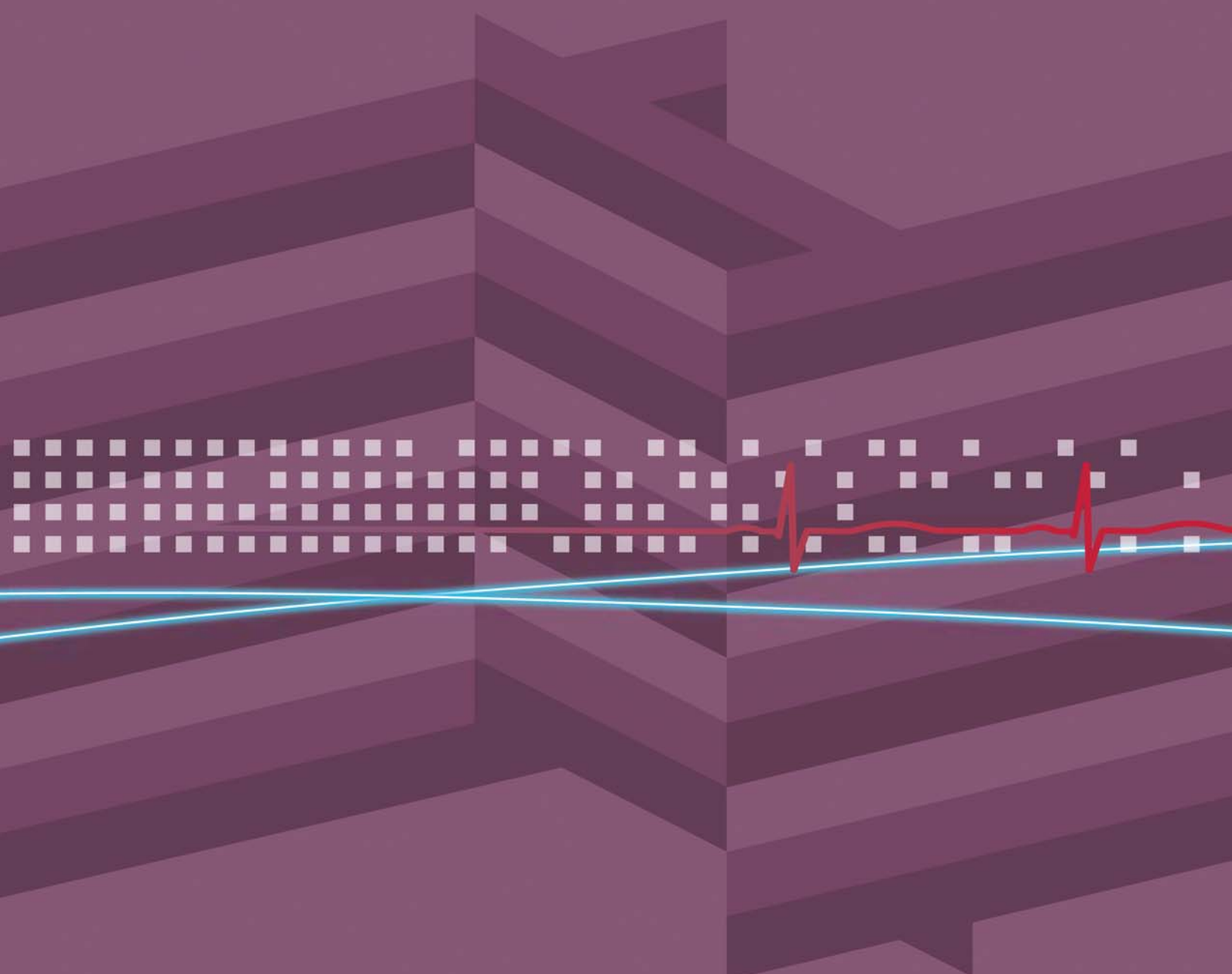


# Telehealth Practice Recommendations for **Diabetic Retinopathy** February 2011



## TABLE OF CONTENTS

I.	Preamble .....	4
II.	Introduction.....	5
III.	Background.....	6
	A.    The Diabetic Retinopathy Study (DRS) .....	6
	B.    Early Treatment Diabetic Retinopathy Study (ETDRS).....	6
	C.    Diabetes Control and Complications Trial (DCCT) /Epidemiology of Diabetes Interventions & Complications (EDIC) .....	6
	D.    The United Kingdom Prospective Diabetes Study (UKPDS).....	7
	E.    The Diabetic Retinopathy Clinical Research Network (DRCR) .....	8
IV.	Principles of a telehealth diabetic retinopathy program .....	8
	A.    Mission.....	8
	B.    Vision.....	8
	C.    Goals .....	9
	D.    Guiding Principles .....	9
V.	Ethics .....	9
VI.	Clinical validation.....	9
	A.    Category 1 .....	10
	B.    Category 2.....	10
	C.    Category 3.....	10
	D.    Category 4.....	10
VII.	Communication.....	11
VIII.	Personnel Qualifications .....	11
	A.    Medical Care Supervision.....	12
	B.    Patient Care Coordinator.....	12
	C.    Image Acquisition.....	12
	C.    Image Review and Evaluation .....	12
	D.    Information Systems .....	12
IX.	Equipment Specifications .....	12
	A.    Interoperability.....	13
	B.    Image acquisition.....	14
	C.    Compression .....	14
	D.    Data Communication and Transmission.....	14
	E.    Computer Display .....	14
	F.    Archiving and Retrieval.....	14
	G.    Security .....	15
	H.    Reliability and Redundancy.....	15
	I.    Documentation.....	15
	J.    Image Analysis.....	15
X.	Legal Requirements .....	15
	A.    Facility Accreditation.....	16
	B.    Health Insurance Portability and Accountability Act (HIPAA) .....	16
	C.    Privileging and Credentialing .....	16
	D.    Stark Act and Self-referrals .....	16
	E.    State Medical Practice Acts / Licensure .....	16

F.	Tort Liability.....	17
G.	Duty.....	17
H.	Standards of Care.....	17
I.	Consent.....	17
XI.	Quality Control.....	18
XII.	Operations.....	18
XIII.	Customer Support.....	18
A.	Originating Site.....	18
B.	Transmission.....	18
C.	Distant Site.....	19
XIV.	Financial Factors.....	19
A.	Reimbursement.....	19
B.	Grants.....	19
C.	Federal Program.....	19
D.	Other Financial Factors.....	20
E.	Equipment Cost.....	20
XV.	Summary.....	20
XVI.	Tables and Figures.....	21
XVII.	Abbreviations.....	24
XVIII.	Glossary.....	25
XIX.	Appendix.....	26
1.	Interoperability.....	26
2.	Digital Imaging and Communications in Medicine (DICOM) Metadata.....	27
3.	Computer-Aided Detection.....	28
4.	Health Insurance Portability and Accountability Act (HIPAA).....	29
5.	Privileging and Credentialing.....	30
6.	Quality Control.....	30
7.	Operational Specifications.....	34
8.	Customer Support.....	37
9.	Reimbursement.....	38
XX.	References.....	40
XXI.	Contributors.....	49

## PREAMBLE

The American Telemedicine Association (ATA), with members from throughout the United States and the world, is the principal organization bringing together telemedicine practitioners, healthcare institutions, vendors and others involved in providing remote healthcare using telecommunications. The ATA is a nonprofit organization that seeks to bring together diverse groups from traditional medicine, academia, technology and telecommunications companies, e-health, allied professional and nursing associations, medical societies, government and others to overcome barriers to the advancement of telemedicine through the professional, ethical and equitable improvement in healthcare delivery.

ATA has embarked on an effort to establish practice guidelines and technical standards for telemedicine to help advance the science and to assure uniform quality of service to patients. Guidelines and standards are developed by panels that include experts from the field and other strategic stakeholders. The guidelines are designed to serve as an operational reference and educational tools to aid providing appropriate care for patients. The guidelines and standards generated by ATA undergo a thorough consensus and rigorous review, with final approval by the ATA Board of Directors. Recommendations will be reviewed and updated periodically.

The practice of medicine is an integration of the science and art of preventing, diagnosing, and treating diseases. It should be recognized that adherence to these guidelines and standards will not guarantee accurate diagnoses or successful outcomes. The purpose of these guidelines and standards is to assist practitioners in pursuing a sound course of action to provide effective and safe medical care founded on current information and evidence-based medicine, available resources, and patient needs. ATA recognizes that safe and effective practice requires specific training, skills, and techniques as described in this document.

The goal of *Telehealth Practice Recommendations for Diabetic Retinopathy* is to support telehealth programs to improve clinical outcomes and promote reasonable and informed patient and care provider expectations. Recommendations are based on reviews of current evidence, medical literature and clinical practice. The recommendations are not intended as strict requirements. They may need to be adapted for patient care or situations where more or less stringent interventions are necessary. Recommendations are also not intended to serve as legal advice or as a substitute for legal counsel. This document does not replace sound medical judgment or clinical decision-making. Supplemental information and examples are included in the appendix.

This document is property of ATA. Any reproduction or modification of the published practice guideline and technical standards must receive prior approval by ATA.

## INTRODUCTION

Telemedicine is the use of medical information exchanged from one site to another via electronic communications. Closely associated with telemedicine is the term telehealth, often used to encompass a broader definition of remote healthcare other than clinical service.<sup>1</sup> Telehealth holds the promise of increased adherence to evidenced-based medicine, improved consistency of care and reduced cost. Goals for an ocular telehealth program include preserving vision, reducing vision loss, and providing better access to medical care.

*Telehealth Practice Recommendations for Diabetic Retinopathy* presents recommendations for designing, implementing, and sustaining an ocular telehealth care program. It also addresses current DR telehealth clinical, technical, and administrative issues that form the basis for evaluating DR telehealth techniques and technologies. These recommendations are consistent with recent federal legislation and industry best practices that emphasize interoperable health information exchange. This document will be reviewed periodically and revised to reflect evolving technologies, regulations, and clinical guidelines.

## BACKGROUND

Diabetes mellitus (DM) is a leading cause of death, disability, and blindness in the United States.<sup>2,3</sup> It afflicts as much as 8% of the American population<sup>4</sup> and the prevalence and incidence of DM is increasing in the United States and worldwide.<sup>5, 6, 7, 8</sup> The World Health Organization (WHO) estimates there are currently more than 285 million people worldwide with DM and predicts 439 million people with DM by 2030.<sup>9</sup> The United States Centers for Disease Control and Prevention estimates 18 million Americans have diagnosed DM and an additional 6.4 million have the disease but have not yet been diagnosed.<sup>5</sup>

DR is a microvascular complication of both type 1 and type 2 DM and develops in nearly all persons with DM. For the past 20 years, DR has remained the most common cause of blindness in working age adult populations in the United States and other developed countries. DR is the most frequently occurring microvascular complication of diabetes, affecting nearly all persons with 15 or more years of diabetes.<sup>10,11</sup> The WHO estimates that after 15 years of DM, approximately 2% of people with DM become blind, while 10% develop severe visual handicap.<sup>12</sup> In the Wisconsin Epidemiologic Study of Diabetic Retinopathy (WESDR), 13% of the study population with DM duration less than 5 years and 90% with duration of DM 10-15 years had some level of DR when onset of DM was prior to age 30 years (presumed to have type 1 DM). For those with onset at age 30 years or older (presumed to have type 2 DM), 40% taking insulin and 24% not taking insulin had some level of DR when the duration of DM was less than 5 years. Eighty-four percent taking insulin and 53% not taking insulin had some level of DR when duration of DM was 15-20 years.<sup>13, 14</sup> In the United Kingdom Prospective Diabetes Study (UKPDS), which enrolled subjects at the time of diagnosis of DM, nearly 40% of participants had some level of DR at diagnosis.<sup>15</sup>

DR affects more than 5.3 million Americans over the age of 18 (2.5% of the US population).<sup>16</sup> Diabetic macular edema (DME) is a manifestation of DR that may occur in any stage of DR and leads to loss of central vision.<sup>17,18,19</sup> The natural course of DME is characterized by chronic retinal vascular leakage and retinal thickening, often with intraretinal lipid deposition.<sup>6</sup> Over a 10-year period in the WESDR, DME was present in 24% of patients. Visually threatening clinically significant macular edema (CSME) was present in 10% of patients. DME is more common in type 2 diabetes patients on insulin than in type 1 diabetes patients, and prevalence increases in both types as the duration of diabetes increases.

The medical, social, and economic ramifications of DR are substantial. Evidence-based treatments in clinical studies spanning 40 years demonstrate ways to virtually eliminate the risk of severe vision loss from proliferative diabetic retinopathy. Treatments are also available to significantly reduce the risk of legal blindness and moderate vision loss. For a variety of reasons, effective treatments such as laser surgery are underutilized.

Clearly defined clinical standards for evaluating and treating DR have been established. Major multi-center clinical trials in the United States and United Kingdom provide the science behind DR clinical management.

#### *The Diabetic Retinopathy Study (DRS)*

The DRS (1971-1975) demonstrated conclusively that scatter (panretinal) laser photocoagulation reduces the risk of severe vision loss from proliferative diabetic retinopathy (PDR) by as much as 60%.<sup>20-22</sup>

#### *Early Treatment Diabetic Retinopathy Study (ETDRS)*

The ETDRS (1979-1990) demonstrated that scatter laser photocoagulation can reduce a person's risk of severe vision loss (best corrected vision of 5/200 or worse) to less than 2%. It also demonstrated that focal laser photocoagulation can reduce the risk of moderate vision loss (a doubling of the visual angle) from diabetic macular edema by 50%. There was no adverse effect on progression of DR or risk of vitreous hemorrhage for patients with DM who take up to 650 mg of aspirin per day.<sup>23-27</sup>

#### *Diabetes Control and Complications Trial (DCCT)/Epidemiology of Diabetes Interventions and Complications (EDIC)*

The DCCT (1983-1993) compared conventional blood glucose control to intensive blood glucose control in patients with type 1 DM and little or no DR. The DCCT conclusively demonstrated that for persons with type 1 DM, intensive control of blood glucose as reflected in measurements of glycosylated hemoglobin A1c:

- Reduces the risk of a three-step progression of DR by 54%
- Reduces the risk of developing severe nonproliferative DR (NPDR) or proliferative DR (PDR) by 47%
- Reduces the need for laser surgery by 56%
- Reduces the risk of diabetic macular edema by 23%<sup>28-34</sup>

Significantly, the EDIC study showed that at 10 years after the completion of the DCCT, subjects in the intensive control group continued to show a substantial decrease in risk of progression of DR compared to the conventional control group, despite a near convergence of hemoglobin A1c levels in both groups.<sup>35,36</sup>

*The United Kingdom Prospective Diabetes Study (UKPDS)*

The UKPDS (1977-1999) demonstrated similar findings to the DCCT for persons with type 2 DM.<sup>37,38</sup> As in the DCCT/EDIC studies, a legacy effect was noted, with subjects with intensive control continuing to have a lower risk of microvascular complications despite convergence of A1c levels in the intensive and conventional control groups.<sup>39</sup>

*The Diabetic Retinopathy Clinical Research Network (DRCR network)*

The DRCR network is a collaborative network funded by the National Eye Institute in the United States to facilitate multicenter clinical research of diabetic retinopathy, diabetic macular edema and associated conditions. The study involves more than 200 centers and 109 active sites nationwide in the US.<sup>40,41</sup> Current and completed studies conducted by the DRCR Network are available on the networks website.

Because DR is often asymptomatic in its early stages, many people do not seek annual retinal examination as recommended by the American Diabetes Association, the American Academy of Ophthalmology, the American Optometric Association, and other professional societies. Others may lack care due to socio-economic factors, geographic or travel restrictions, or ignorance of the need for regular retinal examination for DR. It is estimated that 40 to 50% of adults with DM in the US do not receive recommended eye care to diagnose and treat DR.<sup>42</sup> Studies also show no major improvement in examination rates over the previous five years.

Approximately 26% of patients with type 1 DM and 36 percent with type 2 DM have never had their eyes examined.<sup>43</sup> These patients tend to be older, less educated, and more recently diagnosed than those receiving regular eye care.<sup>43</sup> They are also likely to live in rural areas and receive healthcare from a family or general practitioner.<sup>43</sup> Alarming, in one study, 32% of patients with DM at high risk for vision loss had never received an eye examination.<sup>44</sup> Upon examination, however, almost 61% exhibit DR, cataract, glaucoma, or other ocular manifestations of DM.

The prevalence of DR is high and the incidence is growing in step with worldwide increases in DM. Loss of vision due to DR has a considerable impact on personal and societal resources. In the US, approximately 24,000 persons become blind from DM each year. It is estimated that programs to identify and treat DR could annually save the US healthcare budget nearly \$400 million.<sup>45,46</sup>

DR is readily diagnosed by appropriate examination. Film-based retinal imaging has been a mainstay of DR clinical care and research for many years. Digital retinal imagery is a relatively new tool for assessing patients with DR. Seven standard field photography with digital images has been accepted as the standard for the DRCR network. Digital retinal

imagery as part of a telehealth program has the potential to increase DR diagnoses, resulting in timely treatment and preservation of vision.

DM and its eye complications provide an ideal model for telehealth initiatives. DR care has a firm foundation in evidenced-based medicine. DR is classified by specific retinal lesions, exacts a significant personal and socioeconomic toll, and is a treatable disease. According to the WHO, telehealth programs are “designed to integrate telecommunications systems into the practice of protecting and promoting health,” while “telemedicine programs are designed to integrate telecommunications into diagnostic and therapeutic intervention for the practice of curative medicine.”<sup>47</sup> Ocular telehealth and telemedicine have the potential of delivering eye care to those without access. They can also provide enhanced care to those with readily available ocular care. Telehealth programs can establish and enforce quality of care by linking to national clinical trial scientific data; offer education modules to healthcare professionals, patients, and communities; and facilitate recruitment for clinical trials. The American Diabetes Association recognized the value of fundus imaging in its *2010 Clinical Practice Recommendations* for DR when they noted:

“High-quality fundus photographs can detect most clinically significant diabetic retinopathy. Interpretation of images should be performed by a trained eye care provider. While retinal photography may serve as a screening tool for retinopathy, it is not a substitute for a comprehensive eye exam, which should be performed at least initially and at intervals thereafter as recommended by an eye care professional.”<sup>48</sup>

### **PRINCIPLES OF A TELEHEALTH DIABETIC RETINOPATHY PROGRAM**

Private individuals, public and private institutes, national and international agencies, and individual governments on multiple levels may undertake telemedicine programs for DM and DR. Designing, building, implementing, and sustaining an ocular telehealth DR program requires a clearly defined mission, vision, goals, and guiding principles. The following statements are a guide for leadership and staff in developing and sustaining appropriate and effective programs.

#### *Mission*

Increase access and adherence to demonstrated standards of care among individuals with DM.

#### *Vision*

Ocular telehealth can be an integral component of primary care for patients with DM. Ocular telehealth has the potential to expand access to diabetic retinal examinations for individuals with DM consistent with evidence-based recommendations for diabetic eye care (i.e., ETDRS, DCCT-EDIC, UKPDS, DRCR). Ocular telehealth also has the ability to extend access to diabetes eye care, offer alternative methods for receiving appropriate eye care, and integrate diabetes eye care into patients’ total healthcare.



### *Goals*

- Reduce the incidence of vision loss due to DR
- Improve access to diagnosis and management of DR
- Decrease the cost of identifying patients with DR
- Promote telehealth to enhance the efficiency and clinical effectiveness of evaluation, diagnosis and management of DR
- Promote telehealth to enhance the availability, quality, efficiency and cost-effectiveness of remote evaluation for DR

### *Guiding principles*

Although ocular telehealth programs offer new opportunities to improve access and quality of care for people with DR, programs should be developed for deployment in a safe and effective manner. Program outcomes should be closely monitored to meet or exceed current standards of care for retinal examination.

DM adversely affects most parts of the eye and has a diverse influence on visual function. As a component of informed consent, patients should be aware that a validated teleophthalmology examination of the retina may substitute for a traditional onsite dilated retinal evaluation for DR, but it is not a replacement for a comprehensive eye examination. Until telemedicine programs are developed to include all necessary components of a comprehensive eye exam, a periodic, in-person comprehensive eye examination by a qualified provider continues to be essential.

## **ETHICS**

Regardless of the program, the care of the patient should not be compromised. This responsibility encompasses a broad range of issues including, but not limited to, confidentiality, image quality, data integrity, clinical accuracy and reliability.

## **CLINICAL VALIDATION**

Multicenter, national clinical trials provide evidence-based criteria for clinical guidelines in diagnosing and treating DR. Telehealth programs for DR should define program goals and performance in relationship to accepted clinical standards. In general, the selection of an ocular telehealth system for evaluating diabetic retinopathy should be based on the unique needs of the healthcare setting.

ETDRS thirty-degree, stereo seven-standard field, color, 35 mm slides are an accepted standard for evaluating DR. Although no standard criteria have been widely accepted as performance measurements of digital imagery used for DR evaluation, current clinical trials sponsored by the National Eye Institute have transitioned to digital images for DR assessment. Telehealth programs for DR should demonstrate an ability to compare favorably with ETDRS film or digital photography as reflected in kappa values for agreement of diagnosis, false positive and false negative readings, positive predictive value, negative predictive value, sensitivity and specificity of diagnosing levels of retinopathy and macular edema.<sup>49-51</sup> Inability to obtain or read images should be considered a positive finding and patients with unobtainable or unreadable images should

be promptly re-imaged or referred for evaluation by an eye care specialist. One program reported the majority of patients referred due to unreadable images actually had ocular disease that would have resulted in referral if adequate images had been obtained.<sup>52</sup>

It is recognized that severity levels of DR other than those defined by the ETDRS are used for grading DR (see Table 1 for comparisons between ETDRS levels of DR and the International Clinical Diabetic Retinopathy Disease Severity Scale, and Table 2 for comparisons between ETDRS DME and the International Clinical Diabetic Retinopathy Disease Severity Scale).<sup>53</sup> Protocols should state the reference standard used for validation and relevant datasets used for comparison.

*Telehealth Practice Recommendations for Diabetic Retinopathy* recognizes four categories of validation for DR telehealth programs using ETDRS thirty-degree, stereo seven-standard field, color, 35 mm slides as a reference standard. Validation category signifies a program's overall clinical performance and goal. Information about the program's validation performance should be available to users.

#### *Category 1*

Category 1 validation indicates a system can separate patients into two categories: (a) those who have no or very mild nonproliferative DR (ETDRS level 20 or below), and (b) those with levels of DR more severe than ETDRS level 20. Functionally, Category 1 validation allows identification of patients who have no or minimal DR and those who have more than minimal DR.

#### *Category 2*

Category 2 validation indicates a system can accurately determine if sight-threatening DR is present or not present as evidenced by any level of DME, severe or worse levels of nonproliferative DR (ETDRS level 53 or worse), or proliferative DR (ETDRS level 61 or worse).<sup>25</sup> Category 2 validation allows identification of patients who do not have sight-threatening DR and those who have potentially sight-threatening DR. Patients with sight-threatening DR generally require prompt referral for management.

#### *Category 3*

Category 3 validation indicates a system can identify ETDRS defined levels of nonproliferative DR (mild, moderate, or severe), proliferative DR (early, high-risk), and DME with accuracy sufficient to determine appropriate follow-up and treatment strategies. Category 3 validation allows patient management to match clinical recommendations based on clinical retinal examination through dilated pupils.

#### *Category 4*

Category 4 validation indicates a system matches or exceeds the ability of ETDRS photos to identify lesions of DR to determine levels of DR and DME. Functionally, Category 4 validation indicates a program can replace ETDRS photos in any clinical or research program.

A telehealth program's validation category impacts clinical, business, and operational features. The category influences hardware and software technology, staffing and support, clinical workflow and outcomes, quality assurance and business plan. Equipment cost, technical difficulty and training requirements are likely higher in categories that allow patient management to match clinical recommendations based on clinical retinal examination through dilated pupils.

A telehealth program's goals and desired performance may influence choice of technology and protocol. Some programs use pupil dilation. Others perform imaging with nonmydriatic cameras and undilated pupils. A higher rate of unreadable photographs has been reported through undilated versus dilated pupils.<sup>54,55</sup> Diabetic persons often have smaller pupils and a greater incidence of cataracts, which may limit image quality if performed through an undilated pupil. Pupil dilation is associated with a very small risk of angle-closure glaucoma. Although the risk of inducing angle-closure glaucoma with dilation using 0.5% tropicamide is minimal with no reported cases in a large meta-analysis,<sup>56</sup> programs using pupil dilation should have a defined protocol to recognize and address this potential complication.

Depending on the telehealth program, images may or may not be acquired and reviewed stereoscopically. There has been concern that macular edema may not always be detectable through non-stereo imaging modalities.<sup>57</sup> One approach without stereoscopic evaluation relies on hard exudates in the central macular field or within one disc diameter of the center of the macula as a surrogate marker for macular edema.<sup>58</sup> Clinically significant macular edema is often accompanied by other DR lesions that may also independently trigger referral. It is, therefore, possible that a program without stereoscopic capabilities may be validated to identify macular edema with acceptable sensitivity.

## **COMMUNICATION**

Communication is the foundation of ocular telehealth.<sup>59</sup> Communication should be coordinated and reliable between originating and distant sites, telehealth providers and patients, and telehealth providers and other members of the patient's healthcare team. Providers interpreting retinal telehealth images should render reports in accordance with relevant jurisdictions and community standards.

## **PERSONNEL QUALIFICATIONS**

Telehealth programs for DR depend upon a variety of functions. Distinct individuals may assume these responsibilities or a person may assume several roles.

### *Medical care supervision*

An ophthalmologist with expertise in evaluation and management of DR usually assumes ultimate responsibility for the program and is responsible for oversight of image interpretation and patient well-being. Responsibilities include recommendations for

appropriate care management and providing feedback to the imager and program to ensure images are of appropriate quality.

#### *Patient care coordinator*

The patient care coordinator ensures that each patient receives DR education and complete and appropriate follow-up, especially for those meeting criteria for referral.

#### *Image acquisition*

Image acquisition personnel (“imagers”) are responsible for acquiring retinal images. A licensed eye care professional may not be physically available at all times during a telehealth session. Imagers should possess knowledge and skills for independent imaging or with assistance and consultation by telephone, including:

- Understanding basic ocular telehealth technology and principles
- Demonstrated qualifications for obtaining appropriate image fields of diagnostic quality
- Understanding the clinical appearance of common retinal diseases requiring immediate evaluation
- Communication skills for patient informed consent and education
- Understanding of angle closure glaucoma if pupil dilation is performed, including entry-level skills in screening for shallow anterior chamber and recognition of angle closure signs and symptoms

#### *Image review and evaluation*

Image review and evaluation specialists are responsible for timely grading of images for retinal lesions and determining levels of DR. Only qualified readers should perform retinal image grading and interpretation. Qualification should include academic and clinical training. If a reader is not a licensed eye care provider, specific training is required. Grading skills should be appropriate to technology used in the ocular telehealth program. A licensed, qualified eye care provider with expertise in DR and familiarity with program technology should supervise readers. An adjudicating reader may resolve ambiguous or controversial interpretation. In most cases, an adjudicating reader will be an ophthalmologist with special qualifications in DR by training or experience.

#### *Information systems*

An information systems specialist is responsible for connectivity, data integrity, availability of stored images, and disaster recovery.<sup>60, 61</sup> The specialist should be available in case of system malfunction to solve problems, initiate repairs and coordinate system-wide maintenance.

### **EQUIPMENT SPECIFICATIONS**

Telehealth systems used in the U.S. should conform to Federal Drug Administration regulations. Telehealth systems used inside and/or outside the U.S. should meet applicable international, American, and local statutes, regulations and accepted standards. Elements include:

- Image acquisition hardware (computers, cameras and other peripherals)

- Image transmission, storage and retrieval systems
- Image analysis and clinical workflow management (scheduling follow-up examinations, clinical communication management, and decision support tools)

Equipment specifications will vary with program needs and available technology (Figure 1). Equipment should provide image quality and availability appropriate for clinical needs and guidelines. The diagnostic accuracy of any imaging system should be validated prior to incorporation into a telehealth system.<sup>49-51, 62, 63</sup>

Technologies should be Digital Imaging and Communications in Medicine (DICOM) and Health Level 7 (HL 7) standards compliant. New equipment and periodic upgrades to incorporate expanded DICOM standards should be part of an ongoing quality-control program. DICOM Supplement 91 (Ophthalmic Photography), which addresses ophthalmic digital images, was released in 2004 and updated in 2009.<sup>64</sup>

### *Interoperability*

An integrated digital healthcare system is increasingly an expectation of patients, providers, and regulators. Integration occurs on several levels, leading to more than one definition of “interoperability.” The U.S. Health Information Technology for Economic and Clinical Health (HITECH) Act mandates specific measures of interoperability for electronic health records (EHR). HITECH includes personal health records in broad scale health information exchange.

Interoperability also impacts the ability to interconnect devices and computer applications with EHR and practice management systems. Terminology, hardware, software, and communication standards are required. Harmonization of these standards is needed to allow information exchange between systems and interoperable use of data by devices and software from different vendors. Conformance to standards is increasingly driven by federal regulations and market influences. DR ocular telehealth systems should include non-proprietary interoperability by using components that conform to:

- DICOM<sup>TM65</sup>
- HL7<sup>66</sup>
- Integrating the Healthcare Enterprise (IHE) Eye Care<sup>67 68</sup>
- Systematized Nomenclature of Medicine (SNOMED-CT<sup>TM</sup>)<sup>69</sup>
- Health Information Technology Standards Panel (HITSP)<sup>70</sup>

Conformance to standards does not ensure interoperability. Also, interoperability of electronic medical records, EHR and personal health records may not always be possible or practical. Exchange of DR images and reports can also be accomplished through physical media as provided by HITSP Interoperability Specification, IS 05 - Consumer Empowerment and Access to Clinical Information via Media.<sup>70</sup> Additional information about interoperability is available in Appendix 1.

### *Image acquisition*

Retinal image datasets should adhere to DICOM standards. Patient information, eye and retina characteristics, image type, and other data should be linked to image files as metadata.<sup>64</sup> Retinal evaluation data defining type of retinal examination and the retinal image set should be included and linked to image files as metadata. Additional information such as medical and surgical history, laboratory values, etc. may be included as metadata with an image set (Appendix 2).

### *Compression*

Data compression may facilitate transmission and storage of retinal images. Compression may be used if algorithms have undergone clinical validation. DICOM recognizes JPEG and JPEG2000 for lossy compression of medical images in Supplement 91, Ophthalmic Photography SOP Classes.<sup>64, 71</sup> Compression types and ratios should be periodically reviewed to ensure appropriate clinical image quality and diagnostic accuracy. Eye care providers overseeing image grading are responsible for diagnostic accuracy.

### *Data communication and transmission*

A variety of technologies are available for data communication and transfer. Ocular telehealth programs should determine specifications for transmission technologies best suited to their program. Transmission systems should have robust error checking to ensure no loss of clinical information.<sup>72</sup> Data communications should be compliant with DICOM standards. Ocular telehealth system equipment manufacturers should supply DICOM conformance statements.

If ocular telehealth applications are integrated with existing health information systems, interoperability should incorporate DICOM conformance, interface with HL7 standards, and establish appropriate routing for patient scheduling and report transmission.<sup>73</sup>

### *Computer display*

Monitors and settings should be validated for clinical diagnostic accuracy. Any validated monitor technology can be used (e.g. cathode ray tube, liquid crystal display, gas plasma panel). Retinal images used for diagnosis should be displayed on high-quality monitors of appropriate size and resolution. Displays should be calibrated regularly to ensure fidelity with original validation display conditions. Re-validation should be performed if settings are changed. Ambient light level, reflections and other artifacts should be controlled to ensure standardized viewing.

### *Archiving and retrieval*

Ocular telehealth systems should provide storage capacities in compliance with facility, state and federal medical record retention regulations. Images may be stored at imaging or reading sites, or offsite, and must satisfy all jurisdiction requirements. Past images and reports should be available.

Each facility should have digital image archiving policies and procedures equivalent to existing policies for protecting other data and hardcopy records. Telehealth programs

should also address Health Insurance Portability and Accountability Act (HIPAA) security requirements for data archive and disaster recovery.

### *Security*

Ocular telehealth systems should have network and software security protocols to protect patient confidentiality and identification of image data. Measures should be taken to safeguard and ensure data integrity against intentional or unintentional data corruption. Privacy should be ensured through a minimum 128-bit encryption and two-factor authentication technology. Digital signatures may be used at image acquisition sites. Transmission of retinal imaging studies and study results should conform to HIPAA requirements.

### *Reliability and redundancy*

Written policies and procedures should be in place to ensure continuity of care at levels similar to using hardcopy retinal imaging studies and medical records. Policies and procedures should include internal redundancy systems, backup telecommunications and a disaster plan. Digital retinal images and reports should be retained as part of patient medical records to meet regulatory, facility and medical staff clinical needs.

### *Documentation*

Readers rendering reports on DR level or other ocular abnormalities should comply with standardized diagnostic and management guidelines as established by the American Academy of Ophthalmology<sup>74, 75</sup> or the American Optometric Association.<sup>76</sup> Reports should be based on HL7 and DICOM standards software forms and meets interoperability standards. Forms should allow ocular findings be recorded to accepted standards of defined DR levels. Medical nomenclature should conform to SNOMED CT® standards. Transmission of reports should conform to HIPAA privacy and security requirements.

### *Image analysis*

Computer algorithms to enhance digital retinal image quality or provide automated identification of retinal pathology are emerging technologies. Image analysis tools for enhancing image quality (i.e., histogram equalization, edge sharpening, image deconvolution) or identifying lesions such as hemorrhages or hard exudates can be used to aid retinopathy assessment. Image processing algorithms should undergo rigorous clinical validation. Appendix 3 summarizes computer-aided detection of DR research and development.

## **LEGAL REQUIREMENTS**

Legal and regulatory issues relating to the practice of ocular telehealth are generally the same as other telemedicine modalities and carry the risk management considerations of conventional medical practice.<sup>59, 77, 78</sup> A DR telehealth program should use the same safeguards to mitigate risk.

### *Facility accreditation*

Some hospital telehealth programs fall within regulatory jurisdictions of the Joint Commission (JC) and/or Centers for Medicare and Medicaid Services.<sup>66</sup> The JC and Accreditation Association for Ambulatory Health Care (AAAHC) accredit ambulatory healthcare.<sup>66</sup> Previous and current JC standards indicate accrediting bodies cover telemedicine activities, making regulatory compliance a mandatory component for most hospital based telehealth programs. There are specific references to telemedicine in JC Environment of Care and Medical Staff sections. CMS and AAAHC requirements occur indirectly through related activities, for example, standards for contract care.

### *Health Insurance Portability and Accountability Act (HIPAA)*

Ocular telehealth programs should obtain professional consultation for HIPAA compliance specific to their program. Telehealth programs should consider HIPAA privacy<sup>79</sup> and security<sup>80</sup> regulations in clinical, administrative, and technical operation plans. Privacy and security issues are detailed in Appendix 4.

### *Privileging and credentialing*

Telehealth providers may require formal privileging and credentialing. Providers responsible for interpretation of retinal telehealth images should be credentialed and obtain privileges at originating and distant sites if required by applicable statutes, regulations, and facility bylaws.<sup>81, 82</sup> See Appendix 5 for alternatives per JC guidelines.

### *Stark Act and self-referrals*

Some telemedicine risks are more problematic than others (e.g., anti-trust, fraud and abuse, kickbacks and self-referrals). Self-referrals occur when physicians refer patients to medical facilities in which they have a financial interest. For example, an ophthalmologist places a retinal imaging workstation in a primary care provider's office at deep discount or gratis and reads images at little or no charge. The "anti-kickback" Stark statute may have been violated if patients needing laser treatment are referred to the ophthalmologist. This may be prevented by charging the primary care provider full market value for equipment and services and offering the patient a choice of referral ophthalmologists for treatment.

### *State medical practice acts / licensure*

Many telehealth legal issues assume telemedicine is the practice of medicine. These issues are addressed variably by state medical practice acts, but even in the absence of specific statutory or regulatory definitions, telehealth legal claims would be difficult to defend against otherwise.<sup>59</sup> All states require licensure for rendering medical care to patients located in the state. It could be argued that, with telemedicine, the remote patient "visits" the physician in the physician's state. However, for practical purposes, a physician is considered subject also to the medical practice laws and regulations where the patient is located.<sup>83</sup>

Some states are permissive regarding telemedicine service from physicians residing outside the state while other states provide a special "telemedicine license." Many states are amending medical practice acts to specifically address telemedicine. More than half



require full and unrestricted licensure to render care by telemedicine to a patient residing in the state.

DR telehealth providers should carefully examine telemedicine rules in the states of intended practice. Licensure summaries are provided by the American Medical Association and the Office for the Advancement of Telehealth.<sup>83, 84</sup> A comprehensive review of licensure is available at the American Telemedicine Association State Telemedicine Policy Center website.<sup>66</sup>

### *Tort liability*

Telehealth providers should consult with their professional liability carrier to ensure coverage in both originating and distant sites. Telemedicine may reduce liability risks through improved access and quality of care. However, telemedicine can also complicate traditional tort liability. Issues include which entity or physician owes a duty to the patient, standards of care, jurisdiction, and choice of law.<sup>59</sup> Although telemedicine providers should consult an attorney familiar with telemedicine law, the fundamental aspects of tort law are fairly uniform across jurisdictions:

- A physician has a duty to a patient to act within the accepted standards of care
- Standards of care were violated
- A patient suffered an injury due to the violation of standard of care

### *Duty*

A physician's duty arises from the physician-patient relationship.<sup>85</sup> Telemedicine alters the traditional context of this relationship but a telemedicine encounter is sufficient to establish the relationship.<sup>86</sup>

### *Standards of care*

The telemedicine community is in early stages of establishing standards. The American Medical Association believes medical specialty societies should develop and implement telemedicine "practice parameters," defined as educational tools and patient management strategies, to assist physicians' clinical decision making.<sup>87</sup> Because telemedicine standards of care are not well established, questions could arise regarding appropriateness of a telemedicine DR evaluation, whether appropriate technology was selected (e.g., Validation Category 1, 2, 3 or 4), or whether the outcome was sufficient. An example of a controversial outcome is failure to diagnose nondiabetic retinopathy pathology evident in images (e.g., venous occlusion, choroidal neovascular membrane), or not evident in images (choroidal melanoma anterior to the equator, peripheral retinal tear).

### *Consent*

Ocular telehealth is considered part of a treatment or procedure.<sup>88</sup> Patients have the right to autonomous, informed participation in healthcare decisions.<sup>89</sup> Informed consent is required for clinical treatments and procedures, including those delivered via telemedicine. When treatments or procedures delivered through ocular telehealth are considered low risk and within commonly accepted standards of practice, signature consent may not be required.<sup>81</sup> Ocular telehealth services for DR may satisfy these criteria. Patients should be informed that they have a choice of telehealth and non-

telehealth ocular treatments or procedures. Practitioners should provide patients information about the ocular telehealth program they would reasonably want to know, including:

- Whether the services is novel or experimental
- Differences between care delivered using ocular telehealth and face-to-face
- Benefits and risks of using ocular telehealth in the patient's situation
- Description of what is to be done at the patient's site and the remote site

### **QUALITY CONTROL**

Regardless of an image's origin, providers should ensure the quality of medical images meet specified standards. Policies should be in place to assure patient care and safety,<sup>77, 90, 91</sup> including addressing non-DR eye diseases and findings not specifically related to DM. Telehealth programs should also develop protocols that include policies and procedures for monitoring and evaluating performance.<sup>81</sup> Corrective action of undesired trends and continuing education should be included. Evaluation should be tailored to and include all components, such as image acquisition, transmission and reading. Image acquisition and reading quality assessment and performance improvement are similar to clinical settings. Quality assessment should measure staff performance, data quality and workflow. Reviews of telehealth program outcomes are fundamental to sustained quality care and ongoing performance improvement. Quality assurance should employ peer-review of outcome and identification of fallout cases to guide corrective interventions.<sup>92, 93</sup> Training and education standards should be developed. An example of performance categories, training and quality assurance method is included in Appendix 6.

### **OPERATIONS**

A DR operations manual contains key instructions and operational information. It can also describe quality assurance and staff training procedures. A comprehensive manual enables normal operations during leadership absence. Manuals should be dynamic documents that grow and evolve ever more tailored to the telehealth program. Appendix 7 describes suggested manual components.

### **CUSTOMER SUPPORT**

DR telehealth programs use advanced technology in a range of settings, operated by diverse staff with varying expertise. Support should be tailored to internal and external customers. Support can be categorized by:

#### *Originating site*

- Imager - imaging process, software, recurrent training, recertification
- Imaging device - image acquisition, device faults, preventive maintenance
- Provider/clinical contact - report interpretation, billing

#### *Transmission*

- Connectivity

- Data loss/recovery

#### *Distant site*

- Reader adjudication, recurrent training, recertification
- Diagnostic display equipment and software

Originating and distant sites may be in the same facility with data transmission contained within a single local area network. Support for such systems will be simpler than geographically distributed programs. Technical support can be divided into levels, or tiers, depending on difficulty or urgency. Tiered help desks are common and a convenient way to accommodate program needs. A DR telehealth program should establish standards for addressing customer support needs. Appendix 8 provides examples of levels and support priority.

### **FINANCIAL FACTORS**

Telehealth program sustainability depends on a well-developed business plan. Reimbursement, remuneration and cost are complex issues. Diagnostic and procedural coding, coverage and reimbursed amounts are important considerations. Because of differences between regions, payers, and clinical settings, each program should tailor billing protocols with Medicare, Medicaid, and private insurance intermediaries.

#### *Reimbursement*

Billing code and coverage are pivotal components for reimbursement. Both are needed for effective compensation. Prior to 2011, many DR telehealth programs used the 92250 (Fundus Photography with interpretation and report) Current Procedural Terminology (CPT) code. Billing is usually divided into technical or image capture, and professional or interpretation components. Some programs used CPT 92499 (Unlisted Ophthalmic Service or Procedure), which requires negotiated use with the fiscal intermediary or carrier. The Center of Medicare and Medicaid Services (CMS) approved two new codes specific for remote retinal imaging in the fall of 2010. CPT 92227 and 92228 became effective January 1, 2011. See Appendix 9 for additional information.

#### *Grants*

Grants have been used to establish telemedicine programs for defined circumstances and duration. Although an important method for proof of concept, grants are usually not viable for sustained clinical operation. DR telehealth programs should have business plans that ensure revenue for sustainability, usually through reimbursement for services via Medicare, Medicaid, or private insurance carriers.

#### *Federal programs*

There are several large telemedicine programs that reside within federal agencies and are funded by recurring federal appropriations. Examples include the Indian Health Service and the Veterans Health Administration.

### *Other financial factors*

Non-revenue producing benefits of a DR telehealth program may include cost savings and improved operational efficiencies over traditional care delivery; however, benefits may not be realized by the entity creating them. For example, patients and third-party payers may realize financial savings produced by a DR telehealth program operated by a physician, while physicians funding the program realize no savings. Under current reimbursement policy, DR telehealth may be a better business model in closed systems, such as managed care, where costs and return on investment are realized by the same entity. Government pay-for-performance incentive programs may change the relationship between program funding and reimbursement in the future. Appendix 9 contains information on logistic efficiencies, disease prevention and resource utilization.

### *Equipment cost*

With the decreasing cost of computing and telecommunications, a retinal camera may be the largest capital investment for a DR telehealth program. Imaging costs depends on many factors. Devices range from \$3,500 to over \$65,000 including fundus camera, camera back, lenses, computer, software, and network hardware. Costs can be amortized over several years. Specialty imaging devices are in development with the potential to reduce cost further.

## **SUMMARY**

Ocular telemedicine and telehealth have the potential to decrease vision loss from DR. Planning, execution, and follow up are key factors for success. Telemedicine is complex, requiring the services of expert teams working collaboratively to provide care matching the quality of conventional clinical settings. Improving access and outcomes, however, makes telemedicine a valuable tool for our diabetic patients. Programs that focus on patient needs, consider available resources, define clear goals, promote informed expectations, appropriately train personnel, and adhere to regulatory and statutory requirements have the highest chance of achieving success.

## Tables and Figures

Table 1

International clinical DR scale compared to Early Treatment Diabetic Retinopathy Study (ETDRS) levels of diabetic retinopathy

<b>International Classification Level of DR</b>	<b>ETDRS Level of DR</b>
No apparent retinopathy	Levels 10, 14, 15; DR absent
Mild NPDR	Level 20; very mild NPDR
Moderate NPDR	Levels 35, 43, 47; moderate NPDR
Severe NPDR	Levels 53A-E; severe NPDR, very severe NPDR
PDR	Levels 61,65,71,75,81,85; PDR, high-risk PDR, very severe or advanced PDR

DR = diabetic retinopathy; NPDR = nonproliferative diabetic retinopathy; PDR = proliferative diabetic retinopathy

Table 2

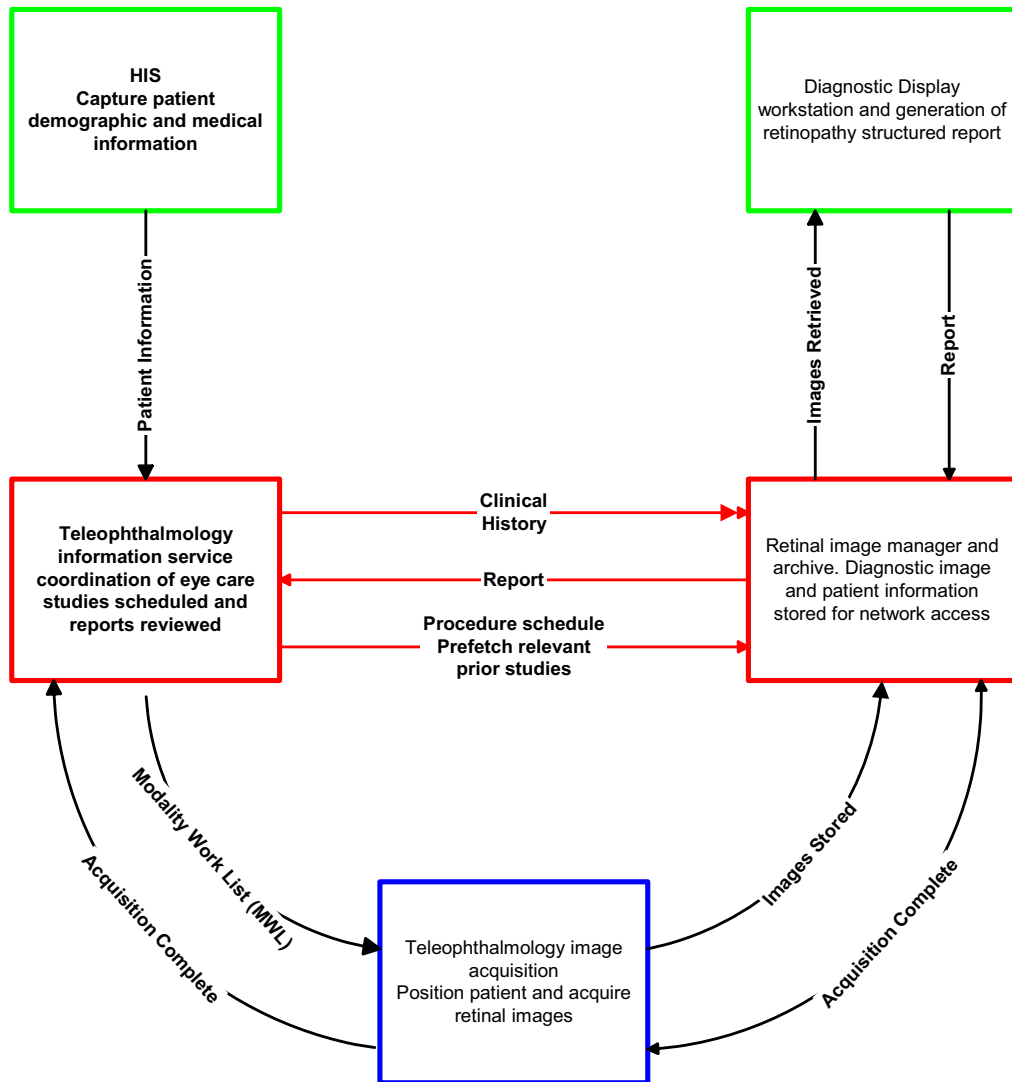
International clinical DME scale compared to ETDRS where noted.

<b>Disease Severity Level</b>	<b>Findings</b>	<b>DME scale</b>
DME apparently absent	No apparent retinal thickening or hard exudates (HE) in posterior pole	
DME apparently present	Some apparent retinal thickening or HE in posterior pole	Mild DME: some retinal thickening or HE in posterior pole but distant from center of the macula <b>(ETDRS: DME but not CSME)</b>
		Moderate DME: retinal thickening or HE approaching the center but not involving the center <b>(ETDRS: CSME)</b>
		Severe DME: retinal thickening or HE involving the center of the macula <b>(ETDRS: CSME)</b>

DME = diabetic macular edema; HE = hard exudates; CSME = clinically significant macular edema

Figure 1

Example of a workflow diagram of an ocular telehealth system based on Integrating the Healthcare Enterprise (IHE) infrastructure for radiology. Despite variations between equipment, communications and information systems, a generalized workflow diagram is possible for any ocular telehealth system.



## Abbreviations

AAAHHC	Accreditation Association for Ambulatory Health Care
ATA	The American Telemedicine Association
CE	Continuing education
CPC	Continuing Professional Competency
CPT	Current Procedural Terminology code
CQI	Continuous Quality improvement
DCCT	Diabetes Control and Complications Trial
DICOM	Digital Imaging and Communications in Medicine
DM	Diabetes mellitus
DME	Diabetic macular edema
DR	Diabetic retinopathy
DRCR	The Diabetic Retinopathy Clinical Research Network
DRS	Diabetic Retinopathy Study
DRVS	Diabetic Retinopathy Vitrectomy Study
EDIC	Epidemiology of Diabetes Interventions and Complications
EHR	Electronic health records
ETDRS	Early Treatment Diabetic Retinopathy Study
FDA	Food and Drug Administration
HITECH	Health Information Technology for Economic and Clinical Health Act
HIPAA	Health Insurance Portability and Accountability Act
HIS	Hospital information system
HITSP	Health Information Technology Standards Panel
HL7	Health Level 7
IHE	Integrating the Healthcare Enterprise
IS	Information specialist
JC	Joint Commission (formerly Joint Commission for the Accreditation of Healthcare Organizations)
JPEG	Joint Photographic Experts Group
M & M	Morbidity and mortality
NPDR	Nonproliferative diabetic retinopathy
OD	Right eye
OS	Left eye
PACS	Picture Archiving Communication System
PDR	Proliferative diabetic retinopathy
SNOMED	Systematized Nomenclature of Medicine
UKPDS	United Kingdom Prospective Diabetes Study
WESDR	Wisconsin Epidemiologic Study of Diabetic Retinopathy
WHO	World Health Organization



## **GLOSSARY**

**DICOM - Digital Imaging and Communication in Medicine**

An international standard for distributing, storing, and viewing medical images.

**HITECH Act - Health Information Technology for Economic and Clinical Health Act**  
Legislation designed to promote the adoption and meaningful use of health information technology.

**HITSP - Health Information Technology Standards Panel**

A public and private sector partnership working toward interoperability between clinical and business health information systems. HITSP is coordinated through the US Department of Health and Human Services' Office of the National Coordinator for Health Information Technology (ONCHIT).

**HL7 - Health Level 7**

An international framework for the electronic exchange of clinical, financial and administrative information among computer systems in hospitals, clinical laboratories, pharmacies, etc.

**IHE - Integrating the Healthcare Enterprise**

A global initiative by healthcare professionals and industry to improve computer sharing of healthcare information through coordinated use of established standards such as DICOM and HL7.

**NHIN - Nationwide Health Information Network**

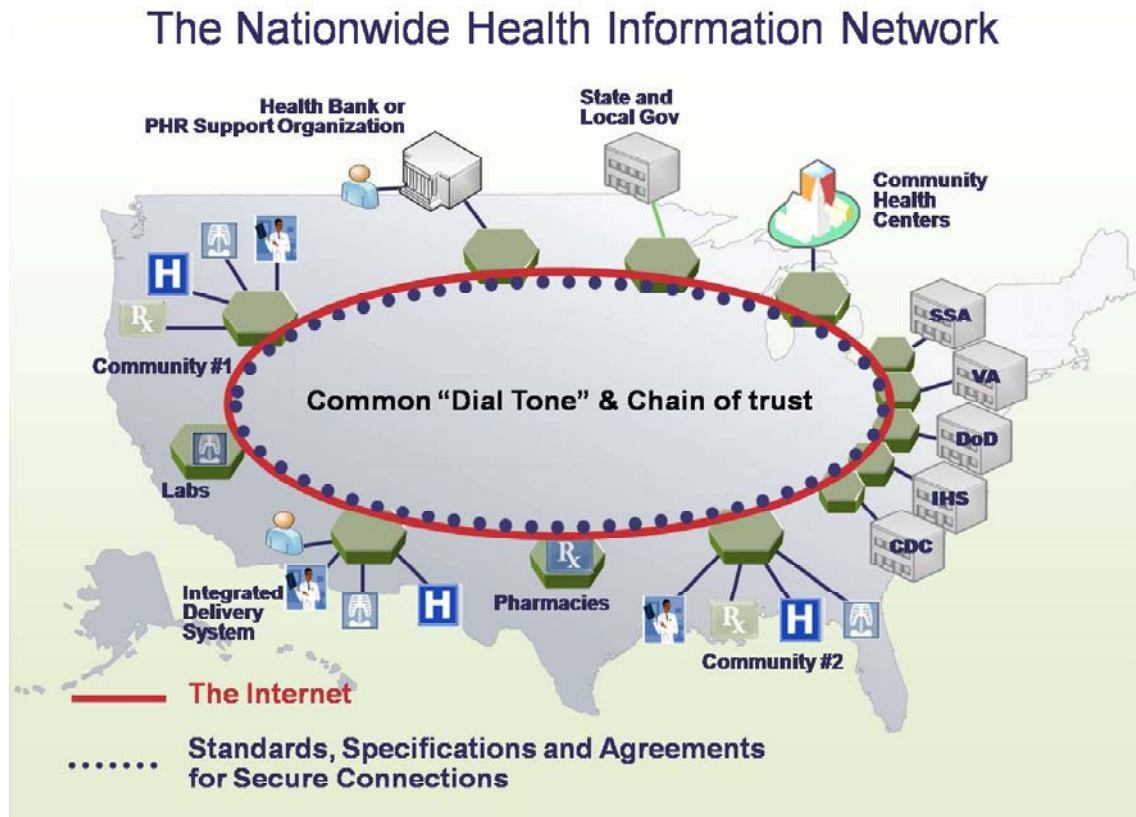
An initiative for improving the quality and efficiency of healthcare through the secure exchange of health information.

**SNOMED CT® - Systematic Nomenclature of Medicine Clinical Terms**

A system of clinical healthcare terminology covering diseases, findings, procedures, microorganisms, pharmaceuticals, etc.

## Appendix 1: Interoperability

Diagram of NHIN and CONNECT.



Interoperability should reduce errors from duplicate manual data entry and aid clinical decision support systems and alerting. In 2004, President Bush signed an executive order creating the Office of the National Coordinator for Health Information Technology (ONC). ONC provides leadership for developing nationwide interoperable health information technology infrastructures and processes. The goal is widespread adoption of electronic health records (EHR) by 2014. In 2009, President Obama signed the American Recovery and Reinvestment Act (ARRA). ARRA includes the Title XIII-Health Information Technology for Economic and Clinical Health Act (HITECH). HITECH mandates interoperability of EHR systems to allow broad health data exchange for timely and accurate access to patient health information.

Consequently, there is increasing focus on interoperability and health information exchange at the national level, especially for "meaningful use" of EHR. To help achieve the goals of the HITECH Act, the Nationwide Health Information Network (NHIN) and the CONNECT NHIN Health Information Exchange (NHIE) Gateway Community Portal are developing technologies to facilitate data exchange.<sup>66</sup> The NHIE Gateway is part of the larger NHIN CONNECT Initiative and will enable federal healthcare agencies and healthcare providers to share patient information efficiently. CONNECT has broad participation including the Centers for Medicare & Medicaid Services (CMS), Veterans

Administration (VA), Department of Defense (DoD), and the Indian Health Service. CONNECT's first user was the Social Security Administration (SSA). While CONNECT was developed to create an NHIE, its long term goal is becoming NHIN's "backbone." CONNECT was first released in December 2008. A schematic describing NHIN and CONNECT is illustrated in Figure 1.<sup>66</sup>

Key NHIN development activities are the Data Use and Reciprocal Support Agreement (DURSA) and other legal agreements for data exchange. These documents will describe information shared, accessed, used, disclosed, and privacy and security. The CONNECT NHIN consortium's November 2009 DURSA addressed many aspects of data exchange and use including consent, obligations, permitted use of data, and data ownership.

Key components include:

- Extension of HIPAA to all participants of NHIN
- HIPAA is the floor for all activities on NHIN but local and state laws that go beyond HIPAA are not preempted
- Limited permitted uses of data (e.g., neither use for research or legal/enforcement is allowed)

All participants must respond to a data request from an NHIN member. Sharing data is not required but the request for data must be acknowledged. Once data is transferred to a recipient, data is owned by the recipient and can be shared/exchanged in conformance to policies.

## **Appendix 2: DICOM Metadata**

DICOM files contain metadata with information about image data. Parenthetical codes refer to DICOM header metadata.

### Demographics

- Patient name (0010, 0010)
- Medical ID number (0010, 0020)
- Patient birth date (0010, 0030)
- Gender (0010, 0040)
- Date and time of examination (0008, 0020) and (0008, 0030)
- Name of facility or institution of acquisition (0008, 0080)
- Accession number (0008, 0050)
- Modality or source equipment that produced the ophthalmic photography series (0008, 0060)
- Referring physician's name (0009, 0090)
- Manufacturer (0008, 0070)
- Manufacturer model name (0008, 1090)
- Software version (0018, 1020)
- Station name (0008, 1010)

### Examination information

- Image type or image identification characteristics (0008, 0008)
- Instance number or image identification number (0020, 0013)
- Mydriatic (pupil dilation) or non-mydriatic (no pupil dilation) imaging. Pupil dilated Yes/No (0022, 000D), dilating agent (0022, 001C)
- Size of field or horizontal field of view in degrees (i.e., 20-degree, 30-degree, 45-degree, 50-degree, 60-degree, and 200-degree) (0022, 000B)
- Identification of single retinal field images, simultaneous or non-simultaneous stereo pairs
- Identification of stereo pairs. Left image sequence (0022, 0021), right image sequence (0122, 0022)
- Monochrome gray scale or color bit depth, ophthalmic photography 8-bit images (0028, 0100, 0028, 0101), 16-bit images (0028, 0102)
- Laterality of eye, right, left, or both eyes; OD, OS, or OU (0020, 0062)
- Retinal region such as Diabetic Retinopathy Study fields 1 to 7 (0008, 0104) .
- Ratio and type (i.e., wavelet or JPEG) of compression, if used. Lossy compression Yes/No (0028, 2112), lossy compression ratio (0028, 2112), lossy compression method (0028, 2114)
- Detector type, CCD or CMOS (0018, 7004)
- Spatial resolution of the image (i.e., 640 x 480, 1000 x 1000, etc.)
- Free text field for retinal imager study comments (i.e., presence of media opacities, poor fixation, poor compliance, etc.)
- Description of any image post-processing
- Measurement data and/or pixel spacing (0028, 0030)

### Appendix 3: Computer-Aided Detection

Different computer-based methods to assist detecting retinopathy have been developed. Most efforts rely on traditional image analysis tools such as region growing, edge detection, and segmentation algorithms to identify features such as the optic disc, retinal vascular tree and vessel crossover. Computers require mathematically exact data definitions to characterize retinal lesions of interest for analysis. Feature extraction is a fundamental step in this process. Algorithms have been developed to segment retinal images,<sup>94-101</sup> assess venous beading<sup>102,103, 104</sup> and retinal thickness,<sup>105-108</sup> quantify lesions<sup>103, 109-120</sup> and maculopathy,<sup>121-123</sup> and detect retinopathy.<sup>103, 110, 124-127</sup> Microaneurysm counting has also shown clinical value as a predictor of retinopathy development. Retinal image analysis to detect and count microaneurysms has relied on morphological, thresholding and segmentation techniques.<sup>94, 126, 128-134</sup>

Lesion detection algorithms generally perform five steps: preprocessing, localization and segmentation of the optic disk, segmentation of the retinal vasculature, localization of the macula and fovea, and localization and segmentation of retinopathy. A variety of outcome measures have been used to validate algorithms.

The use of neural networks to “teach” systems to recognize patterns has shown initial promise but has yet to achieve consistent sensitivity and specificity to be clinically acceptable.<sup>135-137</sup> Results of automated grading to diagnose retinopathy have also been mixed, although some approaches have produced encouraging results.<sup>138-141</sup> Risks in adopting computer-aided retinopathy detection include unknown algorithm lesion sensitivity and specificity, the possibility of false negatives, and/or missed referable cases.<sup>142</sup>

Change analysis assessment is a computer-based approach fundamentally different from feature detection. Instead of identifying specific lesions, change analysis highlights differences between retinal images over time for use in primary and secondary care.<sup>143</sup> This approach leaves decision-making to the readers and avoids ethical issues of software-based misdiagnosis.<sup>141</sup>

Content-based image retrieval has recently been applied to computer retinal image analysis.<sup>144, 145</sup> The concept is based on retrieving related images from a large database of adjudicated retinal images and using pictorial content to provide correct assessment of clinical diagnosis. Resulting content feature lists provide indices for storage, search, and retrieval of related images. The probabilistic nature of content-based image retrieval allows statistically appropriate predictions of presence, severity, and manifestations of common retinal diseases such as diabetic retinopathy. This approach is attractive but, given the diversity of characteristics and spatial locations of retinal lesions, it may suffer performance issues as image libraries grow. The use of fuzzy logic algorithms is being studied.<sup>146</sup>

Computer-assisted retinal image analysis is also being applied to abnormal retinal vasculature and vascular changes such as vessel diameter, tortuosity, angle and retinal vasculature fractal dimension. Retinal image quantitative measures have demonstrated predictive value in diabetic retinopathy and cardiovascular disease.<sup>147-149</sup>

When computer-assisted lesion identification or retinopathy diagnosis decision support is used, the following should be included in the free text field of DICOM headers:

- Algorithm used
- Algorithm functionality
- Algorithm's validated sensitivity and specificity

#### **Appendix 4: Health Insurance Portability and Accountability Act (HIPAA)**

Privacy elements include:

- Covered entity - any organization, person or business associate thereof that transmits protected health information in electronic or paper form<sup>150</sup>
- Covered information - individually identifiable health information in any form or medium used or disclosed by a covered entity
- Voluntary consent - covered entities may obtain a patient's consent before using or disclosing his or her health information

Electronic protected health information security includes:

- Confidentiality - requires authentication and role-based access to data (must have a need to know)
- Integrity - requires methods for assuring no unauthorized altering or destruction of data
- Availability - requires methods for disaster recovery, backup, and access to data under all conditions

## **Appendix 5: Privileging and Credentialing**

Many healthcare facilities use the Joint Commission's (JC) *Elements of Performance* for privileging and credentialing providers, even when not strictly required. At one time, *Elements of Performance* allowed privileging and credentialing telemedicine providers by proxy under specific conditions. This policy allowed the originating site to import privileging and credentialing materials from the distant site if certain criteria were met (i.e., the distant site was JC accredited), staff were privileged for the activity at the distant site, and peer review was conducted and reported to the originating site with other quality information. CMS had planned to require JC to conform to more restrictive criteria effective July 15, 2010, eliminating privileging and credentialing by proxy as described by JC MS.13.01.01. CMS reversed its position in May 2010, however, with a proposed rule defining conditions for a proxy method of privileging and credentialing telemedicine providers.<sup>151</sup> As of October 2010, this decision is still under review.

MS.13.01.01, EP 1 - Licensed independent practitioners responsible for patient care, treatment, and services via telemedicine are credentialed and privileged at the originating site according to standards MS.06.01.03 through MS.06.01.16. Standards allow medical staff to use information another hospital gathers for physician credentialing provided the other hospital is a Medicare-participating hospital. Data includes:

- Licensure
- Training
- Board certification verifications
- National Practitioner Data Bank queries
- Sanction queries
- References
- Other information gathered during the credentialing process

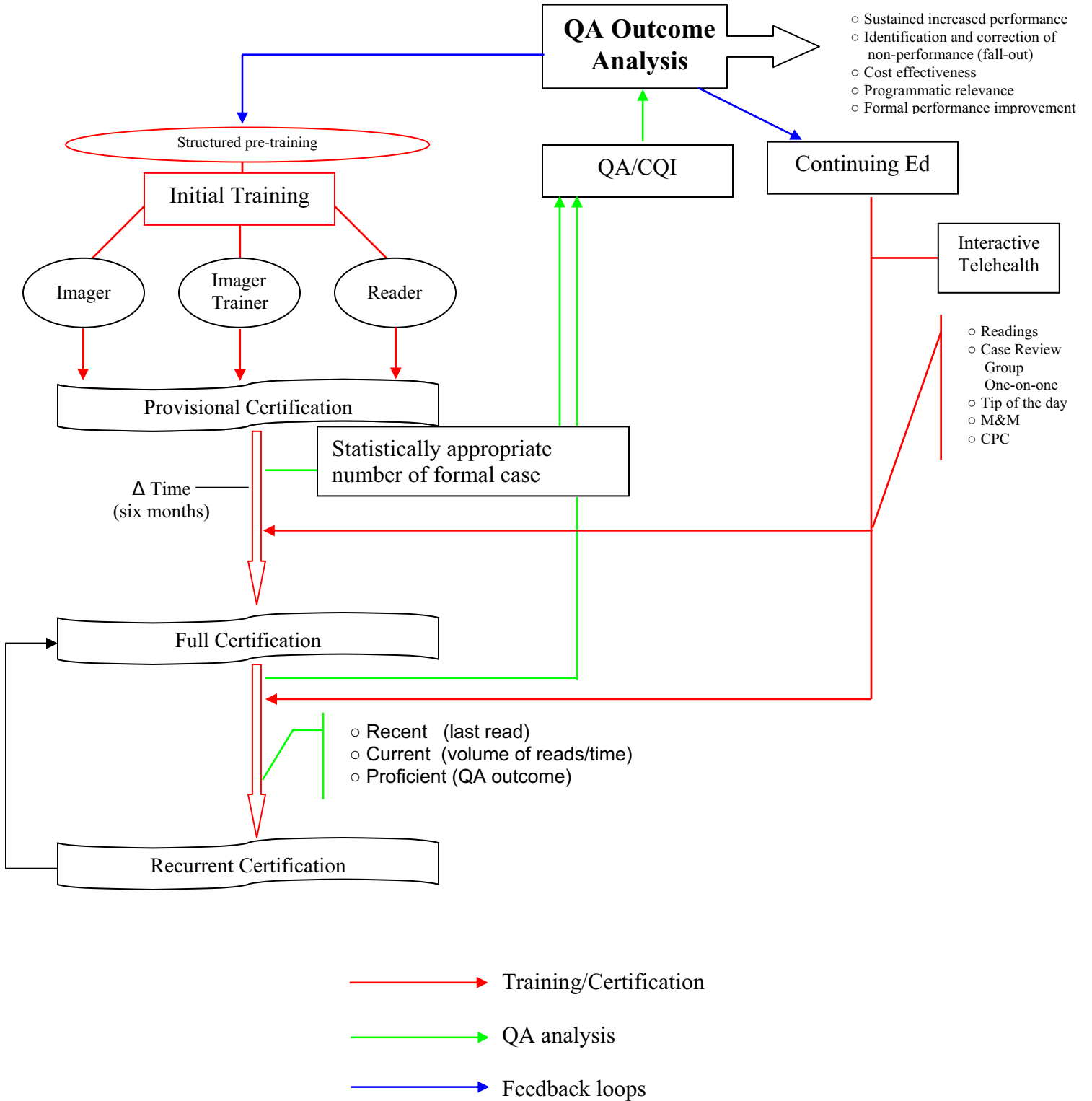
## **Appendix 6: Quality Control**

The following are major categories of performance to be evaluated; evaluation of all categories below may not be applicable to some programs:

- Originating site
  - Administrative
    - Primary care provider and nursing surveys
    - Patient surveys
    - DR surveillance rate for catchment area of the program
    - Successful patient enrollment rate (sustained vs. initial)

- Successful referral completion rate
  - Imager
    - Ungradeable study rate
      - Field definition
      - Focus
      - Stereo
    - Imaging time
    - Continuing Quality Improvement (CQI)
    - Continuing education
  - Equipment
    - Preventative maintenance schedule
    - Out of service rate
- Reading Center
  - Administrative
    - Average acquisition to reading center time
    - Average time to read standard study
    - Average time to read stat study
    - Average acquisition to report delivered time
    - Exception rate and time
  - Technical- network, servers, software, etc
  - Reader
    - Average time per read
    - Assessment program of peer review or over-reads at appropriate levels of detail, and including ungradable and exception rates
      - Inter-observer agreement, if multiple readers are used
      - Intra-observer agreement
      - External review
      - Test set performance
    - Agreement with live exam, if appropriate
    - Assessment fallout/outlier review and evaluation
    - CQI
    - Continuing education

Example of a quality assurance and continuous quality improvement flow diagram.





Multiple feedback loops allow continuing education (CE) programs to adapt to changing conditions. Reviews allow CE performance and cost effectiveness to be continuously enhanced (Figure 2). The following are examples of training and quality assurance method:

Standardized training for imager, imager trainers, readers, and reader trainers.<sup>152</sup>

- Structured, self-study, pre-training of imager and reader to provide minimum background knowledge.
- Structured curriculum training with defined endpoints to assure knowledge and skills proficiency.
- Provisional certification followed by full certification based on experience with a minimum number of patients over a minimum period of time. Experience should demonstrate required levels of proficiency documented by formal quality assurance (QA) review of a fixed number of cases.
- Time-limited certification of imagers and readers. Recertification should be based on the period since last clinical encounter, number of clinical encounters over a period of time, and proficiency as documented by formal QA review. Ocular telehealth programs should create certification methods that are formally defined and relevant to the program.

Ongoing sampling of imager and reader performance by formal criteria based on QA review should be performed. A review of trends in fallout from outcome analyses can be used to assess:

- Proficiency
- Opportunities for program improvement
- Need for changes in initial or recurring training
- Need for additional training of an imager or reader

CE is an important component of any QA/CQI (Continuous Quality Improvement) program and a fundamental method of ensuring current competency<sup>153</sup>. CE should be dynamic and sensitive to patient and staff's changing needs. The following are considerations:

- Adjust CE content by end-to-end program testing through data sampling and outcome analysis
- Adjust CE program to maintain temporal relevance to aggregate and individual clinical populations
- Deliver CE in formats to achieve desired outcome with maximum efficiency and effectiveness. Format examples include periodic self-study curriculum with pre and post-study testing, newsletters and email "Tips of the Day." A variety of interactive CE sessions using telehealth technology are available such as group-based or one-on-one case reviews, morbidity and mortality (M & M) conferences and conferences patterned on Clinical Pathological Conference (CPC) concepts.

Similar guidance comes from broadly distributed programs outside the U.S. The U.K. National Screening Committee adopted digital photography in 2000 for a systematic

national risk reduction program.<sup>154</sup> Their model incorporates trained professionals, recorded outcomes, targets and standards, quality assurance and promotion to increase screening rates. Criteria and minimal/achievable standards were proposed for each quality assurance objective.<sup>155</sup> Other ongoing quality assurance programs are publishing measures and outcomes. For example, a U.K. diabetes center re-graded a percentage of images to determine appropriateness of referrals for clinic examination.<sup>156</sup> Other programs have reported outcomes measuring image quality, intra-grader reliability and percentage of grader-generated reports within 48 hours of grading images.<sup>157</sup>

## **Appendix 7: Operational Specifications**

Possible components of an ocular telehealth operations manual include:

- Company overview & history
- Organization chart
- Mission statement
- Opening procedures
- Closing procedures
- Cash handling
- Daily tasks
- Alarm system operations
- Safe opening and closing procedures
- Contact numbers for emergencies or information
- Employee shift coverage
- Website procedures
- Customer service procedures
- Marketing
- Sales procedures
- Sales quotas
- Commission payments
- Order processing
- Special orders
- Shipping & receiving
- Equipment handling
- Imaging site deployment
- Reading center operations
- Quality assurance
- Privileging and credentialing (originating and distant site)
- Equipment maintenance (office, clinical, it, etc.)
- Security procedures
- Emergency procedures
- Services pricing and discounts

Additional details and examples that may not be applicable to all programs:

- Prospect engagement

- Sales representative receives prospect information from sales/marketing representative
- Contacts prospect via telephone, e-mail, letter, etc.
- Documents contact in CRM software (Customer Relationship Management)
- Update status of prospect after each contact
- Prospect closes and becomes client
- Contract
  - Send approved contract and lease agreement to prospect
  - Customer returns agreement with signature and deposit check or signature and lease document
  - Submit lease document to leasing company
  - Receive notification of lease approval or equipment paid in full
- Setup new customer
  - Complete the Customer Deployment Prerequisite Datasheet
  - Identify static external IP address (may require IT liaison)
  - Identify internal DNS server, Default Gateway and Subnet Mask
  - Identify user details. Assign new unique clinician identifier for each clinician. Assign a new locally unique user id for that user
  - Identify primary care clinician information. Assign new unique clinician identifier for each clinician
- Stage equipment
  - Receive application update CD and shipment checklist and installation checklist from Hosted Solution Management Team
  - Notify admin of expected ship date
  - Admin notifies customer of expected ship date
  - Assemble equipment
    - Camera back
    - Fundus camera
    - Table
    - PC
    - Desktop reference book
  - Apply the software image to the PC
  - Apply the application update to the PC
  - Run a local photography test using the customer photography instructions
  - Disassemble and repack for shipment
- Customer installation & test
  - Validate training personnel and room availability prerequisites & installation date with customer
  - Validate static IP prerequisites
  - Complete relevant parts of installation checklist Carry out install as per installation document)
  - Schedule/conduct deployment test
  - Complete relevant parts of installation checklist
  - Store in appropriate customer folder

- Imager training
  - Ensure that all staff due for training are present.
  - Provide introduction to fundus camera - startup, calibration, configuration, imaging, shutdown.
  - Provide introduction to imaging software- logon, configuration, patient registration, image storage and transmission, shutdown, logoff.
  - Step through photography process using training scripts, tests and real patients.
  - Obtain sign-off from all trained staff. Complete relevant parts of installation checklist and store in appropriate customer folder.
- Photography
  - Administrator
    - Diabetic patient presents at front desk
    - Verify that the patient has not had retinal photographs within 12 months by checking records
    - Add retinal photography to their routing slip for the day
  - Photographer
    - Diabetic patient presents for photography
    - Drop the patient in to today's clinic according to the instructions
  - Monitor messaging
    - Run the "Daily Report" for all locations
    - Inspect the report to validate it conforms to the business rules (see exceptions)
  - Monitor message resend
    - Examine message queue for timed-out message send
    - Resend any outstanding messages, log the ids of the messages that have required a resend
  - Reader
    - Grade according to the protocol
  - Monitor grading throughput
    - Run the "Remote Grading Report" for all locations
    - Inspect for outstanding grading
    - Inspect for disagreements
  - Quality control grading
    - Log on to the application
    - Check the Secondary grading queue for available grading, click the "Grade" button to start grading
    - Conduct secondary Grading according to the protocol
  - Monitor service status
    - Verify service status
    - Verify bandwidth usage
    - Verify nightly logs
    - Verify/action outstanding ticket log
  - Monitor nightly backups

- Open backup log – verify that there were no errors, and that backup files exist

### Appendix 8: Customer Support

Some programs may find it appropriate to consolidate the following example of an ocular telehealth program three-level help desk.

#### Level 1

This is the entry point for most/all initial support requests. Support staff can satisfy image acquisition issues and entry level troubleshooting of software and data transmission. If the request for support is determined to be outside Level 1 scope, the call is triaged to the Level 2 or Level 3 Help Desk.

#### Level 2

For more complex software and data transmission issues a second level support is needed to provide solutions that are more technical in nature.

#### Level 3

A third level support is needed for troubleshooting and resolving proprietary technology. Typically, this concerns the imaging device and associated equipment, e.g., camera backs, relay lenses, software applications, etc.

Configuration may be arranged as to priority:

Impact Level	Definition	Target Call Back Time	Target Resolution Time
1	<b>Critical:</b> Critical System Software is entirely unavailable or severely degraded to the point of un-usability and there is no workaround/alternative	15 minutes	4 hours
2	<b>Major:</b> Non-Critical System Software is entirely unavailable or; Critical System Software is entirely unavailable or severely degraded to the point of un-usability and there is a workaround/alternative	1 hour	8 hours
3	<b>Minor:</b> Part of a System is unavailable.	Not applicable	14 working days
4	<b>Non-urgent user interface issues:</b> System has failed to meet its specification or; Request for information about how to use the System	Not applicable	21 working days
5	<b>Good Will:</b> Anything else, e.g. · State Changes · Letter Changes · Patient Merges · Resetting Grading · Correcting Observations	Not applicable	Approx 180 days

## Appendix 9: Reimbursement

### *Medicare*

CPT 92227: remote imaging for detection of retinal disease (eg, retinopathy in a patient with diabetes) with analysis and report under physician supervision, unilateral or bilateral.

CPT 92228: remote imaging for monitoring and management of active retinal disease (eg, diabetic retinopathy) with physician review, interpretation and report, unilateral or bilateral.

These new remote retinal imaging codes allow for detection of retinal disease (92227) and the monitoring and management of active retinal disease (92228). They specifically address the clinical application of telemedicine modalities for DR.<sup>158</sup> Although the new codes went into effect January 3, 2011, ATA and other national organizations requested a CMS review contending these codes poorly defined the role of telemedicine for DR and undervalued services provided by DR telemedicine applications.<sup>159</sup> In its request, ATA noted 92227's definition does not reflect actual DR remote retinal imaging clinical applications, and that 92228 does not reflect the complexity of care associated with DR remote imaging. CPT 92227 assigns zero RVUs to physicians' work. CPT 92228 significantly undervalues the physician's responsibility and care. ATA also expressed concern that 92228 restricts reimbursement to only patients with active retinal disease. Total RVUs assigned to these new codes are markedly less than the previously used CPT 92250 (fundus photography), although similar equipment, staff and physician effort are involved. ATA joined with many teleophthalmology-DR programs and several specialty professional societies to formally contest the description and reimbursement values of the new codes during the public comment period.

### *Medicaid*

The reimbursement for Medicaid is generally 10% to 20% lower than Medicare.

### *Commercial insurance carrier reimbursement*

Most private and commercial carriers reimbursed DR telehealth programs using CPT code 92250. Some used the level II HCPCS code, S0625 (Retinal Telescreening by Digital Imaging of Multiple Different Fundus Areas to Screen for Vision-Threatening Conditions). Some carriers reimburse for the service but require pupil dilation. Due to this variation among carriers, each must be contacted to determine the requirements for reimbursement. How commercial insurance carriers will treat new CPT codes 92227 and 92228 is currently unknown.

### *Other financial factors*

#### Logistic efficiencies

Geographic disparities in care can result in access to care issues that are costly in terms of time transportation, and missed opportunity. Telemedicine can close these distances electronically with a possible overall savings in costs.

#### Disease prevention

Increasing the surveillance rate of DR through telemedicine contributes to increased treatment and reduction in diabetes-related vision loss.<sup>26, 160</sup> This can result in significant healthcare savings through cost avoidance.<sup>45, 46</sup>

#### Resource utilization

Some DR telehealth programs have shown to be less costly and more effective than convention retinal examinations for the detection of DR.<sup>161</sup> This may allow a reduction in the overall cost of care with the same or expanded scope of services through the re-tasking of costly human resources.

## REFERENCES

1. Telemedicine Defined. Available at <http://www.americantelemed.org/i4a/pages/index.cfm?pageid=3333>.
2. Hogan P, Dall T, Nikolov P. Economic costs of diabetes in the US in 2002. *Diabetes Care* 2003;26:917-32.
3. Diabetes Public Health Resource. Available at <http://www.cdc.gov/diabetes/pubs/factsheets/atwork.htm>.
4. Harris M. Diabetes Mellitus: Summary. National Institutes of Health, 1995:1-36.
5. Centers for Disease Control and Prevention. National diabetes fact sheet: General information and national estimates on diabetes in the United States, 2007. Atlanta, GA: U.S. Department of Health and Human Services, Centers for Disease Control and Prevention.
6. Kempen JH, O'Colmain BJ, Leske MC, et al. The prevalence of diabetic retinopathy among adults in the United States. *Arch Ophthalmol* 2004;122:552-63.
7. Wild S, Roglic G, Green A, Sicree R, King H. Global prevalence of diabetes: estimates for the year 2000 and projections for 2030. *Diabetes Care* 2004;27:1047-53.
8. Saaddine JB, Honeycutt AA, Narayan KM, Zhang X, Klein R, Boyle JP. Projection of diabetic retinopathy and other major eye diseases among people with diabetes mellitus: United States, 2005-2050. *Arch Ophthalmol* 2008;126:1740-7.
9. IDF Diabetes Atlas. Global Burden: Prevalence and Projections, 2010 and 2030. <http://www.diabetesatlas.org/content/diabetes-and-impaired-glucose-tolerance>. 2010.
10. Klein R, Klein BE, Moss SE, Davis MD, DeMets DL. The Wisconsin epidemiologic study of diabetic retinopathy. II. Prevalence and risk of diabetic retinopathy when age at diagnosis is less than 30 years. *Arch Ophthalmol* 1984;102:520-6.
11. Klein R, Klein BE, Moss SE, Davis MD, DeMets DL. The Wisconsin epidemiologic study of diabetic retinopathy. III. Prevalence and risk of diabetic retinopathy when age at diagnosis is 30 or more years. *Arch Ophthalmol* 1984;102:527-32.
12. World Health Organization. Diabetes estimates and projections. WHO. 2008. Last copyright date: 2008. Available from: URL: <http://www.who.int/mediacentre/factsheets/fs312/en/index.html> (last accessed June 2, 2009).
13. Klein R. The epidemiology of diabetic retinopathy: findings from the Wisconsin Epidemiologic Study of Diabetic Retinopathy. *Int Ophthalmol Clin* 1987;27:230-238.
14. Klein R, Klein BE, Moss SE. The Wisconsin epidemiological study of diabetic retinopathy: a review. *Diabetes Metab Rev* 1989;5:559-570.
15. Kohner EM, Aldington SJ, Stratton IM, et al. United Kingdom Prospective Diabetes Study, 30: diabetic retinopathy at diagnosis of non-insulin-dependent diabetes mellitus and associated risk factors. *Arch Ophthalmol* 1998;116:297-303.
16. Aiello LP, Gardner TW, King GL, et al. Diabetic retinopathy. *Diabetes Care* 1998;21:143-156.
17. Ferris FL, 3rd, Patz A. Macular edema. A complication of diabetic retinopathy. *Surv Ophthalmol* 1984;28 Suppl:452-61.



18. Klein R, Klein BE, Moss SE, Cruickshanks KJ. The Wisconsin Epidemiologic Study of Diabetic Retinopathy. XV. The long-term incidence of macular edema. *Ophthalmology* 1995;102:7-16.
19. Klein R, Klein BE, Moss SE, Davis MD, DeMets DL. The Wisconsin epidemiologic study of diabetic retinopathy. IV. Diabetic macular edema. *Ophthalmology* 1984;91:1464-74.
20. Photocoagulation treatment of proliferative diabetic retinopathy. Clinical application of Diabetic Retinopathy Study (DRS) findings, DRS Report Number 8. The Diabetic Retinopathy Study Research Group. *Ophthalmology* 1981;88:583-600.
21. Photocoagulation treatment of proliferative diabetic retinopathy: relationship of adverse treatment effects to retinopathy severity. Diabetic retinopathy study report no. 5. *Dev Ophthalmol.* 1981;2:248-261.
22. Four risk factors for severe visual loss in diabetic retinopathy. The third report from the Diabetic Retinopathy Study. The Diabetic Retinopathy Study Research Group. *Arch Ophthalmol.* 1979;97:654-655.
23. Aspirin effects on mortality and morbidity in patients with diabetes mellitus. Early Treatment Diabetic Retinopathy Study report 14. *JAMA* 1992;268:1292-1300.
24. Fundus photographic risk factors for progression of diabetic retinopathy. ETDRS report number 12. Early Treatment Diabetic Retinopathy Study Research Group. *Ophthalmology* 1991;98:823-833.
25. Early photocoagulation for diabetic retinopathy. ETDRS report number 9. Early Treatment Diabetic Retinopathy Study Research Group. *Ophthalmology* 1991;98:766-785.
26. Photocoagulation for diabetic macular edema. Early Treatment Diabetic Retinopathy Study report number 1. Early Treatment Diabetic Retinopathy Study research group. *Arch Ophthalmol* 1985;103:1796-1806.
27. Focal photocoagulation treatment of diabetic macular edema. Relationship of treatment effect to fluorescein angiographic and other retinal characteristics at baseline: ETDRS report no. 19. Early Treatment Diabetic Retinopathy Study Research Group. *Arch Ophthalmol.* 1995;113:1144-1155.
28. Retinopathy and nephropathy in patients with type 1 diabetes four years after a trial of intensive therapy. The Diabetes Control and Complications Trial/Epidemiology of Diabetes Interventions and Complications Research Group. *N Engl J Med* 2000;342:381-389.
29. Epidemiology of Diabetes Interventions and Complications (EDIC). Design, implementation, and preliminary results of a long-term follow-up of the Diabetes Control and Complications Trial cohort. *Diabetes Care* 1999;22:99-111.
30. Early worsening of diabetic retinopathy in the Diabetes Control and Complications Trial. *Arch Ophthalmol.* 1998;116:874-886.
31. Lifetime benefits and costs of intensive therapy as practiced in the diabetes control and complications trial. The Diabetes Control and Complications Trial Research Group. *JAMA* 1996;276:1409-1415.
32. The absence of a glycemic threshold for the development of long-term complications: the perspective of the Diabetes Control and Complications Trial. *Diabetes* 1996;45:1289-1298.

33. The relationship of glycemic exposure (HbA1c) to the risk of development and progression of retinopathy in the diabetes control and complications trial. *Diabetes* 1995;44:968-983.
34. Progression of retinopathy with intensive versus conventional treatment in the Diabetes Control and Complications Trial. *Diabetes Control and Complications Trial Research Group. Ophthalmology* 1995;102:647-661.
35. Babaya N. Effect of intensive therapy on the microvascular complications of type 1 diabetes mellitus. *JAMA* 2002;287:2563-9.
36. White NH, Sun W, Cleary PA, et al. Prolonged effect of intensive therapy on the risk of retinopathy complications in patients with type 1 diabetes mellitus: 10 years after the Diabetes Control and Complications Trial. *Arch Ophthalmol* 2008;126:1707-15.
37. Intensive blood-glucose control with sulphonylureas or insulin compared with conventional treatment and risk of complications in patients with type 2 diabetes (UKPDS 33). UK Prospective Diabetes Study (UKPDS) Group. *Lancet* 1998;352:837-53.
38. Tight blood pressure control and risk of macrovascular and microvascular complications in type 2 diabetes: UKPDS 38. UK Prospective Diabetes Study Group. *Bmj* 1998;317:703-13.
39. Holman RR, Paul SK, Bethel MA, Matthews DR, Neil HA. 10-year follow-up of intensive glucose control in type 2 diabetes. *N Engl J Med* 2008;359:1577-89.
40. Diabetic Retinopathy Clinical Research Network. *DRCRnet Manual of Operations*. Version 3.0. Tampa, FL: Jaeb Center for Health Research Foundation; April 8, 2005.
41. Diabetic Retinopathy Clinical Research Network (DRCRnet) public web site. Available at [http://drcrnet.jaeb.org/ViewPage.aspx?PageName=Home\\_Page](http://drcrnet.jaeb.org/ViewPage.aspx?PageName=Home_Page).
42. National Committee for Quality Assurance's 2009 State of Health Care Quality Report. Available at <http://www.ncqa.org/tabid/836/Default.aspx>.
43. Witkin SR, Klein R. Ophthalmologic care for persons with diabetes. *JAMA* 1984;251:2534-2537.
44. Sprafka JM, Fritsche TL, Baker R, Kurth D, Whipple D. Prevalence of undiagnosed eye disease in high-risk diabetic individuals. *Arch Intern Med* 1990;150:857-861.
45. Javitt JC, Aiello LP, Chiang Y, Ferris FL, III, Canner JK, Greenfield S. Preventive eye care in people with diabetes is cost-saving to the federal government. Implications for health-care reform. *Diabetes Care* 1994;17:909-917.
46. Javitt JC, Aiello LP. Cost-effectiveness of detecting and treating diabetic retinopathy. *Ann Intern Med* 1996;124:164-169.
47. World Health Organization W. Telehealth and telemedicine will henceforth be part of the strategy for health for all, 2002.
48. Standards of medical care in diabetes--2009. *Diabetes Care* 2009;32 Suppl 1:S13-61.
49. Fransen SR, Leonard-Martin TC, Feuer WJ, Hildebrand PL. Clinical evaluation of patients with diabetic retinopathy: accuracy of the Inoveon diabetic retinopathy-3DT system. *Ophthalmology* 2002;109:595-601.
50. Bursell SE, Cavallerano JD, Cavallerano AA, et al. Stereo nonmydriatic digital-video color retinal imaging compared with Early Treatment Diabetic Retinopathy Study

- seven standard field 35-mm stereo color photos for determining level of diabetic retinopathy. *Ophthalmology* 2001;108:572-85.
51. Rudnisky CJ, Hinz BJ, Tennant MT, de Leon AR, Greve MD. High-resolution stereoscopic digital fundus photography versus contact lens biomicroscopy for the detection of clinically significant macular edema. *Ophthalmology* 2002;109:267-74.
  52. Cavallerano AA, Cavallerano JD, Katalinic P, Tolson AM, Aiello LP, Aiello LM. Use of Joslin Vision Network digital-video nonmydriatic retinal imaging to assess diabetic retinopathy in a clinical program. *Retina* 2003;23:215-23.
  53. Wilkinson CP, Ferris FL, 3rd, Klein RE, et al. Proposed international clinical diabetic retinopathy and diabetic macular edema disease severity scales. *Ophthalmology* 2003;110:1677-82.
  54. Scanlon PH, Foy C, Malhotra R, Aldington SJ. The influence of age, duration of diabetes, cataract, and pupil size on image quality in digital photographic retinal screening. *Diabetes Care* 2005;28:2448-53.
  55. Murgatroyd H, Ellingford A, Cox A, et al. Effect of mydriasis and different field strategies on digital image screening of diabetic eye disease. *Br J Ophthalmol* 2004;88:920-4.
  56. Pandit RJ, Taylor R. Mydriasis and glaucoma: exploding the myth. A systematic review. *Diabet Med* 2000;17:693-9.
  57. Whited JD. Accuracy and reliability of teleophthalmology for diagnosing diabetic retinopathy and macular edema: a review of the literature. *Diabetes Technol Ther* 2006;8:102-11.
  58. Bresnick GH, Mukamel DB, Dickinson JC, Cole DR. A screening approach to the surveillance of patients with diabetes for the presence of vision-threatening retinopathy. *Ophthalmology* 2000;107:19-24.
  59. Blum J. Internet medicine and the evolving legal status of the physician-patient relationship. *J Leg Med* 2003;4:413-55.
  60. Kasztelowicz P. Security of medical data transfer and storage in Internet Cryptography, antiviral security and electronic signature problems, which must be solved in nearest future in practical contest. *Pol J Pathol* 2003;3:209-14.
  61. Brebner E, Bebnar J, Ruddick-Bracken H, Wootton R, Ferguson J. The importance of setting and evaluating standards of telemedicine training. *J Telemed Telecare* 2003;9:7-9.
  62. Lee P. Telemedicine: opportunities and challenges for the remote care of diabetic retinopathy. *Arch Ophthalmol* 1999;117:1639-40.
  63. Zeimer R, Zou S, Meeder T, al. e. A fundus camera dedicated to the screening of diabetic retinopathy in the primary-care physician's office. *Invest Ophthalmol* 2002;43:1581-87.
  64. Digital Imaging and Communications in Medicine (DICOM) Supplement 91: Ophthalmic Photography Image SOP Classes. Available at: [ftp://medical.nema.org/medical/dicom/final/sup91\\_ft2.pdf](ftp://medical.nema.org/medical/dicom/final/sup91_ft2.pdf) (Last accessed on December 8, 2009).
  65. DICOM™ Digital IMaging and Communications in Medicine. Available at <http://medical.nema.org/>.
  66. Health Level Seven International (HL7) . Available at <http://www.hl7.org/>.

67. IHE Eye Care (EYECARE) Technical Framework Volume 1 (EYECARE TF-1) Integration Profiles. Revision 3.7 – Final Text February 15, 2010. Available at [http://www.ihe.net/Technical\\_Framework/upload/ihe\\_eyecare\\_tf\\_rev3-7\\_vol1\\_Final\\_Text\\_2010-02-15.pdf](http://www.ihe.net/Technical_Framework/upload/ihe_eyecare_tf_rev3-7_vol1_Final_Text_2010-02-15.pdf).
68. IHE Eye Care (EYECARE) Technical Framework Volume 2 (EYECARE TF-1) Transactions. Revision 3.7 – Final Text February 15, 2010. Available at [http://www.ihe.net/Technical\\_Framework/upload/ihe\\_eyecare\\_tf\\_rev3-7\\_vol2\\_Final\\_Text\\_2010-02-15.pdf](http://www.ihe.net/Technical_Framework/upload/ihe_eyecare_tf_rev3-7_vol2_Final_Text_2010-02-15.pdf).
69. SNOMED Terminology Solutions. Available at [http://www.cap.org/apps/docs/snomed/sts/snomed\\_ct.html](http://www.cap.org/apps/docs/snomed/sts/snomed_ct.html).
70. HITSP. Available at <http://www.hitsp.org/>.
71. Digital Imaging and Communications in Medicine (DICOM) Supplement 61: JPEG 2000 Transfer Syntaxes. Available at: [ftp://medical.nema.org/medical/dicom/final/sup61\\_ft.pdf](ftp://medical.nema.org/medical/dicom/final/sup61_ft.pdf) (Last accessed on December 8, 2009).
72. Kuzmak PM, Dayhoff RE. Minimizing Digital Imaging and Communications in Medicine (DICOM) Modality Worklist patient/study selection errors. J Digit Imaging 2001;14:153-7.
73. Csipo D, Dayhoff RE, Kuzmak PM. Integrating Digital Imaging and Communications in Medicine (DICOM)-structured reporting into the hospital environment. J Digit Imaging 2001;14:12-6.
74. American Academy of Ophthalmology Preferred Practice Pattern®. Diabetic Retinopathy. September 2008. <http://one.aao.org/CE/PracticeGuidelines/PPP.aspx?sid=ab789157-5312-4bbe-86ed-8d164ffa9567>.
75. American Academy of Ophthalmology Summary Benchmarks for Preferred Practice Pattern® Guidelines Retinopathy. November 2008. Available at <http://one.aao.org/CE/PracticeGuidelines/default.aspx?sid=783ead7d-3e06-433b-8f17-2971eb3816f8>.
76. Optometric Clinical Practice Guideline: Care of the Patient with Diabetes Mellitus. Available at <http://www.aoa.org/documents/CPG-3.pdf>.
77. Silverman R. Current legal and ethical concerns in telemedicine and e-medicine. J Telemed Telecare 2003;9:7-9.
78. Bilimoria N. Telemedicine: laws still need a dose of efficiency. J Med Pract Manage 2003;6:289-94.
79. U.S. Department of Health and Human Services. Health information privacy: summary of HIPAA Privacy Rule. Last updated: May 2003. Accessed at <http://www.hhs.gov/ocr/privacy/hipaa/understanding/summary/index.html>
80. National Center for Chronic Disease Prevention and Health Promotion. Diabetes Public Health Resource: Age-Adjusted Rates of Seeing a Health Professional for Diabetes in the Last Year per 100 Adults with Diabetes, United States, 2001. Center for Disease Control and Prevention. 2001. Accessed at <http://www.cdc.gov/diabetes/statistics/preventive/fig4g.htm>.
81. Eliasson A, Poropatich R. Performance improvement in telemedicine: the essential elements. Mil Med 1998;8:530-5.

82. Comprehensive Accreditation Manual for Hospitals: The Official Handbook. MS.4.120. MS.4.130.: The Joint Commission on the Accreditation of Healthcare Organizations, 2008.
83. AMA, Physician Licensure: An update of trends. <http://www.ama-assn.org/ama/pub/about-ama/our-people/member-groups-sections/young-physicians-section/advocacy-resources/physician-licensure-an-update-trends.shtml>.
84. OAT Telemedicine Licensure Report. <ftp://ftp.hrsa.gov/telehealth/licensure.pdf>
85. Rigelhaupt, JL., Annotation, What Constitutes Physician-Patient Relationship for Malpractice Purposes, 17 A.L.R. 4th 132 (1982).
86. E-Health and Antitrust. Telemedicine and E-Health Law, Rel 7. Ch7,1.04[3a]. Law Journal Press, 2008.
87. AMA, Policy Statement H-480.968, Telemedicine. [http://www.ama-assn.org/ama1/pub/upload/mm/16/a08\\_hod\\_resolutions.pdf](http://www.ama-assn.org/ama1/pub/upload/mm/16/a08_hod_resolutions.pdf)
88. VA. VHA Informed Consent for Clinical Treatments and Procedures: VA, 2003.
89. Stanberry B. Legal ethical and risk issues in telemedicine. Comput Methods Programs Biomed 2001;64:225-33.
90. Iserson K. Telemedicine: a proposal for an ethical code. Camb Q Healthc Ethics 2000;9:404/6.
91. Mossman K. Medical testing: issues and ethics. Forum Appl Res Public Policy 1997;3:90-101.
92. Borkowski W, Mielniczuk H. Telemedical management system of structured clinical documentation-application for quality assurance and multicenter clinical trials. Pol J Pathol 2003;3:193-5.
93. McCrossin R. Managing risk in telemedicine. J Telemed Telecare 2003;9:6-9.
94. Spencer T, Olson JA, McHardy KC, Sharp PF, Forrester JV. An image-processing strategy for the segmentation and quantification of microaneurysms in fluorescein angiograms of the ocular fundus. Comput Biomed Res 1996;29:284-302.
95. Fleming AD, Goatman KA, Philip S, Olson JA, Sharp PF. Automatic detection of retinal anatomy to assist diabetic retinopathy screening. Phys Med Biol 2007;52:331-45.
96. Usher D, Dumskyj M, Himaga M, Williamson TH, Nussey S, Boyce J. Automated detection of diabetic retinopathy in digital retinal images: a tool for diabetic retinopathy screening. Diabet Med 2004;21:84-90.
97. Staal J, Abramoff MD, Niemeijer M, Viergever MA, van Ginneken B. Ridge-based vessel segmentation in color images of the retina. IEEE Trans Med Imaging 2004;23:501-9.
98. Lowell J, Hunter A, Steel D, et al. Optic nerve head segmentation. IEEE Trans Med Imaging 2004;23:256-64.
99. Martinez-Perez ME, Hughes AD, Thom SA, Bharath AA, Parker KH. Segmentation of blood vessels from red-free and fluorescein retinal images. Med Image Anal 2007;11:47-61.
100. Solouma NH, Youssef AB, Badr YA, Kadah YM. A new real-time retinal tracking system for image-guided laser treatment. IEEE Trans Biomed Eng 2002;49:1059-67.
101. Smith RT, Chan JK, Nagasaki T, et al. Automated detection of macular drusen using geometric background leveling and threshold selection. Arch Ophthalmol 2005;123:200-6.

102. Gregson PH, Shen Z, Scott RC, Kozousek V. Automated grading of venous beading. *Comput Biomed Res* 1995;28:291-304.
103. Hejlesen O, Ege B, Englmeier KH, Aldington S, McCanna L, Bek T. TOSCA-Imaging--developing Internet based image processing software for screening and diagnosis of diabetic retinopathy. *Stud Health Technol Inform* 2004;107:222-6.
104. Kozousek V, Shen Z, Gregson P, Scott RC. Automated detection and quantification of venous beading using Fourier analysis. *Can J Ophthalmol* 1992;27:288-94.
105. Asrani S, Zeimer R, Goldberg MF, Zou S. Application of rapid scanning retinal thickness analysis in retinal diseases. *Ophthalmology* 1997;104:1145-51.
106. Sadda SR, Tan O, Walsh AC, Schuman JS, Varma R, Huang D. Automated detection of clinically significant macular edema by grid scanning optical coherence tomography. *Ophthalmology* 2006;113:1187 e1-12.
107. van Dijk HW, Kok PH, Garvin M, et al. Selective loss of inner retinal layer thickness in type 1 diabetic patients with minimal diabetic retinopathy. *Invest Ophthalmol Vis Sci* 2009;50:3404-9.
108. Cabrera Debuc D, Somfai GM. Early detection of retinal thickness changes in diabetes using optical coherence tomography. *Med Sci Monit* 2010; 16(3): MT 15-21.
109. Larsen M, Godt J, Larsen N, et al. Automated detection of fundus photographic red lesions in diabetic retinopathy. *Invest Ophthalmol Vis Sci* 2003;44:761-6.
110. Englmeier KH, Schmid K, Hildebrand C, et al. Early detection of diabetes retinopathy by new algorithms for automatic recognition of vascular changes. *Eur J Med Res* 2004;9:473-8.
111. Larsen N, Godt J, Grunkin M, Lund-Andersen H, Larsen M. Automated detection of diabetic retinopathy in a fundus photographic screening population. *Invest Ophthalmol Vis Sci* 2003;44:767-71.
112. Fleming AD, Philip S, Goatman KA, Williams GJ, Olson JA, Sharp PF. Automated detection of exudates for diabetic retinopathy screening. *Phys Med Biol* 2007;52:7385-96.
113. Junge J, Boon KL, Ottens DE. Automatic information processing in diabetic retinopathy. *Ophthalmologica* 1975;171:3-6.
114. Muller W, Spittel U, Gobel H. First results of the computer-aided classification of electroretinograms. *Ophthalmologica* 1986;192:228-33.
115. Guo W, Marshall G. ORDMKV: a computer program fitting proportional odds model for multi-state Markov process. *Comput Methods Programs Biomed* 1995;46:257-63.
116. Marshall G, Guo W, Jones RH. MARKOV: a computer program for multi-state Markov models with covariables. *Comput Methods Programs Biomed* 1995;47:147-56.
117. Zana F, Klein JC. A multimodal registration algorithm of eye fundus images using vessels detection and Hough transform. *IEEE Trans Med Imaging* 1999;18:419-28.
118. Zahlmann G, Kochner B, Ugi I, et al. Hybrid fuzzy image processing for situation assessment. *IEEE Eng Med Biol Mag* 2000;19:76-83.
119. Niemeijer M, van Ginneken B, Staal J, Suttorp-Schulten MS, Abramoff MD. Automatic detection of red lesions in digital color fundus photographs. *IEEE Trans Med Imaging* 2005;24:584-92.

120. Chanwimaluang T, Fan G, Fransen SR. Hybrid retinal image registration. *IEEE Trans Inf Technol Biomed* 2006;10:129-42.
121. Walter T, Klein JC, Massin P, Erginay A. A contribution of image processing to the diagnosis of diabetic retinopathy--detection of exudates in color fundus images of the human retina. *IEEE Trans Med Imaging* 2002;21:1236-43.
122. Sanchez CI, Garcia M, Mayo A, Lopez MI, Hornero R. Retinal image analysis based on mixture models to detect hard exudates. *Med Image Anal* 2009;13:650-8.
123. Goldberg RE, Varma R, Spaeth GL, Magargal LE, Callen D. Quantification of progressive diabetic macular nonperfusion. *Ophthalmic Surg* 1989;20:42-5.
124. Abramoff MD, Niemeijer M, Suttorp-Schulten MS, Viergever MA, Russell SR, van Ginneken B. Evaluation of a system for automatic detection of diabetic retinopathy from color fundus photographs in a large population of patients with diabetes. *Diabetes Care* 2008;31:193-8.
125. Bouhaimed M, Gibbins R, Owens D. Automated detection of diabetic retinopathy: results of a screening study. *Diabetes Technol Ther* 2008;10:142-8.
126. Sinthanayothin C, Boyce JF, Williamson TH, et al. Automated detection of diabetic retinopathy on digital fundus images. *Diabet Med* 2002;19:105-12.
127. Gonzalez Villalpando C, Gonzalez Villalpando ME, Martinez Diaz S, et al. A diabetic retinopathy screening program as a strategy for blindness prevention. *Arch Med Res* 1997;28:129-35.
128. Goatman KA, Cree MJ, Olson JA, Forrester JV, Sharp PF. Automated measurement of microaneurysm turnover. *Invest Ophthalmol Vis Sci* 2003;44:5335-41.
129. Hipwell JH, Strachan F, Olson JA, McHardy KC, Sharp PF, Forrester JV. Automated detection of microaneurysms in digital red-free photographs: a diabetic retinopathy screening tool. *Diabet Med* 2000;17:588-94.
130. Cree MJ, Olson JA, McHardy KC, Sharp PF, Forrester JV. A fully automated comparative microaneurysm digital detection system. *Eye (Lond)* 1997;11 ( Pt 5):622-8.
131. Baudoin CE, Lay BJ, Klein JC. Automatic detection of microaneurysms in diabetic fluorescein angiography. *Rev Epidemiol Sante Publique* 1984;32:254-61.
132. Fleming AD, Philip S, Goatman KA, Olson JA, Sharp PF. Automated microaneurysm detection using local contrast normalization and local vessel detection. *IEEE Trans Med Imaging* 2006;25:1223-32.
133. Spencer T, Phillips RP, Sharp PF, Forrester JV. Automated detection and quantification of microaneurysms in fluorescein angiograms. *Graefes Arch Clin Exp Ophthalmol* 1992;230:36-41.
134. Sleightholm MA, Arnold J, Aldington SJ, Kohner EM. Computer-aided digitisation of fundus photographs. *Clin Phys Physiol Meas* 1984;5:295-301.
135. Frame AJ, Undrill PE, Cree MJ, et al. A comparison of computer based classification methods applied to the detection of microaneurysms in ophthalmic fluorescein angiograms. *Comput Biol Med* 1998;28:225-38.
136. Sivakumar R, Ravindran G, Muthayya M, Lakshminarayanan S, Velmurughendran CU. Diabetic Retinopathy Analysis. *J Biomed Biotechnol* 2005;2005:20-27.
137. Nayak J, Bhat PS, Acharya R, Lim CM, Kagathi M. Automated identification of diabetic retinopathy stages using digital fundus images. *J Med Syst* 2008;32:107-15.



138. Niemeijer M, van Ginneken B, Cree MJ, et al. Retinopathy online challenge: automatic detection of microaneurysms in digital color fundus photographs. *IEEE Trans Med Imaging*;29:185-95.
139. Abramoff MD, Reinhardt JM, Russell SR, et al. Automated Early Detection of Diabetic Retinopathy. *Ophthalmology*.
140. Niemeijer M, van Ginneken B, Russell SR, Suttorp-Schulten MS, Abramoff MD. Automated detection and differentiation of drusen, exudates, and cotton-wool spots in digital color fundus photographs for diabetic retinopathy diagnosis. *Invest Ophthalmol Vis Sci* 2007;48:2260-7.
141. Philip S, Fleming AD, Goatman KA, et al. The efficacy of automated "disease/no disease" grading for diabetic retinopathy in a systematic screening programme. *Br J Ophthalmol* 2007;91:1512-7.
142. Abramoff MD, Niemeijer M, Russell SR. Automated detection of diabetic retinopathy: barriers to translation into clinical practice. *Expert Rev Med Devices*;7:287-96.
143. Hill, N.R. and D.R. Matthews, Automated Retinal Image Differencing (ARID) - Abstracts of the 40th Annual Meeting of the European Association for the Study of Diabetes. Munich, Germany, 5-9 September 2004. *Diabetologia*, 2004. 47 Suppl 1: p. A1-489.
144. Tobin KW, Abdelrahman M, Chaum E, Govindasamy V, Karnowski TP. A probabilistic framework for content-based diagnosis of retinal disease. *Conf Proc IEEE Eng Med Biol Soc* 2007;2007:6744-7.
145. Chaum E, Karnowski TP, Govindasamy VP, Abdelrahman M, Tobin KW. Automated diagnosis of retinopathy by content-based image retrieval. *Retina* 2008;28:1463-77.
146. LeBlanc K, Saffiotti A. Multirobot object localization: a fuzzy fusion approach. *IEEE Trans Syst Man Cybern B Cybern* 2009;39:1259-76.
147. Nguyen TT, Wong TY. Retinal vascular changes and diabetic retinopathy. *Curr Diab Rep* 2009;9:277-83.
148. Bayu Sasongko M, Wang JJ, Donaghue KC, et al. Alterations in Retinal Microvascular Geometry in Young Type 1 Diabetes. *Diabetes Care*.
149. Grauslund J, Green A, Kawasaki R, Hodgson L, Sjolie AK, Wong TY. Retinal Vascular Fractals and Microvascular and Macrovascular Complications in Type 1 Diabetes. *Ophthalmology*.
150. Covered Entity Charts. Centers for Medicare and Medicaid Services. Accessed at [http://www.cms.hhs.gov/HIPAAGenInfo/06\\_AreYouaCoveredEntity.asp](http://www.cms.hhs.gov/HIPAAGenInfo/06_AreYouaCoveredEntity.asp).
151. <http://edocket.access.gpo.gov/2010/pdf/2010-12647.pdf>.
152. Blignault I, Kennedy C. Training for telemedicine. *J Telemed Telecare* 1999;5 Suppl 1:S112-4.
153. Comprehensive Accreditation Manual for Hospitals: The Official Handbook. EC6.10. MS.5.10: The Joint Commission on the Accreditation of Healthcare Organizations, 2004.
154. Scanlon PH. The English national screening programme for sight-threatening diabetic retinopathy. *J Med Screen* 2008;15:1-4.
155. Garvican L, Clowes J, Gillow T. Preservation of sight in diabetes: developing a national risk reduction programme. *Diabet Med* 2000;17:627-34.



156. Arun CS, Young D, Batey D, et al. Establishing ongoing quality assurance in a retinal screening programme. *Diabet Med* 2006;23:629-34.
157. Schneider S, Aldington SJ, Kohner EM, et al. Quality assurance for diabetic retinopathy telescreening. *Diabet Med* 2005;22:794-802.
158. Centers for Medicare and Medicaid Services, Department of Health and Human Services. Medicare Program; Payment Policies Under the Physician Fee Schedule and Other Revisions to Part B for CY 2011. <http://www.regulations.gov/#!documentDetail:D=CMS-2010-0205-1982>. Accessed on 2-6-2011.
159. Ocular Special Interest Group, American Telemedicine Association. American Telemedicine Association comments on Diabetic Retinopathy Codes. <http://hub.americantelemed.org/AMERICANTELEMED/AMERICANTELEMED/Resources/ViewDocument/Default.aspx?DocumentKey=8fd1a5c2-9158-46a5-a0a4-e2843fd0cd96&MessageKey=dae865a4-f48c-4cee-87ce-c815c7e06bc1>. Accessed on 2-6-2011.
160. Early photocoagulation for diabetic retinopathy. ETDRS report number 9. Early Treatment Diabetic Retinopathy Study Research Group. *Ophthalmology* 1991;98:766-85.
161. Whited JD, Datta SK, Aiello LM, et al. A modeled economic analysis of a digital tele-ophthalmology system as used by three federal health care agencies for detecting proliferative diabetic retinopathy. *Telemed J E Health* 2005;11:641-51.

## CONTRIBUTORS

### *Second Edition*

Editorial Committee: Helen K. Li, MD (Chair), Mark Horton, OD, MD (Co-chair), Sven-Erik Bursell, PhD, Jerry Cavallerano, OD, PhD, Ingrid Zimmer-Galler, MD, Mathew Tennant, MD.

### Writing Committees

Clinical: Jerry Cavallerano, OD, PhD (Chair), Ingrid Zimmer-Galler, MD

Technology: Sven-Erik Bursell, PhD (Chair), Michael Abramoff, MD, PhD, Edward Chaum, MD, PhD, Debra Cabrera DeBuc, PhD.

Operations: Mark Horton, OD, MD (Chair), Helen K. Li, MD, Tom Leonard-Martin, PhD, MPH, Mathew Tennant, MD, Marc Winchester, BA.

### Other contributors

Reviewers [R], ATA Standard and Guidelines Committee Members [SG], ATA Staff [S]  
 Nina Antoniotti, RN, MBA, PhD [Chair, SG]  
 Jordana Bernard, MBA [S]  
 David Brennan, MSBE [SG]  
 Anne Burdick, MD, MPH [SG]  
 Jerry Cavallerano, OD, PhD [SG]

Brian Grady, MD [SG]  
Tom Hirota, DO [SG]  
Elizabeth Krupinski, PhD [Vice Chair, SG]  
Cindy K. Leenknecht, MS, APRN-CS, CCRP [SG]  
Jonathan Linkous, MPA [S]  
Lou Theurer [SG]  
Jill Winters, PhD, RN [SG]

*First Edition*

American Telemedicine Association, Ocular Telehealth Special Interest Group, and the National Institute of Standards and Technology Working Group

American Telemedicine Association Executive Committee:

Jonathan D. Linkous, Richard Bakalar, MD, Adam Darkins, MD, Col. Ronald K. Poropatich MD

American Telemedicine Association Ocular Telehealth Special Interest Group:

Jerry Cavallerano, OD, PhD (Chair), Mary G. Lawrence, MD, MPH (Vice Chair)

Editorial Committee:

Helen K. Li, MD (Co-chair), Mathew Tennant, MD (Co-chair), Sven Bursell, PhD, Jerry Cavallerano, OD, PhD, Mark Horton, OD, MD, Richard Bakalar, MD

Writing Committees:

Clinical: Jerry Cavallerano, OD, PhD (Chair), Mary G. Lawrence, MD, MPH, Ingrid Zimmer-Galler, MD, COL Wendall Bauman, MD

Technology: Sven Bursell, PhD (Chair), W. Kelly Gardner

Operations: Mark Horton, OD, MD (Chair), Lloyd Hildebrand, MD, Jay Federman, MD

National Institute of Standards and Technology:

Lisa Carnahan

Veterans Administration:

Peter Kuzmak, John M. Peters, Adam Darkins, MD

At Large Group Participants:

Jehanara Ahmed, MD, Lloyd M. Aiello, MD, Lloyd P. Aiello, MD, PhD, Gary Buck, Ying Ling Chen, PhD, Denise Cunningham, CRA, RBP, MEd, Eric Goodall, Ned Hope, Eugene Huang, PhD, Larry Hubbard, MAT, Mark Janczewski, MD, J.W.L. Lewis, PhD, Hiro Matsuzaki, COL Francis L. McVeigh, OD, Jordana Bernard, Diane Parker-Taillon, Robert Read, Peter Soliz, PhD, Bernard Szirth, PhD, COL Robert A. Vigersky, MD, COL Thomas Ward, MD

American Telemedicine Association Administrative Contributor:

Catherine Diver