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Table S1. The current IOTN awards.

Project	Project Title	Award Organization	PIs
Number	, and the second	_	(Lead PI in bold)
UG3CA244697	Intercepting Progression from Pre-	Weill Medical College of	N.K. Altorki, A.
	invasive to Invasive Lung	Cornell University	Borczuk,
	Adenocarcinoma		O.Elemento, T.
			Mcgraw, V. Mittal
U01CA233097	Epithelium-Derived Alarmins Role in	Massachusetts General	S. Demehri
	Breast Cancer Immunoprevention	Hospital	
UG3CA244687	Recurrent Tumor-specific Alternately	University of Minnesota	D. Largaespada
	Processed Transcripts as a Source of		
	Neoantigens for NF1-associated		
	Malignant Peripheral Nerve Sheath Tumor Immunoprevention		
U01DE029255	Robust Immuno-prevention Strategies	University of Michigan	Y. L. Lei, J.C.
UU1DE029233	for High-risk Oral Epithelial Dysplasia	University of Wilchigan	Brenner, N. Neamati
U01CA233056	Neoantigen Vaccination for Lynch	Weill Medical College of	S. Lipkin, E. Vilar-
001CA233030	Syndrome Immunoprevention	Cornell University / MD	Sanchez
	Syndrome minimioprevention	Anderson Cancer Center	Sunchez
U01CA244452	Mechanisms of Exosome Driven	University of California,	R. Blelloch, L. Fong
	Immunoregulation of Cancer Progression	San Francisco	
U01CA239258	Enhancing Cell Therapy for Brain	Children's Research	C. Bollard, C.R.
	Tumors	Institute / Johns Hopkins	Cruz, R. Jones, B.
		University / University of	Savoldo
		North Carolina at Chapel	
		Hill	
U01CA233096	Cytokine Immunotherapies for	Yale University	M. Bosenberg, A.
	Melanoma		Ring
U01CA232758	B Cell-Dependent Anti-Tumor Immunity	H. Lee Moffitt Cancer	J. Conejo-Garcia
	in Ovarian Cancer	Center & Research	
U01CA233078	N.Cl. and Istina and Issues and Issues Com-	Institute	M. Demetriou
U01CA2330/8	N-Glycosylation and Immunotherapy for Cancer	University of California, Irvine	M. Demetriou
U01CA244314	Human CD3epsilon Co-potentiation to	University of Missouri-	D. Gil Pagés
0010/1244514	Boost Immunotherapy	Columbia	D. On 1 ages
U01CA244291	Optimizing Myeloma-specific Immunity	Fred Hutchinson Cancer	G. Hill
001011211291	After Autologous Stem Cell	Research Center	O. IIII
	Transplantation		
U01AA027681	Immunosuppressive Mechanisms	University of California,	M. Karin, A. El-
	Responsible for Development of Non-	San Diego / University of	Khoueiry, S.
	Viral Liver Cancer and Control of its	Southern California	Shalapour, H.
	Response to Immune Checkpoint		Tsukamoto
	Inhibitors		
U01CA233084	MUC1-C is a Target for Reversing	Dana-Farber Cancer	D. Kufe, K.K. Wong
	Immune Evasion and Resistance to	Institute / New York	
******	Immunotherapies	University	7.17
U01CA233102	Immunomodulation of the Tumor	University of Wisconsin –	Z. Morris, J.
	Microenvironment with Molecular	Madison	Weichert
	Targeted Radiotherapy to Facilitate an		
	Adaptive Anti-Tumor Immune Response to Combined Modality Immunotherapies		
U01CA233085	Reprograming the Tumor	Roswell Park	K. Odunsi, A.
0010A233003	Microenvironment to Overcome	Comprehensive Cancer	Gambotto, D.B.
	Multiple Primary and Acquired Immune	Comprehensive Cancer	Kozbor
	intercipie i ilmary and Acquired ilmilule		1102001

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	Resistance Mechanisms in Ovarian	Center / University of	
	Cancer	Pittsburgh	
U01DE028227	Stimulating Neo-Antigen Specific T Cell	La Jolla Institute for	S. Schoenberger,
	Responses in Head and Neck Cancers	Allergy and Immunology /	J.S. Gutkind, A. Rao
		University of California,	
		San Diego	
U01DE028233	Targeting the Immunosuppressive	Baylor College of	A. Sikora, A.
	Tumor Microenvironment to Enhance	Medicine	Annapragada
	Efficacy of Radiotherapy and Immuno-		
	Radiotherapy for Oral Cancer		
U01DE029188	Defining Mechanisms of Immunotherapy	Dana-Farber Cancer	R. Uppaluri, D.
	Resistance in Head and Neck Squamous	Institute	Barbie, R. Haddad
	Cell Carcinomas		
U01CA233100	Molecular and Immune Drivers of	Dana-Farber Cancer	E. Van Allen, L.
	Immunotherapy Responsiveness in	Institute / University of	Fong
	Prostate Cancer	California, San Francisco	-
U01CA233074	Targeting Alternative Splicing for TCR	University of California,	O. Witte, G. Crooks,
	Discovery in Small Cell Carcinomas	Los Angeles / Children's	Y. Xing
		Hospital of Philadelphia	
U54CA244719	Nano-immuno-oncology Approaches to	University of Texas	J. Gao, Z. Chen
	Overcome Tumor Immune Evasion	Southwestern Medical	
		Center	
U54CA244711	Engineering the Next Generation of T	University of	C. June, G. Linette,
	Cells	Pennsylvania	M. Milone
U54CA244438	UCSF Center for Synthetic Immunology:	University of California,	W. Lim, T. Desai,
	Tools to Reprogram the Immune System	San Francisco	K. Roybal
	to Combat Cancer		
U54CA244726	Biomaterials to Create T Cell Immunity	Harvard University /	D. Mooney, F. Hodi,
		Dana-Farber Cancer	D. Scadden, W.
		Institute	Shih, C. Wu
U01AR077511	Identification of Pathways to Mitigate	National Jewish Health /	D.Y. Leung, J.A.
	Immune-Related Adverse Events with	Memorial Sloan Kettering	Kern, M. Lacouture
	Cancer Immunotherapy	Cancer Center	
U01CA247573	Engineering Immunotherapeutic	Columbia University	T. Danino
	Probiotics to Mitigate irAEs		
U24CA232979	IOTN: Data Management and Resource-	Roswell Park	A. Hutson, S. Liu,
	Sharing Center	Comprehensive Cancer	M. Morgan, K.
		Center	Odunsi
U24CA233032	IOTN: Cellular Immunotherapy Data	Medical College of	M. Pasquini
	Resource (CIDR)	Wisconsin	

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Table S2. The IOTN initiatives.

FOAs	IOTN Initiative	Length of Initiative	Goal
RFA-CA-17-045/ RFA-CA-19-015	Cancer Immunotherapy Research Projects (U01)	5 years	Improve cancer treatment with immune-based approaches by enhancing our understanding of factors that contribute to immunosuppression, including within the immune cells, tumor cells, or tumor microenvironments.
RFA-CA-17-046/ RFA-CA-19-014	Cancer Immunoprevention Research Projects (U01)	5 years	Identify the earliest exploitable changes in the carcinogenic process and develop preclinical immune-based interventions for prevent the development of cancers.
RFA-CA-17-047	Data Management and Resource-Sharing Center (DMRC) (U24)	5 years	Provide overall support, and facilitate collaboration, data management, and data sharing among the IOTN-funded components, as well as promote scientific outreach with other Cancer Moonshot initiatives and the larger scientific community.
RFA-CA-17-048	Cellular Immunotherapy Data Resource (CIDR) (U24)	5 years	Support a data registry for collecting outcomes of patients receiving cellular immunotherapies, that could be utilized for observational studies or inform subsequent pre-clinical studies and clinical trials.
RFA-CA-19-012	Cancer Immunoprevention Research Projects (UG3/UH3)	UG3 - 2 years UH3 - 3 years	Perform exploratory studies aimed at immune target identification and validation, followed by the development and preclinical testing of specific immunoprevention interventions.
RFA-CA-19-013	Immuno-engineering to Improve Immunotherapy (i3) Centers (U54)	5 years	Incorporate innovative engineering solutions to accelerate the preclinical development of immunotherapeutic and immunopreventive interventions that are more effective, safer, and that could benefit a wider patient population.
RFA-CA-19-044	Advancing Cancer Immunotherapy by Mitigating Immune- Related Adverse Events (irAE) (U01)	5 years	Establish a deeper understanding of the origins and activation pathways leading to inflammatory or autoimmune adverse events that currently limit the use of various immunotherapy regimens in patients.