#### **CURRICULUM VITAE**

Date Updated: October 7, 2021

### **Part I: General Information**

Name: Eleftheria Maratos-Flier

Position: Professor Emerita, Harvard Medical School

Director, Translational Medicine, NIBR

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**Education:** 

B.A. New York University, New York, New York

M.D. Mount Sinai School of Medicine

New York, New York

# **Licensure and Certification:**

Massachusetts License Registration American Board of Internal Medicine

# **Academic Appointments:**

1982-1987	Instructor in Medicine	Harvard Medical School
1987-1999	Assistant Professor of Medicine	Harvard Medical School
1999-2010	Associate Professor of Medicine	Harvard Medical School
2010-2018	Professor of Medicine	Harvard Medical School
2018-	Professor Emerita	Harvard Medical School

# **Current Position:**

2018- Director, Translation Medicine

Cardiovascular and Metabolism Translational Medicine

Novartis Institutes for Biomedical Research

# **Hospital or Affiliated Institution Appointments:**

1982-1999 Associate Physician Brigham and Women's Hospital

Boston, Massachusetts.

1987- Active Provisional Staff Beth Israel Deaconess Medical

Center Boston, Massachusetts.

	1982-1983	Research Associate	Joslin Diabetes Center Section of Cellular and Molecular Physiology Boston, MA
	1984-2000	Investigator	Joslin Diabetes Center Section of Cellular and Molecular Physiology Boston, MA
	2000- 2004	Head, Section on Obesity Senior Investigator	Joslin Diabetes Center, Boston, Massachusetts
Other	Professional I	Positions:	
	2001-2018	Affiliate Faculty	Program in Neuroscience Harvard Medical School
	2002-2010	Society Research Fellow	Peabody Society Harvard Medical School
	2005-2018	Associate Advisor	Peabody Society Harvard Medical School
	2006-2010	Director, Role of Discovery in Medicine, Year I course	Harvard Medical School
	2011-2018	Associate Member	Broad Institute
	2018-	Director, Translational Medicine	NIBR, Cambridge
Hospital and Health Care Organization Service Responsibilities:			
	1982-1990 1982-1996 1982-2000 2000-2004 2004-	Diabetes Attending Diabetes Attending Diabetes Outpatient Clinic Diabetes Outpatient Clinic Staff Physician	Joslin Diabetes Center Brigham and Women's Hospital Brigham and Women's Hospital Joslin Diabetes Center Beth Israel Deaconess

# Major Administrative Responsibilities:

1984-1986	Co-Organizer Longwood Medical Area Diabetes Seminar Series
1991-1994	Co-Organizer Longwood Medical Area Diabetes Seminar Series
1994-2002	Director, Molecular Core, Joslin Diabetes Center

1999-2002	Organizer, Longwood Medical Area Diabetes Seminar Series
2002-2004	Director, Animal Physiology Core, Joslin Diabetes Center
2004-2012	Director, Animal Physiology Core, Division of Endocrinology
	Beth Israel Deaconess Medical Center
2006-2008	Associate Director, Endocrine Fellowship, BIDMC
2009-2018	Director, Office for Academic Careers and Faculty Development
	Beth Israel Deaconess Medical Center

# **Major Committee Assignments:**

# **Professional Societies:**

1984-	American Diabetes Association	Member
1987-	The Peabody Society, Harvard Medical School	Member
1996-	The Endocrine Society	Member
2003	Association of American Physicians (AAP)	Elected
2003-2010	NAASO	Member
2005-2016	Society for Neuroscience	Member
2007-2016	American Physiological Society	Member

# **Community Service Related to Professional Work:**

1983-2007	Volunteer science teaching in elementary and high school
1993-2004	Co-manager of academic discussion list on healthcare reform, "Healthre"

# **Editorial Boards:**

1999-2003	Endocrinology
2002-2007	Obesity Research, Associate Editor
2005-2010	Diabetes, Editorial Board
1998-	Ad hoc: Nature Medicine, Cell Metabolism, Proceedings of the National
	Academy of Science, Journal of Clinical Investigation, American Journal
	of Physiology.
2013-	Editorial Board, Molecular Metabolism

#### **Awards and Honors:**

1972	Phi Beta Kappa
1980	Postdoctoral Fellowship, American Cancer Society
1981-1982	Mary K. Iacocca Research Fellowship, Joslin Diabetes Center
1998, 2002	Merck Senior Fellows Award to Maratos-Flier's Fellows
2003	Annual Patricia Usher Memorial Lecture, Beth Israel Deaconess
	Medical Center
2012	Saul Horowitz Distinguished Alumni Award, Mount Sinai School
	of Medicine
2020	Endocrine Society Roy O, Greep Award for Outstanding Research
	Contributions

#### PART II: Research, Teaching and Clinical Contributions

#### A. Narrative Report:

### Report of Research

Overview of Major Research Interests:

- Fibroblast growth factor 21: Regulation of energy expenditure, hepatic metabolism and pancreatic function.
- o Melanin Concentrating Hormone: Physiology and Mechanism of Action
- Hypothalamic peptides and appetite regulation/interaction of leptin with other regulators of eating behavior
- Human translational studies

Obesity is a significant medical problem which is a risk factor for the development multiple illnesses including type II diabetes, cardiovascular disease, peripheral vascular disease, liver disease and several cancers. While obesity results from excess caloric intake in relation to energy expenditure, the underlying causes remain poorly understood. Energy balance is known to be regulated by complex interactions between the periphery and the central nervous system; however, energy balance is also regulated by diet. My major research interest is defining the molecular mediators which regulate inter-relationships between the brain, the periphery and diet thus contributing to the obese state and to the pathologic consequences of obesity.

1: **FGF21**: *Past observations:* Fibroblast growth factor 21 is a member of the endocrine FGF family. It is expressed in several metabolically active tissues including liver, white adipose tissue, brown adipose tissue and pancreas. I first became interested in FGF21 when my group found that hepatic expression was massively induced in mice on a ketogenic diet (KD). Using a gene discovery approach, we identified FGF21 as a critical mediator of fatty acid oxidation. We were first to report this and to show that adenovirus mediated knockdown led to marked fatty liver and serum hypertriglyceridemia in mice eating KD, as mice fail to increase fatty acid oxidation and ketogenesis appropriately. We further demonstrated that FGF21 is required for the normal metabolic response (weight loss and increased energy expenditure) to the diet. We confirmed the role of FGF21 by showing that FGF21 deficient mice show none of the metabolic effects of KD, gaining rather than losing weight and also failing to increase energy expenditure.

My group since demonstrated direct action of FGF21 on the liver, FGF21 resistance in obesity and FGF21 induced browning in inguinal adipose tissue that mimics cold exposure. We were also the first to report on the link between serum FGF21 and non-alcoholic fatty liver disease in humans (NAFLD).

#### Work on FGF21 and MCH:

**FGF21** and **Inflammation:** After finding that FGF21 correlated with NAFLD in humans we speculated that FGF21 might play a role in the progression of NAFLD to non-alcoholic-steatosis and cirrhosis. Using a non-obese model of NAFLD (generated by feeding mice a methionine choline deficient diet) we found that mice lacking FGF21 (FGF21KO) show accelerated progression of NAFLD and develop both increased fibrosis and inflammation. Furthermore, replacement of FGF21 in FGFKO mice consuming the steatotic diet prevents the adverse effects of the diet on the liver.

My group also found that FGF21deletion is associated with pancreatic inflammation. To induce obesity in mice, we use a standard high fat/high sucrose diet. As expected, WT mice become obese and insulin resistant and develop islet cell hyperplasia. However the histology of the acinar pancreas appears normal. When littermate mice lacking FGF21 are placed on the same diet, they develop the same degree of obesity and insulin resistance as the wild type mice. However we found remarkable infiltration of the acinar pancreas in the periductal area with lymphocytes. Histologically these are large, uniform appearing infiltrates that are positive for CD3 antigens, indicating a T-cell infiltrate. Future work in this area would involve immune-characterization of the infiltrates as a prerequisite to understanding the molecular mechanisms involved.

FGF21 and Carbohydrate Metabolism: My group defined a critical role for FGF21 in sugar metabolism and pursued this in both mouse and human studies. In FGF21-KO mice we described a rapid deterioration in mice consuming either high sucrose or high fructose diets, independent of the fat content of the diet. As carbohydrate consumption induces ChREBP we are examining the relationship between this transcription factor and FGF21. This finding led directly to clinical studies in humans. Prior to our work that in normal human subjects, fructose ingestion leads to acute robust (average four fold) increases of FGF21 which are maximal after two hours of ingestion, no acute regulation of FGF21 had been discovered in humans. In obese subjects and in those with metabolic syndrome, baseline levels are increased however there is an additional acute increase after fructose loading leading to a higher area under the curve. The emerging data suggest that FGF21 may serve as a fructose sensor and regulate fructose disposal.

2. **Melanin Concentrating Hormone:** *Past Observations:* My research defined the role of the neuropeptide (MCH), which is exclusively expressed in the lateral hypothalamus, as an orexigenic peptide. I then reported that deletion of MCH was associated with a lean, hypermetabolic phenotype and that overexpression was associated with increased susceptibility to obesity. My group defined intracellular signaling pathways mediated by Gq and Gi coupled receptors and were the first to note that MCH played a role in regulating dopaminergic tone in the accumbens nucleus. We recently generated an MCH receptor-cre mouse which we used to map the expression of the rodent melanin concentrating hormone receptor (MCHR1). We also generated a mouse that co-expresses MCHR1 with MCHR2, (MCHR2 is seen is only expressed in higher vertebrates) and found that MCHR2 acts to limit diet induced obesity and mediates actions distinct from MCHR1. My lab focused on a series of studies aimed at mapping outputs of

the MCH neurons in the lateral hypothalamus. Although neuronal populations with the highest expression of MCH receptor are in the striatum, especially the nucleus accumbens, the synaptic connectivity between the lateral hypothalamus is not particularly active and reports on direct MCH effects on accumbens neurons are inconsistent. We found an active connection between MCH neurons in the lateral hypothalamus and the lateral septum, an area of the brain that is involved in anxiety and reward functions. This work is being continued by a colleague focusing on using an optogenetic and DREADD approach in defining MCH actions in this area.

3. **Human Studies:** As an endocrinologist I am interested in translational studies in humans. Prior to joining Novartis I was awarded investigator initiated funding from Astra-Zeneca to study the mechanisms of weight loss of the diabetes drug exenatide (Byetta) in obese non-diabetic women, focusing on the effects of exenatide on energy expenditure. My second area of interest, noted above, is understanding human FGF21 physiology and the relationship between the FGF21 response to fructose and diet composition, metabolic syndrome and obesity. The work on FGF21 in humans was funded using discretionary funding.

#### **Current Work as TME at Novartis Institutes for Biomedical Research:**

As a member of the cardiovascularmetabolic (CVM) group my work focuses on assets aimed at targeting obesity, T2 Diabetes and cardiovascular disease. I currently direct two active studies that have progressed to human subjects. For one asset I hve been the TA responsible to close out a SAD studu. I was also the director responsible for formulating the clinical study protocol for an upcoming PoC study. This involved multiple aspects of input from line functions such as PK and PCS amd regulatory, interactions with principle investigators at CROs. The protocol was recently approved by the internal review board. A successful update of the investigators brochure was just finalized. The protocol will be submitted shortly to the FDA. I work on a daily basis with the clinical trial operational lead on aspects of data transfer and other issues required. We are aiming for FPFV in December, 2021. A second, earlier stage protocol will move into FIH in late 2022. I also closed out a study that was a successful proof of concept trial. This asset is being out-licensed and a manuscript on the results is in review with the journal Science and Translational medicine. For clinical studies I lead teams which include members with diverse expertise.. I am responsible for evaluating and integrating information relevant to conduct of the studies including safety and efficacy. I am also responsible for obtaining internal board approvals for conduct design, assembling IND to regulatory authorities, writing protocols and investigators brochures, managing incoming data from clinical studies, writing final study reports. I also am responsible for direct interactions with principle investigators at CROs as well as influencing choice of CROs.

My other significant role involves is my regular interaction with pre-clinical scientists in the CVM space. Given my academic background I function as a local KOL providing information on the physiologic and mechanistic contributors to obesity and metabolic dysregulation. I contribute to decisions to on choice of targets to evaluate. I am also a member of the pre-clinical teams evaluating drug candidates in pre-clinical models to ensure adequate information has been acquired to facilitate transition to clinical study. In this role I am active on two early teams with assets aimed at the treatment of metabolic disease.

# **Professional Development and Teaching:**

**Academic Careers and Faculty Development:** In late 2009 I took on the role of directing this office at Beth Israel Deaconess. This was a new office, mirroring a trend in other hospitals, aimed at providing structured opportunities for faculty. I was one of several senior level faculty who applied and I was offered the position after interviews with members of the BIDMC center for education and after submitting a proposal describing my plans for "growing" a program.

**Programs:** In this role I put in place multiple programs; these ranged from 60-90 minute single sessions to day long programs. A sample of topics covered included:

(Myself as presenter, 2-3 times a year)

- Giving a Lecture
- Making it to the next step for Postdocs
- Making it to the next step for Instructors
- Publishing your paper
- Assembling RO1 (three sessions)

#### Other presenters:

- Negotiating
- Hiring and Firing
- Promotion to Assistant Professor
- Promotion to Associate Professor
- Promotion to Professor
- Taxes for Foreign Nationals
- Work Life Balance
- Making the Most out of a Meeting as Chair or Participant

#### Workshops with multiple presentations:

- Junior Faculty Leadership Workshop (one day)
- Assembling a K award (one day, twice a year)

On a yearly basis, 500 post-doctoral fellows and faculty attend these session. The overall rating for quality and utility averages a 4.3 on a scale of 1-5 where 5 is the best and 1 is the worst.

#### **Guideline Development:**

In 2014 I was asked by Dr. Richard Schwartzstein to convene a task force to develop guidelines for post-doctoral fellows training at Beth Israel Deaconess Medical Center, including MDs, PhD, citizens and foreign nationals. I convened a committee of faculty which included a department chair, a division chief, and a range of senior to junior faculty to develop these guidelines. We solicited input from research administration, BIDMC legal and human resources. In addition to guidelines involving responsibilities and benefits we also assembled the forms and procedures required to meet the guidelines such as annual review forms, salary forms, appointment extension forms and vacation tracking forms. The guidelines will be presented to BIDMC leadership in March for approval, pending feedback from the leadership group.

Once the guidelines are approved, I planned assembled a ½ day postdoctoral fellow orientation which would be offered three times a year and will required attendance for incoming postdoctoral fellows. The next project in this area was to developed a set of guidelines for mentors/mentees aimed at improving this occasionally difficult relationship. These planned activities ended when I transitioned to Novartis.

# **Advising:**

From 2010 until 2018 I consulted with faculty for career advising. I saw an average of six faculty members per month for "one on one" advice. The faculty queries range from a review of the Harvard format Curriculum Vitae to specific concerns about readiness for promotion to seeking advice about problematic mentor relationships.

# **Teaching:**

Between 2007 and 2010, I was involved in developing and co-directing a new, one month, required course for first year HMS students, "the Role of Discovery in Medicine". The goal of this course is to highlight the critical importance of scholarly work in changing medical paradigms and medical practice and to examine methodology essential to the process of discovery. This course has evolved and is now an elective for students planning to submit proposals for funding from HMS for either bench or clinical/translational research. When I initially worked on the course, two diseases, Obesity/Type II diabetes and Breast Cancer were used as disease paradigms in seminars aimed at assisting students in assembling high quality research proposals. A series of lectures in the fall covered such topics such as approaches to clinical versus bench research, finding a lab and writing a research proposal.

I also provided ongoing supervision of 4-6 post-doctoral research fellows in my laboratory. I have supervised senior thesis of three seniors at Harvard college. I mentor and acted as a senior advisor to two junior level investigators. I also participate as an invited speaker in the IHP course at Harvard Medical School.

I regularly teach in the upper level Endocrinology course organized by Dr. Monty Krieger at MIT.

# **Clinical Contributions:**

I see outpatients with diabetes one afternoon monthly a diabetes obesity specialist. This clinical activity takes place at Beth Israel Deaconess Medical Center. Approximately 90% of my patients involve long term follow up. Each afternoon clinic generates an addition two to three hours of follow up time in the form of patient phone calls and communication with referring physicians. Diabetes care requires particularly intensive phone follow up for the purpose of blood sugar monitoring.

<b>B. Research Funding Information</b> (entirely for bench research).				
Years	Source	<u>PI</u>	Grant Title	

Completed:

1983-1984	Juvenile Diabetes Foundation 82R609		PI	Viral Interactions with Beta Cells
1984-1989	NIH Clinical Investigator Award 5K08AM1252		PI	Viral Pathogenesis in Endocrine Cells
1984-1986	American Diabetes Associ	ation	PI	Cellular Processing of Reovirus and Infection of Islet Cells
1989-1992	American Heart Association	PI	Viral Cells	Receptors on Endocrine
1990-1995	NIH R01 A128971-05	PI		cular Characterization Reovirus Receptor
1994-1999	Markey Charitable Trust	PI		Sequencing & Synthesis Core
1995-1996	Boston Obesity Research Center NIH 5P30DK46200-04	PI	Hypo	fication of Novel thalamic Genes that are rentially Regulated in Obesity
1996-1997	American Diabetes Association	PI		nin Concentrating Hormone: A vel Orexigenic Neuropeptide
1996-1998	Eli Lilly and Company	PI	•	directed to Obesity, Energy eostasis and Metabolic Control
1996-1997	NIH 5P30DK36836-11	PI		: A Novel Regulator of eding Behavior
1998-2003	NIH 1R01DK53978	PI		omic and Physiologic acterization of MCH Action
1999-2004	NIH 1R01DK56113	PI		Targeting Approaches to MCH nergy Balance
2000-2010	NIH 1P01DK56116-06	PI	B. Ka	am Project Grant with hn PI PI on Project IV and Core C
2007-2010	Amylin Pharmaceuticals	PI	Loss,	Effect of Exenatide on Weight Energy Expenditure and er in Obese Women Without

# Diabetes

2005-2011	NIH R01DK069983-05	PI	Actions of MCH in the Brain
2011-2012	Jeffrey B Picower Foundation	PI	Molecular mediators of FGF21 action
Ongoing:			
2011-2019	NIH R01DK028082-29	PI	Metabolic Actions of FGF21
2012-2016	Amylin Pharmaceuticals	PI	Patterns, predictors and mechanisms Of weight loss with exenatide treatment in overweight and obese women without diabetes.

# C. Report on Current Research Activities:

# Project:

My current research activities focus on translational studies related to obesity, type 2 diabetes and lipid disorders at the Novartis Institutes for Biomedical Research. I serve as translational medicine expert leading teams to transition potential drugs from pre-clinical studies into humans.

# **Report on Teaching**

# **Local Contributions**

a. Medical School Courses

1982-1996	Teaching of residents and fellows while Attending Physician at Brigham and Women's Hospital, Boston, Massachusetts.
1982- 2004.	Supervised research training of 1-3 M.D. Postdoctoral Fellows, medical and college students and Research Assistants at the Joslin Diabetes Center, Boston, Massachusetts
2004-2018	Supervised research training of 4-6 MD or PhD postdoctoral fellows, medical students and undergraduates in the Endocrine Division of Beth Israel Deaconess Medical Center
1984-1988	First year medical tutor at Harvard Medical School, Boston, Massachusetts.
1988-1992	Tutor in Patient/Doctor I, Harvard Medical School This course meets weekly throughout the academic year. In addition to a two hour tutorial approximately two additional weeks of time outside the classroom for observation of students,

	with patients, discussions and feedback were required.
1991-1994	Advisor, Peabody Society in Pilot Advisory Program, Harvard Medical School
1995-2005	Participated as lecturer and tutor in CBC, 5 hours each time
2005-2006	Tutor, Prevention and Nutrition
2006-2009	Co-director, Role of Discovery in Medicine, required course for first year Harvard Medical Students
April 2007	Thesis Committee, Jacob Newman Marcus, Orexin Receptors and the Central Autonomic System, Harvard Medical School
Dec. 2007	Thesis Committee, Thaddeus John Unger, Examination of Deficits in Energy Balance and Affective Behavior Following Central or Hypothalamic Depletion of Brain-Derived Neurotrophic Factor, Tufts University
May 2013	2013 Leadership and Faculty Development Conference & 2013 Minority Health Policy Annual Meeting, Session Commenter

# b. Advisory and Supervisory Responsibilities

1982-1996 While an attending physician at the Brigham I met with fellows and residents three times per week while on service (two months of the year.) The sessions were devoted to presentation of patients and discussion and teaching. (They were not used for provision of clinical services.)

1994- Supervise research for 2-5 research fellow approximately 350 hrs/yr.

2005-2008 Co-direct, Endocrine Fellowship program with Dr. Evan Rosen

#### c. Advisees and Trainees:

1986-1989 Eric Verdin (postdoctoral fellow), Professor of Medicine, UCSF

1989-1991 Louise Montgomery (postdoctoral fellow), Associate Professor of Biology, Chairman Department of Biology, Marymount University, Arlington, VA

1991-1993 Ali El-Ghorr (postdoctoral fellow), Faculty Division of Biomedicine and Clinical Laboratory Sciences, University of Edinburgh

1994-1997 Daqing Qu, (postdoctoral fellow), Research Scientist, Millipore Corporation

1997-1999 David Ludwig (postdoctoral fellow), Associate Professor, Harvard Medical School, Children's Hospital, Boston

1997-2001 Nicholas Tritos, (postdoctoral fellow), Instructor in Medicine, Massachusetts General Hospital, Boston, MA

2002-2005 Richard Bradley (postdoctoral fellow), Private sector, business development.

1997-2000 Jason Mastaitis (pre-doc), Ph.D., Department of Neurobiology, Mount Sinai School of Medicine, New York, New York

2000-2003 Daniel Trombly (pre-doc), Graduate student, Department of Neuroscience, Northwestern University, Chicago, Illinois

2001-2002 Jason Karamachandran (senior thesis) Pathology Resident, Stanford University, San Francisco, California

2002-2003 Neha Jadeja, (senior thesis), Medical Student, Harvard Medical School, Boston

2001-2003 Gabriella Segal-Lieberman (postdoctoral fellow) Assistant Professor of Medicine, Sheba Medical Center Institute of Endocrinology, Ramat-Gan, Israel

2001-2004 Efi Kokkotou, (postdoctoral fellow), Assistant Professor of Medicine, Harvard Medical School, Division of Gastroenterology, Beth Israel Deaconess Medical Center, Boston

2002-2006 Pavlos Pissios, (postdoctoral fellow), Asssistant Professor of Medicine, Harvard Medical School, Division of Endocrinology, Beth Israel Deaconess Medical Center, Boston

2003-2007 Justin Y. Jeon, (postdoctoral fellow), Assistant Professor, Exercise Physiology, Korea

2003-2007 Adam R. Kennedy, (postdoctoral fellow), Scientist, Cytokinetics, San Francisco, California.

2005-2011 F. Martin Fisher, (postdoctoral fellow), Instructor, Beth Israel Deaconess Medical Center

2006-2008 Jody Dushay (postdoctoral fellow), Instructor, Beth Israel Deaconess Medical Center

2007-2011 Andrew C. Adams, (postdoctoral fellow), Staff Scientist, Lilly Pharmaceuticals

2009-2011 Patricia Chui, (postdoctoral fellow), Attending Physician, New York University Medical School

2011-2013 Co-mentored Tahereh Ghorbani, (postdoctoral fellow), Attending Physician, Joslin Diabetes Center

2012-2014 Elena Toschi, (postdoctoral fellow), Attending Physician, Joslin Diabetes Center

2009-2015 Nicholas Douris, (postdoctoral fellow), Scientist Alkeremes Coorporation, Waltham, MA

2009-2015 Melissa Chee, (postdoctoral fellow), Assistant Professor of Neurobiology, Carlton University, Ontario, Canada.

2013-2017 Darko Stevanovic, post-doctoral fellow, Global Director of Medical and Scientific Affairs - Sanvita Medical at Nova Biomedical

2011-2017 Garima Singhal, post-doctoral fellow, Scientist Dicerna Pharmaceuticals

2015-2018 Bhavna Desai, post-doctoral fellow, Consultant, Simon-Kucher & Partners

# 2. Regional, National and International Contributions:

# Symposium Lecturer:

# Regional:

May 2000	Department of Physiology Seminar, University of Massachusetts Medical School
Oct. 2000	Inaugural Obesity Research Center Symposium Boston University, Invited Speaker
May 2003	Pat Usher Memorial Lecture, Division of Endocrinology Beth Israel Deaconess Medical Center, Boston
Oct. 2006	Beth Israel Deaconess Research Day, Invited Speaker
Oct. 2008	Brigham and Women's/Children's Hospital, Endocrine Grand Rounds
Dec. 2008	Longwood Medical Area Diabetes Seminar, Joslin Diabetes Center
July 2009	Boston Obesity and Nutrition Research Center Annual Symposium
Nov. 2010	Endocrine Grand Rounds, Massachusetts General Hospital, Boston
Mar. 2011	Medical Grand Rounds, Beth Israel Deaconess Medical Center, Boston

Oct. 2011	Endocrine	Grand Rounds.	Rhode Island	Hospital.	Providence

Oct. 2013 Frontiers in Medicine, Beth Israel Deaconess Medical Center, Boston

# National:

Aug. 1996	Aspen Lipid Conference
Feb. 1997	Maui Meeting on Diabetes and Obesity
June 1997	Endocrine Society, 79th Annual Meeting, Symposium Speaker
Sep. 1997	National Institute of Health, NIDDK,
	"The Brain and the Adipocyte"
Mar. 1998	Pennington Biomedical Research Center
	"Nutrition, Genetics and Obesity"
June 1998	American Diabetes Association, 58th Annual Meeting,
	Symposium Speaker "Neuroendocrine Regulation of Energy Balance and Satiety"
Oct. 1998	Banbury Conference/Cold Spring Harbor
	The Molecular Physiology of Weight Regulation and Obesity
June 1999	Endocrine Society, 81st Meeting, Symposium Speaker
Feb. 2000	Keystone Symposium, Invited Speaker
Feb. 2001	Keystone Symposium, Invited Speaker
Jan. 2002	Keystone Symposium, Molecular Control of Adipogenesis and
	Obesity, Invited Speaker
July 2002	Gordon Research Conference, Mechanism of Hormone Action,
	Invited Speaker
Feb. 2003	AAAS Symposium Speaker
June 2004	8 <sup>th</sup> Neuroendocrine Workshop, Invited Speaker
June 2004	Endocrine Society 86 <sup>th</sup> Annual Meeting, Symposium Speaker
Jan. 2005	Keystone Symposium, Invited Speaker
May 2006	Appetite and Obesity, Cold Spring Harbor Symposium, Invited Speaker
May 2008	New York Academy of Sciences, Invited Speaker
June 2008	Endocrine Society, Annual Meeting, Symposium Speaker
Jan. 2009	Keystone Symposium, Invited Speaker
Aug. 2009	Kern Aspen Lipid Conference, Invited Speaker
April 2010	Presidential Symposium, American Society for Nutrition,
-	Invited Speaker
June 2010	American Diabetes Association, Annual Meeting, Invited Speaker
June 2010	Endocrine Society, Annual Meeting, Invited Speaker
Feb. 2011	NIDDK, "Toward a Clinical Definition of Leptin Resistance",
	Invited Participant
Mar. 2011	Endocrine Grand Rounds, University of Pennsylvania
Nov. 2011	Endocrine Grand Rounds, Brown University
Dec. 2011	Endocrine Grand Rounds, Columbia University

Jan. 2012	Keystone Symposium, Invited Speaker, Plenary Session
June 2012	Endocrine Society, Annual Meeting, Invited Speaker
June 2013	ADA, Invited Speaker
Jan. 2014	Vanderbilt University, Department of Physiology Seminar Series
Nov. 2014	Invited Lecturer, Dept. Pharmacology, U of Pennsylvania
June 2015	Iconoclasm Meeting, Prior to ADA, Boston, MA
June 2016	Lecturer at University of Iowa Carver College of Medicine
	Diabetes & Obesity

# International:

July 1997	International Diabetes Foundation, Helsinki Meeting
Aug. 1998	Adipo- Science Meeting, Osaka, Japan
Nov 1999	Institut Pasteur, Euroconference "Obesity: Genetics,
	Pathophysiology and Therapeutics," Invited Speaker
March 2004	Days of Molecular Medicine, Cambridge, England, Invited
	Symposium Speaker
Jan. 2012	Keystone Symposium, Genetic and Molecular Basis of Obesity
	and Body Weight Regulation, Invited Speaker
Aug. 2012	Australian Diabetes Association, Brisbane Australia, Annual
C	Meeting
Mar. 2013	Keystone Symposium, Neuronal Control of Appetite, Metabolism
	and Weight, Invited Speaker
Mar. 2015	Keystone Symposium, Obesity and the Metabolic Syndrome:
	Mitochondria and Energy Expenditure, Invited Speaker
Nov. 2015	Invited Speaker, Karolinska Institute, Stockholm, Sweden
Sep. 2016	Invited Speaker, 4 <sup>th</sup> Helmholtz-Nature Medicine Diabetes
-	Conference, Munich, Germany
Feb. 2017	Invited Speaker, Zydus-Cadila Symposium, Ahmedabad, India
Apr. 2017	Invited Speaker, International Symposium on Insulin Receptor and
	Insulin Action
July 2017	Society for Ingestive Behavior, invited Lecturer, Montreal, Canada
Nov. 2017	Invited Lecturer, University of Montreal, CHUM Research Center
Nov 2017	Invited Lecturer, Australian New Zealand Diabetes Meeting
Mar 2019	Symposium Speaker, Annual Endocrine Society Meeting
ership Roles	

# Leadership Roles

2000-2003	Endocrine Society, Meeting Organizing Committee
2007	Keystone Symposium, Organizer: Obesity, Peripheral and Central
	Pathways Regulating Energy Homeostasis
2015	Keystone Symposium, Obesity and the Metabolic Syndrome:
	Mitochondria and Energy Expenditure, Co-Organizer

# PART III: Bibliography

# **Original Reports**

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- 2. **Maratos E**, Taub RN, Bramis J. Amelioration of streptozotocin-induced diabetes in mice by the implantation of pancreatic islets in diffusion chambers. Mt Sinai J Med NY 1976; 43:415-422.
- 3. Flier JS, **Maratos-Flier E**, McIsaac D, Pallotta J. Endogenous digitalis-like activity circulates in the plasma of the toad, bufo marinus. Nature 1979; 279:342-344.
- 4. **Maratos-Flier E**, Spriggs DR, Fields BN, Kahn CR. Specific plasma membrane receptor for reovirus on rat pituitary cells in culture. J Clin Invest 1983; 72:617.
- 5. **Maratos-Flier E**, Goodman MJ, Fields BN, Kahn CR. Differential effects of viral infection on islet and pituitary cells in culture. Endocrinology 1985; 116:2430-2437.
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- 8. Verdin EM, **Maratos-Flier E**, Kahn CR, Sodoyez JC, Sodoyez-Goffaux F, De Vos CJ, Lynn SP, Fields BN. Visualization of viral clearance in the living animal. Science 1987; 236:439-442
- 9. **Maratos-Flier E**, Kao CY-Y, Verdin EM, King GL. Receptor mediated vectorial transcytosis of epidermal growth factor by MDCK cells. J Cell Biol 1987; 105:1595-1601.
- 10. Verdin EM, Lynn, SP, Fields BN, **Maratos-Flier**, E. Uptake of reovirus serotype I by the lungs from the bloodstream is mediated by the viral hemaglutinin. J Virol 1988; 62:545-551.
- 11. Verdin EM, King GL, **Maratos-Flier E**. Characterization of a common high-affinity receptor for reovirus serotypes 1 and 3 on endothelial cells. J Virol 1989; 63:1318-1323.
- 12. Montgomery LB, Kao C-Y-Y, Verdin E, Cahill C, **Maratos-Flier E**. Infection of a polarized epithelial cell line with wild type reovirus leads to viral persistence and altered cellular function. J Gen Virol 1991; 72:2939-2946.
- 13. El-Ghorr AA, Gordon D, George K, and **Maratos-Flier E**. Regulation of expression of the reovirus on differentiated HL60 cells. J Gen Virol 1992; 73:1961-1968.
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