Designs and Methodologies Used in Landmark Clinical Trials of Glaucoma: Implications for Future Big Data Mining and Actionable Disease Treatment

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INTRODUCTION

A widely agreed upon definition of glaucoma with clear diagnostic criteria to classify disease presence and status remains elusive in both clinical and research settings. For decades, the diagnosis of glaucoma was primarily based on documenting visual field changes through static or kinetic perimetry and correlating these findings with structural changes at the optic nerve head (1, 2). Recently, the 10th World Glaucoma Association (WGA) Consensus Meeting supported the use of optic nerve structural endpoints alone to provide sufficient information for a diagnosis of glaucoma, even in the absence of visual field changes, a state termed pre-perimetric glaucoma (3). The National Institute for Health and Care Excellence (NICE), the national body tasked with issuing clinical guidelines for England and Wales, does not provide strict criteria to diagnose glaucoma (4). There is scarce guidance on criteria to definitively diagnose glaucoma which leads to difficulties in research efforts focused on phenotyping ocular imaging/diagnostic data as a first step toward identifying and predicting disease to enhance clinical care.

LANDMARK TRIALS IN GLAUCOMA

The field of glaucoma has the second largest number of published randomized controlled trials in all of ophthalmology, the majority of which evaluate glaucoma treatments (5, 6). Landmark randomized controlled trials have shaped the practice of glaucoma care and are commonly used to teach medical learners the basics of treating patients with suspected or diagnosed glaucomatous optic neuropathy (7–10). While it is clear that these trials have provided a wealth of information for practical patient care, the added benefit of these data sets is the ability to mine information that can guide the planning of future studies. This can take the form of feeding large labeled data sets into novel machine learning algorithms which potentially can produce new findings to inform patient care (11, 12). The large data sets from previous landmark trials are often published in top tier journals and receive a great deal of attention. However, the building blocks of these studies; including methodologies, patient selection criteria and diagnostic operating procedures, are usually published in a separate document and receive less rigorous attention (1, 2, 13). The aim of this brief report is to clarify the diagnostic criteria used by landmark glaucoma trials with a focus on design and methodology.

We reviewed diagnostic criteria used by landmark glaucoma clinical trials commonly cited in ophthalmic textbooks and review articles (7–10). We also supplemented the mentioned trials by...
TABLE 1 | Landmark glaucoma clinical trials and each of the functional, structural, and intraocular pressure criteria used in glaucoma diagnosis or enrollment.

| Trial | Year | IOP (mmHg) criteria | Visual field criteria | Structural criteria |
|-------|------|---------------------|-----------------------|-------------------|
| The Glaucoma Laser Trial (GLT) (1) | 1991 | >21 | Glaucomatous visual field defect/deterioration on Program 32 (1) | None |
| | | ≥27 | - | cup/disc ratio disparity ≥0.3 |
| | | ≥31 | - | cup/disc ratio ≥0.8 |
| The Advanced Glaucoma Intervention Study (AGIS) (2) | 1994 | >21 | Visual field defect score of at least (25) | - |
| Collaborative Normal-Tension Glaucoma Study (CNTGS) (19) | 1998 | 17–21 | Visual field deterioration | Disc rim deterioration |
| The Collaborative Initial Glaucoma Treatment Study (CIGTS) (13) | 1999 | ≥20 | At least three contiguous points on the total deviation probability plot at the <2% level and a Glaucoma Hemifield Test result that is "outside normal limits," (25) | Glaucomatous disc judged by physicians |
| | | 20–26 | At least two contiguous points in the same hemifield on the total deviation probability plot at the <2% level (25) | Glaucomatous optic disc |
| | | ≥27 | - | Glaucomatous optic disc |
| Ocular Hypertension Treatment Study (OHITS) (20) | 1999 | ≥24 and ≤32* | Corrected pattern standard deviation <0.05 OR glaucoma hemifield test outside normal limits (26) | Stereoscopic optic disc photographs showing a change in the position of vessels (greater than expected by eye movement), development of notch, pit, or development of thinning or pallor in the neural rim. |
| Early Manifest Glaucoma Trial (EMGT) (21) | 1999 | <30* | Glaucoma hemifield test outside normal limits (26) | - |
| | | | Glaucoma hemifield test borderline (28) | Glaucomatous optic disc features correspond to visual field |
| The European Glaucoma Prevention Study (EGPS) (22) | 2002 | >21 to ≤29* | Deterioration from baseline (22) | - |
| Low-Pressure Glaucoma Treatment Study (LoGTS) (23) | 2005 | ≤21* | At least 3 contiguous points depressed more than 8 decibels or 2 contiguous points depressed more than 10 decibels (23) | Glaucomatous optic disc consistent with visual field |
| UK Glaucoma Treatment Study (UKGTS) (18) | 2013 | <30* | Reduction in sensitivity at 2 or more contiguous points with P < 0.01 loss or more, or more contiguous points with P < 0.05 loss or more, or a 10-dB difference across the nasal horizontal midline at 2 or more adjacent points in the total deviation plot (27). | Cup-to-disc ratio of ≥0.7, focal narrowing of the neural rim, or both |

*These intraocular pressure thresholds were used for enrollment, rather than for establishing the diagnosis.

an advanced PubMed search for design and methodology articles published for glaucoma clinical trials where we used the following search strategy:

((Design[Title]) OR (Method[Title])) AND (Glaucoma [Title])

We included trials that were concerned mainly with open angle glaucoma, as this represents the majority of the effort in this space. Studies that had design protocols published in a separate “design and methodology” article were of particular interest.

Fourteen trials were identified in our assessment, however, five of them were surgical trials focused on outcome comparisons and used criteria previously utilized in larger randomized clinical trials (14–18). As a result, we focused on nine clinical trials (1, 2, 13, 19–24). Glaucoma diagnosis was based on one or more of the following criteria: functional criteria in terms of visual field performance, structural criteria in terms of optic disc features, and/or intraocular pressure (IOP). Table 1 details the specific glaucoma diagnostic criteria adopted by each of the landmark clinical trials. Visual field criterion was a pre-requisite diagnostic criterion for most clinical trials, although they varied in their definition for what qualified as a glaucomatous visual field (1, 13, 19, 22, 23, 25, 26). The Glaucoma Laser Trial (GLT) and the Collaborative Initial Glaucoma Treatment Study (CIGTS) did not require visual fields for patients with an IOP of 27 or higher, where only structural evidence of glaucomatous optic disc damage was required (1, 13). More recent trials, including the European Glaucoma Prevention Study (EGPS) (22), Low-Pressure Glaucoma Treatment Study (LoGTS) (23), and UK...
Glaucoma Treatment Study (UKGTS) (18), required structural glaucomatous features and/or visual field glaucomatous features. Figure 1 shows how the different combinations of functional, structural and IOP criteria were utilized across these studies.

**DISCUSSION**

Detailing the presence and/or progression of glaucomatous optic neuropathy is based on the functional and structural characteristics of the optic nerve, relying on the combination of both subjective and objective data obtained from clinical examination and output from various diagnostic modalities. While large randomized clinical trials have provided insights into the treatment of glaucoma, the criteria used for enrollment and interventions in each trial are disparate, making application of findings in the clinical setting difficult at best. With the emergence of artificial intelligence and machine learning techniques that seek to decipher new learnings from these large datasets, a deep and nuanced understanding of the designs and methodologies used is key to unlocking even more data to enhance patient care. We have provided an overview of diagnostic criteria used in landmark randomized controlled trials of open angle glaucoma. These criteria differ in the diagnostic weight placed on subjective visual field studies, objective structural changes of the optic nerve, as well as the use of IOP metrics. Outlining these criteria in a single resource may act as a starting point for discussions on proper methods of mining past data sets while also reaching some consensus for implementing a commonly agreed upon set of diagnostic criteria in future studies to facilitate broader analyses. The ultimate goal is to make findings from large randomized clinical trials more actionable in a real-world clinical setting by leveraging big data sets toward predictive output and guidance for when to observe and when to intervene with escalating care.

**AUTHOR CONTRIBUTIONS**

SA, ME, LS, and MK contributed in research conception, literature review, and manuscript writing. All authors contributed to the article and approved the submitted version.
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