

Adoptive Cellular Immunotherapy

Mark B. Faries, MD, FACS

Professor of Surgery

Chief, Donald L. Morton Melanoma Research Program

Director, Therapeutic Immunology



Cell Therapy: Overview

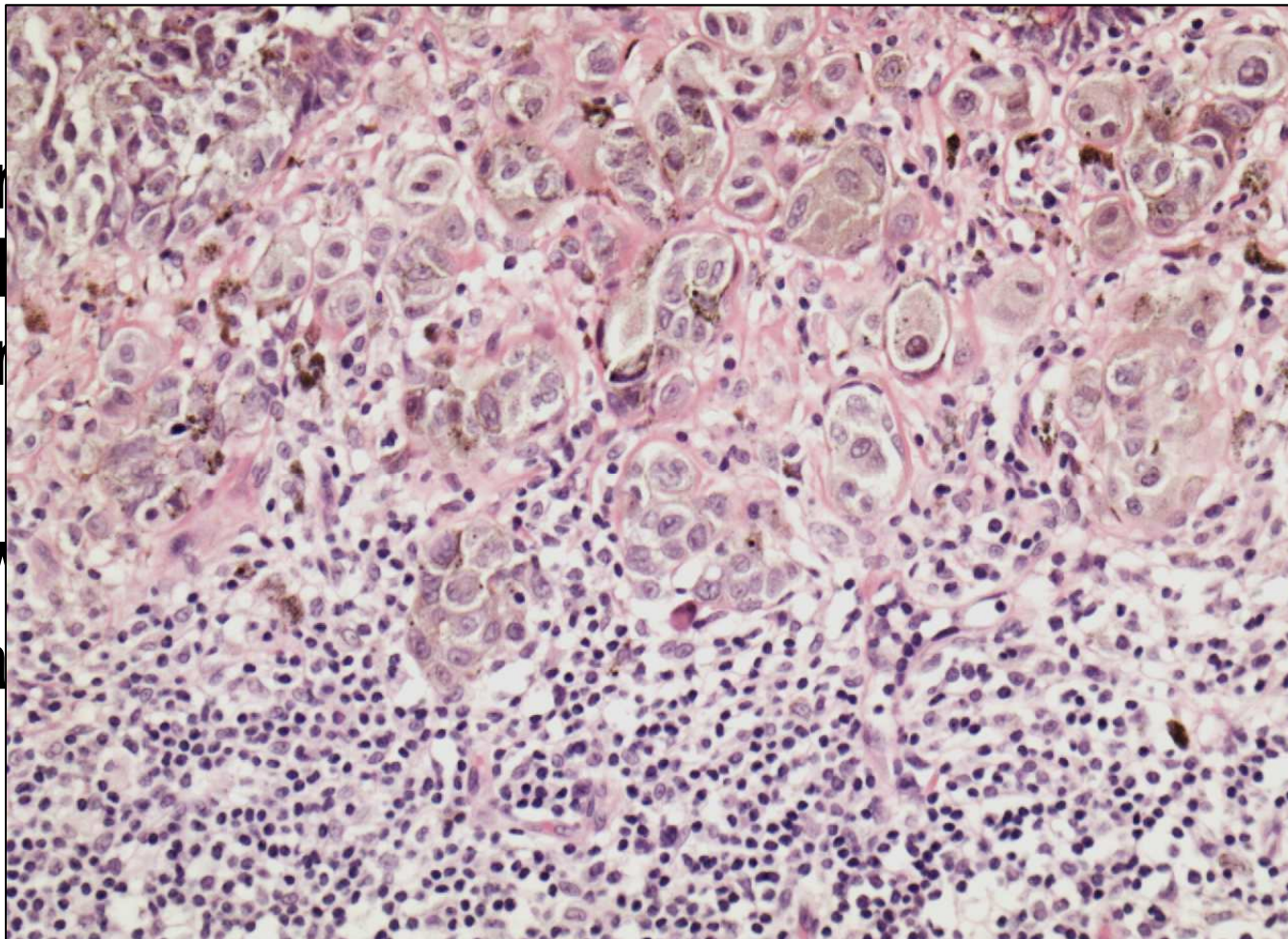
- **Background: Lymphocytes and Cancer**
- **History of ACT**
- **Current Programs and Recent Discoveries**
- **Outstanding issues and questions**
 - **Challenges**
 - **Opportunities**

Tumor-Infiltrating Lymphocytes

- **Endogenous immune responses to cancer are described in numerous tumor types and are strongly related to outcomes**
- **New immune therapies that activate those cells in vivo (e.g. checkpoint blockade) have shown dramatic effects**
- **Adoptive Cell Transfer has demonstrated proof of concept in melanoma and other cancers**

Background

- Natural immune recognition of melanoma is common



- In the in na,
- v a ne

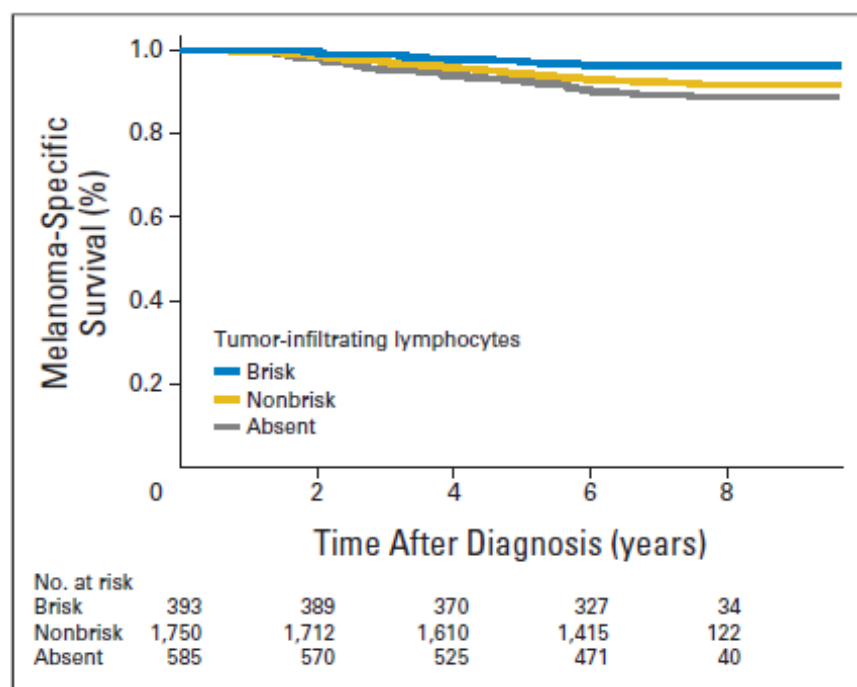
TIL in Melanoma

Tumor-Infiltrating Lymphocyte Grade in Primary Melanomas Is Independently Associated With Melanoma-Specific Survival in the Population-Based Genes, Environment and Melanoma Study

Nancy E. Thomas, Klaus J. Busam, Lynn From, Anne Krickler, Bruce K. Armstrong, Hoda Anton-Culver, Stephen B. Gruber, Richard P. Gallagher, Roberto Zanetti, Stefano Rosso, Terence Dwyer, Alison Venn, Peter A. Kanetsky, Pamela A. Groben, Honglin Hao, Irene Orlow, Anne S. Reiner, Li Luo, Susan Paine, David W. Ollila, Homer Wilcox, Colin B. Begg, and Marianne Berwick

JOURNAL OF CLINICAL ONCOLOGY

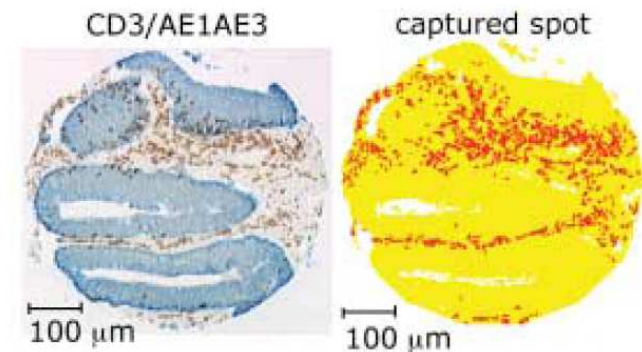
NOVEMBER 20 2013



TIL in Colon: Immunoscore

Effector Memory T Cells, Early Metastasis, and Survival in Colorectal Cancer

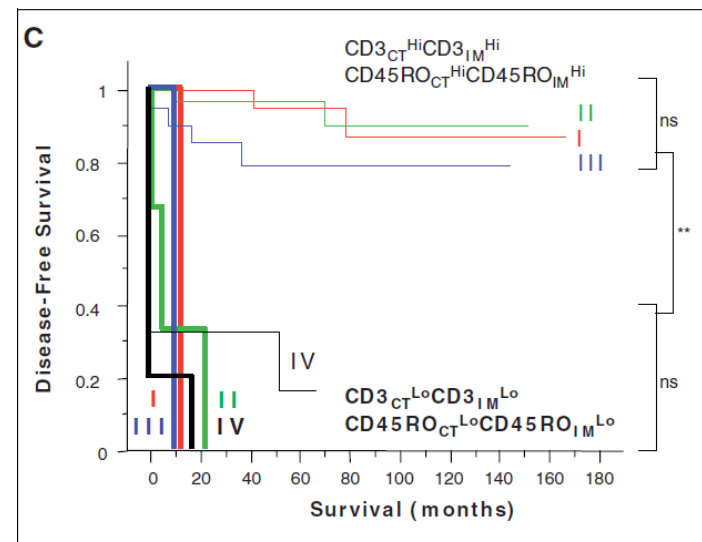
Franck Pagès, M.D., Ph.D., Anne Berger, M.D., Ph.D., Matthieu Camus, M.Sc.,
Fatima Sanchez-Cabo, Ph.D., Anne Costes, B.S., Robert Molitor, Ph.D.,
Bernhard Mlecnik, M.Sc., Amos Kirilovsky, M.Sc., Malin Nilsson, B.S.,
Diane Damotte, M.D., Ph.D., Tchao Meatchi, M.D., Patrick Bruneval, M.D., Ph.D.,
Paul-Henri Cugnenc, M.D., Ph.D., Zlatko Trajanoski, Ph.D.,
Wolf-Herman Fridman, M.D., Ph.D., and Jérôme Galon, Ph.D.
The NEW ENGLAND JOURNAL of MEDICINE DECEMBER 22, 2005



Type, Density, and Location of Immune Cells Within Human Colorectal Tumors Predict Clinical Outcome

Jérôme Galon *et al.*

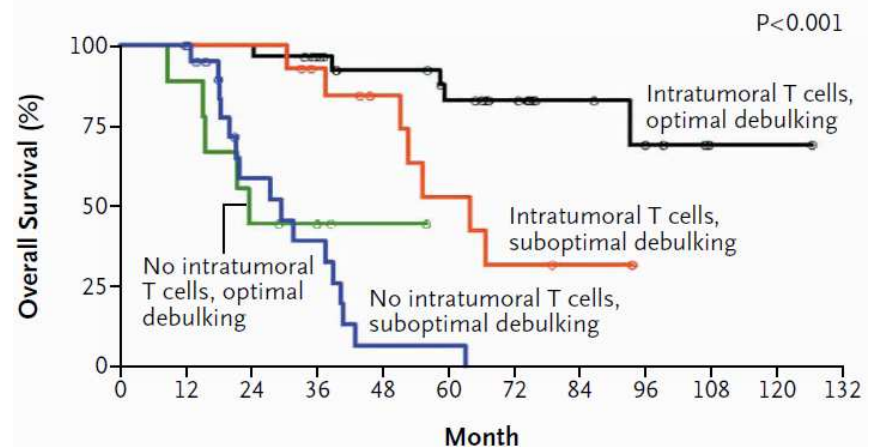
Science **313**, 1960 (2006);



TIL in Ovarian

Intratumoral T Cells, Recurrence, and Survival in Epithelial Ovarian Cancer

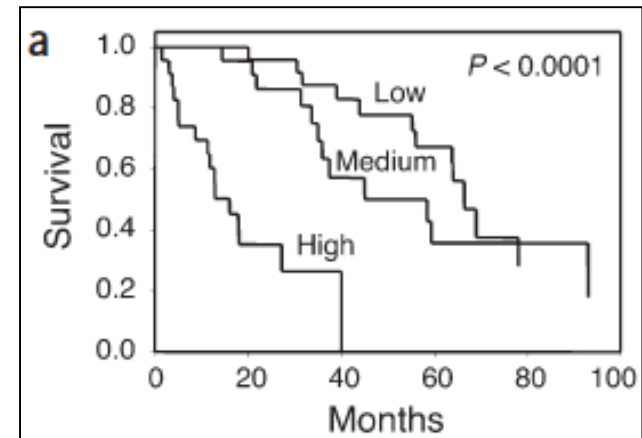
Lin Zhang, M.D., Jose R. Conejo-Garcia, M.D., Ph.D.,
 Dionyssios Katsaros, M.D., Ph.D., Phyllis A. Gimotty, Ph.D.,
 Marco Massobrio, M.D., Giorgia Regnani, M.D.,
 Antonis Makrigiannakis, M.D., Ph.D., Heidi Gray, M.D.,
 Katia Schlienger, M.D., Ph.D., Michael N. Lieberman, Ph.D.,
 Stephen C. Rubin, M.D., and George Coukos, M.D., Ph.D.
The NEW ENGLAND JOURNAL of MEDICINE
 JANUARY 16, 2003



Specific recruitment of regulatory T cells in ovarian carcinoma fosters immune privilege and predicts reduced survival

Tyler J Curiel¹, George Coukos², Linhua Zou¹, Xavier Alvarez¹, Pui Cheng¹, Peter Mottram¹,
 Melina Evdemon-Hogan¹, Jose R Conejo-Garcia², Lin Zhang², Matthew Burow¹, Yun Zhu¹, Shuang Wei¹,
 Ilona Kryczek¹, Ben Daniel¹, Alan Gordon³, Leann Myers¹, Andrew Lackner¹, Mary L Disis⁴, Keith L Knutson⁴,
 Lieping Chen⁵ & Weiping Zou¹

VOLUME 10 | NUMBER 9 | SEPTEMBER 2004 *NATURE MEDICINE*

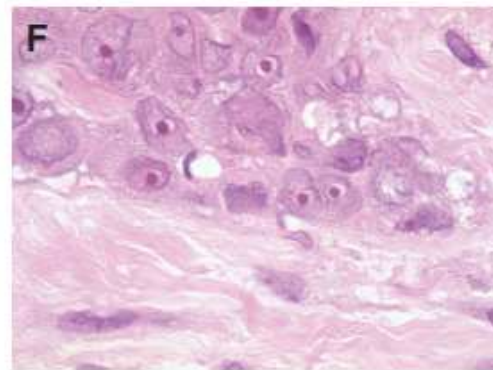
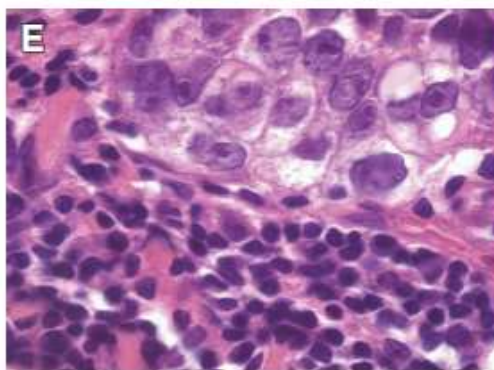
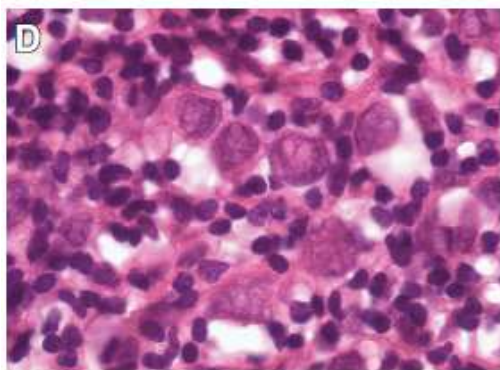
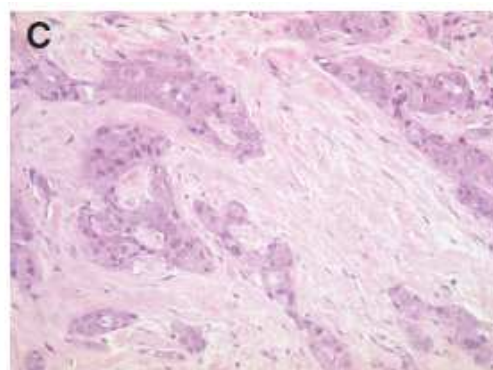
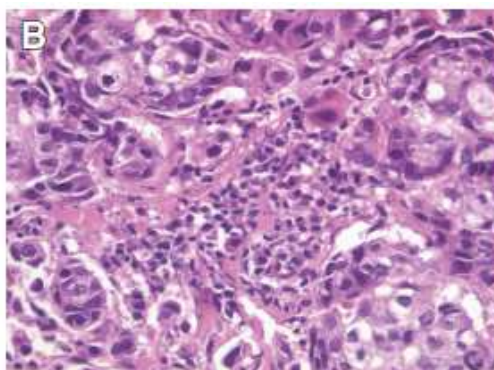
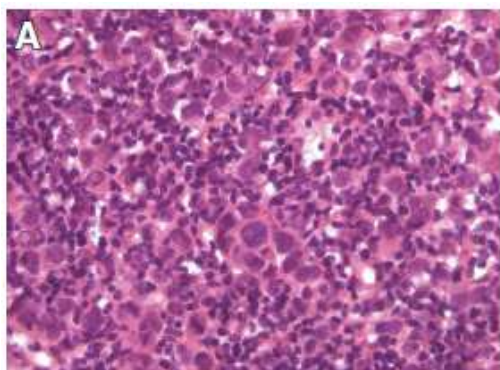


TIL in Breast Cancer

Tumor-Associated Lymphocytes As an Independent Predictor of Response to Neoadjuvant Chemotherapy in Breast Cancer

Carsten Denkert, Sibylle Loibl, Aurelia Noske, Marc Roller, Berit Maria Müller, Martina Komor, Jan Budczies, Silvia Darb-Esfahani, Ralf Kronenwett, Claus Hanusch, Christian von Törne, Wilko Weichert, Knut Engels, Christine Solbach, Iris Schrader, Manfred Dietel, and Gunter von Minckwitz

JANUARY 1 2010 *J Clin Oncol* 28:105-113.

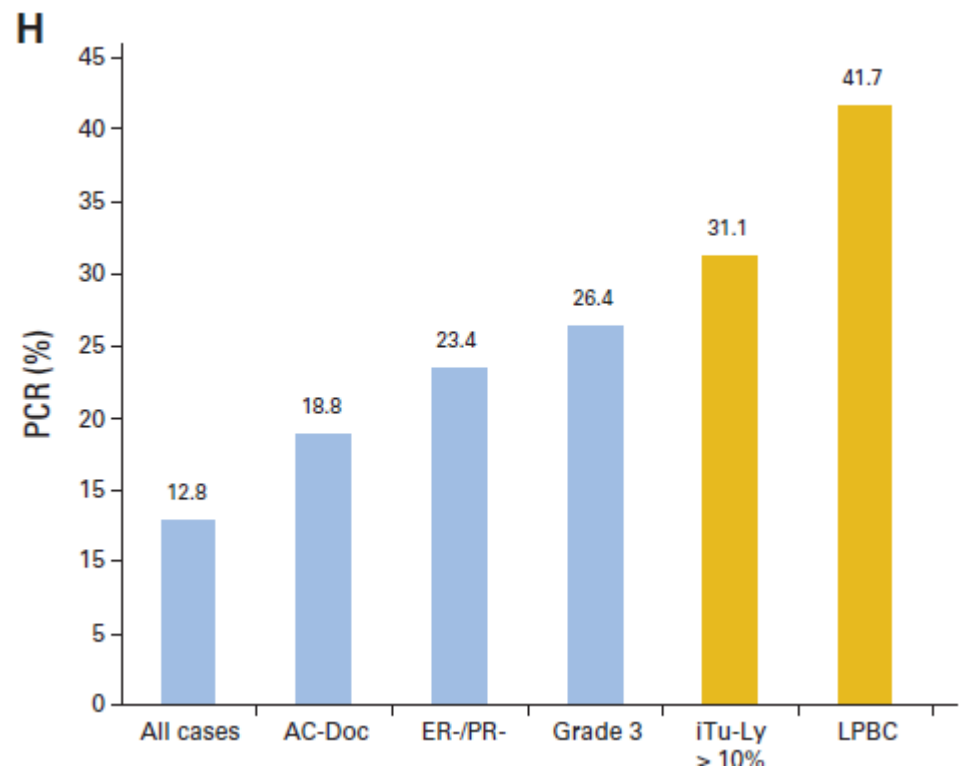
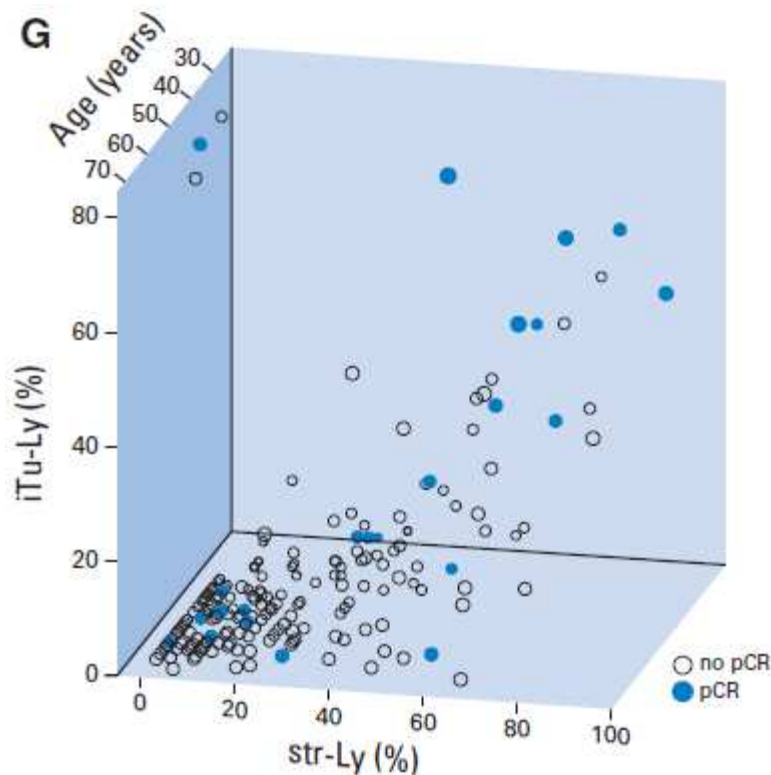


TIL in Breast Cancer

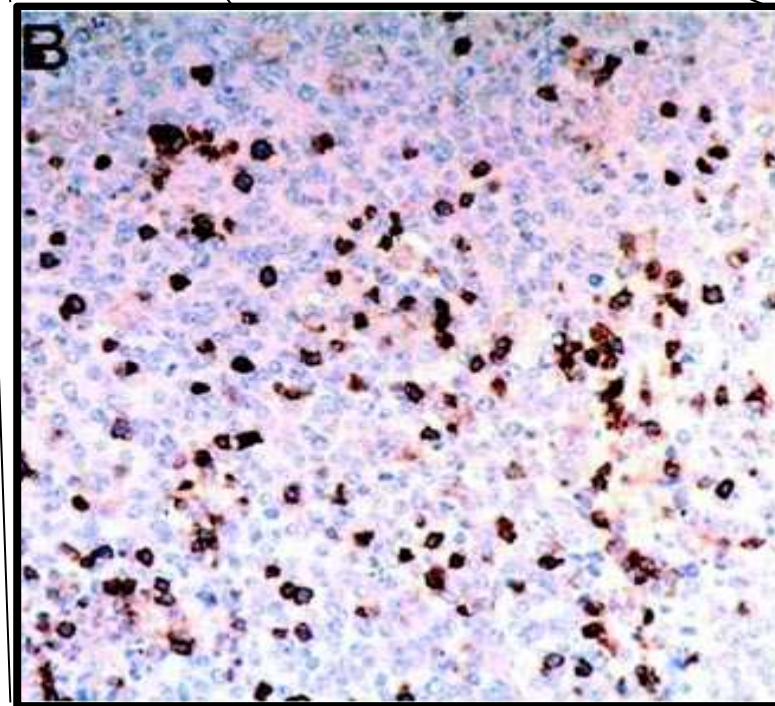
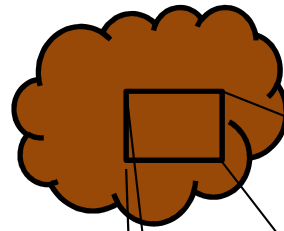
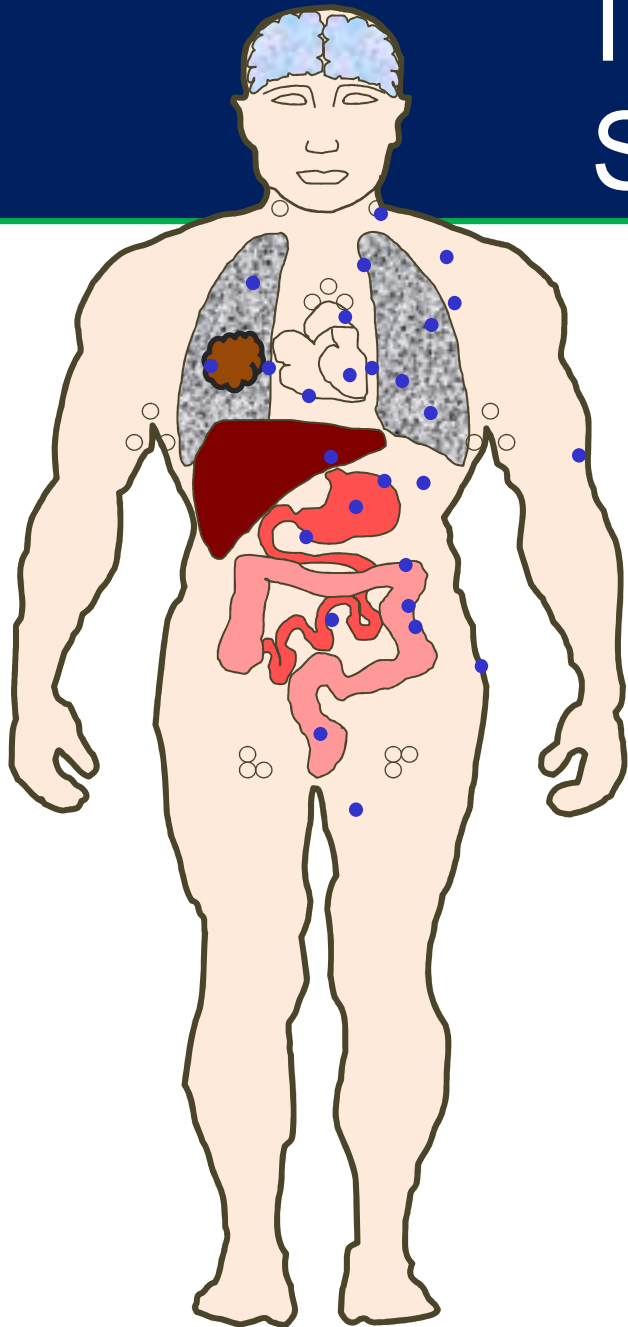
Tumor-Associated Lymphocytes As an Independent Predictor of Response to Neoadjuvant Chemotherapy in Breast Cancer

Carsten Denkert, Sibylle Loibl, Aurelia Noske, Marc Roller, Berit Maria Müller, Martina Komor, Jan Budczies, Silvia Darb-Esfahani, Ralf Kronenwett, Claus Hanusch, Christian von Törne, Wilko Weichert, Knut Engels, Christine Solbach, Iris Schrader, Manfred Dietel, and Gunter von Minckwitz

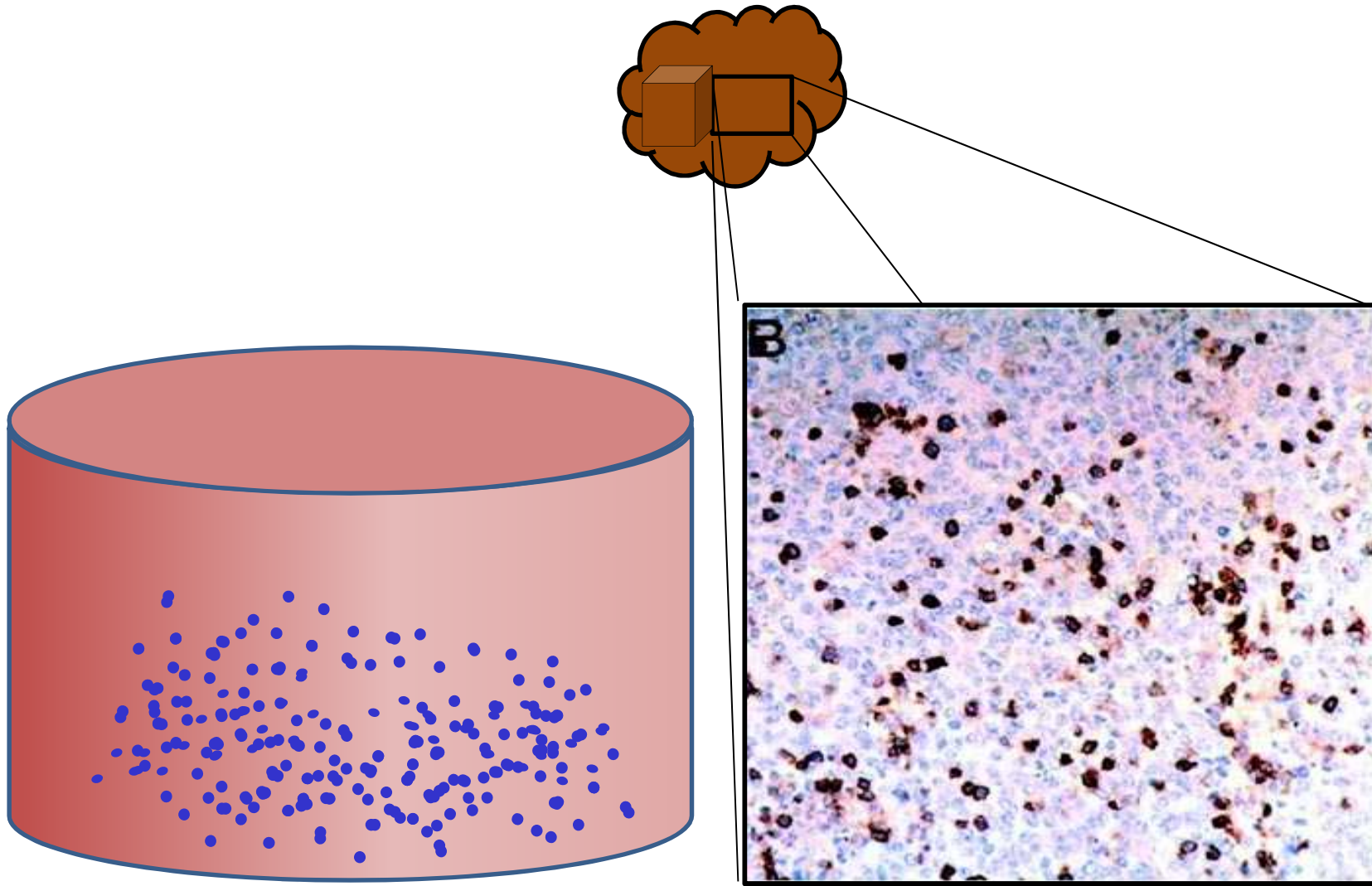
JANUARY 1 2010 *J Clin Oncol* 28:105-113.



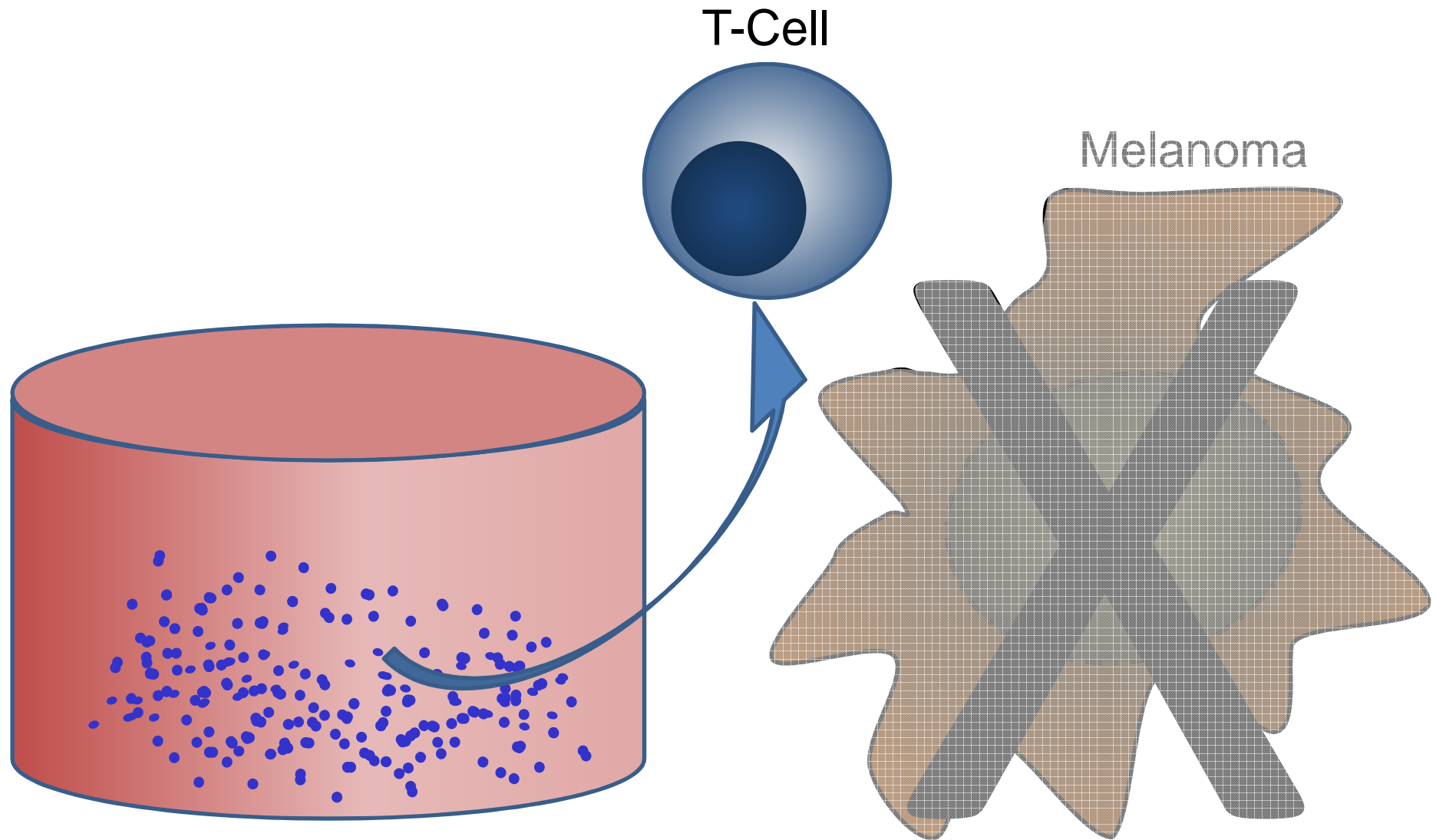
TIL: Current Standard Procedure



TIL: Current Standard Procedure

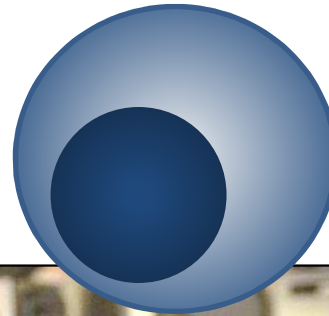


TIL: Current Standard Procedure



TIL: Current Standard Procedure

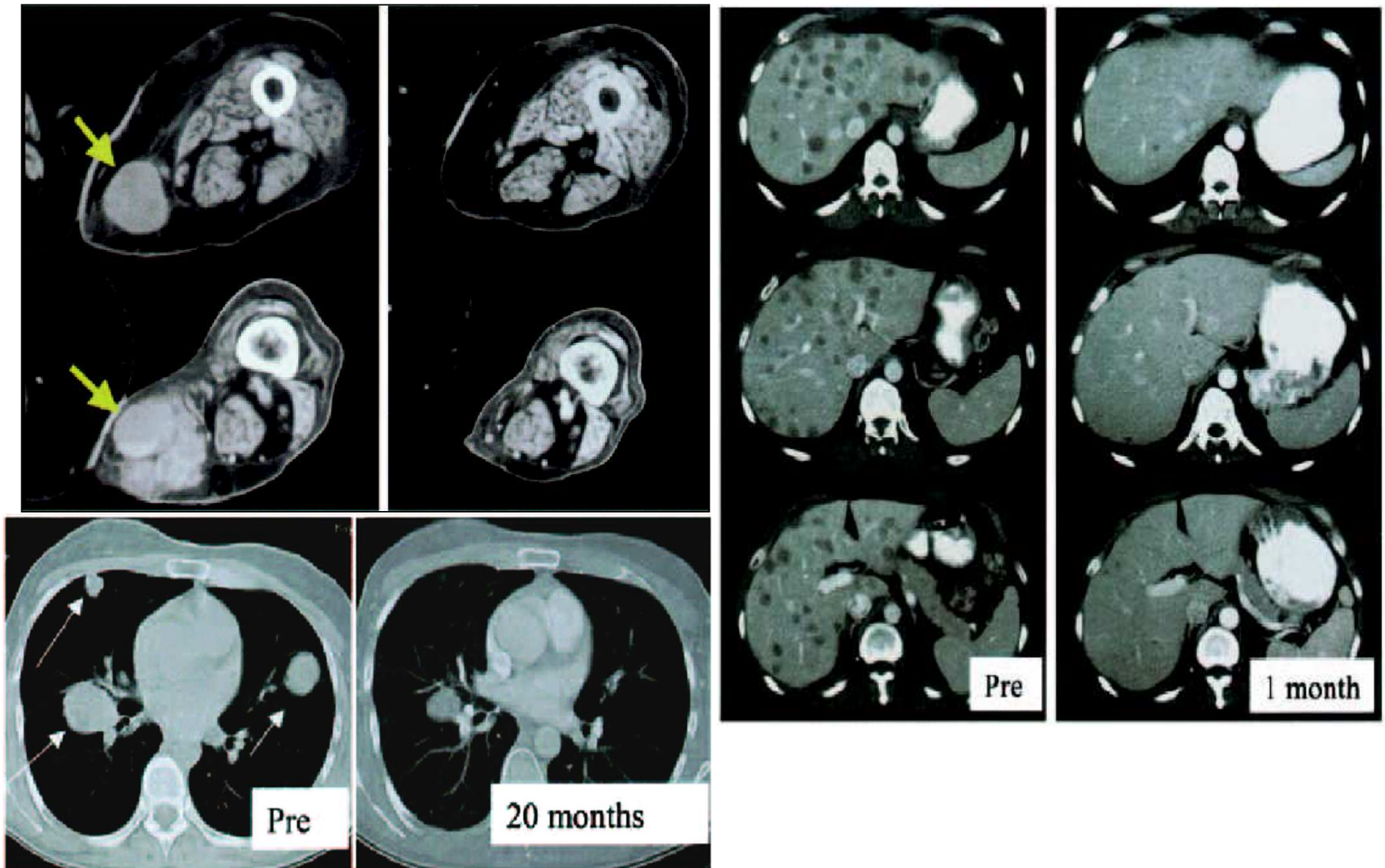
T-Cell



RAPID EXPANSION PROTOCOL



TIL: Clinical Responses



TIL: Early Clinical Data

Treatment of Patients With Metastatic Melanoma With Autologous Tumor-Infiltrating Lymphocytes and Interleukin 2

Journal of the National Cancer Institute,
Vol. 86, No. 15, August 3, 1994

Steven A. Rosenberg, John R.
Yannelli, James C. Yang, Suzanne
L. Topalian, Douglas J.
Schwartzentruber, Jeffrey S.
Weber, David R. Parkinson,
Claudia A. Seipp, Jan H. Einhorn,
Donald E. White*

Table 2. Number of patients responding to treatment with TILs plus IL-2 and duration of response*

	Response to treatment											
	No cyclophosphamide				Plus cyclophosphamide				Total			
	No. of patients				No. of patients				No. of patients			
	Total	CR	PR	% CR + PR	Total	CR	PR	% CR + PR	Total	CR	PR	% CR + PR
Prior IL-2	11	1	2	27	17	0	6	35	28	1	8	32
No prior IL-2	18	3	3	33	40	1	13	35	58	4	16	34
Total	29	4	5	31	57	1	19	35	86	5	24	34

	Duration of response, mo			
	No cyclophosphamide		Plus cyclophosphamide	
	CR	PR	CR	PR
Prior IL-2	23	4, 1	—	8, 7, 6, 5, 5, 1
No prior IL-2	46+, 38, 21+	7, 4, 2	20	53+, 9, 7, 7, 4, 4

*CR = complete response. PR = partial response.

TIL: Early Clinical Data

Treatment of Patients With Metastatic Melanoma With Autologous Tumor-Infiltrating Lymphocytes and Interleukin 2

Steven A. Rosenberg, John R. Yannelli, James C. Yang, Suzanne L. Topalian, Douglas J. Schwartzentruber, Jeffrey S. Weber, David R. Parkinson, Claudia A. Seipp, Jan H. Einhorn, Donald E. White*

Journal of the National Cancer Institute,
Vol. 86, No. 15, August 3, 1994

Table 3. Treatment with TILs plus IL-2: characteristics of TILs associated with response and site of tumor harvest

Characteristics of TILs associated with response	Responder*	Nonresponder*	P
Time in culture, d	33 ± 1	43 ± 2	.0001
Doubling time, d	2.4 ± 0.2	3.5 ± 0.4	.03
% lysis (effector-to-target cell ratio of 40:1)			
Autologous tumor target	25 ± 4	10 ± 2	.0008
Daudi lymphoma target	11 ± 6	6 ± 2	.6
Phenotype, % of cells			
CD3 ⁺	96 ± 1	95 ± 1	.9
CD4 ⁺	24 ± 8	32 ± 5	.3
CD8 ⁺	71 ± 8	62 ± 5	.3
CD56 ⁺	9 ± 3	13 ± 2	.3

Site of tumor harvest	No. of patients	
	Responder	Nonresponder
Lymph node†	6	29
Subcutaneous tumor nodule†	18	19
Muscle	0	3
Lung	1	2
Intraperitoneal mass	0	2
Pleural effusion	0	1
Colon	1	0
Liver	1	0
Spleen	0	1
Ovary	1	0
Bone	1	0
Total	29	57

*Values = means ± SEM.

†Difference in response rates comparing lymph node and subcutaneous harvest sites, $P = .006$.

TIL: Early Clinical Data

Cancer Regression and Autoimmunity in Patients After Clonal Repopulation with Antitumor Lymphocytes

25 OCTOBER 2002 VOL 298 SCIENCE

Mark E. Dudley,¹ John R. Wunderlich,¹ Paul F. Robbins,¹
James C. Yang,¹ Patrick Hwu,¹ Douglas J. Schwartzentruber,¹
Suzanne L. Topalian,¹ Richard Sherry,¹ Nicholas P. Restifo,¹
Amy M. Hubicki,¹ Michael R. Robinson,² Mark Raffeld,³
Paul Duray,³ Claudia A. Seipp,¹ Linda Rogers-Freezer,¹
Kathleen E. Morton,¹ Sharon A. Mavroukakis,¹ Donald E. White,¹
Steven A. Rosenberg^{1*}

Table 1. Patient demographics, treatments received, and clinical outcomes.

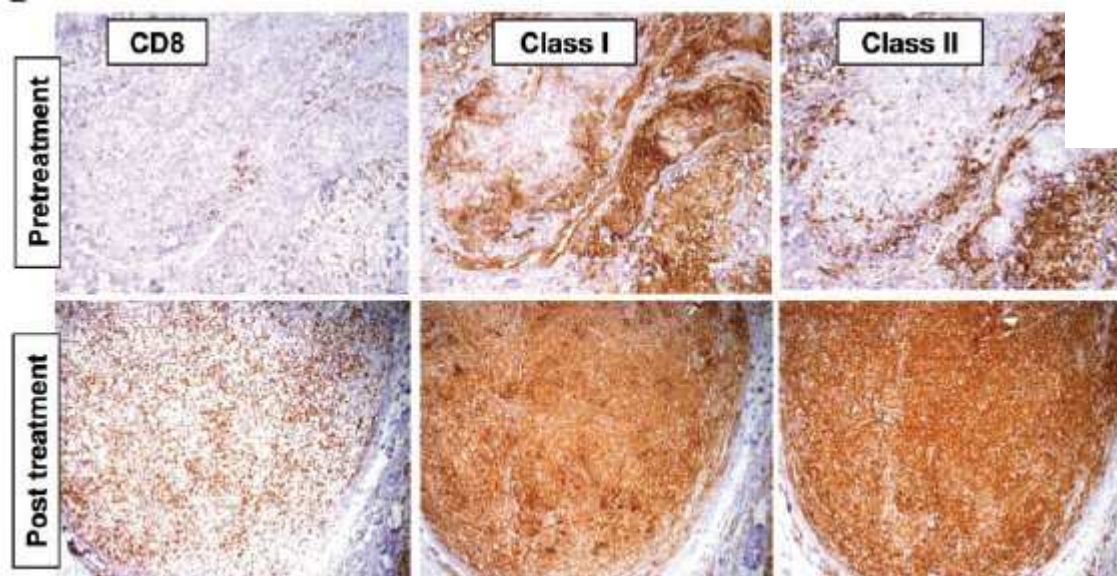
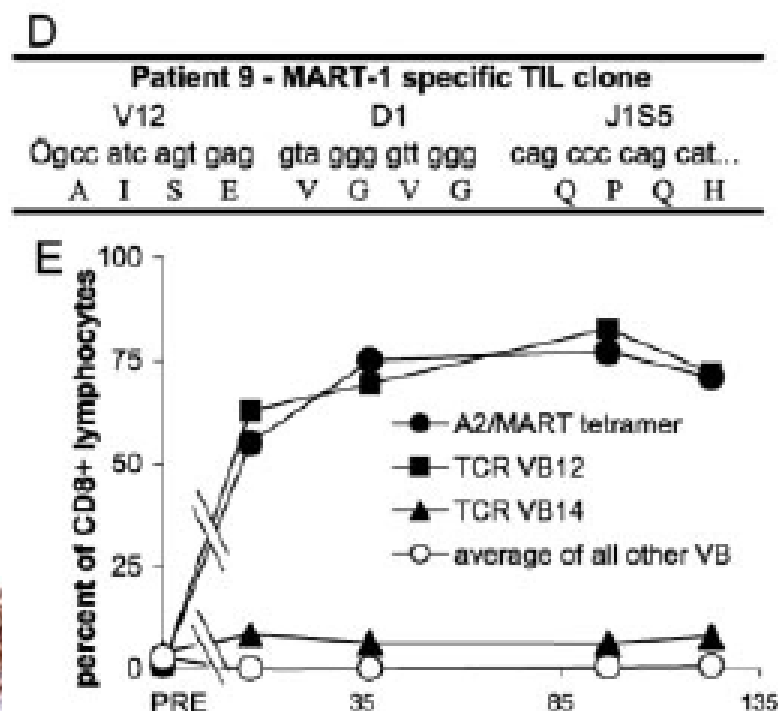
Patient	Age/sex	Treatment*				Sites of evaluable metastases	Response duration (months)	Autoimmunity
		Cells infused† ($\times 10^{-10}$)	CD8/CD4 phenotype‡ (%)	Antigen specificity§	IL-2 (doses)			
1	18/M	2.3	11/39	Other	9	Lymph nodes (axillary, mesenteric, pelvic)	PR¶ (24+)	None
2	30/F	3.5	83/15	MART-1, gp100	8	Cutaneous, subcutaneous	PR (8)	Vitiligo
3	43/F	4.0	44/58	gp100	5	Brain, cutaneous, liver, lung	NR	None
4	57/F	3.4	56/52	gp100	9	Cutaneous, subcutaneous	PR (2)	None
5	53/M	3.0	16/85	Other	7	Brain, lung, lymph nodes	NR-mixed	None
6	37/F	9.2	65/35	Other	6	Lung, intraperitoneal, subcutaneous	PR (15+)	None
7	44/M	12.3	61/41	MART-1	7	Lymph nodes, subcutaneous	NR-mixed	Vitiligo
8	48/M	9.5	48/52	gp100	12	Subcutaneous	NR	None
9	57/M	9.6	84/13	MART-1	10	Cutaneous, subcutaneous	PR (10+)	Vitiligo
10	55/M	10.7	96/2	MART-1	12	Lymph nodes, cutaneous, subcutaneous	PR¶ (9+)	Uveitis
11	29/M	13.0	96/3	MART-1	12	Liver, pericardial, subcutaneous	NR-mixed	Vitiligo
12	37/F	13.7	72/24	MART-1	11	Liver, lung, gallbladder, lymph nodes	NR-mixed	None
13	41/F	7.7	92/8	MART-1	11	Subcutaneous	NR	None

TIL Clinical Effects

Cancer Regression and Autoimmunity in Patients After Clonal Repopulation with Antitumor Lymphocytes

Mark E. Dudley,¹ John R. Wunderlich,¹ Paul F. Robbins,¹
James C. Yang,¹ Patrick Hwu,¹ Douglas J. Schwartzentruber,¹
Suzanne L. Topalian,¹ Richard Sherry,¹ Nicholas P. Restifo,¹
Amy M. Hubicki,¹ Michael R. Robinson,² Mark Raffeld,³
Paul Duray,³ Claudia A. Seipp,¹ Linda Rogers-Freezer,¹
Kathleen E. Morton,¹ Sharon A. Mavroukakis,¹ Donald E. White,¹
Steven A. Rosenberg^{1*}

25 OCTOBER 2002 VOL 298 SCIENCE



TIL: Early Clinical Data RCC

Adoptive Immunotherapy With Tumor-Infiltrating Lymphocytes and Interleukin-2 in Patients With Metastatic Malignant Melanoma and Renal Cell Carcinoma: A Pilot Study

By Peter S. Goedegebuure, Linda M. Douville, Hong Li, Glenn C. Richmond, Deric D. Schoof, Marybeth Scavone, and Timothy J. Eberlein

Journal of Clinical Oncology, Vol 13, No 8 (August), 1995: pp 1939-1949

Table 4. Clinical Responses to TIL and Low-Dose IL-2

Response	Melanoma		RCC		Total	
	No.	%	No.	%	No.	%
CR	3	19	0	0	4	20
MR	0	0	2	50	2	10
NR	9	56*	2	50	10	50
PD	4	25	0	0	4	20
Assessable	16	80	4	20	20	100

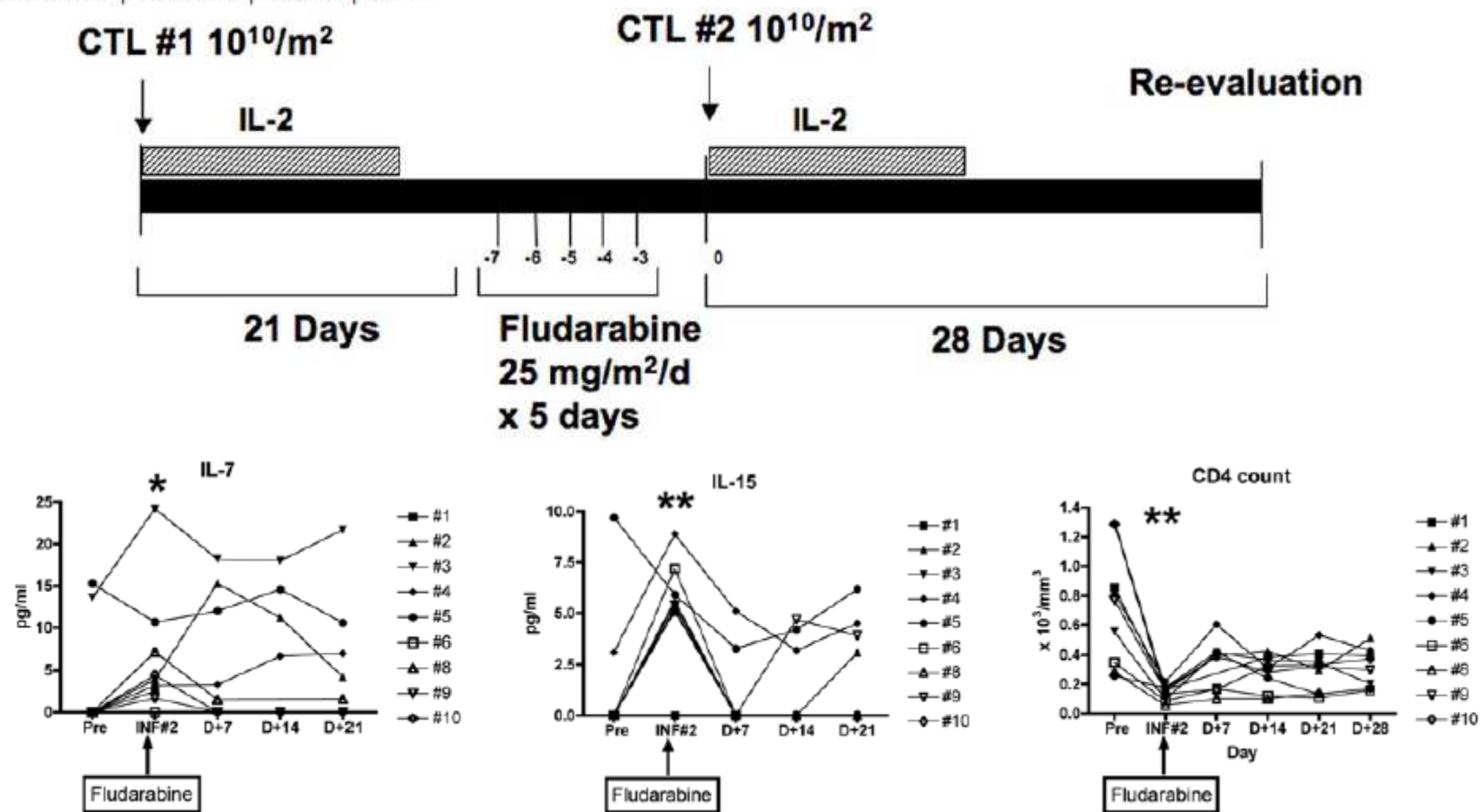


Lymphodepletion

Fludarabine Modulates Immune Response and Extends *In Vivo* Survival of Adoptively Transferred CD8 T Cells in Patients with Metastatic Melanoma

Herschel Wallen*, John A. Thompson, J. Zachary Reilly, Rebecca M. Rodmyre, Jianhong Cao, Cassian Yee

March 2009 | Volume 4 | Issue 3 | e4749

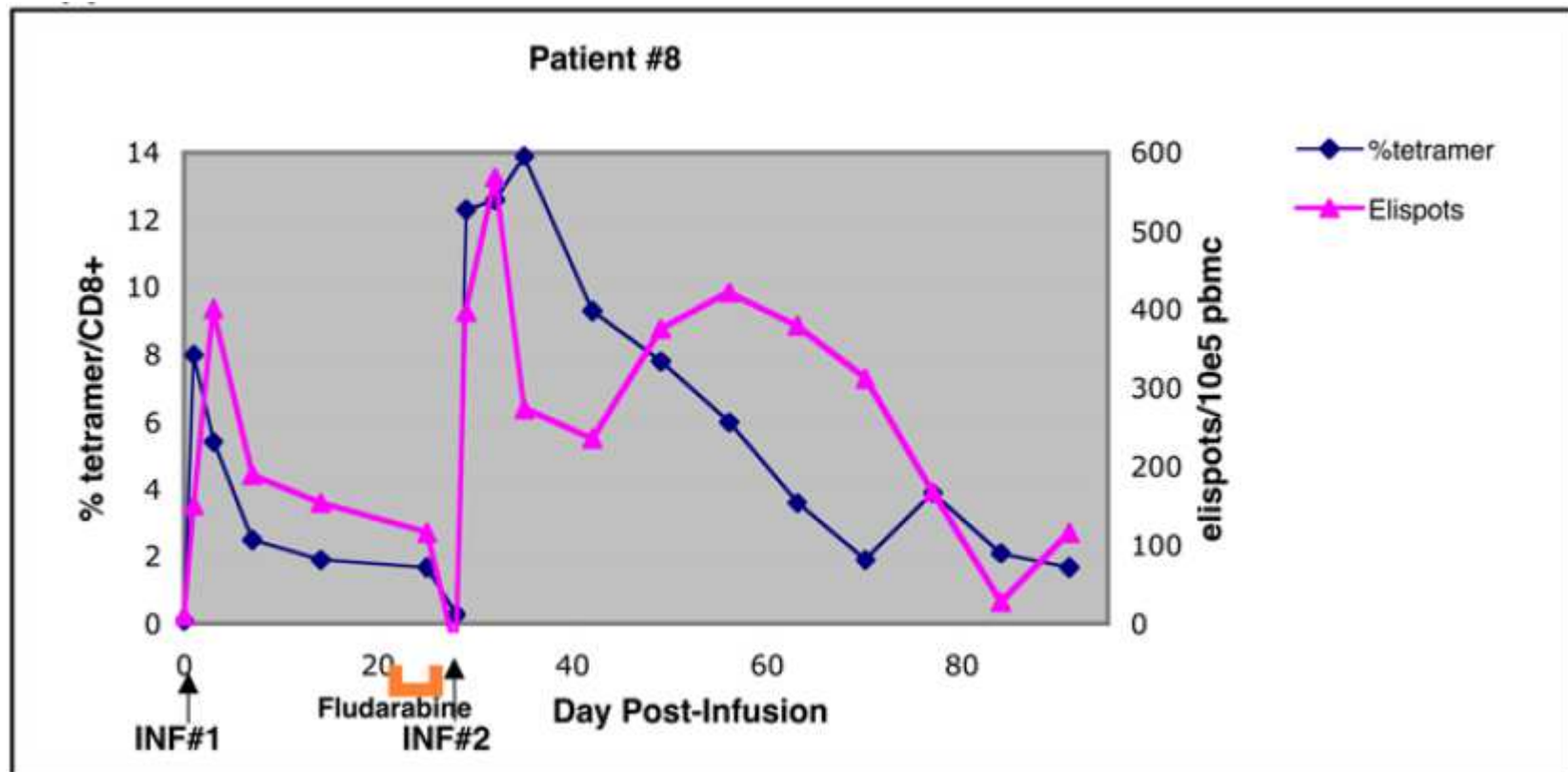


Lymphodepletion

Fludarabine Modulates Immune Response and Extends *In Vivo* Survival of Adoptively Transferred CD8 T Cells in Patients with Metastatic Melanoma

Herschel Wallen*, John A. Thompson, J. Zachary Reilly, Rebecca M. Rodmyre, Jianhong Cao, Cassian Yee

March 2009 | Volume 4 | Issue 3 | e4749



Lymphodepletion

Adoptive Cell Therapy for Patients With Metastatic Melanoma: Evaluation of Intensive Myeloablative Chemoradiation Preparative Regimens

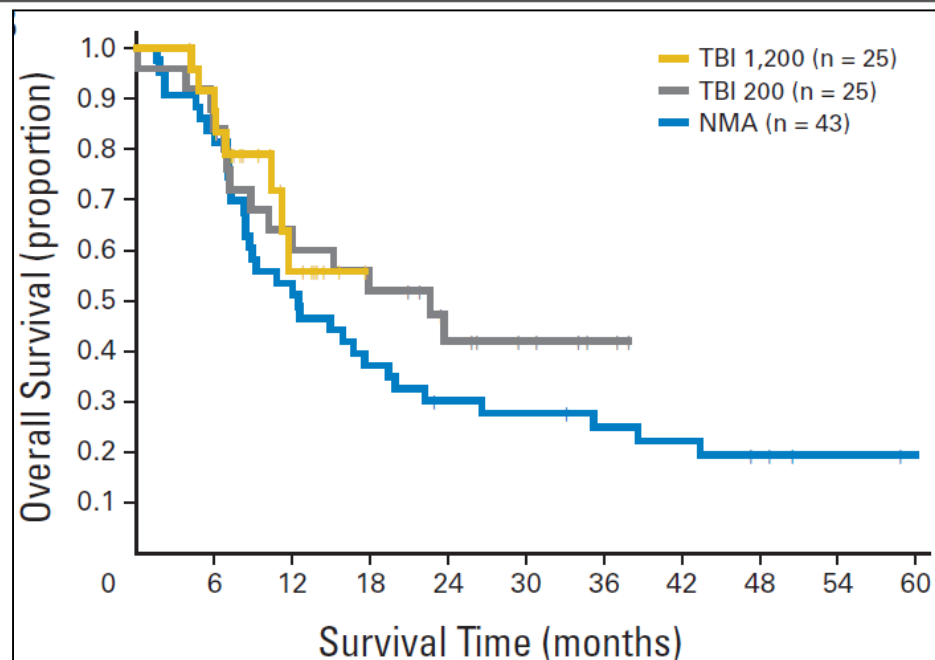
Mark E. Dudley, James C. Yang, Richard Sherry, Marybeth S. Hughes, Richard Royal, Udai Kammula, Paul F. Robbins, JianPing Huang, Deborah E. Citrin, Susan F. Leitman, John Wunderlich, Nicholas P. Restifo, Armen Thomasian, Stephanie G. Downey, Franz O. Smith, Jacob Klapper, Kathleen Morton, Carolyn Laurencot, Donald E. White, and Steven A. Rosenberg

JOURNAL OF CLINICAL ONCOLOGY

VOLUME 26 • NUMBER 32 • NOVEMBER 10 2008

Day of treatment	-7	-6	-5	-4	-3	-2	-1	0	1	2	3
Non-myeloablative	Cy	Cy	Flu	Flu	Flu	Flu	Flu	TIL	IL-2	IL-2	IL-2
Ablative (200cGy)		Cy Flu	Cy Flu	Flu	Flu	Flu	TBI	TIL	IL-2	IL-2 CD34+	IL-2
Ablative (1,200cGy)	Cy Flu	Cy Flu	Flu	Flu	Flu TBI	TBI	TBI	TIL	IL-2 CD34+	IL-2	IL-2

VOLUME 26 • NUMBER 32 • NOVEMBER 10 2008



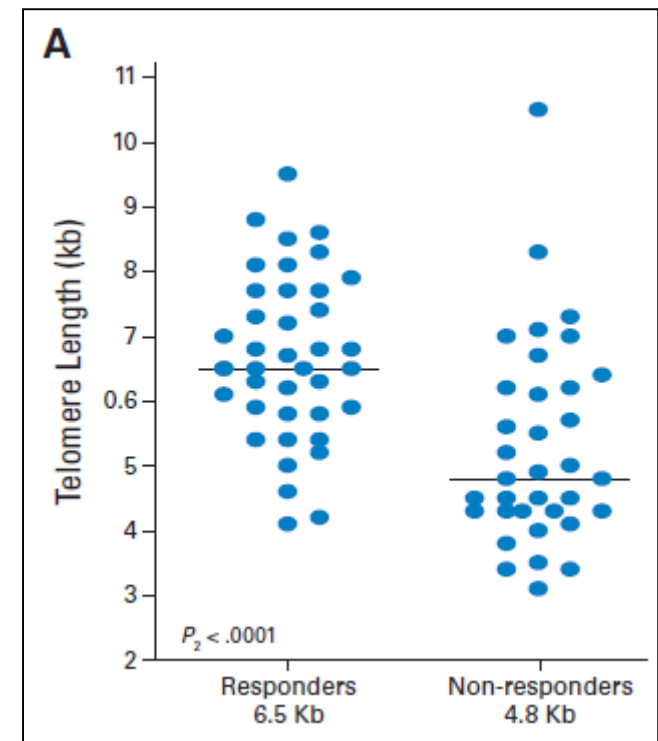
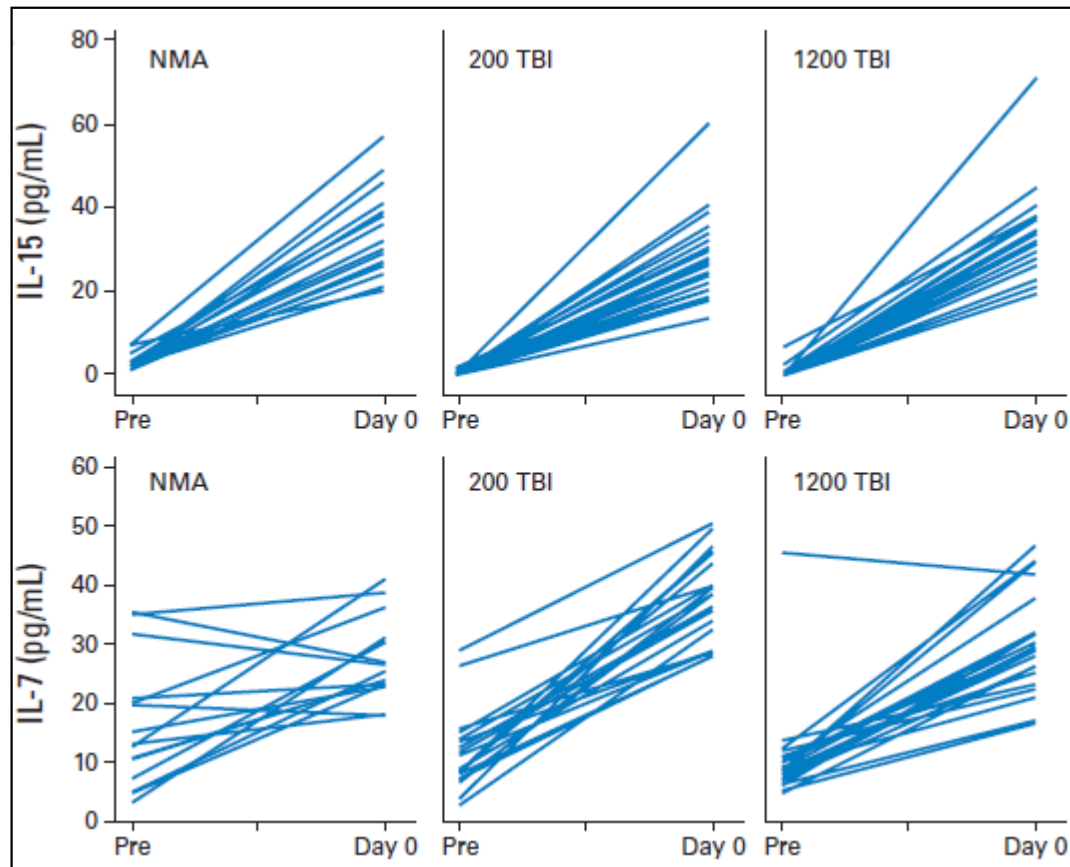
Lymphodepletion

Adoptive Cell Therapy for Patients With Metastatic Melanoma: Evaluation of Intensive Myeloablative Chemoradiation Preparative Regimens

Mark E. Dudley, James C. Yang, Richard Sherry, Marybeth S. Hughes, Richard Royal, Udai Kammula, Paul F. Robbins, JianPing Huang, Deborah E. Citrin, Susan F. Leitman, John Wunderlich, Nicholas P. Restifo, Armen Thomasian, Stephanie G. Downey, Franz O. Smith, Jacob Klapper, Kathleen Morton, Carolyn Laurencot, Donald E. White, and Steven A. Rosenberg

JOURNAL OF CLINICAL ONCOLOGY

VOLUME 26 · NUMBER 32 · NOVEMBER 10 2008

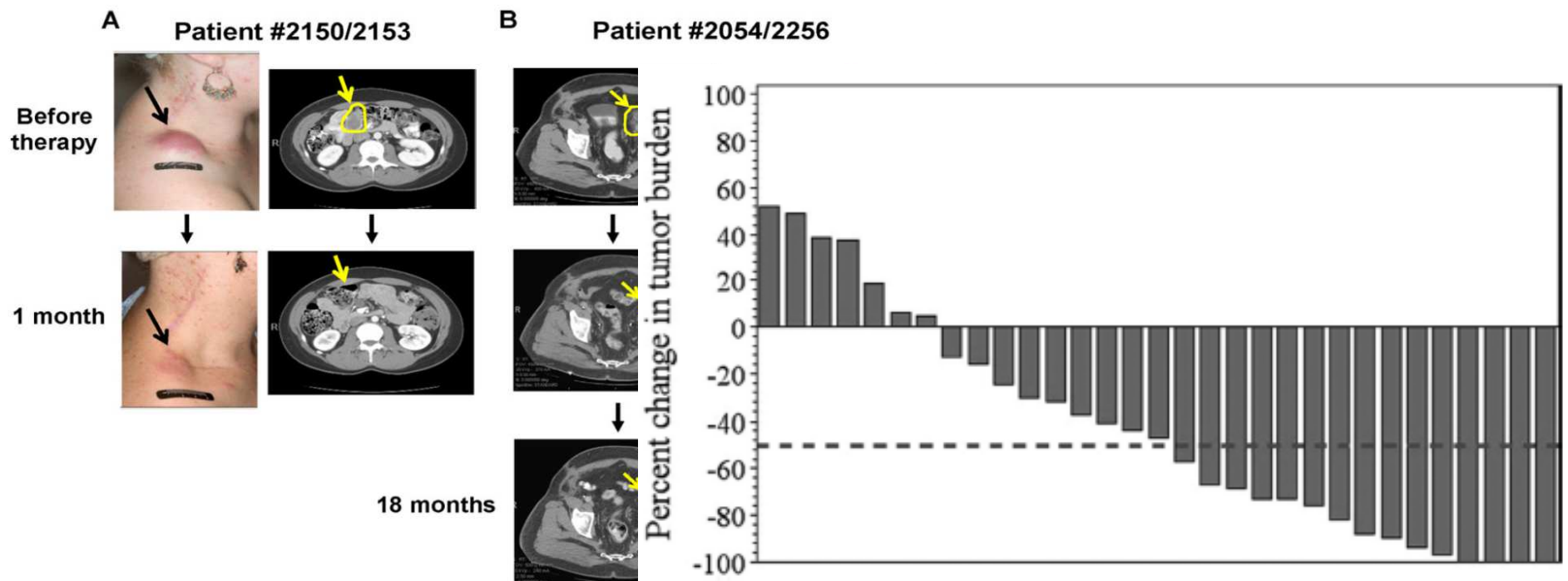


Predicting Response

Specific lymphocyte subsets predict response to adoptive cell therapy using expanded autologous tumor-infiltrating lymphocytes in metastatic melanoma patients

Laszlo G. Radvanyi^{1,*}, Chantale Bernatchez¹, Mingying Zhang¹, Patricia S. Fox², Priscilla Miller¹, Jessica Chacon¹, Richard Wu¹, Gregory Lizee¹, Sandy Mahoney¹, Gladys Alvarado¹, Michelle Glass¹, Valen E. Johnson², John D. McMannis³, Elizabeth Shpall³, Victor Prieto⁴, Nicholas Papadopoulos¹, Kevin Kim¹, Jade Homsy¹, Agop Bedikian¹, Wen-Jen Hwu¹, Sapna Patel¹, Merrick I. Ross⁵, Jeffrey E. Lee⁵, Jeffrey E. Gershenwald⁵, Anthony Lucci⁵, Richard Royal⁵, Janice N. Cormier⁵, Michael A. Davies¹, Rahmatu Mansaray^{1,3}, Orenthial J. Fulbright^{1,3}, Christopher Toth^{1,3}, Renjith Ramachandran^{1,3}, Seth Wardell^{1,3}, Audrey Gonzalez^{1,3}, and Patrick Hwu^{1,*}

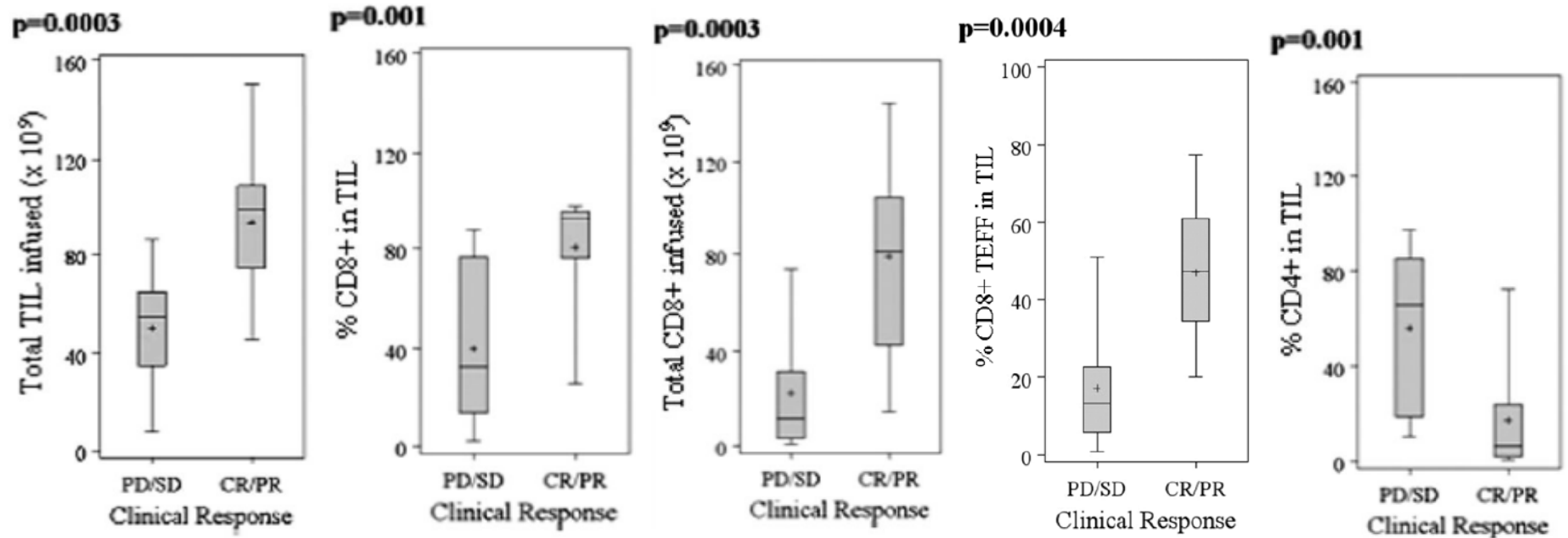
Clin Cancer Res. 2012 December 15; 18(24): 6758–6770.



Predicting Response

Specific lymphocyte subsets predict response to adoptive cell therapy using expanded autologous tumor-infiltrating lymphocytes in metastatic melanoma patients

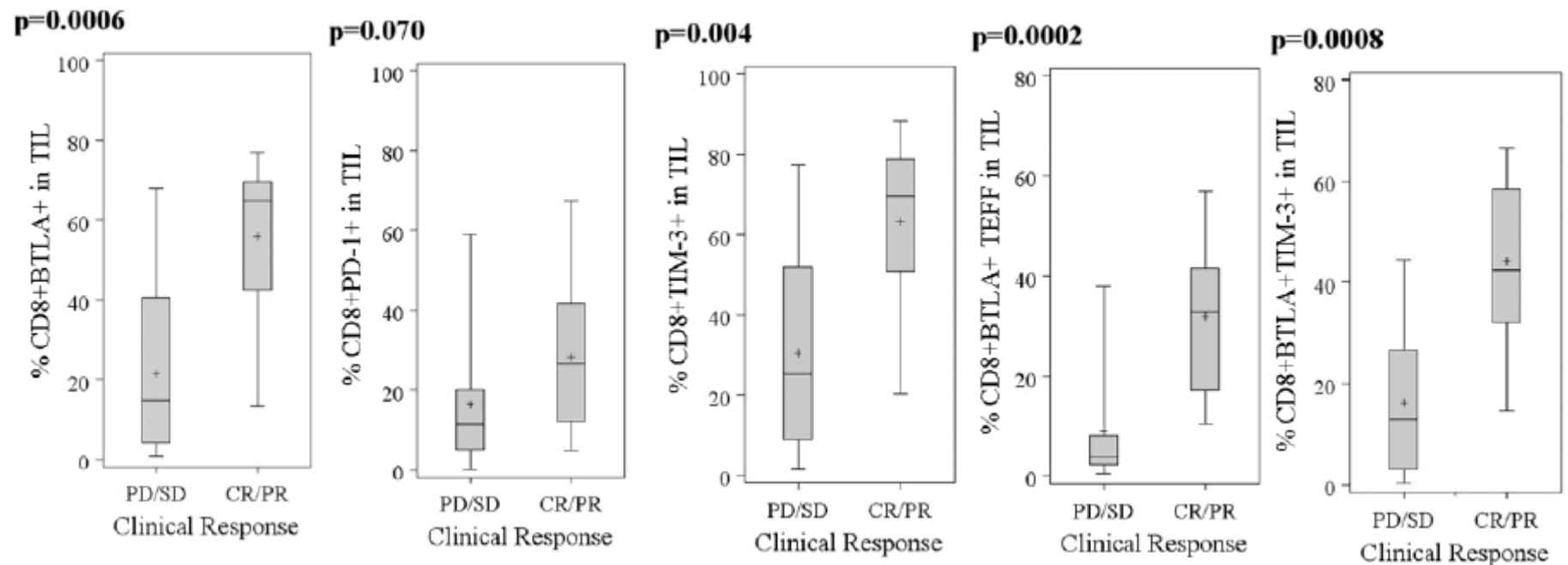
Clin Cancer Res. 2012 December 15; 18(24): 6758–6770.



Predicting Response

Specific lymphocyte subsets predict response to adoptive cell therapy using expanded autologous tumor-infiltrating lymphocytes in metastatic melanoma patients

Clin Cancer Res. 2012 December 15; 18(24): 6758–6770.



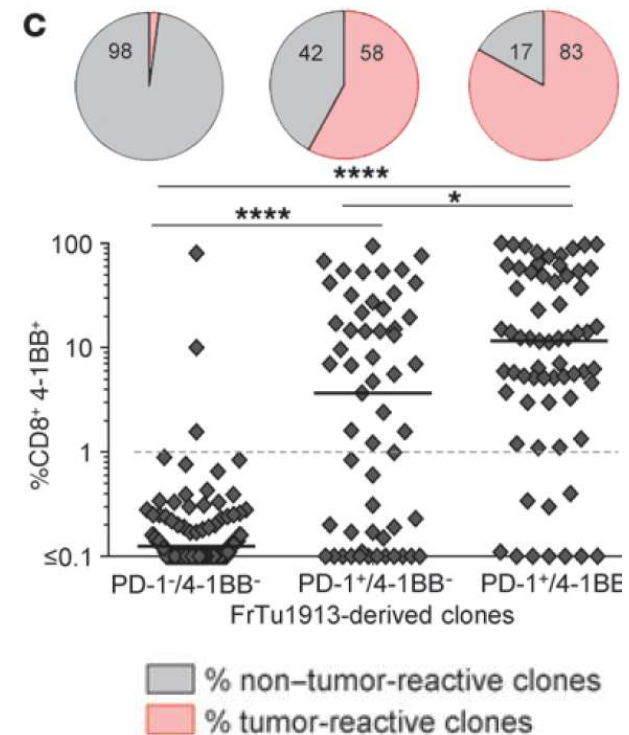
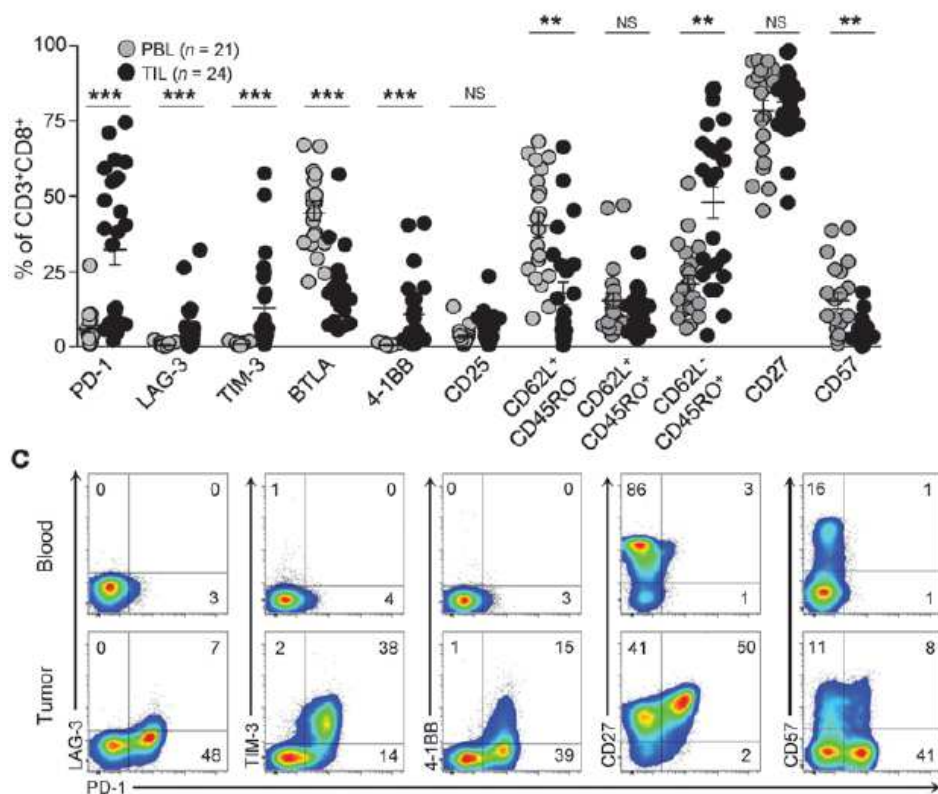
Predicting Response

PD-1 identifies the patient-specific CD8⁺ tumor-reactive repertoire infiltrating human tumors

Alena Gros,¹ Paul F. Robbins,¹ Xin Yao,¹ Yong F. Li,¹ Simon Turcotte,¹ Eric Tran,¹ John R. Wunderlich,¹
Arnold Mixon,¹ Shawn Farid,¹ Mark E. Dudley,¹ Ken-ichi Hanada,¹ Jorge R. Almeida,² Sam Darko,²
Daniel C. Douek,² James C. Yang,¹ and Steven A. Rosenberg¹

The Journal of Clinical Investigation

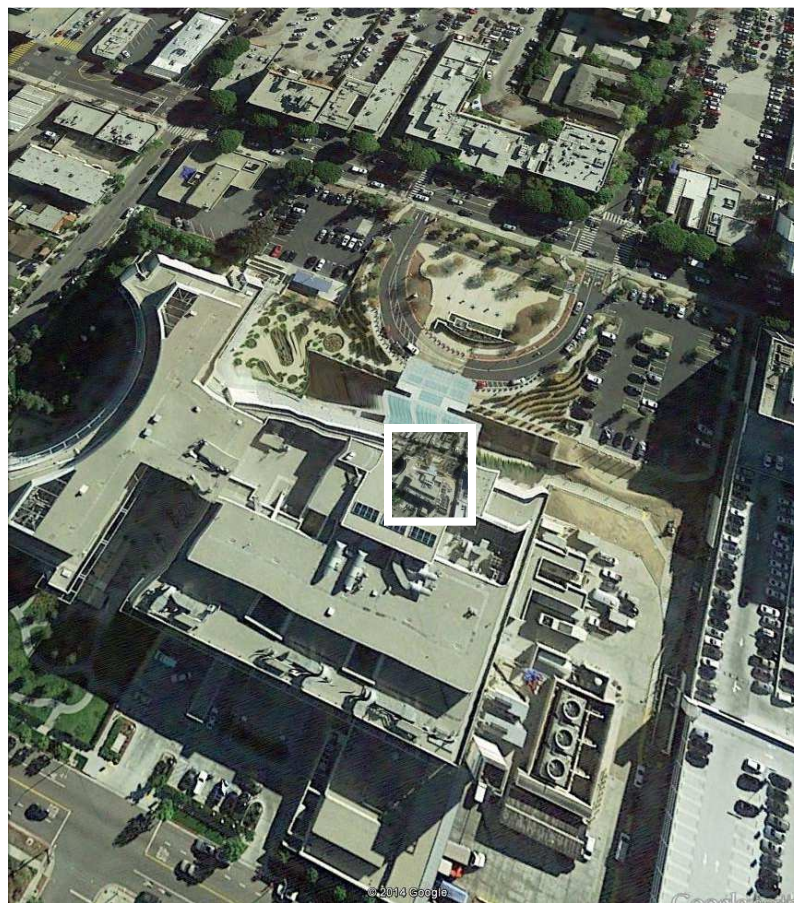
May 2014



Tumor-Infiltrating Lymphocytes: Challenges

- **Technical challenges: (“GMP”)**
- **Timeline**
- **Tumor reactivity**
- **Interleukin-2**

An aerial photograph of a large university campus. The campus features a variety of buildings, including large lecture halls, smaller administrative buildings, and a prominent central building with a distinctive architectural design. A large, oval-shaped track is visible on the left side of the image. The campus is surrounded by green spaces and trees, and a residential area is visible in the upper right corner. A north arrow is located in the top right corner of the image.



Cell Production Facility



Cell Production Facility



Tumor-Infiltrating Lymphocytes: Challenges

- Technical challenges: (“GMP”)
- **Timeline**
- Tumor reactivity
- Interleukin-2

Tumor-Infiltrating Lymphocytes: Challenges

- Technical challenges: (“GMP”)

- **Timeline**



Tumor Removal

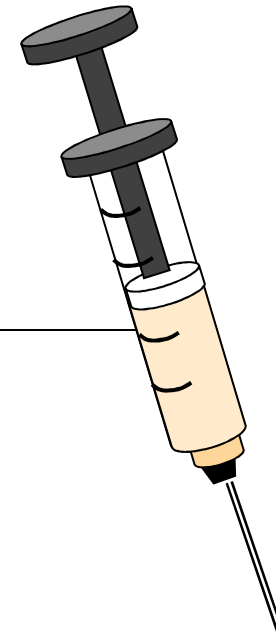
2-4 weeks

Testing



2 weeks

5-8 weeks



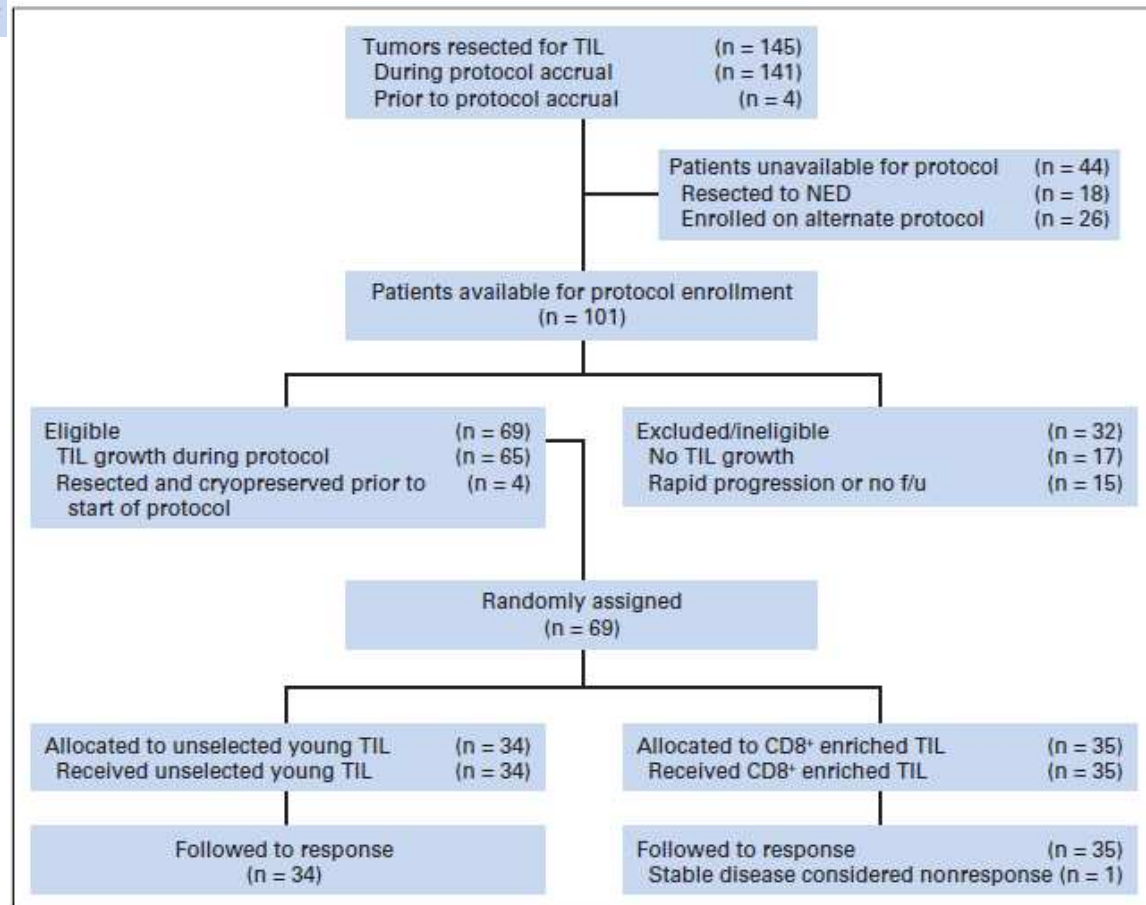
Young TIL

Randomized Selection Design Trial Evaluating
CD8⁺-Enriched Versus Unselected Tumor-Infiltrating
Lymphocytes for Adoptive Cell Therapy for Patients
With Melanoma

JUNE 10 2013

Mark E. Dudley, Colin A. Gross, Robert P.T. Somerville, Young Hong, Nicholas P. Schaub, Shannon F. Rosati,
Donald E. White, Debbie Nathan, Nicholas P. Restifo, Seth M. Steinberg, John R. Wunderlich,
Udai S. Kammula, Richard M. Sherry, James C. Yang, Giao Q. Phan, Marybeth S. Hughes,
Carolyn M. Laurencot, and Steven A. Rosenberg

JOURNAL OF CLINICAL ONCOLOGY



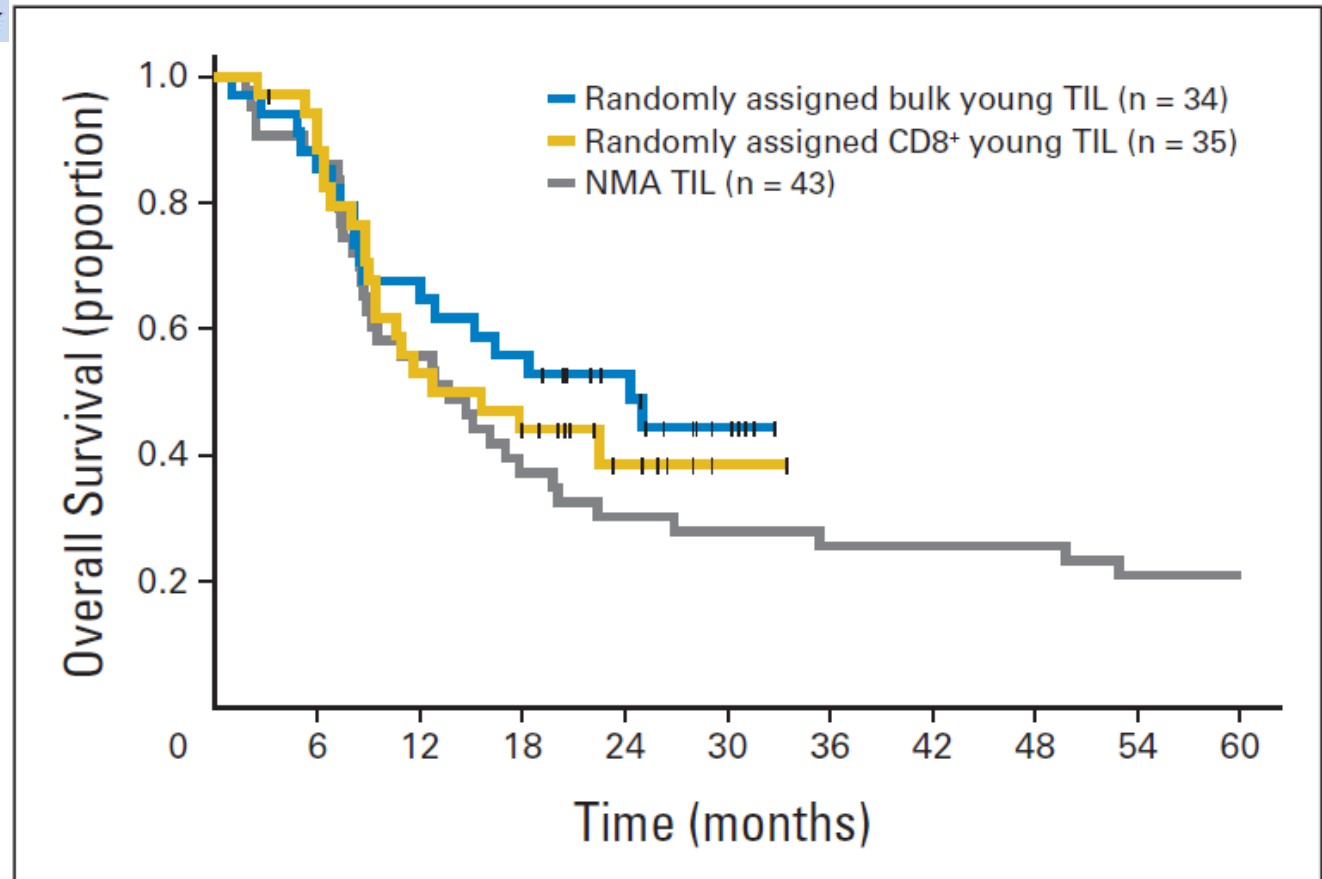
Young TIL

Randomized Selection Design Trial Evaluating
CD8⁺-Enriched Versus Unselected Tumor-Infiltrating
Lymphocytes for Adoptive Cell Therapy for Patients
With Melanoma

JUNE 10 2013

Mark E. Dudley, Colin A. Gross, Robert P.T. Somerville, Young Hong, Nicholas P. Schaub, Shannon F. Rosati,
Donald E. White, Debbie Nathan, Nicholas P. Restifo, Seth M. Steinberg, John R. Wunderlich,
Udai S. Kammula, Richard M. Sherry, James C. Yang, Giao Q. Phan, Marybeth S. Hughes,
Carolyn M. Laurencot, and Steven A. Rosenberg

JOURNAL OF CLINICAL ONCOLOGY



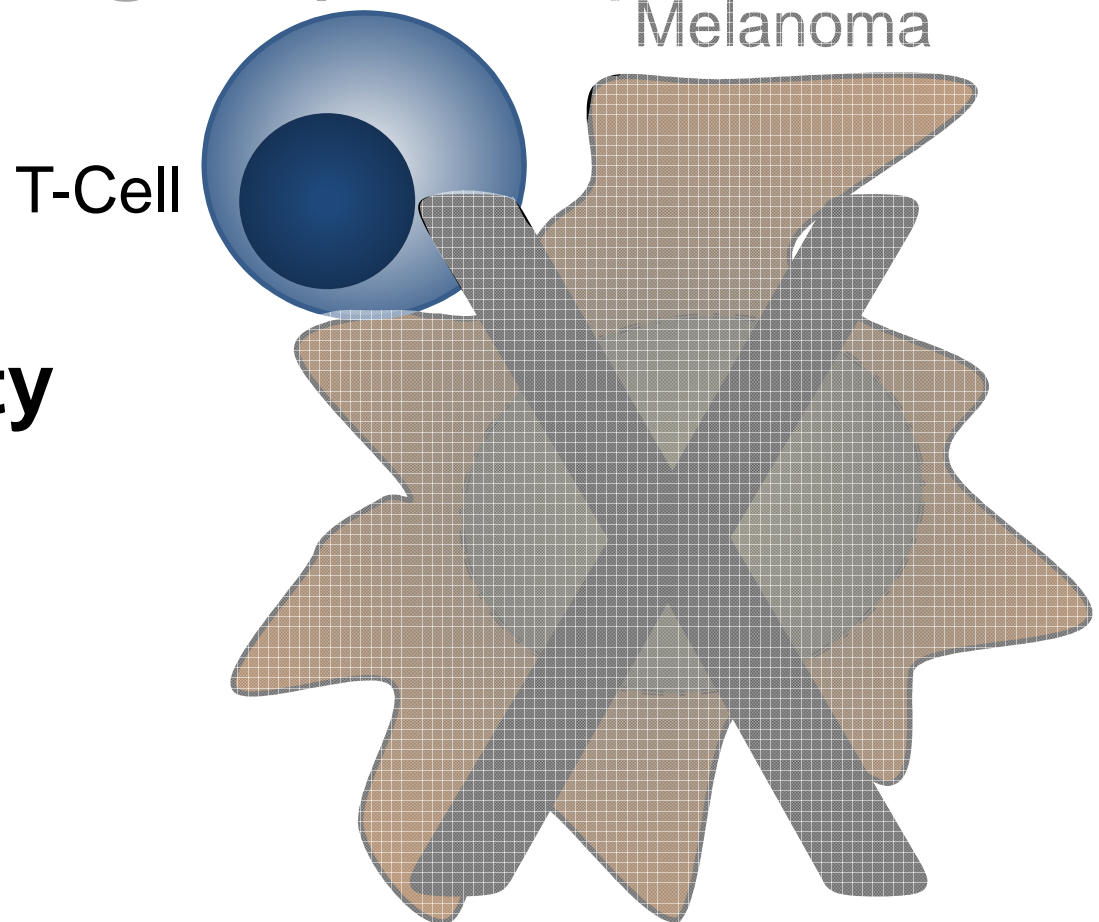
Tumor-Infiltrating Lymphocytes: Challenges

- Technical challenges: (“GMP”)

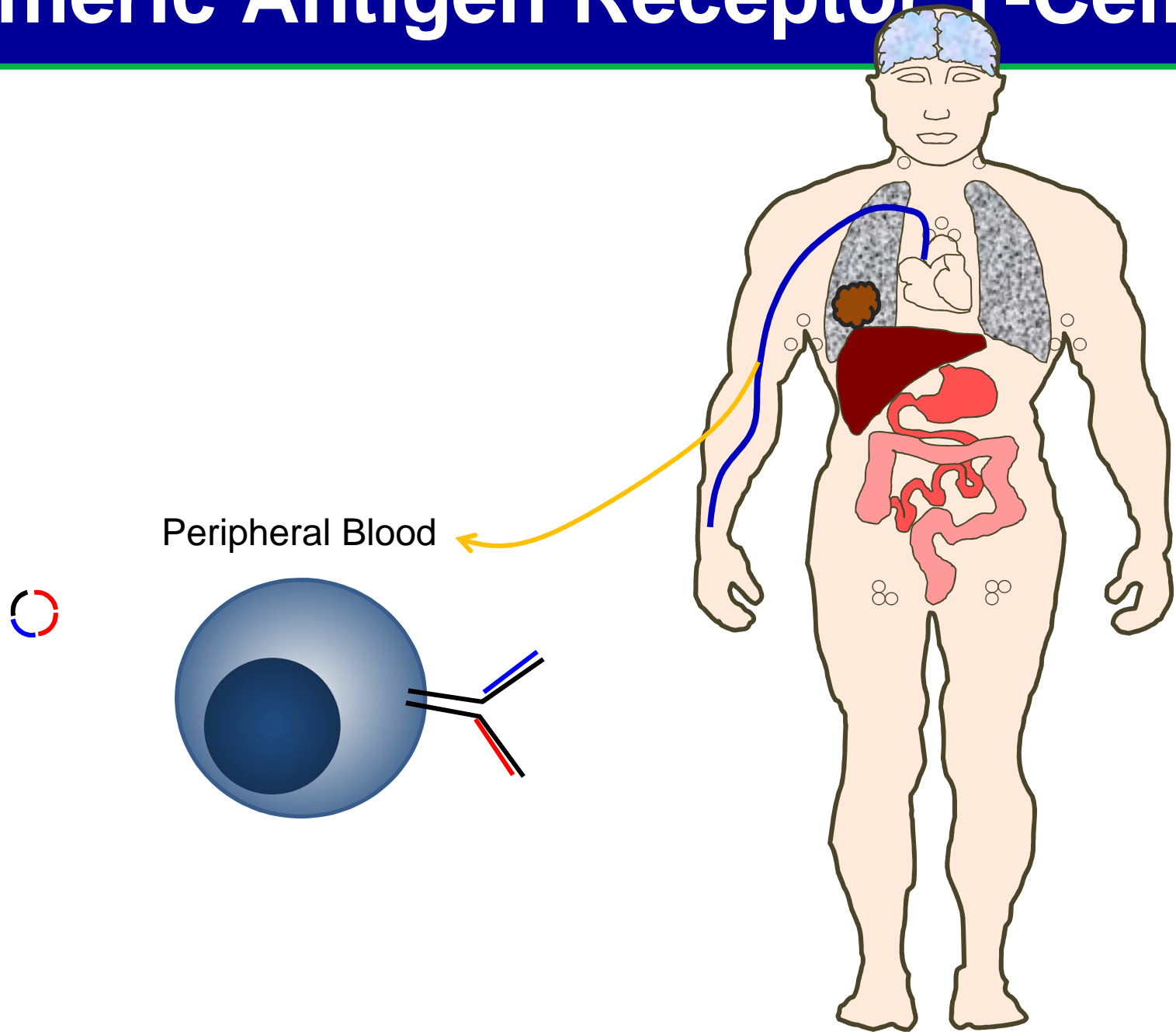
- Timeline

- **Tumor reactivity**

- Interleukin-2



Chimeric Antigen Receptor T-Cells



Chimeric Antigen Receptor T-Cells

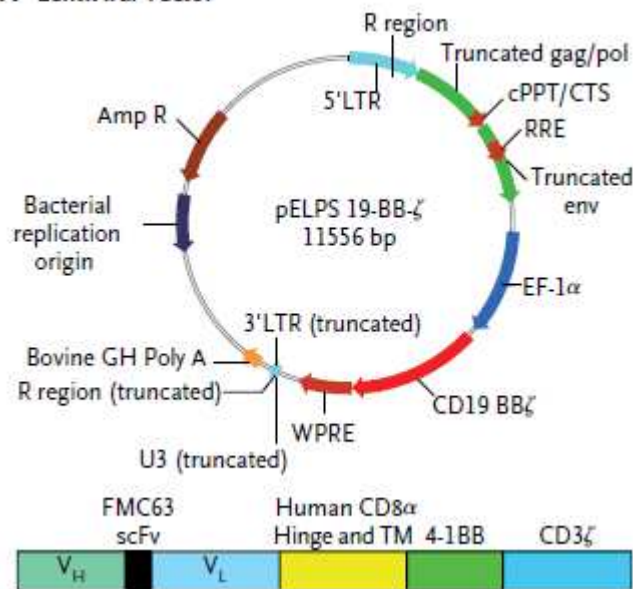
Chimeric Antigen Receptor–Modified T Cells in Chronic Lymphoid Leukemia

David L. Porter, M.D., Bruce L. Levine, Ph.D., Michael Kalos, Ph.D.,
Adam Bagg, M.D., and Carl H. June, M.D.

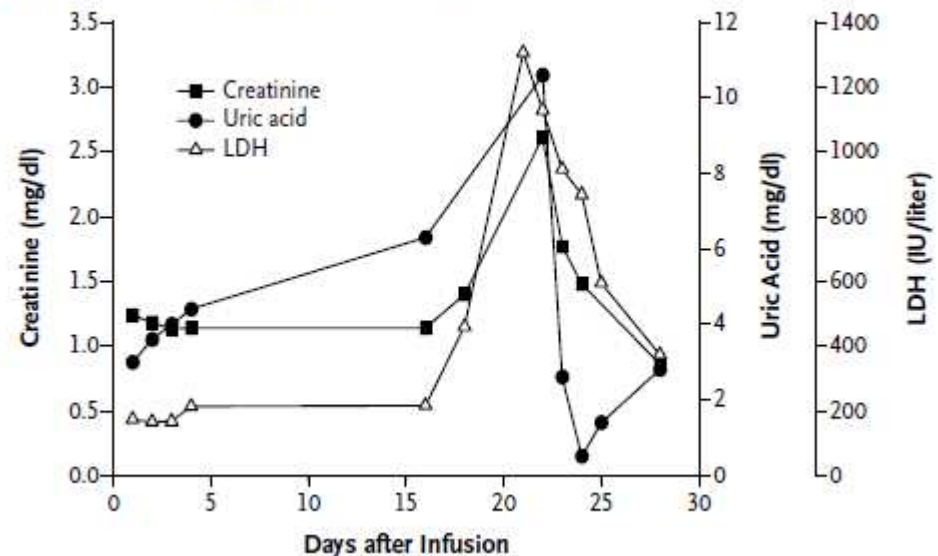
The NEW ENGLAND JOURNAL of MEDICINE

August 10, 2011,

A Lentiviral Vector



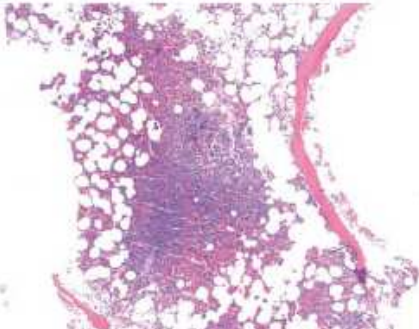
B Serum Creatinine, Uric Acid, and LDH



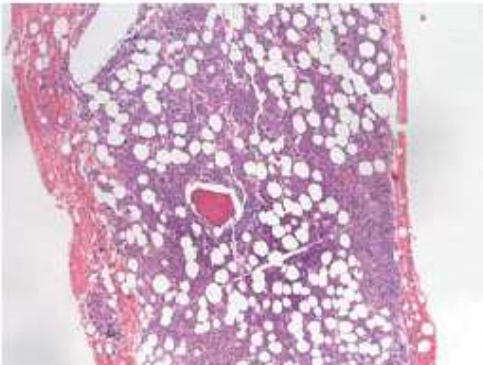
Chimeric Antigen Receptor T-Cells

C Bone Marrow–Biopsy Specimens

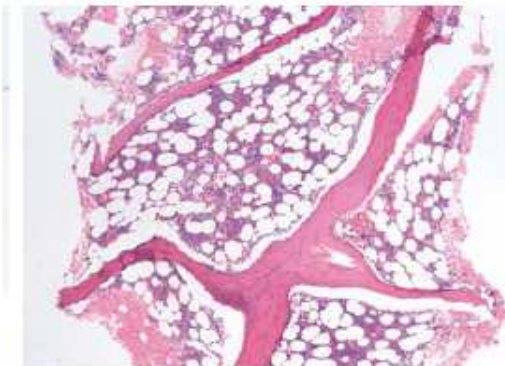
Day –1 (baseline)



Day 23



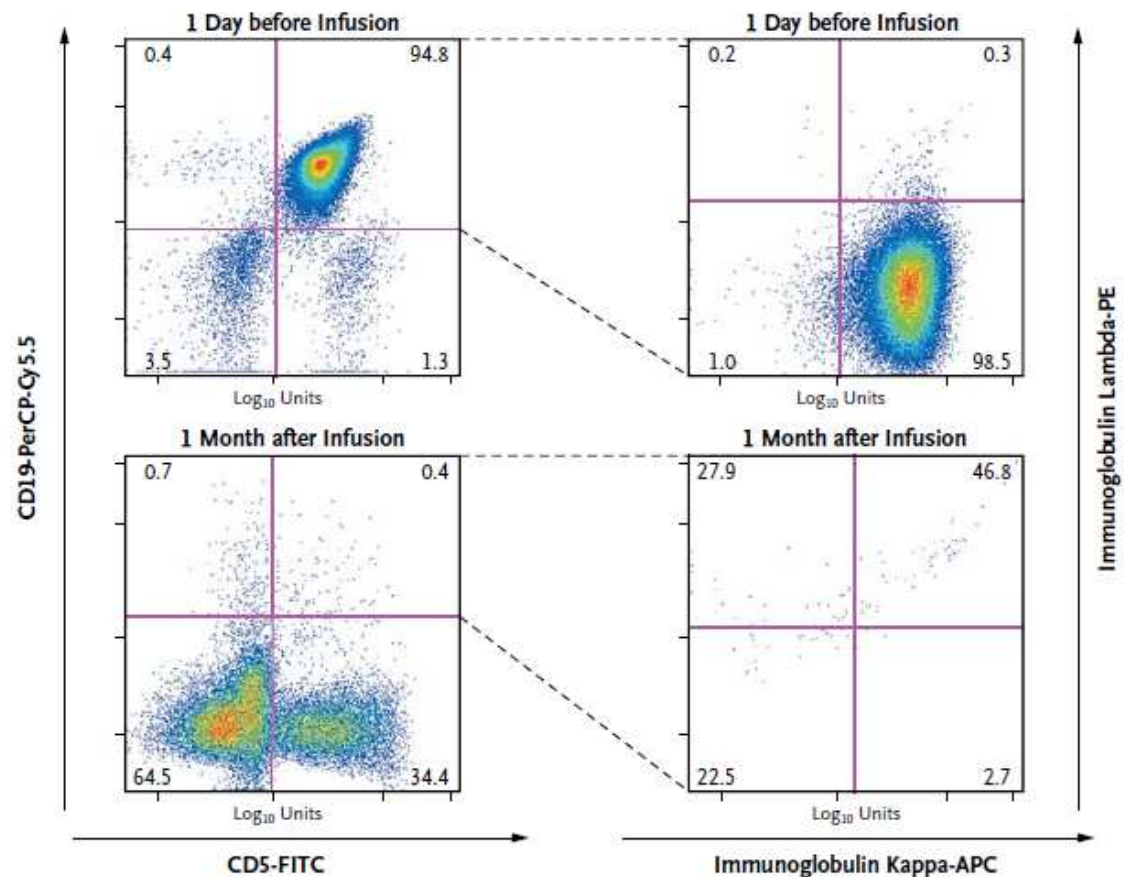
6 Mo



Chimeric Antigen Receptor–Modified T Cells in Chronic Lymphoid Leukemia

The NEW ENGLAND JOURNAL of MEDICINE

August 10, 2011,



Chimeric Antigen Receptor T-Cells

Chemotherapy-Refractory Diffuse Large B-Cell Lymphoma
and Indolent B-Cell Malignancies Can Be Effectively
Treated With Autologous T Cells Expressing an Anti-CD19
Chimeric Antigen Receptor

*James N. Kochenderfer, Mark E. Dudley, Sadik H. Kassim, Robert P.T. Somerville, Robert O. Carpenter,
Maryalice Stetler-Stevenson, James C. Yang, Giao Q. Phan, Marybeth S. Hughes, Richard M. Sherry,
Mark Raffeld, Steven Feldman, Lily Lu, Yong F. Li, Lien T. Ngo, Andre Goy, Tatyana Feldman,
David E. Spaner, Michael L. Wang, Clara C. Chen, Sarah M. Kranick, Avindra Nath, Debbie-Ann N. Nathan,
Kathleen E. Morton, Mary Ann Toomey, and Steven A. Rosenberg*

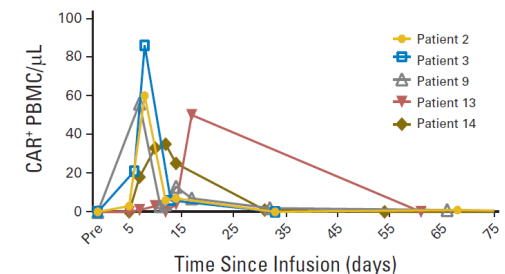
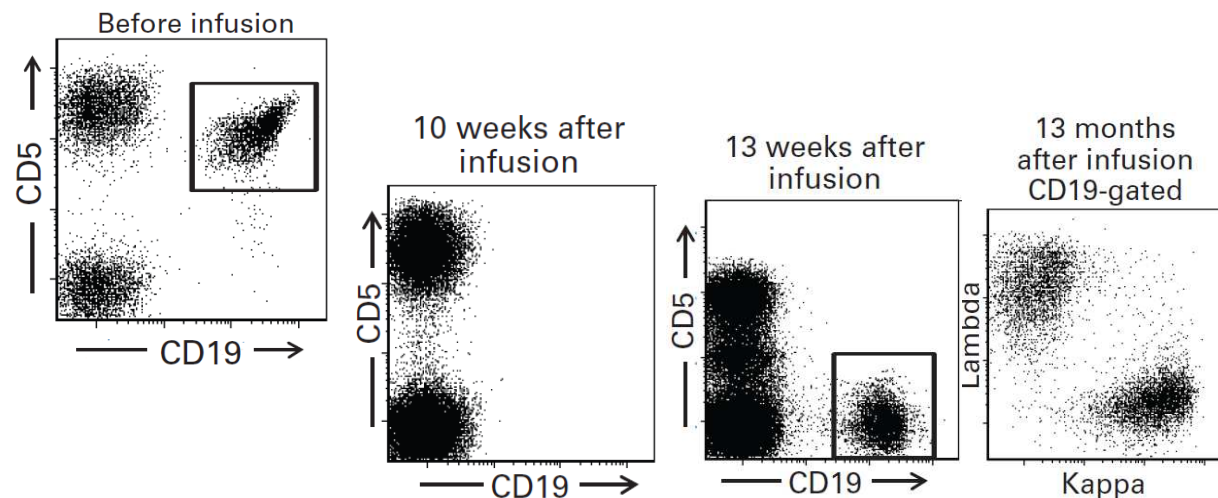
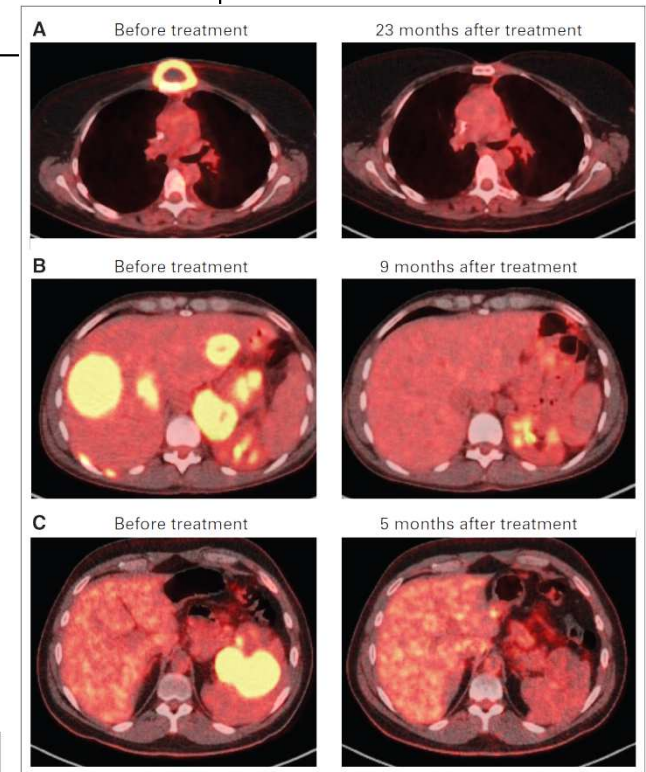
JOURNAL OF CLINICAL ONCOLOGY Published Ahead of Print on August 25, 2014



Chimeric Antigen Receptor T-Cells

Chemotherapy-Refractory Diffuse Large B-Cell Lymphoma and Indolent B-Cell Malignancies Can Be Effectively Treated With Autologous T Cells Expressing an Anti-CD19 Chimeric Antigen Receptor

- 15 patients (9 DLBCL, 6 indolent B-cell malign.)
- 1 Death and several neurologic toxicities
- 4/7 DLBCL with CR, 2 PR, 1 SD
- Overall RR: 12/13 (92%) (9 responses ongoing)



Tumor-Infiltrating Lymphocytes: Modifications

5/9/2014

Patient's Cells Deployed to Attack Aggressive Cancer - NYTimes.com

The New York Times | <http://nyti.ms/1kPowfQ>

BELLE
NOW PLAYING
GET TICKETS

HEALTH | NYT NOW

Patient's Cells Deployed to Attack Aggressive Cancer

By DENISE GRADY MAY 8, 2014

Doctors have taken an important step toward a long-sought goal: harnessing a person's own immune system to fight cancer.

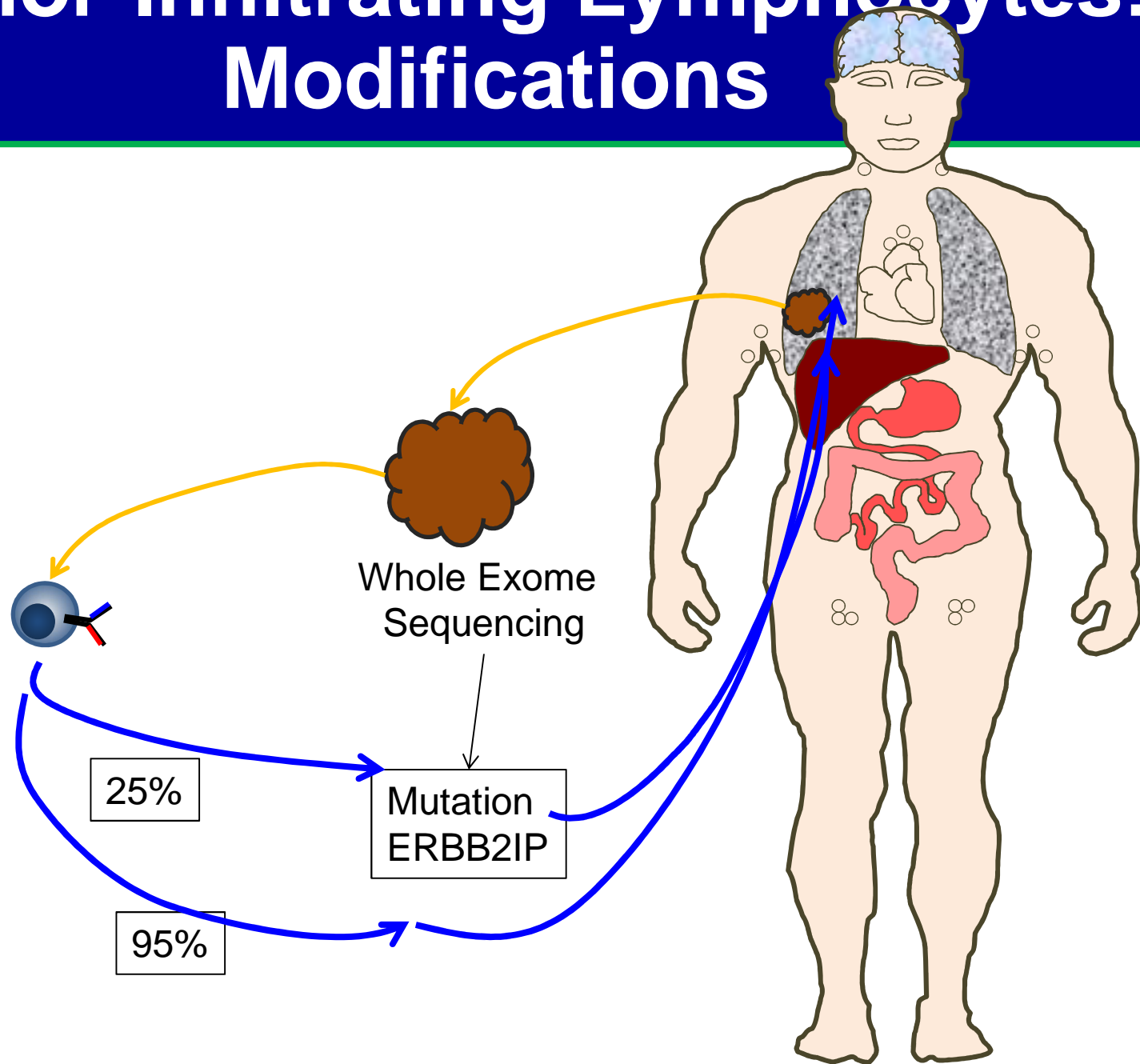
An article published Thursday in the journal Science describes the treatment of a 43-year-old woman with an advanced and deadly type of cancer that had spread from her bile duct to her liver and lungs, despite chemotherapy.

Researchers at the National Cancer Institute sequenced the genome of her cancer and identified cells from her immune system that attacked a specific mutation in the malignant cells. Then they grew those immune cells in the laboratory and infused billions of them back into her bloodstream.

The tumors began "melting away," said Dr. Steven A. Rosenberg, the senior author of the article and chief of the surgery branch at the cancer institute.

The woman is not cured: Her tumors are shrinking, but not gone. And an experiment on one patient cannot determine whether a new treatment works. But the report is noteworthy because it describes an approach that

Tumor-Infiltrating Lymphocytes: Modifications



TIL Opportunities: Combination

- TIL Induction
 - BRAFi
 - XRT
 - Vaccination

BRAF Inhibition Is Associated with Enhanced Melanoma Antigen Expression and a More Favorable Tumor Microenvironment in Patients with Metastatic Melanoma

Microenvironment in Patients with Metastatic Melanoma

Dennie T. Frederick¹, Adriano Piris³, Alexandria P. Cogdill¹, Zachary A. Cooper¹, Cecilia Lezcano⁶, Cristina R. Ferrone¹, Devarati Mitra⁴, Andrea Boni¹, Lindsay P. Newton¹, Chengwen Liu⁷, Weiyi Peng⁷, Ryan J. Sullivan², Donald P. Lawrence², F. Stephen Hodi⁵, Willem W. Overwijk⁷, Gregory Lizée⁷, George F. Murphy⁶, Patrick Hwu⁷, Keith T. Flaherty², David E. Fisher⁴, and Jennifer A. Wargo¹

Clin Cancer Res; 19(5); 1225–31.

January 10, 2013;

