

CHARBONNEAU CANCER INSTITUTE

Mission: To meet the cancer challenge.

Goal: To decrease cancer in the population, improve cancer treatment, and improve the patient experience by conducting biomedical, clinical, health systems, and population health research, from prevention to life after cancer.

Objectives:

- To foster relationships with academic, health services, and industry partners;
- To secure philanthropy for research initiatives with significant potential;
- To increase the competitiveness of the Institute for external funding;
- To build modern infrastructure that will enable our researchers to discover;
- To provide exceptional education and learning opportunities for trainees;
- ◆ To recruit, support, and retain the best and brightest cancer researchers; and
- To facilitate transformative national and international collaborations.

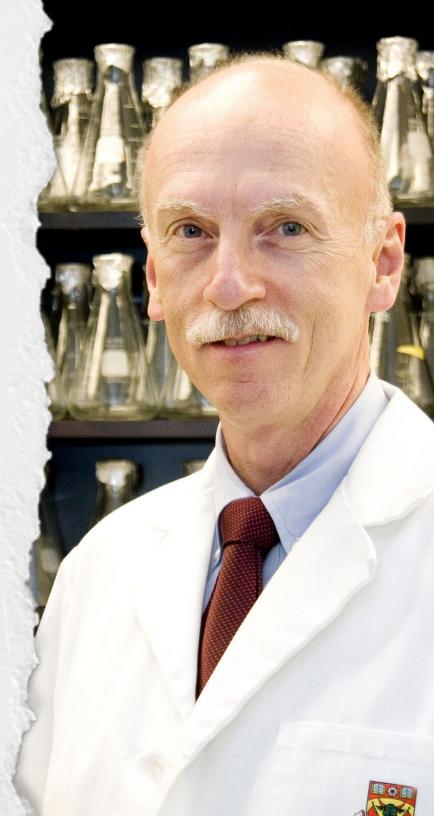


Message from the Director

In 2018, the Cancer Institute marked its 15th anniversary. Initially known as the Southern Alberta Cancer Research Institute, we became Charbonneau in 2014, a change that marked a commitment to develop a research institute of significance at the University of Calgary. This inaugural annual research report highlights several significant achievements over the past twelve months, including the opening of the new Childhood Cancer and Blood Disorders Laboratories and rapidly increasing success in tri-council funding. Again the Institute has supported its research centres and programs at the Cumming School of Medicine, Calgary's teaching hospitals, and the Tom Baker Cancer Centre by recruiting top research talent from across the globe and building key infrastructure. This report celebrates our scientists, donors, partners, and staff for their dedication to research aimed at meeting the cancer challenge and improving cancer treatment, decreasing cancer in the population, and improving the patient experience.

J. 197 G.

J. Gregory Cairncross, MD Director, Charbonneau Cancer Institute Professor, Clinical Neurosciences University of Calgary



Decreasing Cancer in the Population

Early Cancer Detection



EXOSOME-BASED DEVICE FOR BLADDER CANCER

Drs. Amir Sanati Nezhad and **Don Morris** and their team have developed a point-of-care device to collect and analyze exosomes (components of the cell that are released into the blood and have tumour-specific characteristics). The device can accurately detect changes in exosomes that are predictive of cancer, with an initial application in bladder cancer. The team is now refining the device and clinically testing its use.



METABOLOMIC TEST FOR COLORECTAL CANCER

Dr. Oliver Bathe and team (**Drs. Hans Vogel, Robert Hilsden,** and **Karen Kopciuk**) have developed a new metabolomic test to detect colon cancer. Current approaches, such as FIT testing, are effective but not widely adopted, due to their inconvenience. Dr. Bathe's metabolomic test is convenient, less invasive, and was highly accurate in early phase testing. Further testing and validation of the test are now underway.

Cancer Prevention & Epidemiology

Dr. Christine Friedenreich, an Adjunct Professor with the Department of Oncology and Scientific Director of the Department of Cancer Epidemiology and Prevention Research (AHS), is leading the Canadian population attributable risk of cancer (ComPARe) study, which aims to estimate how many cancer cases can be prevented in Canada through the year 2042. In collaboration with **Dr. Darren Brenner** and several Ontario- and Quebec-based investigators, the team is examining cancer risk factors, such as tobacco, diet, energy imbalance, infectious diseases, hormonal therapies, and environmental factors (i.e., air pollution and residential radon). The results of this study will inform cancer prevention and public health initiatives, as well as public policy.



Above: Dr. Amir Nezhad (top); Dr. Oliver Bathe (bottom); Right: Dr. Christine Friedenreich and Dr. Darren Brenner

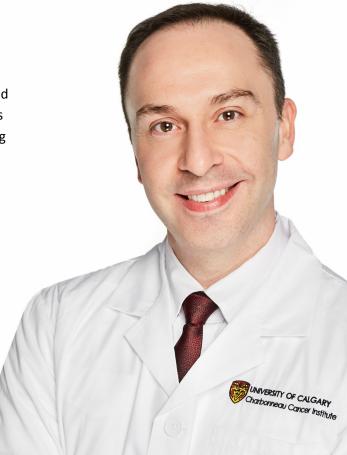


Evict Radon

Dr. Aaron Goodarzi, Canada Research Chair for Radiation Exposure Disease and member of the **Robson DNA Science Centre**, is studying ionizing radiation, such as the alpha particles emitted by radon gas, and how this type of radiation impacts DNA damage and repair. His work is improving our knowledge of cancer development and the human aging process, as well as informing efforts to educate the public about the harms of ionizing radiation and how to avoid or reduce exposure.

Radon is the leading cause of lung cancer in non-smokers and can further increase the risk of lung cancer in smokers. Work published by Dr. Goodarzi in 2017 revealed that 1 in 8 homes in southern Alberta have cancer-causing levels of radon. Yet, with appropriate mitigation, radon can be eliminated in homes.

Dr. Goodarzi now leads a grass-roots, citizen science initiative, **Evict Radon**, to increase awareness about radon testing and mitigation. The Evict Radon campaign has evolved into a national research initiative to test thousands of homes across Canada for radon. Learn more at www.evictradon.ca.



Improving Cancer Treatment

Brain Tumour Biology and Therapeutics

Clark Smith Brain Tumour Centre scientists and clinician researchers have made notable contributions to the understanding and treatment of brain cancer.



Dr. V. Wee Yong spearheaded a brain immunotherapy program and has shown that immune cells in brain tumours can be activated by the off-label use of existing drugs, such as the antifungal drug amphotericin B, to suppress cancer growth in animal model systems. He is leading a CIHR-funded trial, *Mobilizing anti-tumor microglia/macrophages with niacin to improve the prognosis of glioblastoma*, which will study the role of niacin (vitamin B3), a more tolerable drug than amphotericin B, in boosting immune cells within the brain to suppress tumour growth in patients with glioblastoma.



Drs. Donna Senger and Steve Robbins have shown how brain tumour cells are able to influence the adjacent brain environment to ensure their own growth and survival. Using patient-derived brain tumour models, they are working to identify ways to exploit the active interplay between tumour cells, the surrounding normal cells, and the extracellular matrix within the brain to mitigate tumour proliferation, invasion, and therapeutic resistance. Through their work on the Terry Fox Research Institute Translational Cancer Research Project, Modeling and therapeutic targeting of the clinical and genetic diversity of glioblastoma, they identified a specific subset of immune cells that could be targetable in patients with glioblastoma.



Dr. Marco Gallo, Canada Research Chair in Brain Cancer Epigenomics, is interested in the epigenome of brain cancer stem cells. He and his team have generated high resolution 3D genome maps using glioblastoma cells from three patients. These 3D maps have been integrated with epigenetic and transcriptional profiles to understand how individual genes are regulated in different patients. He has also shown that cancer stem cells from glioblastoma activate a group of immune-related genes that allow them to escape the immune system. The protein product of one of these genes is targetable using an experimental compound to which glioblastoma cancer stem cells are especially sensitive.

Above: Dr. V. Wee Yong (top); Dr. Donna Senger (middle); Dr. Marco Gallo (bottom)

Precision Oncology in Multiple Myeloma

Drs. Paola Neri and Nizar Bahlis are conducting genomic studies to identify novel drug targets for multiple myeloma (MM), based on the individual genetic profile of the patient's tumour cells. Using this approach, they are able to deliver personalized and tailored therapeutics for patients with MM and define mechanisms of drug resistance to novel anti-MM agents. By performing single cell immune profiling in patients treated with immunotherapy, they have found that the expansion of central memory T-cells is important to achieve deep and durable responses in patients. Moreover, by analyzing the repertoire of T-cells through single cell sequencing, they are working to design efficient CAR-T cells targeting MM-specific neo-antigens.



Above: Dr. Paola Neri (left); Dr. Doug Mahoney (right)



Oncolytic Virus Therapy

Dr. Douglas Mahoney and colleagues published a paper in *Nature Communications* in 2017, which showed that oncolytic rhabdoviruses and smac (second mitochondrial activator of caspase) mimetic compounds work synergistically to promote anticancer immunity by driving anticancer T-cell activity and reinvigorating exhausted T-cells in immuno-suppressed tumour microenvironments.

Building on his expertise designing therapeutic viruses and engineering immune cells, Dr. Mahoney is collaborating with **Dr. Jennifer Chan** on a new CAR-T cell program in cancer immunotherapy. The **ACTION** (Advanced Cellular Therapy & Immune Oncology)

Initiative will include the identification of new immunotherapy targets, the development of innovative CAR T-cell technologies, preclinical studies in patient-derived and genetically-engineered mouse models, and an innovative clinical trials component at the Alberta Children's Hospital. ACTION strives to position Calgary as a Canadian hub for CAR-T cell innovation and enhance access in Alberta to these new immune-based cancer therapies.

Improving the Patient Experience



Improving Social Adjustment in Survivors of Childhood Cancer

Dr. Fiona Schulte is an Assistant Professor in the Division of Psychosocial Oncology and a psychologist in the Hematology, Oncology and Transplant program at the Alberta Children's Hospital (AHS). Her work has broadly explored psychological outcomes in survivors of pediatric cancers and specifically social adjustment in children who were treated for brain cancer. Her research has provided evidence for the negative impact of

treatments directed at the brain, such as cranial radiation therapy, on social adjustment and cognitive impairment. She recently received funding from Alex's Lemonade Stand Foundation that will allow her and her team to use peer data, multi-informant assessments, and neuroimaging to examine these relationships further. Dr. Schulte's future research will focus on children with acute lymphoblastic leukemia, the most common form of childhood cancer, to examine factors associated with poor social adjustment in this vulnerable population



National Impact: Improving Cancer Related Sexual Dysfunction

Dr. Lauren Walker, an Adjunct Assistant Professor within the Division of Psychosocial Oncology and psychologist at the Tom Baker Cancer Centre (AHS), has identified how to recognize and treat sexual dysfunction within the context of cancer. Over the course of about 10 years, Dr. Walker's research has transformed the way patients with cancer-related sexual dysfunction are supported, including the development of integrated intimacy work-shops, as part of routine care for prostate cancer. She has collaborated to develop a national sexual recovery e-clinic and health care provider training program

based on her research. Dr. Walker's team developed a one-of-a-kind educational program for patients receiving androgen deprivation therapy for prostate cancer. It consists of a professionally facilitated class and a book she co-authored, titled Androgen Deprivation Therapy: An Essential Guide for Prostate Cancer Patients and Their Loved Ones. The program is available online and has been disseminated internationally, with 14 sites having served over 1000 patient/partner participants to date. Ultimately, Dr. Walker hopes that her work will help patients who tend to get overlooked, whether it's those who don't respond to traditional sexual medicine treatments or those who experiencing side effects that no one likes to talk about.

Improving Care Delivery



Toward Better Care: CHORD Consortium

Dr. Winson Cheung, an Associate Professor in the Department of Oncology and the Medical Director of Health Services Research at CancerControl Alberta (AHS), chairs the Cancer Health Outcomes Research Database (CHORD) Consortium, a pan-Canadian collaboration to develop a national cancer data repository for rectal cancer and other malignancies. Having demonstrated the feasibility of a national data sharing collaboration, Dr. Cheung led a study to examine whether adjuvant chemotherapy, which is associated with toxicity and increased healthcare costs, improves outcomes in patients with early stage rectal cancer. The study

included patients from several provinces, including Newfoundland, Ontario, Alberta, and British Columbia. Published in the *American Journal of Clinical Oncology* in 2018, his study showed that post-surgical chemotherapy does not improve outcomes and its use should be carefully considered. By reducing the use of post-surgical chemotherapy, patients may be able avoid some unnecessary toxicity and long-term side effects from treatment.

Palliative Care Early and Systematic (PaCES Study)

In 2018, **Drs. Jessica Simon** and **Aynharan Sinnarajah** (Tom Baker Cancer Centre, AHS) were awarded a grant from CIHR and Alberta Health to develop, implement, and evaluate the impact of an early, systematic and integrated palliative care pathway for advanced colorectal cancer on patient, caregiver, and health system outcomes. The PaCES Study seeks to increase the number of patients with advanced colorectal cancer who receive integrated palliative care and, in doing so, improve both patient-reported and health system outcomes. Initial PaCES activities focused on understanding why palliative care services are accessed late and the challenges that are associated with integrating palliative care into clinical practice. The team discovered that health care providers and patients perceive benefit in receiving palliative care at the same time as cancer-modifying treatments, but both are challenged by misperceptions that palliative care means that end of life is imminent. They also found that healthcare providers believe that a palliative care approach

is part of their role as clinicians but struggle with knowing how best to provide it.

The team is now working to close these gaps through the routine identification of those likely to benefit; improved patient-clinician communication about care preferences; ensuring the essential components of palliative care are provided in a systematic way; on-going liaising between specialists, family doctors, and palliative care providers; and more timely access to palliative care.

Above: Dr. Winson Cheung; Right: Drs. Jessica Simon and Aynharan Sinnarajah

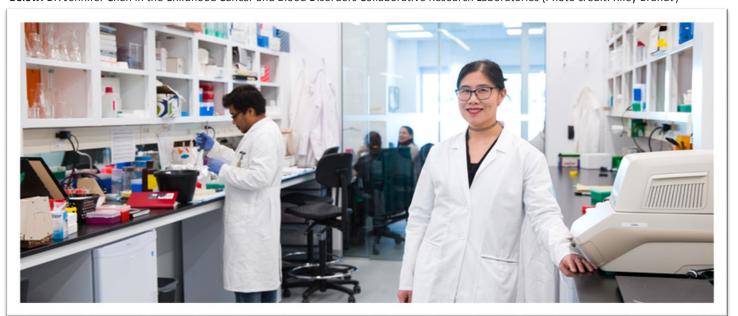
New Childhood Cancer and Blood Disorders

Collaborative Research Laboratories

Thanks to generous gifts from the Alberta Children's Hospital Foundation and Kids Cancer Care Foundation, in January 2018 the Institute opened its new **Childhood Cancer and Blood Disorders Collaborative Research Laboratories** in the Heritage Medical Research Building. This modern, open-concept facility was designed to enable the sharing of equipment and research staff and provide collision space for researchers to share new ideas and plan future research endeavours. The facility was planned and overseen by **Dr. Jennifer Chan** and now accommodates nine principal investigators (**Drs. Faisal Khan, Jan Storek, Aru Narendran, Marco Gallo, Paola Neri, Nizar Bahlis, Sorana Morrissy, Edwin Wang, and Jennifer Chan**) and their trainees and staff.

The facility supports research to: understand the cellular and molecular basis for childhood cancer initiation and progression, innovate and test new therapeutic approaches for the treatment of childhood cancers, and increase the availability of patient-derived samples and experimental models of childhood cancer. It also features a hematology translational laboratory (**Drs. Faisal Khan and Jan Storek**), which was accredited by the College of Physicians and Surgeons of Alberta to provide clinical testing for patients across the province undergoing bone marrow transplant, including Next-Generation Sequencing-based somatic variant results.

Below: Dr. Jennifer Chan in the Childhood Cancer and Blood Disorders Collaborative Research Laboratories (Photo credit: Riley Brandt)



Tri-Council Grant Recipients (2017-2018)

Natural Sciences and Engineering Research Council of Canada:

Dr. Shirin Bonni

Dr. Sarah Childs

Dr. Jennifer Cobb

Dr. Jason de Koning

Dr. Darren Derksen

Dr. Marco Gallo

Dr. Aaron Goodarzi

Dr. Savraj Grewal

Dr. Ebba Kurz

Dr. Haocheng Li

Dr. Jonathan Lytton

Dr. Gregory Moorhead

Dr. Derrick Rancourt

Dr. Karl Riabowol

Dr. Kristina Rinker

Dr. Amir Sanati Nezhad

Dr. David Schriemer

Dr. Hans Vogel

Dr. Edwin Wang

Dr. Gareth Williams

Canadian Institutes of Health Research (CIHR):

Spring 2018 Project Grant:

Dr. Gareth Williams

Dr. Hans Vogel

Fall 2017 Project Grant:

Dr. Jennifer Cobb

Dr. Christine Friedenreich

Dr. Marco Gallo

Dr. Garnette Sutherland

Dr. Voon Wee Yong

Special Operating | Catalyst | Supporting Patient-Oriented Research | Other Grants:

Dr. Oliver Bathe

Dr. Darren Brenner

Dr. Gregory Cairncross

Dr. Michael Monument

Dr. Derrick Rancourt

Drs. Jessica Simon and Aynharan Sinnarajah

Noteworthy Publications by Institute Members (2017-2018)



Smac mimetics and oncolytic viruses synergize in driving anticancer T-cell responses through complementary mechanisms

Dae-Sun Kim^{1,2,3}, Himika Dastidar^{1,2,3}, Chunfen Zhang^{1,2}, Franz J. Zemp^{1,2,3}, Keith Lau^{1,2,3,4}, Matthias Ernst^{1,2} Andrea Rakic^{1,2,5}, Saif Sikdar^{1,2,3}, Jahanara Rajwani^{1,2}, Victor Naumenko^{1,2,3,4}, Dale R. Balce⁶, Ben W. Ewanchuk⁷, Pankaj Tailor⁶, Robin M. Yates 6, 6,7, Craig Jenne^{3,4}, Chris Gafuik^{1,2,3} & Douglas J. Mahoney^{1,2,3,7}

Identification of nine new susceptibility loci for endometrial cancer

Tracy A. O'Mara o et al. (Christine Friedenreich)

Identification of the fungal ligand triggering cytotoxic PRR-mediated NK cell killing of Cryptococcus and Candida

Shu Shun Li^{1,2}, Henry Ogbomo^{1,2}, Michael K. Mansour³, Richard F. Xiang^{1,2}, Lian Szabo⁴, Fay Munro⁵, Priyanka Mukherjee⁵, Roy A. Mariuzza⁶, Matthias Amrein⁵, Jatin M. Vyas 6, Stephen M. Robbins^{7,8} & Christopher H. Mody (1) 1,2,4

DOI: 10.1038/s41467-018-04743-2

Quantitative in vivo whole genome motility screen reveals novel therapeutic targets to block cancer metastasis

Konstantin Stoletov¹, Lian Willetts¹, Robert J. Paproski¹, David J. Bond¹, Srijan Raha¹, Juan Jovel^{2,3}, Benjamin Adam⁴, Amy E. Robertson¹, Francis Wong¹, Emma Woolner¹, Deborah L. Sosnowski¹, Tarek A. Bismar⁵, Gane Ka-Shu Wong^{2,3,6}, Andries Zijlstra⁰ ⁷ & John D. Lewis⁰

DOI: 10.1038/±41467-018-06654-8 OPEN

TGF-β-associated extracellular matrix genes link cancer-associated fibroblasts to immune evasion and immunotherapy failure

Ankur Chakravarthy^{1,2}, Lubaba Khan^{3,4,5}, Nathan Peter Bensler^{3,4,5}, Pinaki Bose^{3,4,5,6} & Daniel D. De Carvalho (5) 1,2

nature

Nature, 2017 September 14; 549(7671): 227-232. doi:10.1038/nature23666

Fate mapping of human glioblastoma reveals an invariant stem cell hierarchy

Xiaoyang Lan^{1,2,3}, David J. Jörg^{4,5}, Florence M. G. Cavalli^{1,2}, Laura M. Richards^{12,13}, Long V. Nguyen⁷, Robert J. Vanner^{1,2,3}, Paul Guilhamon^{12,13,14}, Lilian Lee^{1,2}, Michelle Kushida^{1,2}, Davide Pellacani^{7,8}, Nicole I. Park^{1,2,3}, Fiona J. Coutinho^{1,2,3}, Heather Whetstone^{1,2}, Hayden J. Selvadurai^{1,2}, Clare Che^{1,2}, Betty Luu^{1,2}, Annaick Carles⁹, Michelle Moksa⁹, Naghmeh Rastegar^{1,2}, Renee Head^{1,2}, Sonam Dolma^{1,2,11}, Panagiotis Prinos^{13,20}, Michael D. Cusimano 17,18, Sunit Das 17,18, Mark Bernstein 16,18, Cheryl H. Arrowsmith 13,20, Andrew J. Mungall 10, Richard A. Moore 10, Yussanne Ma 10, Marco Gallo 19, Mathieu Lupien 12, 13, 14, Trevor J. Pugh^{12,13}, Michael D. Taylor^{1,2,11,15,18}, Martin Hirst^{9,10}, Connie J. Eaves^{7,8}, Benjamin D. Simons^{4,5,6,*}, and Peter B. Dirks^{1,2,3,15,18,*}

medicine

Article | Published: 11 September 2017

A mouse model for embryonal tumors with multilayered rosettes uncovers the therapeutic potential of Sonichedgehog inhibitors

Julia E Neumann, Annika K Wefers, Sander Lambo, Edoardo Bianchi, Marie Bockstaller, Mario M Dorostkar, Valerie Meister, Pia Schindler, Andrey Korshunov, Katja von Hoff, Johannes Nowak, Monika Warmuth-Metz, Marlon R Schneider, Ingrid Renner-Müller, Daniel J Merk, Mehdi Shakarami, Tanvi Sharma, Lukas Chavez, Rainer Glass, Jennifer A Chan, M Mark Taketo, Philipp Neumann, Marcel Kool & Ulrich Schüller™

ARTICLES

s://doi.org/10.1038/s41590-017-0030-x

nature immunology

Aged polymorphonuclear leukocytes cause fibrotic interstitial lung disease in the absence of regulation by B cells

Jung Hwan Kim^{1,2}, John Podstawka^{1,2}, Yuefei Lou^{1,2}, Lu Li^{1,2}, Esther K. S. Lee^{1,2}, Maziar Divangahi³, Björn Petri^{4,5}, Frank R. Jirik⁶, Margaret M. Kelly^{1,7} and Bryan G. Yipp ^{⊙1,2*}

ENDOCRINOLOGY

Molecular profiling of thyroid nodule fine-needle aspiration cytology

Markus Eszlinger, Lorraine Lau, Sana Ghaznavi, Christopher Symonds, Shamir P. Chandarana, Moosa Khalil & Ralf Paschke Nature Reviews Endocrinology 13, 415-424 (2017)



Nat Struct Mol Biol. 2018 June ; 25(6): 482-487. doi:10.1038/s41594-018-0065-1

Dissection of DNA double-strand break repair using novel single-molecule forceps

Jing L. Wang^{1,*}, Camille Duboc^{1,*}, Qian Wu^{2,*}, Takashi Ochi^{2,8}, Shikang Liang², Susan E. Tsutakawa³, Susan P. Lees-Miller⁴, Marc Nadal¹, John A. Tainer^{3,5}, Tom L. Blundell², and Terence R. Strick^{1,6,7,†}

genetics

Nat Genet. 2017 May ; 49(5): 680-691. doi:10.1038/ng.3826.

Identification of twelve new susceptibility loci for different histotypes of epithelial ovarian cancer

A full list of authors and affiliations appears at the end of the article. (Martin Kobel)

nature nanotechnology

Nat Nanotechnol. 2017 Jul;12(7):701-710. doi: 10.1038/nnano.2017.56. Epub 2017 Apr 24

Peptide-MHC-based nanomedicines for autoimmunity function as T-cell receptor microclustering devices.

Singha S¹, Shao K¹, Yang Y¹.², Clemente-Casares X¹, Solé P³, Clemente A³, Blanco J³, Dai Q⁴, Song F⁴,

Liu SW⁵, Yamanouchi J¹, Umeshappa CS¹, Nanjundappa RH¹, Detampel P⁶, Amrein M⁶, Fandos C³, Tanguay Rˀ,

Newbigging S⁵, Serra P³, Khadra A⁵, Chan WCW⁴, Santamaria P¹.³.

Cell

A Hematogenous Route for Medulloblastoma Leptomeningeal Metastases

Livia Garzia, Noriyuki Kijima, A. Sorana Morrissy, Pasqualino De Antonellis, Ana Guerreiro-Stucklin, Borja L. Holgado, Xiaochong Wu, Xin Wang, Michael Parsons, Kory Zayne, Alex Manno, Claudia Kuzan-Fischer, Carolina Nor, Laura K. Donovan, Jessica Liu, Lei Qin, Alexandra Garancher, Kun-Wei Liu, Shella Mansouri, Betty Luu, Yuan Yao Thompson, Vijay Banaswamy, John Peacock, Hamza Farooq, Patryk Skowron, David J.H. Shih, Angela Li, Sherine Ensan, Clinton S. Robbins, Myron Cybulsky, Siddhartha Mittra, Yussanne Ma, Richard Moore, Andy Mungali, Yoon-Jae Cho, William A. Weiss, Jennifer A. Chan, Cynthia E. Hawkins, Maura Massimino, Nada Jabado, Michal Zapotocky, David Sumerauer, Eric Bouffet, Peter Dirks, Uir Tabori, Poul H.B. Sorense, Priscilla K. Brastianos, Kenneth Aldape, Steven J.M. Jones, Marco A. Marra, James R. Woodgett, Robert J. Wechsler-Reya, Daniel W. Fults, and Michael D. Taylor'

Cancer Cell

Comparative Molecular Analysis of Gastrointestinal Adenocarcinomas

Yang Liu, ^{1,2,22} Nilay S. Sethi, ^{1,2,22} Toshinori Hinoue, ^{3,22} Barbara G. Schneider, ^{4,22} Andrew D. Cherniack, ^{1,2} Francisco Sanchez-Vega, ⁵ Jose A. Seoane, ⁶ Farshad Farshidfar, ⁷ Reanne Bowlby, ⁸ Mirazul Islam, ^{1,2} Jaegil Kim, ¹ Walid Chatlia, ⁸ Rehan Akbani, ¹⁰ Rupa S. Kanchi, ¹⁰ Charles S. Rabkin, ¹¹ Joseph E. Willis, ¹² Kenneth K. Wang, ¹³ Shannon J. McCall, ¹⁴ Lopa Mishra, ¹⁵ Akinyemi I. Ojesina, ^{16,21} Susan Bullman, ² Chandra Sekhar Pedamallu, ² Alexander J. Lazar, ¹⁷ Ryo Sakai, ¹⁸ The Cancer Genome Atlas Research Network, Vésteinn Thorsson, ^{19,23,4} Adam J. Bass, ^{1,2,20,23,4} and Peter W. Laird ^{3,22,24,4}

Cancer Cell. 2017 August 14; 32(2): 185-203.e13. doi:10.1016/j.ccell.2017.07.007.

Integrated Genomic Characterization of Pancreatic Ductal Adenocarcinoma

The Cancer Genome Atlas Research Network* (Oliver Bathe)



The NEW ENGLAND JOURNAL of MEDICINE

Overall Survival with Palbociclib and Fulvestrant in Advanced Breast Cancer

N.C. Turner, D.J. Slamon, J. Ro, I. Bondarenko, S.-A. Im, N. Masuda, M. Colleoni, A. DeMichele, S. Loi, S. Verma, H. Iwata, N. Harbeck, S. Loibl, F. André, K. Puyana Theall, X. Huang, C. Giorgetti, C. Huang Bartlett, and M. Cristofanilli

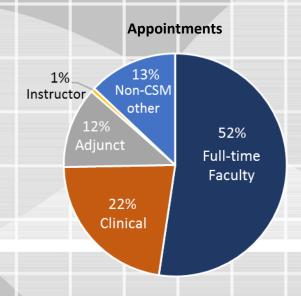
Venetoclax–Rituximab in Relapsed or Refractory Chronic Lymphocytic Leukemia

J.F. Seymour, T.J. Kipps, B. Eichhorst, P. Hillmen, J. D'Rozario, S. Assouline, C. Owen, J. Gerecitano, T. Robak, J. De la Serna, U. Jaeger, G. Cartron, M. Montillo, R. Humerickhouse, E.A. Punnoose, Y. Li, M. Boyer, K. Humphrey, M. Mobasher, and A.P. Kater

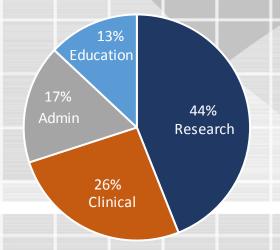
Obinutuzumab for the First-Line Treatment of Follicular Lymphoma

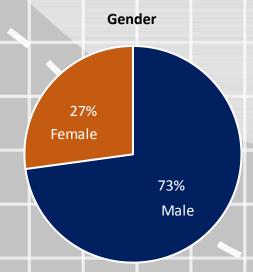
R. Marcus, A. Davies, K. Ando, W. Klapper, S. Opat, C. Owen, E. Phillips, R. Sangha, R. Schlag, J.F. Seymour, W. Townsend, M. Trněný, M. Wenger, G. Fingerle-Rowson, K. Rufibach, T. Moore, M. Herold, and W. Hiddemann

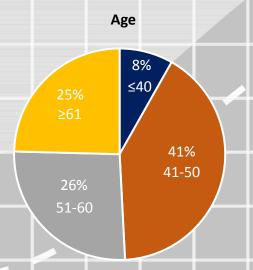
Institute Metrics



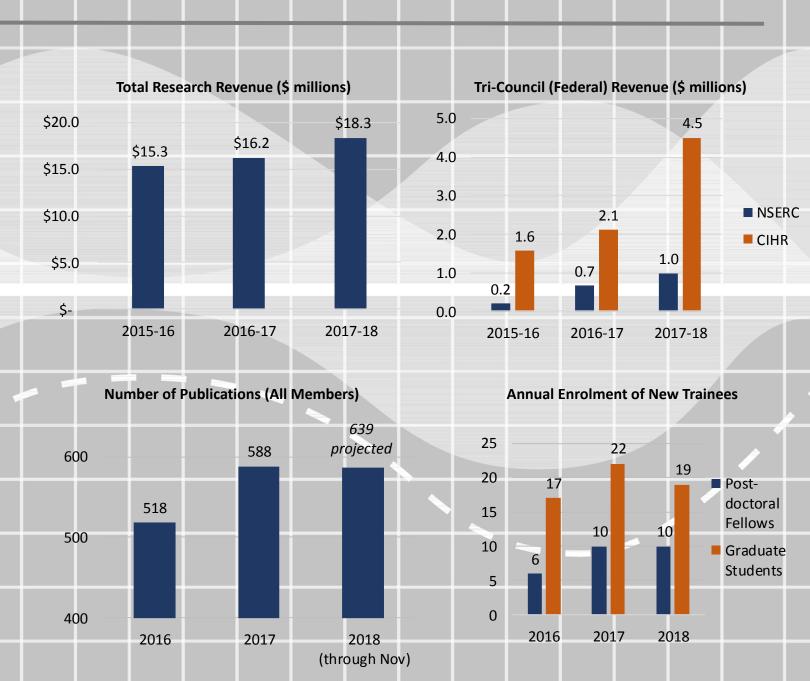
Activity Profile of Full Members (Full-Time Faculty)



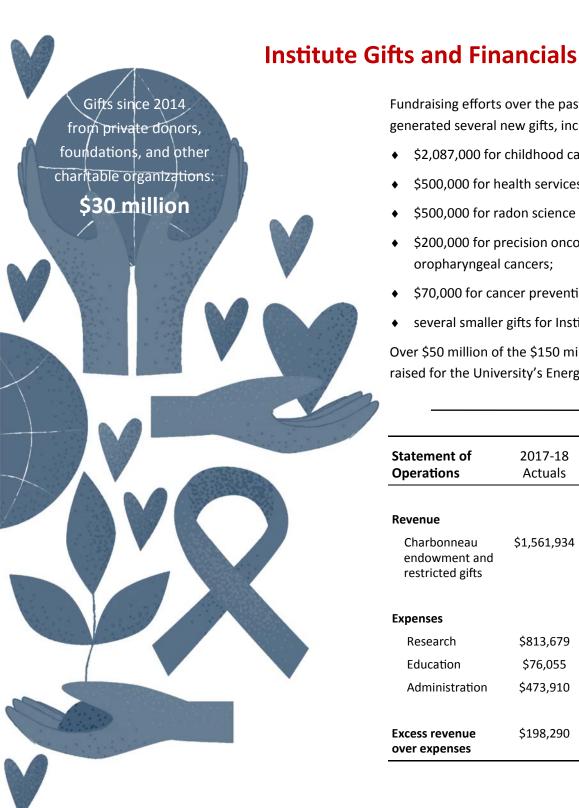




Source: Office of Faculty Analytics, Cumming School of Medicine; Charbonneau Cancer Institute internal administrative data



Notes: Total research revenue includes grants, gifts, and donations received by Institute members; Tri-Council (Federal) revenue, as a portion of the total research revenue, includes grants from the Canadian Institutes of Health Research (CIHR) and the Natural Sciences and Engineering Research Council (NSERC).



Fundraising efforts over the past year (2017-2018) have

- \$2,087,000 for childhood cancer research;
- \$500,000 for health services research;

generated several new gifts, including:

- \$500,000 for radon science (Evict Radon campaign);
- \$200,000 for precision oncology focused on oropharyngeal cancers;
- \$70,000 for cancer prevention research; and
- several smaller gifts for Institute programs and researchers.

Over \$50 million of the \$150 million cancer target has been raised for the University's Energize Campaign.

Statement of	2017-18	2018-19	2019-20
Operations	Actuals	Budget	Forecast
Revenue			
Charbonneau endowment and restricted gifts	\$1,561,934	\$1,763,484	\$1,188,123
Expenses			
Research	\$813,679	\$1,258,526	\$652,523
Education	\$76,055	\$118,500	\$118,500
Administration	\$473,910	\$462,000	\$427,000
Excess revenue over expenses	\$198,290	(\$75,542)	(\$9,900)

Institute Members

Full Members

Amrein, Matthias
Anderson, Ronald
Bahlis, Nizar
Barbera, Lisa
Bathe, Oliver
Beattie, Tara
Beaudry, Paul
Bebb, Gwyn
Bismar, Tarek
Bonni, Shirin
Bose, Pinaki
Brenner, Darren
Brown, Chris
Bultz, Barry

Cairncross, J. Gregory
Campbell, Tavis
Card, Cynthia
Carlson, Linda
Chan, Jennifer
Cheng, Tina
Cheung, Winson
Cobb, Jennifer
Craighead, Peter
Demetrick, Douglas
Doll, Corinne M
Donnelly, Bryan
Dort, Joseph
Dowden, Scot
Duggan, Maire

Friedenreich, Christine Fujita, Don Gallo, Marco Ghatage, Prafull Giese-Davis, Janine Goodarzi, Aaron Grewal, Savraj

Hao, Desirée Hilsden, Robert Johnston, Randal Kelly, John Khan, Faisal M.

Koebel, Martin

Hamilton, Mark

Kopciuk, Karen Kurz, Ebba Lau, Harold Lee, Ki-Young Lees-Miller, Susan

Lewis, Victor Mahoney, Douglas Monument, Michael

Morris, Don Morrissy, Sorana Moules, Nancy Narendran, Aru Nelson, Gregg Neri, Paola Olivotto, Ivo A Owen, Carolyn Paschke, Ralf Peters, Cheryl Quan, May Lynn Rancourt, Derrick Riabowol, Karl

Robinson, John W Ruether, Dean Schriemer, David C Schulte, Fiona Senger, Donna

Rinker, Kristina

Robbins, Stephen

Simon, Jessica Sinclair, Shane Smith, Wendy Stewart, Douglas Storek, Jan

Sutherland, Garnette

Tang, Patricia Temple-Oberle, Claire Tremblay, Alain

Tremblay, Alain
Trpkov, Kiril
Truong, Tony
Verma, Sunil
Vogel, Hans
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Arnie Charbonneau Cancer Institute

Health Research Innovation Centre

3280 Hospital Drive NW

Calgary, AB T2N 4Z6

403.210.3934

charbonneau@ucalgary.ca

www.charbonneau.ucalgary.ca