

EDITORIAL

Bringing Access to the Full Spectrum of Cancer Research: A Call for Papers

The *PLOS Medicine* Editors*

* medicine_editors@plos.org



 OPEN ACCESS

Citation: The *PLOS Medicine* Editors (2015) Bringing Access to the Full Spectrum of Cancer Research: A Call for Papers. *PLoS Med* 12(4): e1001817. doi:10.1371/journal.pmed.1001817

Published: April 17, 2015

Copyright: © 2015 The *PLOS Medicine* Editors. This is an open access article distributed under the terms of the [Creative Commons Attribution License](http://creativecommons.org/licenses/by/4.0/), which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited.

Funding: The authors are each paid a salary by the Public Library of Science, and they wrote this editorial during their salaried time.

Competing Interests: The authors' individual competing interests are at <http://www.plosmedicine.org/static/editorsinterests.action>. PLOS is funded partly through manuscript publication charges, but the *PLOS Medicine* Editors are paid a fixed salary (their salaries are not linked to the number of papers published in the journal).

Abbreviations: AACR, American Association for Cancer Research; FDA, Food and Drug Administration; HPV, human papilloma virus; ICGC, International Cancer Genome Consortium; LMICs, low- and middle-income countries; TCGA, The Cancer Genome Atlas.

Provenance: Written by editorial staff; not externally peer reviewed

Cancer is a keyword in innumerable stories of both loss and hope. In 2012, 14.1 million people were diagnosed with cancer worldwide, with an estimated 8.2 million deaths annually [1]. The disease burden is daunting, yet long-term investment of resources in research is now yielding cause for optimism. For example, the United States Food and Drug Administration (FDA) has approved over 30 cancer drugs within the past three years [2], many of which were designed in response to insights gained from genomic investigations and developed to specifically target a gene or protein that is required for tumor growth, spread, or survival.

Understanding the clonal evolution of a cancer and the mutations that accrue over time could result in even more precise therapies. Recent findings from the cancer genomics field have also revealed the extent to which cancers are not just heterogeneous among people but also within an individual [3,4]. Propelled by improved technologies such as single-cell sequencing, longitudinal studies on an individual can trace a cancer's development over time and through metastases, providing guidance for potentially individualized therapies.

As an open-access journal committed to publishing outstanding research and commentaries on the major challenges to human health worldwide, *PLOS Medicine* endeavors to provide a prominent venue for publishing global cancer research. With growing and aging populations, low- and middle-income countries (LMICs) are disproportionately affected by the increasing numbers of cancers worldwide, with more than 60% of the world's total cancer cases and 70% of the deaths occurring in Africa, Asia, and Central and South America [1]. In lower-resource settings, the situation is made worse by the lack of early detection and access to treatment. Recent Policy Forum articles in *PLOS Medicine* have outlined key components to the establishment of national childhood cancer strategies in LMICs [5] and strategies to improve cancer care for cervical cancer in LMICs, including scale-up of human papilloma virus (HPV) vaccination and integration of care and prevention services with HIV and maternal services [6]. Given that disparities also exist in access to cancer care within high-income countries (e.g., for lung cancer treatment [7]), *PLOS Medicine* welcomes submissions of research that provides solutions to access to care within both low- and high-resource settings.

Reducing the burden of cancer means addressing the full spectrum of contributing conditions. Accordingly, in recent months, *PLOS Medicine* has published research on topics ranging from identifying and improving environmental and lifestyle-related exposures [8,9,10] to optimizing the benefits and reducing the harms of screening [11,12,13,14], as well as translating basic science towards state-of-the-art treatments [3,4,15,16] and ensuring that such interventions are available to all who need them [7,17].

Within this spectrum, the editorial team believes that translational genomics represents a “state of the art” wavelength, able to illuminate both basic pathophysiology and therapeutic

The *PLOS Medicine* Editors are Clare Garvey, Thomas McBride, Linda Nevin, Larry Peiperl, Amy Ross, and Paul Simpson.

decisions and options. The editorial team is actively engaging with the cancer genomics community, both through our academic editors, such as Andrew Beck (Dana-Farber/ Harvard Cancer Center) who discussed the importance of open access to cancer genomics data in a recent editorial [18], and also via our attendance at key conferences in 2015. Two consortia that provide broad public access to cancer sequence data are The Cancer Genome Atlas (TCGA, <http://cancergenome.nih.gov/>) and International Cancer Genome Consortium (ICGC, <https://icgc.org/>). We are very pleased to feature an accompanying interview blog [19] with Francis Ouellette, Associate Director of Informatics and Biocomputing at the Ontario Institute for Cancer Research, who discusses the remit of the ICGC and TCGA projects, how these projects have generated a tidal wave of data that has reshaped how people consider analyzing such datasets, and his hopes for how these findings will translate into clinical applications.

To coincide with the meeting of the American Association for Cancer Research (AACR), to be held in Philadelphia from April 18 to 22, 2015, *PLOS Medicine* is launching a Cancer Research Collection [20], an open-access collection of recently published articles representing the full spectrum of clinically relevant cancer research and commentary, from translational to clinical to epidemiological. To expand the collection, and to support the mission we share with the AACR to conquer cancer through research and education, we are issuing a call to the clinical genomics and cancer research community for papers that provide novel insights into cancer heterogeneity, progression, and translational and clinical medicine, with strong potential to advance patient care, public policy, or clinical research agendas. Papers submitted in response to the call for papers will be included in the collection if accepted for publication. Please submit a presubmission inquiry to the editorial team at <http://www.editorialmanager.com/pmedicine/default.aspx>.

Author Contributions

Wrote the first draft of the manuscript: CG. Wrote the paper: CG LN TM AR LP PS. Agree with manuscript results and conclusions: CG LN TM AR LP PS. All authors have read, and confirm that they meet, ICMJE criteria for authorship.

References

1. Stewart BW, Wild CP, editors (2014) World Cancer Report 2014. <http://www.iarc.fr/en/publications/books/wcr/index.php>. International Agency for Research on Cancer.
2. US FDA (2015) Hematology/Oncology (Cancer) Approvals & Safety Notifications. <http://www.fda.gov/Drugs/InformationOnDrugs/ApprovedDrugs/ucm279174.htm>. Accessed 17 March 2015.
3. Schwarz RF, Ng CKY, Cooke SL, Newman S, Temple J, et al. (2015) Spatial and Temporal Heterogeneity in High-Grade Serous Ovarian Cancer: A Phylogenetic Analysis. *PLoS Med* 12(2): e1001789. doi: [10.1371/journal.pmed.1001789](https://doi.org/10.1371/journal.pmed.1001789) PMID: [25710373](https://pubmed.ncbi.nlm.nih.gov/25710373/)
4. Mroz EA, Tward AM, Hammon RJ, Ren Y, Rocco JW (2015) Intra-tumor Genetic Heterogeneity and Mortality in Head and Neck Cancer: Analysis of Data from The Cancer Genome Atlas. *PLoS Med* 12(2): e1001786. doi: [10.1371/journal.pmed.1001786](https://doi.org/10.1371/journal.pmed.1001786) PMID: [25668320](https://pubmed.ncbi.nlm.nih.gov/25668320/)
5. Gupta S, Rivera-Luna R, Ribeiro RC, Howard SC (2014) Pediatric Oncology as the Next Global Child Health Priority: The Need for National Childhood Cancer Strategies in Low- and Middle-Income Countries *PLoS Med* 11: e1001656. doi: [10.1371/journal.pmed.1001656](https://doi.org/10.1371/journal.pmed.1001656) PMID: [24936984](https://pubmed.ncbi.nlm.nih.gov/24936984/)
6. Singhrao R, Huchko M, Yamey G (2013) Reproductive and Maternal Health in the Post-2015 Era: Cervical Cancer Must Be a Priority. *PLOS Med* 10: e1001499. doi: [10.1371/journal.pmed.1001499](https://doi.org/10.1371/journal.pmed.1001499) PMID: [23966840](https://pubmed.ncbi.nlm.nih.gov/23966840/)

7. Forrest LF, Adams J, Wareham H, Rubin G, White M (2013) Socioeconomic Inequalities in Lung Cancer Treatment: Systematic Review and Meta-Analysis. *PLoS Med* 10(2): e1001376. doi: [10.1371/journal.pmed.1001376](https://doi.org/10.1371/journal.pmed.1001376) PMID: [23393428](https://pubmed.ncbi.nlm.nih.gov/23393428/)
8. Kitahara CM, Flint AJ, Berrington de Gonzalez A, Bernstein L, Brotzman M, et al. (2014) Association between Class III Obesity (BMI of 40–59 kg/m²) and Mortality: A Pooled Analysis of 20 Prospective Studies. *PLoS Med* 11(7): e1001673. doi: [10.1371/journal.pmed.1001673](https://doi.org/10.1371/journal.pmed.1001673) PMID: [25003901](https://pubmed.ncbi.nlm.nih.gov/25003901/)
9. Ulucanlar S, Fooks GJ, Hatchard JL, Gilmore AB (2014) Representation and Misrepresentation of Scientific Evidence in Contemporary Tobacco Regulation: A Review of Tobacco Industry Submissions to the UK Government Consultation on Standardised Packaging. *PLoS Med* 11(3): e1001629. doi: [10.1371/journal.pmed.1001629](https://doi.org/10.1371/journal.pmed.1001629) PMID: [24667150](https://pubmed.ncbi.nlm.nih.gov/24667150/)
10. Zheng W, McLerran DF, Rolland BA, Fu Z, Boffetta P, et al. (2014) Burden of Total and Cause-Specific Mortality Related to Tobacco Smoking among Adults Aged ≥45 Years in Asia: A Pooled Analysis of 21 Cohorts. *PLoS Med* 11(4): e1001631. doi: [10.1371/journal.pmed.1001631](https://doi.org/10.1371/journal.pmed.1001631) PMID: [24756146](https://pubmed.ncbi.nlm.nih.gov/24756146/)
11. Singal AG, Pillai A, Tiro J (2014) Early Detection, Curative Treatment, and Survival Rates for Hepatocellular Carcinoma Surveillance in Patients with Cirrhosis: A Meta-analysis. *PLoS Med* 11(4): e1001624. doi: [10.1371/journal.pmed.1001624](https://doi.org/10.1371/journal.pmed.1001624) PMID: [24691105](https://pubmed.ncbi.nlm.nih.gov/24691105/)
12. Castañón A, Landy R, Cuzick J, Sasieni P (2014) Cervical Screening at Age 50–64 Years and the Risk of Cervical Cancer at Age 65 Years and Older: Population-Based Case Control Study. *PLoS Med* 11(1): e1001585. doi: [10.1371/journal.pmed.1001585](https://doi.org/10.1371/journal.pmed.1001585) PMID: [24453946](https://pubmed.ncbi.nlm.nih.gov/24453946/)
13. Tammemägi MC, Church TR, Hocking WG, Silvestri GA, Kvale PA, et al. (2014) Evaluation of the Lung Cancer Risks at Which to Screen Ever- and Never-Smokers: Screening Rules Applied to the PLCO and NLST Cohorts. *PLoS Med* 11(12): e1001764. doi: [10.1371/journal.pmed.1001764](https://doi.org/10.1371/journal.pmed.1001764) PMID: [25460915](https://pubmed.ncbi.nlm.nih.gov/25460915/)
14. Ross-Innes CS, Debiram-Beecham I, O'Donovan M, Walker E, Varghese S, et al. (2015) Evaluation of a Minimally Invasive Cell Sampling Device Coupled with Assessment of Trefoil Factor 3 Expression for Diagnosing Barrett's Esophagus: A Multi-Center Case–Control Study. *PLoS Med* 12(1): e1001780. doi: [10.1371/journal.pmed.1001780](https://doi.org/10.1371/journal.pmed.1001780) PMID: [25634542](https://pubmed.ncbi.nlm.nih.gov/25634542/)
15. Kim JH, Sohn BH, Lee H-S, Kim S-B, Yoo JE, et al. (2014) Genomic Predictors for Recurrence Patterns of Hepatocellular Carcinoma: Model Derivation and Validation. *PLoS Med* 11(12): e1001770. doi: [10.1371/journal.pmed.1001770](https://doi.org/10.1371/journal.pmed.1001770) PMID: [25536056](https://pubmed.ncbi.nlm.nih.gov/25536056/)
16. Martinez-Torres A-C, Quiney C, Attout T, Boulet H, Herbi L, et al. (2015) CD47 Agonist Peptides Induce Programmed Cell Death in Refractory Chronic Lymphocytic Leukemia B Cells via PLCγ1 Activation: Evidence from Mice and Humans. *PLoS Med* 12(3): e1001796. doi: [10.1371/journal.pmed.1001796](https://doi.org/10.1371/journal.pmed.1001796) PMID: [25734483](https://pubmed.ncbi.nlm.nih.gov/25734483/)
17. Eng A, McCormack V, dos-Santos-Silva I (2014) Receptor-Defined Subtypes of Breast Cancer in Indigenous Populations in Africa: A Systematic Review and Meta-Analysis. *PLoS Med* 11(9): e1001720. doi: [10.1371/journal.pmed.1001720](https://doi.org/10.1371/journal.pmed.1001720) PMID: [25202974](https://pubmed.ncbi.nlm.nih.gov/25202974/)
18. Beck AH (2015) Open Access to Large Scale Datasets Is Needed to Translate Knowledge of Cancer Heterogeneity into Better Patient Outcomes. *PLoS Med* 12(2): e1001794. doi: [10.1371/journal.pmed.1001794](https://doi.org/10.1371/journal.pmed.1001794) PMID: [25710538](https://pubmed.ncbi.nlm.nih.gov/25710538/)
19. *PLOS Medicine* Editors (2015 Apr 17) Cancer genomics: data, data and more data. In: Speaking of Medicine Blog. Available: <http://blogs.plos.org/speakingofmedicine/2015/04/17/interview-francis-ouellette/>
20. *PLOS Medicine* Editors (2015) Cancer Research Collection homepage. Available: www.ploscollections.org/plosmedicinecancerresearch