivity reaction -Changed RAL to DTG for less pill burden -Stopped ETV due to interaction with DTG
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	7	No	N/A
	8	No	N/A
5	9	Yes	Clinical history with reported resistance to multiple classes of medications (M41L, V188L, M184V, T215F, K103N)
	10	Yes	
	11	Yes	

HIV D+/R+ ART

Donor	Donor ART	Recipient	Recipient Post-transplant ART*	Organ
1	3TC DTG RTV DRV	1	FTC TAF DTG MVC	Liver
		2	FTC TAF DTG MVC	Kidney
2	ATV RTV TDF FTC	3	RAL ABC 3TC	Kidney
		4	ABC 3TC DTG	Kidney
3	FTC+TAF ATV/c	5	MVC DTG DRV	Kidney
4	EVG/c/TAF/FTC	6	3TC AZT RAL	Kidney
		7	FTC TDF EFV	Kidney
		8	DTG FTC TAF	Liver
5	ATV DTG RTV TDF+FTC	9	DTG RPV TAF FTC	Kidney
		10	DTG RPV TAF FTC	Kidney
		11	DRV/r DTG TAF FTC	Liver

*FTC = emtricitabine; TAF = tenofovir alafenamide fumarate; DTG = dolutegravir; MVC = maraviroc; RAL = raltegravir; ABC = abacavir; 3TC = lamivudine; DRV = darunavir; AZT = zidovudine; TDF = tenofovir disoproxil fumarate; EFV = efavirenz; ATV/c = atazanavir; EVG = elvitegravir; c = cobicistat

HIV- donor characteristics

Characteristic	HIV- donor (N=22)
Suspected False Positive Cases	6 (27%)
HIV Ab+/NAT negative	4 (18%)
HIV Ab-/NAT positive	2 (9%)
Hepatitis C Antibody Positive; N (%)	5 (23%)
Hepatitis C NAT Positive; N (%)	6 (27%)
Hepatitis B Surface Ag Positive; N (%)	1 (5%)
Hepatitis B Core Antibody Positive; N (%)	2 (9%)

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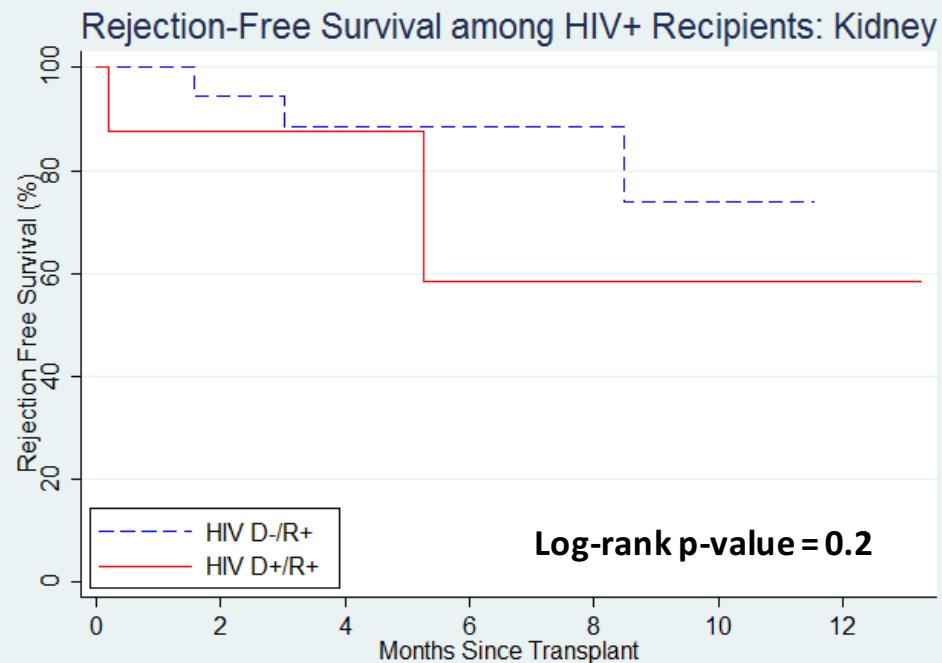
Post-transplant follow-up

	Overall (N=41)	HIV D-/R+ (N=30)	HIV D+/R+* (N=11)
Med (IQR) months	5.4 (2.9-8.8)	5.6 (4.4-9.2)	2.9 (1.1-5.3)
Range of follow-up	0.2 – 13.3	0.3-11.6	0.2-12.3

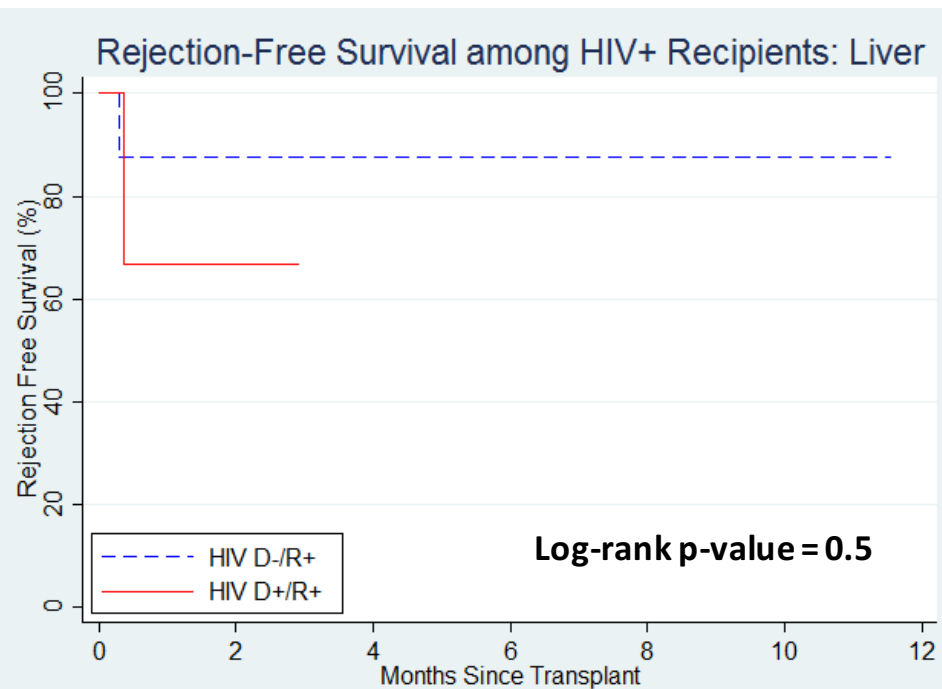
Post-transplant outcomes

	HIV D-/R+	HIV D+/R+
	N=30	N=11
Rejection episodes; N (%)	4 (13.3%)	3 (27.3%)
Opportunistic Infections; N (%)	0 (0%)	0 (0%)
Breakthrough viremia; N (%)	0 (0%)	0 (0%)
Malignancy; N (%)	0 (0%)	0 (0%)
Graft failure; N (%)	0 (0%)	1 (9.1%)
Death; N (%)	0 (0%)	0 (0%)

Among those who
received kidneys:



Among those who
received livers:



Serious Adverse Event Rates

	Patients (N)	Number of SAEs	Person- years	SAEs per person-year	p-value
HIV D-/R+	30	16	16.67	0.96	0.7
HIV D+/R+	11	6	5.13	1.17	

	HIV D+	HIV D-
Acute cellular rejection	0	2
Acute kidney injury	0	1
Altered mental status	1	0
CMV viremia	0	1
Hydronephrosis of transplanted kidney	0	1
Hyperglycemia	1	2
Hypotension	0	1
Lymphocele	1	0
Neutropenia	0	1
Oral ulcers	0	1
PD catheter malfunction	0	1
Pneumonia	0	1
Possible bleed after biopsy	1	0
Renal vein thrombosis	1	0
Tacrolimus toxicity	1	0
Upper respiratory infection	0	1
Urine leak	0	1
UTI	0	1
Wound dehiscence	0	1

HOPE *IN ACTION* → **TO DATE**

HIV-to-HIV Transplants Save Lives

**LIVES SAVED BY
TRANSPLANT** **26**

11 **ORGAN DONOR
HEROES**

**TRANSPLANT
CENTER
PARTNERS** **19**

15 **ORGAN
PROCUREMENT
ORGANIZATIONS**

Visit transplantepi.org for more about HOPE
Register to be an organ donor at registerme.org

HOPE

IN ACTION →

Learn if the use of HIV+ deceased donors in the is
safe and effective in the US

- Parent/Pilot transplant trial
- NIH multicenter trial

NIH U01: HIV D+/R+ Kidney Transplant

- 15 US Transplant Centers
- Deceased donor kidney transplant
- Compare HIV D-/R+ and HIV D+/R+, **N = 80 per arm**
- Anticipated start in fall



Program Officer:
Jonah Odum, MD PhD



National Institute
of Allergy and
Infectious Diseases



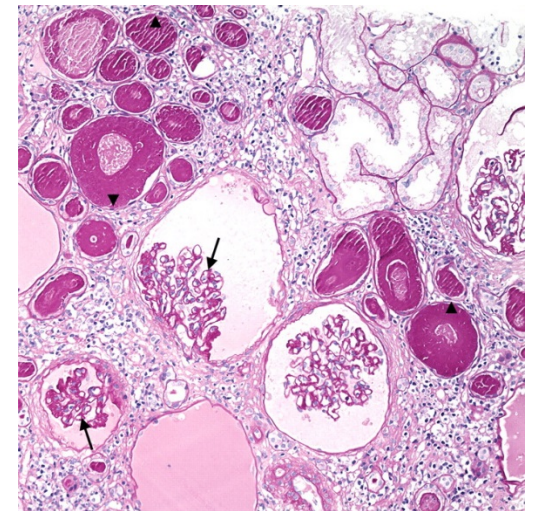
Project Manager:
Natasha Watson, MSN

Renal Disease

Allograft rejection

HIV associated renal diseases

- HIV associated nephropathy
- HIV associated immune complex disease
- Non-collapsing FSGS
- Thrombotic microangiopathy



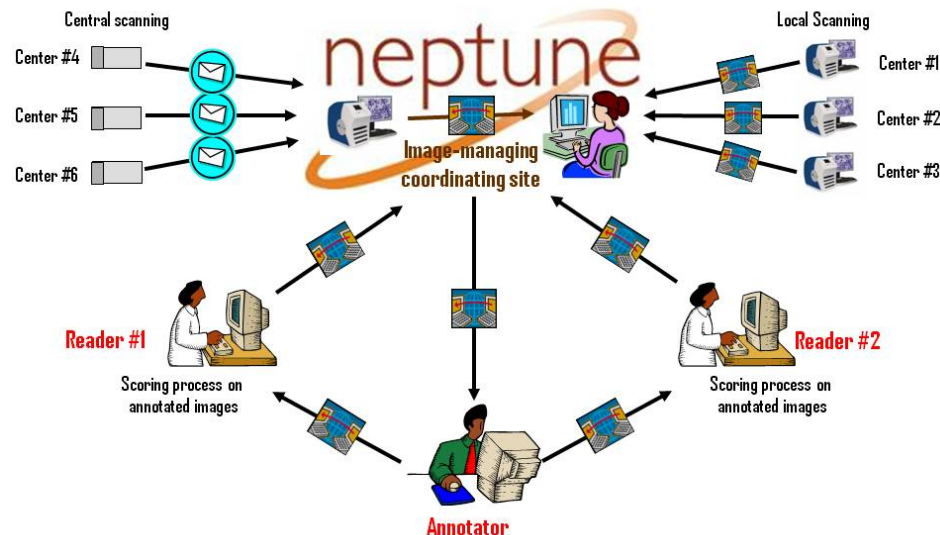
Digital Repository of Biopsies

Stephen Hewitt, MD, PhD

Head of Experimental Pathology



NEPTUNE Digital Pathology Protocol



HOPE
IN ACTION

Genetic Risk: Donors and Recipients

- ApoL1 protein variants in donors and recipients



Jeffrey Kopp, MD

NIDDK

Chief, Kidney Diseases Branch



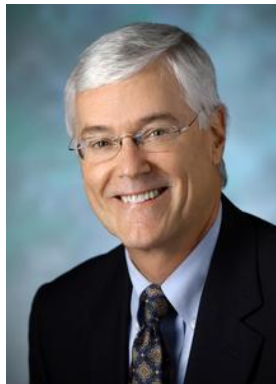
Cheryl Winkler, PhD

NCI

Head, Molecular Genetic
Epidemiology

Virology Studies

- Incidence of HIV-superinfection in blood
- Incidence of HIV-superinfection in tissue by laser capture microdissection
- Changes in HIV latent reservoir



Andrew Redd, PhD NIAID/JHU
Tom Quinn, MD NIAID/JHU
Aaron Tobian, MD PhD JHU

HIV Reservoir and Implications for Cure Efforts

HOPE
IN ACTION →

The Journal of Clinical Investigation

BRIEF REPORT

Rapamycin-mediated mTOR inhibition uncouples HIV-1 latency reversal from cytokine-associated toxicity

Alyssa R. Martin,¹ Ross A. Pollack,² Adam Capoferri,^{2,3} Richard F. Ambinder,¹ Christine M. Durand,² and Robert F. Siliciano^{2,3}

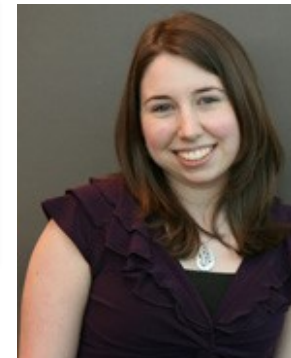


Alyssa Martin, PhD

- HIV PCR based assays are problematic but viral outgrowth assays (gold standard), costly, not always feasible

Defective proviruses rapidly accumulate during acute HIV-1 infection

Katherine M Bruner¹, Alexandra J Murray¹, Ross A Pollack¹, Mary G Soliman¹, Sarah B Laskey¹, Adam A Capoferri^{1,2}, Jun Lai¹, Matthew C Strain³, Steven M Lada⁴, Rebecca Hoh⁵, Ya-Chi Ho¹, Douglas D Richman^{3,4,6}, Steven G Deeks⁵, Janet D Siliciano¹ & Robert F Siliciano^{1,2}



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Christine Durand, MD
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Susan Lerner, MD

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Elizabeth Ferry, RN
Farzan Saeed
Nicole Turgeon, MD
Dasia Webster

Duke University

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Cameron Wolfe, MBBS

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Jonathan Bromberg, MD PhD

Massachusetts General Hospital

Margaret Thomas, CCRC
David Wojciechowski, DO

Weill Cornell Medical College

Thangamani Muthukumar, MD
Benjamin Samstein, MD
Catherine Small, MD

Methods: HIV testing

OPTN: FDA-Approved HIV Screening for Donor Products

Assay Name	Assay Type ^a	Maker	Sensitivity	Specificity	FP Rate
HIV Antibody assays					
Genetic Systems HIV-1/HIV-2 Plus O	EIA	Bio-Rad	100%	99.89%	0.11%
ABBOTT PRISM HIV O Plus	ChLIA	Abbott	100%	99.94%	0.06%
HIVAB HIV-1/HIV-2 (rDNA) EIA	EIA	Abbott	100%	99.90%	0.1%
HIV NAT assays					
COBAS AmpliScreen HIV-1 Test, v1.5	PCR	Roche	99.1%	99.7%	0.3%
Procleix HIV-1/HCV	TMA	Gen-Probe	99.8%	99.87%	0.13%
Procleix Ultrio	TMA	Gen-Probe	99.6%	99.8%	0.2%
COBAS TaqScreen MPX Test	PCR	Roche	100%	99.98%	0.02%
Procleix Ultrio Plus	TMA	Gen-Probe	99.72%	100%	0%

Donor HIV testing

Donor	OPTN Ab	Result	OPTN NAT	Result	HOPE qPCR
1	Genetic systems HIV 1/2/O EIA	Pos	Procleix Ultrio Plus	Neg	Neg
2	Genetic systems HIV 1/2/O EIA	Pos	COBAS Taq screen MTX v2	Neg	Neg
3	Abbott Prism HIV O plus	Pos	Procleix Ultrio Plus	Neg	Neg
4	Genetic systems HIV 1/2/O EIA	Neg	Procleix Ultrio Plus	Pos	Neg
5	Genetic systems HIV 1/2/O EIA	Neg	COBAS Taq screen MTX v2	Pos	Neg
6	Genetic systems HIV 1/2/O EIA	Pos	Procleix Ultrio Plus	Neg	Neg



South Africa: HIV-to-HIV

27 recipients

- CD4 > 200, VL < 50
- Median age 41
- 56% male
- 96% African
- 94% HIV-associated nephropathy

15 donors

- Median age 30
- Cause of death: trauma, overdose, SAH
- 14 treatment naïve, 1 on antiretroviral therapy

The NEW ENGLAND JOURNAL of MEDICINE

ORIGINAL ARTICLE

HIV-Positive-to-HIV-Positive Kidney Transplantation — Results at 3 to 5 Years

Elmi Muller, M.B., Ch.B., M.Med., Zunaid Barday, M.B., Ch.B.,
Marc Mendelson, M.D., Ph.D., and Delawir Kahn, M.B., Ch.B., Ch.M.

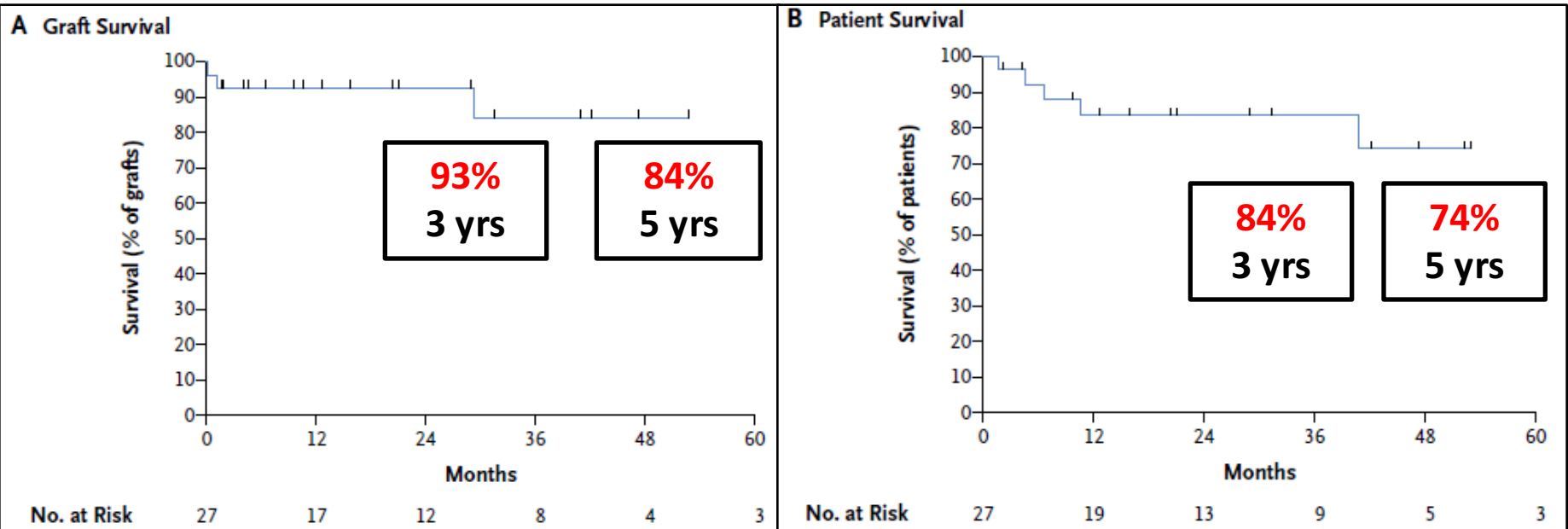
South Africa: HIV-to-HIV

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N =27

Muller et al, NEJM; 2015; 372:613-20.

South Africa: HIV-to-HIV

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HIV disease and infections

- 100% HIV suppression maintained
- One case of extrapulmonary TB, one of aspergillus

Rejection (all received ATG/thymoglobuline)

- 8% - 1 year
- 22% - 3 years

Recurrent HIV-AN?

- 3/27 early histologic changes with normal eGFR