

# Eric Edward Bryant, PhD

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## Education & Experience

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- 2011—'18     **PhD, Columbia University**  
*Department:* [GSAS Biological Sciences](#)  
*Thesis:* [Systems genetics of DNA damage tolerance — cisplatin, RAD5 & CRISPR-mediated nonsense](#)  
*Mentor:* [Rodney Rothstein](#)  
*Co-Mentor:* [Alberto Ciccia](#)  
*Committee:* [Songtao Jia](#), [Elizabeth Miller](#) & [Matthew Weitzman](#)  
*Keywords:* Systems genetics / Genetic interaction networks / Landscape enrichment analysis / Synthetic lethality / DNA replication, recombination & repair / Chromosome mobility / CRISPR iSTOP / Bioinformatics / R programming
- 2009—'11     **Gene Oracle, Research Assistant**  
*Role:* De-novo gene synthesis & degenerate codon library construction. Pipeline management and development.  
*Manager:* [Shannon Phan](#)
- 2006—'09     **BS, University of California Los Angeles**  
*Department:* [UCLA Microbiology, Immunology & Molecular Genetics](#)
- 2004—'06     **West Valley College**  
*Department:* [WVC Biology](#)

## Software

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- 2019     **[screenmill R package](#): capture, annotate, quantify, review and analyze time-series colony growth.** This package was used for colony quantification and interaction analysis in Bryant et al. [2019](#). A nice example analysis using screenmill can be found in [Figure 1D](#).
- 2017     **[iSTOP R package](#): design guides to introduce stop codons with CRISPR-mediated base editors.** This package was written to facilitate guide design for Billon et al. [2017](#). iSTOP can be configured to generate any desired missense mutation using any hypothetical base editor.

## Teaching

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- 2013     **Columbia University, Teaching Assistant for Molecular Biology**  
*Professors:* [Songtao Jia](#) & [Ron Prywes](#)
- 2012     **Columbia University, Teaching Assistant for Cell Biology**  
*Professors:* [Elizabeth Miller](#) & [Chloë Bulinski](#)

## Awards

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- 2018     Departmental distinction for PhD dissertation defense
- 2017—'18     TL1 NIH training grant, clinical and translational research
- 2016—'17     T32 NIH training grant, cancer biology
- 2013—'15     T32 NIH training grant, biological sciences
- 2013     James Howard McGregor award (student with unusual promise as a teacher of zoology)

## Member

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- 2012—'18     [New York Academy of Sciences: Genome Integrity Discussion Group](#)
- 2012—'18     [Genetics Society of America](#)
- 2006—pres.     [NOLS Alumni](#)

## Publications

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### Lead contribution

- 2019-09-26 Rad5 dysregulation drives hyperactive recombination at replication forks resulting in cisplatin sensitivity and genome instability.  
**Bryant EE**, Šunjevarić I, Berchowitz L, Rothstein R, Reid RJD.  
*Nucleic Acids Research*. 2019 Sep 26;47(17):9144–9159  
[PMID: 31350889](#) – [PMCID: PMC6753471](#) – [DOI: 10.1093/nar/gkz631](#)
- 2019-01-09 Systems genetics of DNA damage tolerance – Cisplatin, *RAD5* & CRISPR-mediated nonsense.  
**Bryant EE**.  
*Columbia University*.  
[DOI: 10.7916/d8-k1do-kb09](#)
- 2017-09-21 CRISPR-mediated base editing enables efficient disruption of eukaryotic genes through induction of STOP codons.  
Billon P\*, **Bryant EE\***, Joseph SA, Nambiar TS, Hayward SB, Rothstein R, and Ciccio A.  
*Molecular Cell*. 2017 Sep 21;67(6):1068–1079.e4  
[PMID: 28890334](#) – [PMCID: PMC5610906](#) – [DOI: 10.1016/j.molcel.2017.08.008](#)  
\*co-first authors

### Supporting contribution

- 2019-10-01 DNA damage triggers increased mobility of chromosomes in G1 phase cells.  
Smith MJ, **Bryant EE**, Joseph FJ, Rothstein R.  
*Molecular Biology of the Cell*. 2019 Oct 1;30(21):2620–2625  
[PMID: 31483739](#) – [PMCID: PMC6761769](#) – [DOI: 10.1091/mbc.E19-08-0469](#)
- 2018-09-01 Increased chromosomal mobility after DNA damage is controlled by interactions between the recombination machinery and the checkpoint.  
Smith MJ, **Bryant EE**, Rothstein R.  
*Genes & Development*. 2018 Sep 1;32(17-18):1242–1251  
[PMID: 30181361](#) – [PMCID: PMC6120718](#) – [DOI: 10.1101/gad.317966.118](#)
- 2016-10-01 A synthetic dosage lethal genetic interaction between CKS1B and PLK1 is conserved in yeast and human cancer cells.  
Reid RJD, Du X, Šunjevarić I, Rayannavar V, Dittmar J, **Bryant EE**, Maurer M, and Rothstein R.  
*Genetics*. 2016 Oct 1;204(2):807–819  
[PMID: 27558135](#) – [PMCID: PMC5068864](#) – [DOI: 10.1534/genetics.116.190231](#)