**eMERGE Network Proposal for Analysis**

Project/Manuscript Concept Sheet

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| **Reference Number** | NT260 |
| **Submission Date** | 10/20/2017 |
| **Project Title** | Association of *CYP2D6* genotypes in the eMERGE PGx Cohort with EHR-derived phenotypes. |
| **Tentative Lead Investigator (first author)** | John Logan Black III MD  |
| **Tentative Senior Author (last author)** | Iftikhar Kullo MD |
| **All other authors**  | Hugues Sicotte PhD, Sandra Peterson, Laura Rasmussen-Torvik, Cindy Prows, Adam Gordon, Teri Manolio, Josh Denny, Dan Roden and other interested eMERGE investigators |
| **Sites Involved** | All eMERGE PGx sites |
| **Background and Significance** | * CYP2D6 is difficult to genotype from short read NGS data due to a large number of CNV-related conditions and star alleles that are present with variable haplotypes
* We have generated a novel approach to determining CNV of CYP2D6 using NGS short read technology based upon an understanding of the CYP2D locus
* This novel approach has been encoded into a software solution called CNVAR. The software not only identified CNVs (deletions, duplications, multiplications, hybrid genes) but also provides star allele genotypes
* CNVAR is in clinical production at Mayo Clinic
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| **Outline of Project** | Aim I. Determine CYP2D6 genotype of samples in the eMERGE PGx dataset that were sequenced using PGRN\_SEQv1 and estimate allele frequencies, noting novel alleles where possible.Aim II. Investigate the association of CYP2D metabolizer status with adverse drug response phenotypes and conduct PheWAS of metabolizer status. |
| **Desired****Variables (essential for analysis****indicated by \*)** | Age, sex, ADR codes, phecodes for PheWAS. |
| **Desired data** | We require and have access to the required sequencing files. |
| **Planned Statistical Analyses** | Simple descriptive analysis of allele frequency and number of samples that cannot be genotyped using CNVAR. Results will have unique identifier, genotype (\*allele), phenotype data (metabolizer status). A genotype to phenotype look up table will be provided. |
| **Ethical considerations** | None |
| **Target Journal** | * TBD
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| **Milestones** | October 2017: Proposal submissionNovember 2017: Initiate analysis of sequencing filesNovember 15, 2017: Possible poster submission to Pacific Symposium on Biocomputing (stretch goal)December 2017: Complete analysis of all samplesJanuary 2018: Return results to eMERGEMarch 2018: Manuscript preparationMay 2018: Manuscript submission |