Inclusion of intraocular pressure data into the University of California Health Data Warehouse

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Funding sources: This work is supported by National Institutes of Health/National Eye Institute Grants (P30EY022589, UL1TR001442, DP50D029610, R01MD014850) and an unrestricted grant from Research to Prevent Blindness (New York, NY). The sponsor or funding organization had no role in the design or conduct of this research.

Financial Disclosure: Sally L. Baxter reported grants from the National Institutes of Health during the study conduction, grants from Research to Prevent Blindness and the University of California; equipment from Optomed and Topcon; and nonfinancial support from the University of California Office of the President Grant outside the submitted work. No other disclosures were reported.

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ABSTRACT

Background

Efforts to bring ophthalmology data into OMOP have often been behind those of other fields [1]. There is ongoing work by the Eye Care and Vision Working Group to improve this, through vocabulary gap analysis, data transformation design and institutional efforts to implement ETLs [2]. This work documents a cumulation of such efforts, where intraocular pressure, a key ophthalmology datapoint, was loaded into the University of California Health Data Warehouse (UCHDW), an OMOP data warehouse that aggregates data from multiple UC health systems.

Intraocular pressure (IOP) is considered an ophthalmic "vital sign" and is the only modifiable risk factor of glaucoma [3], the world's leading cause of irreversible blindness. IOP has been frequently used either for cohort identification [4] or outcome evaluation [5] in many landmark clinical trials in ophthalmology. As of now, all glaucoma therapeutic approaches (i.e., hypotensive medications and surgical and laser techniques) aim to lower IOP. Without representation of IOP, it would be impossible to study the outcomes of glaucoma care. Evaluation of IOP is also important for monitoring post-operative or post-procedural outcomes (e.g., after cataract surgery or intravitreal injections) and potential adverse effects related to ocular or systemic medications.

Methods

All patients with IOP measurement records in the UCHDW as of February 2024 were included. The process involved mapping IOP terms from each site's Epic Kaleidoscope EHR to OMOP standard concepts, transforming data into the OMOP model, and ensuring data quality. The value_as_number field was generated using a text-to-float SAFE_CAST, that cast any non-numeric values to NULL. Analyses included examining sampling characteristics across patients, comparison of the IOP measurement distribution with a reference standard, characterization of outliers, and a review of sampling over time.

Results

95,700 patients had IOP measurements, with a total of 1,255,300 individual measurement events recorded (1,158,200 when null records were excluded). All loaded values were confirmed to be numeric; no string or text values (e.g., "soft") were present. The mean (SD) IOP after outlier exclusion was 15.27 (3.72) mmHg, which is consistent with previously published population-based studies. There were 1500 (0.13%) measurements that qualified as outliers (IOP>50 mmHg). The study highlighted data quality issues, such as anomalous entries (e.g., non-physiologic values) and significant artifacts in recorded measurement dates. A notable finding was that 35% of records were listed as occurring on a single day, due to EHR system transition (Figure 1).

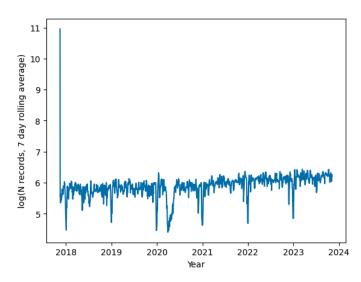


Figure 1: Distribution of IOP records over time, demonstrating an erroneous spike at the beginning of records. Note that y axis is a log scale.

Conclusion: The successful inclusion of IOP data into the UCHDW demonstrates a significant milestone in operationalizing vision data in OMOP. It underscores the importance of close communication between clinical/research teams and IT teams, and the necessity of rigorous data quality checks. The study also identifies areas for improvement in future data transformation efforts, such as handling non-numeric values and validating data against source records. The OHDSI community can use this as a case study for lessons from incorporating new specialty domains, and should be aware of the new influx of previously unavailable vision data, as similar work is being repeated across multiple institutions.

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