Virtual clinics in glaucoma care: face to face versus remote decision making

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# Synopsis

The study demonstrates that virtual decision making, by senior medical staff, for stable glaucoma has reasonable agreement with face to face clinical decisions. There are no significant safety issues with virtual clinics.

#### Abstract

**Background/aims:** To examine the agreement in clinical decisions of glaucoma status made in a virtual glaucoma clinic with those made during a face-to-face consultation.

**Methods:** A trained nurse and technicians entered data prospectively for 204 patients into a proforma. A subsequent face-to-face clinical assessment was completed by either a glaucoma consultant or fellow. Proformas were reviewed remotely by one of two additional glaucoma consultants and 12 months later, by the clinicians who had undertaken the original clinical examination. The inter-observer and intra-observer decision making agreement of virtual assessment versus standard care were calculated.

**Results:** We identified adverse disagreement between face to face and virtual review in 7/204 (3.4%, 95% CI: 0.9%, 5.9%) patients, where virtual review failed to predict a need to accelerated follow-up identified in face to face review. Mis-classification events were rare, occurring in 1.9% (95% CI: 0.3% and 3.8%) of assessments. Inter-observer kappa [95% confidence intervals; CI] showed only fair agreement (0.24 [0.04 to 0.43]); this improved to moderate agreement when only consultant decisions were compared against each other (k = 0.41)

[0.16 to 0.65]). The intra-observer agreement kappa [95% CI] for the consultant was 0.274 [0.073 to 0.476], and for the fellow was 0.264 [0.031 to 0.497].

**Conclusion:** The low rate of adverse mis-classification, combined with the slowly progressive nature of most glaucoma, and the fact that patients will all be regularly reassessed, suggests that virtual clinics offer a safe, logistically viable option for selected glaucoma patients.

### Introduction

Chronic disease management is a particular challenge facing the United Kingdom (UK) National Health Service (NHS). [1] It is estimated the UK population is growing at a rate of 0.7% per annum, and will have reached approximately 72 million by 2031. As with most Western countries, the UK has an increasingly ageing population; in 2003, there were 20 million individuals over the age of 50 years and this number is predicted to increase by 36% by 2031. [2] Older people are disproportionately heavy users of healthcare services, with implications for both resource funding and management. [3] The NHS is also under increasing pressure to make efficiency savings. [4 5]

Current models of care for many chronic diseases require that patients make repeated visits to hospitals and clinics in order to undergo assessments and tests. For an efficient, consultant led service, clinically relevant information should be presented in such a way to allow rapid decision making. Stable patients are then quickly identified, allowing consultants more time to discuss difficult management decisions with unstable patients. This requires the segregation of patients with the same disease process into different categories depending on clinical risk.

Glaucoma, as an example of a chronic eye condition, is the commonest cause of irreversible blindness worldwide, [6] but blindness can be prevented by early diagnosis and treatment. [7] Chronic diseases often require lifelong monitoring with periodic treatment adjustment. It has been estimated that in England around half a million people suffer with glaucoma, with up to 1.2 million suffering from ocular hypertension or suspected glaucoma; which approximates to 2.4 million NHS outpatient visits per year. [8] There are increasing capacity problems coping with the number of glaucoma related out-patient appointments within the NHS hospital eye service (HES). Modelling predicts that this is set to worsen with current methods of NHS glaucoma patient management. [9]

It has been suggested that a 'virtual clinic' model of community-based monitoring of stable glaucoma disease may help tackle capacity issues in the UK HES. For example, clinical test data can be collected by technical staff, uploaded to a server and presented to a consultant ophthalmologist logging on remotely.

Whilst there are now a number of ophthalmology services across the UK that provide versions of these 'virtual clinic' services, there is currently no evidence that this method of clinical management is at least equivalent to face-to-face consultations in safety and robustness of decision-making. The purpose of our study was to examine the agreement between the decisions made by clinicians reviewing patient data remotely with those made by clinicians undertaking a face-to-face consultation (i.e standard outpatient care). There were two research objectives: firstly to establish the inter-observer agreement between remote review clinicians and those delivering standard outpatient care, and secondly to evaluate the intra-observer agreement, by comparing the decisions made by clinicians who provided standard care with the decisions they made when reviewing patient data remotely.

### Methods

Consecutive adult glaucoma patients with a planned, greater than six monthly follow up frequency were considered by the Glaucoma Service to be suitable for this type of clinic.. Patients with poor mobility, poor visual field technique, or poor quality optic disc imaging or other diagnosis not suited to remote assessments were not considered suitable. Only patients with open angles (including pseudoexfoliation and pigment dispertion syndrome) were considered. The Hospital's Performance and Information team provided the details of patients' follow up schedules and over a 7 month period from March to September 2011 for two adult glaucoma outpatient clinics.

The 'virtual clinic' was set up in the Hospital's Clinical Research Facility and was staffed by 3 technicians and an ophthalmic nurse. Patients underwent the same routine investigations and examinations that were to be conducted in their clinic appointment. The flow of the clinic is a follows: each technician manned a designated workstation: visual acuity measurement, visual field measurement (Humphrey Field Analyzer, Zeiss, CA), optic disc imaging. The latter consisted of disc and macular photography (TRC NW8, Topcon Medical Systems, Oakland, USA) and scanning laser ophthalmoscopy with the Heidelberg Retina Tomograph (HRT; Heidelberg Engineering, Heidelberg, Germany). A questionnaire was used to capture aspects of the patient's symptoms and history, which was reviewed by the nurse. The nurse also measured and recorded intraocular pressure (IOP) data using a Goldmann Applanation Tonometer and made a slit-lamp assessment of the anterior segment. The nurse then completed a data capture proforma that contained aspects of the clinical examination from that day, including any comment on the patient's symptoms where relevant. Advice was offered by the nurse on common anterior segment complaints, such as blepharitis. Significant drop related side effects were recorded. The nurse also observed drop technique and offered advice where appropriate. The patient then went on to see a consultant (JC) or senior fellow (RP) for their face-to-face consultation. The clinicians' observations of the patient's glaucoma status and change in management were recorded. The follow up outcomes were subsequently risk stratified to best fit the categories required in the remote reviews and shown in Table 1.

The audit department authorized the study team (audit code: CA15/GL/25) to collect data on the patients for remote assessment by members as described below.

Patient details including ethnicity, diagnosis, general health status, angle assessment, central corneal thickness measurement, maximum IOP over the course of their follow up in the service, results of any phasing examinations and drop usage were recorded on a separate paper proforma, along with a summary of the clinical assessment at their preceding outpatient appointment. These data, along with all current and previous visual field plots and current HRT results (including trend analysis where available) were available for 'remote review'. Consultants undertaking the remote review (PF, KB) also had access to digital optic disc photographs.

Remote reviewers were asked to indicate when the next follow up visit should be, based on the patient's risk of developing significant visual loss over the follow up interval (Table 1). They were also advised that they could comment upon the glaucoma status and whether they noted any other factor in the examination that warranted further investigation.

Very low risk	Low risk	Medium risk	High risk	Emergency
Stable review	Stable review	Consultant	Consultant	Immediate
appointment	appointment 6	clinic	clinic	referral to
12 months	months	appointment	appointment	glaucoma
		within 4	within 2 weeks	clinic or A&E
		months		

Table 1: risk stratification for remote assessors

Twelve months after the patients' visits, the two clinicians who conducted the face-to-face assessment (JC and RP) were asked to perform the remote review process in order to determine whether their management decisions would be the same as that made when they undertook the face-to-face assessment. The time frame was an attempt to prevent the clinicians from recalling their original decisions.

Following collection of data, the characteristics of the clinical findings that led to the clinical conclusion of progressive change were presented to the co-authors. Agreement between the authors that there was evidence of disease progression was then achieved before data analysis was completed.

#### Data analysis

Cohen's Kappa statistic used to assess both interobserver (i.e. face to face clinicians' versus remote review clinicians') agreement and intraobserver (same clinicians' face to face versus their deferred/later remote review) agreement.

Using the face-to-face review as the reference standard, the sensitivity and specificity of the ability of the virtual clinic assessment to detect unstable disease was also calculated. We also looked at agreement between specific clinicians.

#### Results

Two hundred and eighty-six patients reviewed over a 6 month period and screened for eligibility for inclusion, with 217 meeting the criteria for having their data remotely reviewed. Data for 13 patients were irrevocably damaged following a flooding incident; thus complete data were available for 204 patients.

The primary diagnoses were: primary open angle glaucoma n=88 (43%), primary ocular hypertension n= 59 (29%), glaucoma suspects n= 33 (16%), secondary glaucoma n = 18 (9%), secondary OHT n = 4 (2%) and primary angle closure n = 2 (1%). The average [standard deviation; range] visual field mean deviation in the better and worse eyes were - 1.63 [2.66; -15.14 to 3.35] dB and -4.96 [5.84; -27.94 to 1.99] dB respectively.

At the face-to-face assessment, 21 (10%) patients were identified as being 'unstable', in that they required a change in management. In the virtual clinic assessment, patients deemed as 'very low' or 'low risk' were classed as 'stable', whilst those deemed 'medium – high risk' or 'emergency' were recorded as 'unstable'. Where the reviewer had indicated that the classification was the result of a non-glaucomatous issue, and that the glaucoma itself was stable, patients were marked as 'stable'.

## Inter-observer agreement

All 5 authors reviewed the hospital medical records for patients considered unstable on face to face assessment, and stable on virtual assessment (N= 14). Thus, consensus agreement was ensured for the face-to-face clinical suspicion of "unstable" glaucoma or OHT. This consensus opinion led to 7 patients deemed 'unstable' at their face-to-face review as being reclassified 'stable'. The kappa table is shown in Table 2. The kappa statistic for the inter-observer agreement, comparing face-to-face with virtual assessment, was 0.320 [95% confidence intervals, CI, 0.112 to 0.529], indicating 'fair agreement'. The sensitivity and specificity [95% CI] of the virtual clinic was 50.0 [23.0 to 77.0]% and 91.6[86.7 to 95.1]%, respectively.

	Face-to-face Face-to-face		Total	
	'stable'	'unstable'		
Remote	174	7	181	
'stable'				
Remote	16	7	23	
'unstable'				
Total	190	14	204	

## Table 2: Kappa table for interobserver agreement analysis.

Face to face reviews were performed by either a consultant or senior glaucoma fellow (*JC*,*RP*) and remote reviews were performed by one of 2 consultants (*PF*, *KB*). Agreement was 'fair' between face to face and remote review.

Furthermore, we examined the agreement between consultants only, excluding any data from senior trainees (fellows). Table 3 illustrates the agreement between consultant face-to-face review and remote assessment. This suggests that consultant virtual clinic decisions agree well with consultant face-to-face management decisions.

Grade	Remote decision	% Sensitivity	% Specificity
	agreement kappa	[95% CI]	[95% CI]
	[95% CI]		
Consultant	0.406	75.0	89.1
	[0.161 to 0.651]	[34.9 to 96.8]	[81.7 to 94.2]

Table 3: Agreement, sensitivity and specificity between virtual clinic decisions andclinical management decisions made by consultant.The data suggest that consultantsundertaking remote reviews agree well with the consultant undertaking face-to-face review.

The 7/204 (3.4%, 95% CI: 0.9%, 5.9%) patients who were "misclassified" as stable during the virtual clinic assessment, but 'unstable' at the face-to-face review, comprised of one patient with POAG, 2 glaucoma suspects and 4 patients with glaucoma secondary to either pseudoexfoliation or pigment dispersion syndrome. Further details of their diagnoses and visual status are reported in Table 4. The characteristics suggest that 5 of the patients had 'early' disease based on the depth of their visual field defect. [10] The other 2 with more advanced disease were recommended to re-attend in 6 months in the virtual clinic assessment.

Diagnosis	Age	RVA	LVA	RMD	LMD	Recommended
	(years)	(Snellen)	(Snellen)	(dB)	(dB)	follow up
						remote review
PXF	84	6/24	6/12	-1.59	-0.84	12 months
glaucoma						
PXF	74	6/6	6/6	-0.74	-2.79	12 months
glaucoma						
PXF	61	6/6	6/6	-3.16	-3.74	6 months
glaucoma						
PDG	66	6/6	6/9	-4.11	-15.81	6 months
POAG	80	6/12	6/6	-22.16	-12.76	6 months
Suspect	53	6/5	6/6	0.22	0.38	12 months
Suspect	65	6/5	6/5	-1.08	-3.44	6 months

**Table 4:** Characteristics of patients who were not identified as progressing at the remote review. *Key:* PXF = pseudoexfoliation, PDG = pigment dispersion glaucoma, POAG =primary open angle glaucoma, VA = visual acuity, MD = visual field mean deviation.

## Intraobserver agreement

The consultant and fellow undertaking clinical examinations performed 'virtual clinic' assessments for 194 of the original 204 patients. The intraobserver agreement kappa [95% CI] for the consultant was 0.274 [0.073 to 0.476], and for the fellow was 0.264 [0.031 to 0.497]. The sensitivity [95% CI] and specificity [95% CI] of 'virtual clinic' management decisions were 75.0 [34.9 to 96.8] % and 81.0 [72.1 to 88.0] % for the consultant and 60.0 [26.2 to 87.8] % and 78.9 [67.6 to 87.7] % for the fellow, respectively.

In total, 6 patients were deemed as 'stable' during the virtual clinic assessment (4 by the Fellow and 2 by the Consultant), and were deemed unstable during the face-to-face examination. During the remote review, the disc imaging of the two patients progressing in the Consultants' clinical assessments was not available.

As expected, co-pathology was identified in the clinical assessment. Potentially significant symptoms were successfully identified during the remote assessment, but were rare (diplopia 1%, amaurosis fugax 0.5%, photopsia 0.5%). Appropriate, additional investigations were recommended for all of these patients. One (0.5%) leaking trabeculectomy bleb was identified clinically but not via the remote assessment. Common, non-sight threatening anterior segment disorders (episcleritis, blepharitis, drop allergy) that required additional treatment as a result of the clinical assessment, were identified remotely in only 1 of 6 (16.6%) of patients, through the patient questionnaire. New retinal findings (dry ARMD, epiretinal membrane) were identified in 4 (2%) of patients on clinical assessment and half of these were identified with the remote assessment from fundus photography.

### Discussion

Our study aimed to evaluate agreement between ophthalmologists' management decisions made during the standard outpatient visit and a 'virtual clinic'. Intra-observed agreement for both Consultant and Fellow grades were similar (0.27 versus 0.26) and only moderate and the classification categories may be too arbitrary, particularly with the separation of six and twelve months into low and very low risk. The results suggest that the virtual clinic has low sensitivity at detecting unstable disease. However, closer inspection shows that agreement and sensitivity are dependent on the grade of the clinician, with non-consultants adopting a more cautious and less consistent approach in their management decisions. Decision making between remote and face-to-face assessments are likely to differ, but the identification of progressing disease is critical. The wide range in confidence intervals for sensitivity suggest that far larger and long-term studies are required to confirm virtual clinic sensitivity for detecting progressing disease. The number of progressing patients is expected to be small in this population of patients.

A similar study was conducted by Gupta et al in a tertiary ophthalmic referral centre in India. [11] Agreement was assessed between ophthalmologists diagnosing and managing patients during their clinical examination with those made via teleophthalmology review. The kappa statistic for the diagnosis and management of glaucoma cases was 0.52 and 0.53, respectively. Another study examining the agreement between a district hospital ophthalmologist and a specialist glaucomatologist undertaking teleophthalmology reviews found similar levels of agreement in diagnosing glaucoma in African eyes.[12] The relatively low level of agreement in management decisions not unusual. A single ophthalmologist showed some disagreement with their own management decisions when presented with cases a period after their initial clinical examination. [13] There exists a diversity of clinical opinion amongst glaucomatologists on the best management options for individual patients. [14]

Decisions based on multiple investigations, all open to interpretation, are likely to lead to variations in recommended outcomes. In our study, the more experienced the clinicians were prepared to accept more uncertainty and recommend longer follow up periods. The study design forced clinicians to classify the patient's risk of glaucoma progression. The arbitrary distinction between stable ( $\geq 6$  months follow up) and unstable (<6 months follow up) may have led to over representation of poor stability identified in the study (table 2).

All of the patients remained under the hospitals care. This continuation of care would give an opportunity for 'instability' to be detected at subsequent visits. Glaucoma progression can be difficult to detect particularly if based on clinician judgement of paper-based visual field printouts. [15] With the inherent noise within visual field test results, clinicians may reasonably adopt a 'wait and see' approach to managing suspected disease progression. [16] In this study, the two patients (2/204, 1.9%, 95% CI: 0.3, 3.8) in whom there was relatively advanced visual field loss and disagreement between clinical assessment and virtual assessment were recommended to be seen again in 6 months. Progression seen at that virtual assessment would instigate a change in management. Large scale modelling studies have shown that patients with early disease, defined as less than -6dB of mean deviation sensitivity loss,[10] are at low risk of progressing to blindness once they are in the ophthalmology healthcare system. [17] Thus, perhaps the traditional outpatient consultation should remain for patients with worse disease status, who are at a higher risk of developing significant visual loss over their lifetime.

Whilst concerning symptoms, suggestive of new onset and non-glaucoma associated eye disease, were identified remotely; other slowly changing processes, such as age related macular degeneration were not well recognised and patients should be advised of the limitations and recommended to undertake regular community optometry assessments. Suitable correspondence between HES and community optometrists is important to clarify the limitations of virtual review clinics. Local side effects to treatment and common ocular surface diseases, such as blepharitis, are poorly identified on remote assessments and a suitably trained ophthalmic nurse present at the data gathering assessments will identify and manage most of these issues. The nurse can also make recommendations on the extent of visually significant lens opacity.

In summary, our study suggests that the use of a virtual clinic in the management of stable glaucoma patients has some limitations, and may not provide the level of sensitivity at detecting unstable disease that is found in the standard outpatient environment. However, our study identified no serious safety concerns, and suggests that significant mis-classification events are rare, occurring between in 0.3% and 3.8% of assessments. The slowly progressive nature of most glaucoma and the periodic re-assessments, suggest that virtual clinics offer a safe, logistically viable option for selected patients. We recommend patients enrolled in such schemes be those at low risk of progression to significant visual loss over each follow up

interval. Remote assessments are most reliably performed by consultant level glaucoma specialist. Patients should be encouraged to see their optometrist routinely to assess for new, unrelated eye disease.

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## Contributions

JC: Study design, data collection, data interpretation, manuscript preparation

RP: Study design, data collection, data interpretation, manuscript critique

AK: Data analysis, data interpretation, manuscript preparation

PF: Study design, data collection, data interpretation, manuscript critique

KB: Study design, data collection, data interpretation, manuscript critique

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