

2018 MARCH[®] Vision Care Kansas Provider Reference Guide



keeping an eye on your health[®]

Provider Reference Guide | **Disclaimer**

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1.1 About the Provider Reference Guide

MARCH[®] is committed to working with our contracted providers and their staff to achieve the best possible health outcomes for our members. This guide provides helpful information about MARCH[®] eligibility, benefits, claim submission, claim payments, and much more. For easy navigation through this guide, click on the Table of Contents to be taken to the section of your choice.

This version of the Provider Reference Guide was revised on September 26, 2018. Reviews and updates to this guide are conducted as necessary and appropriate. Update notifications are distributed as they occur through provider newsletters. A current version of this guide is always available on our website at www.marchvisioncare.com. To request a current copy of the Provider Reference Guide on CD, please contact our Provider Relations Department at the appropriate state-specific phone number (below).

MARCH[®] would like to thank our providers for their participation in the delivery of quality vision care services to our members.

1.2 Contact Information

Phone Number	(844) 50-MARCH or (844) 506-2724
Fax Number	(877) MARCH-88 or (877) 627-2488
General Website	www.marchvisioncare.com
Provider Website	providers.eyesynergy.com
Mailing Address	MARCH [®] Vision Care 6701 Center Drive West, Suite 790 Los Angeles, CA 90045
Lab and Contact Lens Orders	providers.eyesynergy.com

1.3 eyeSynergy[®]

MARCH[®] is proud to offer eyeSynergy[®], our web-based solution for electronic transactions. With eyeSynergy[®], providers can:

- Verify member eligibility and benefits.
- Generate confirmation numbers for services (for the definition of “confirmation number”, refer to section 2.1).
- Submit claims.
- Obtain detailed claim status including check number and paid date.
- Access online resources such as a current copy of the Provider Reference Guide.
- Submit lab orders for eyeglasses and contact lenses.

eyeSynergy[®] is provided free of charge to all MARCH[®] participating providers. To access eyeSynergy[®], log onto our website at www.marchvisioncare.com and click on the orange and blue eyeSynergy[®] link located at the top of the page.

Registration

First time users must register before accessing eyeSynergy[®]. Please be prepared to enter the provider’s name or group name, office phone number, and tax identification number during registration. Once verified, you will immediately be provided with a temporary password to log in.

Logging In

Once registered, providers may log into eyeSynergy[®] with their user name and password. Please note that passwords are case-sensitive. As a security feature, the provider will be asked to renew their password every 60 days. After 5 failed log-in attempts, the provider is required to call MARCH[®] Vision Care to reset their password.

Once logged in, you may access the eyeSynergy[®] User Guide located on the Resources menu. This guide includes step-by-step instructions for completing various transactions within eyeSynergy[®].

1.4 Interactive Voice Recognition (IVR) System

Our Interactive Voice Recognition (IVR) System provides responses to the following inquiries twenty-four (24) hours per day, seven (7) days per week:

- Eligibility and benefits.
- Confirmation numbers.
- Claim status.

The IVR System may be accessed by calling (844) 506-2724. Select the provider option and follow the prompts to verify eligibility and benefits, request a confirmation number, or check claim status.

Registration

First-time users must register before accessing the IVR System. Please be prepared to enter your office phone number, office fax number and tax identification number during registration. Once verified, you will be prompted to select a 4-digit PIN for your account.

Logging In

Once registered, you may log into the IVR System using your 10-digit ID and 4-digit PIN. The 10-digit ID is the office phone number provided during registration. The 4-digit PIN is the number designated by your office during registration.

1.5 Electronic Funds Transfer (EFT)

MARCH[®] Vision Care is pleased to offer electronic funds transfer (EFT) and electronic remittance advices (ERAs) as the preferred methods of payments and explanations. EFT is the electronic transfer, or direct deposit, of money from MARCH[®] Vision Care directly into your bank account. ERAs are electronic explanations of payment (EOPs). MARCH partners with PaySpan Health, Inc.[®] (PaySpan) – a solution that delivers EFTs, ERAs/Vouchers, and much more.

The service is free to MARCH[®] providers. The solution delivers ERAs via their website allowing straightforward reconciliation of payments to empower our providers to reduce costs, speed secondary billings, improve cash flow, and help the environment by reducing paper usage.

MARCH[®] offers you the option to receive payments according to preference: electronically deposited into a bank account, or by traditional paper check.

Provider Benefits

As a provider, you gain immediate benefits by signing up for electronic payments from MARCH[®] Vision Care through PaySpan Health:

- Improve cash flow – Electronic payments can mean faster payments, leading to improvements in cash flow.
- Maintain control over bank accounts – You keep TOTAL control over the destination of claim payment funds. Multiple practices and accounts are supported.
- Match payments to advice/vouchers – You can associate electronic payments quickly and easily to an advice/voucher.
- Manage multiple payers – Reuse enrollment information to connect with multiple payers. Assign different payers to different banks.

Signing up for electronic payments is simple, secure, and will only take 5-10 minutes to complete. To complete the registration process, please visit the PaySpan website at www.payspanhealth.com or contact them directly at (877) 331-7154.

1.6 Provider Change Notification

Please help us to ensure your current information is accurately displayed in our provider directory. When possible, please report changes concerning your provider information to us in advance. All changes should be reported to MARCH[®] in writing. Failure to report changes related to your billing address and/or tax identification number may delay claim payments. Examples of changes that need to be reported to MARCH[®] in writing, include, but are not limited to:

- Practice phone and fax number.
- Practice address.
- Billing address (requires W9).
- Tax identification number (requires W9).
- Office hours.
- Practice status regarding the acceptance of new members, children, etc.
- Providers added to practice/providers leaving practice.
- Provider termination.

Please report all changes via mail or fax to:

MARCH[®] Vision Care
Attention: Provider Relations Department
6701 Center Drive West, Suite 790
Los Angeles, CA 90045
Fax: (877) MARCH-88 or (877) 627-2488

Email: providerdemographics@marchvisioncare.com

The Centers for Medicare & Medicaid Services (CMS) requires all providers to verify the accuracy of their information included in the health plan's provider directory on a quarterly basis. MARCH[®] encourages our providers to verify their demographic information through our provider web portal, ^{eye}Synergy[®]. When logging into your ^{eye}Synergy[®] account, you will see a banner on the top of your screen regarding your demographic information. You will click on that banner to be redirected to the demographic verification page where you can quickly verify your information and submit the form electronically. The online verification option is only available to registered and active ^{eye}Synergy[®] users.

2.1 Eligibility and Benefit Verification



MARCH[®] strongly recommends verification of member eligibility and benefits prior to rendering services. Please do not assume the member is eligible if they present a current ID card. Eligibility and benefits should be verified on the date services are rendered.

Confirmation Numbers

A confirmation number is an 11-digit identification number generated when the provider office verifies member eligibility for requested benefits and services through MARCH[®]. Verification is obtained by speaking with a Call Center Representative, or by accessing the IVR or ^{eye}Synergy[®] web portal. Confirmation numbers affirm member eligibility for requested benefits and services. However, confirmation numbers are not required for all services. Providers are strongly encouraged to verify benefits and eligibility on the date services will be rendered.

Benefits that generally require confirmation numbers include, but are not limited to:

- Replacement frames and lenses.
- Medically necessary contact lenses for Medicaid members.
- Two pairs of glasses in lieu of bifocals.
- Prescription sunglasses.

The confirmation request process requires the provider to attest that a member meets the defined benefit criteria, as outlined in the state specific Provider Reference Guide, when applicable. Upon attestation, a confirmation number is generated.

Example: A member is diagnosed with keratoconus and requires contact lenses. The provider is required to request a confirmation and attest to the documented exam findings and/or diagnosis. The submitted claim must include the diagnosis of keratoconus. Provided the member is eligible on the date services were rendered, payment is issued.

The following are examples of instances in which a confirmation number does not guarantee payment of a claim:

- The member is not eligible on the date of service.
- The member's benefit exhausted prior to claim submission.

IMPORTANT: MARCH[®] performs retrospective random chart audits on claims submitted for services requiring attestation.

Covered Benefits

A listing of covered benefits may be accessed by:

- Logging into ^{eye}Synergy[®] at <https://providers.eyesynergy.com>. Click on the Resources menu and select Provider Reference Guide. Benefits may be accessed by selecting the desired state from the drop-down menu. Providers may also access current benefits by plan or patient including the patient's current benefit availability from the Benefits and Eligibility menu in ^{eye}Synergy[®];
- Accessing our website at www.marchvisioncare.com. Click on Doctors and Office Staff and then Provider Resources. Benefits may be accessed by selecting the desired state from the drop-down menu.

Covered benefits include details such as benefit frequency, copayment amount, allowance amount, benefit limitations and benefit criteria.

Methods of Verification

You may access ^{eye}Synergy[®] or the Interactive Voice Recognition System to verify member eligibility, benefits, and to request a confirmation number.

2.2 Non-Covered Services

The Centers for Medicare and Medicaid Services (CMS) prohibits providers from billing or seeking compensation from Medicaid beneficiaries for the provision of services that are covered benefits under their Medicaid plans. However, there are certain circumstances in which a member requests services that are not covered or fully covered under their Medicaid plans.

In these circumstances, the provider must inform the member PRIOR to rendering the non-covered service that the service is not covered and that member will be financially responsible. **Failure to do so may result in the provider being financially responsible for those services even if the member verbally agreed to those services or paid for them up-front.**

Acceptable Waivers

A general waiver stating “the member is responsible for all services not covered by insurance” is not a valid waiver, as it does not specifically define which services are not covered and the amount the member is expected to pay.

The provider is required to have the member sign a waiver form that clearly explains that the specific service/procedure is not covered and that the member acknowledges that he/she will be responsible for the cost of the service(s).

MARCH[®] recommends using the MARCH[®] Non-Covered Service Fee Acceptance Form (available in both Spanish and English) in Exhibit A, but it is not required. If the provider chooses to use another form in place of the MARCH[®] Non-Covered Service Fee Acceptance Form, it must contain the following elements:

- Documentation of the specific services provided (including dates of service, description of procedure/service, amount charged).
- The member’s signed acknowledgement that he/she understands the service is not covered and he/she is financially liable for the services provided.

Once the waiver is signed, the member must receive a copy of the signed waiver. A copy of the signed waiver must also be placed in the member’s medical chart.

3.1 Claim Submission

Preferred Method

MARCH[®] prefers to receive claims electronically via ^{eye}Synergy[®], our web-based solution for electronic transactions. ^{eye}Synergy[®] helps reduce claim errors resulting in faster processing times.

Clearinghouse Submissions

MARCH[®] has a direct agreement with Optum to accept electronic claims. Our payor ID for Optum is 52461.

Paper Claims

Paper claims should be submitted on a red CMS-1500 form and mailed to:

MARCH[®] Vision Care
6701 Center Drive West, Suite 790
Los Angeles, CA 90045

Handwritten and/or faxed claims may delay claim payment.

Clean Claim Definition

MARCH[®] defines a clean claim as a bill from a health care provider that can be processed without obtaining additional information from the provider of service or from a third party. An unclean claim is defined as any claim that does not meet the definition of a clean claim. State specific exceptions to the MARCH[®] clean claim definition are provided below.

Claims submitted for payment should include the following:

- Member name, ID number, date of birth and gender.
- Provider and/or facility name, address and signature.
- Billing name, address and tax identification number.
- The rendering and billing National Provider Identifier (NPI).
- Date of service.
- Current and appropriate ICD-10 codes.
- Service units.
- Current and appropriate CPT/HCPCS codes.
- Current and applicable modifier codes.
- Place of service.
- Usual and customary charges.

MARCH[®] has the right to obtain further information from a provider's office upon request when a submitted claim has errors or when MARCH[®] or the health plan has reasonable grounds for suspecting possible fraud, misrepresentation or unfair billing practices.

Unclean claims are processed in accordance with applicable laws and regulations.

IMPORTANT: Please submit corrected claims on a red CMS-1500 form and clearly indicate on the claim that the submission is a **corrected claim**. This ensures the corrected information will be considered during claims processing and will help prevent payment delays.

3.2 American Medical Association CPT Coding Rules

MARCH[®] reaffirms its adoption of CPT coding rules established by the American Medical Association, Medicaid, and Medicare Regulations, and applicable law:

- For an initial examination of a new patient, providers can use a new eye examination billing code. A provider may also bill for a new member examination if a member has not been examined for 3 consecutive years by that provider/group.
- A routine examination for an established patient in subsequent years can be billed as a follow up examination.
- Providers can continue to bill this way unless the member has not been examined for 3 consecutive years at that office, at which time the service may be billed with a new member examination code as indicated above.

- A medical examination may be billed if the member has the benefit as indicated in MARCH®'s State Specific Provider Reference Guide.
- Follow up examinations for the same medical condition noted above may be billed based on the acuteness of the condition and the documented services provided.
- According to **Medicare Carriers Manual Section 15501.1 H**, if more than one evaluation and management (face-to-face) service is provided on the same day to the same patient, whether by the same provider or more than one provider in the same specialty in the same group, only one evaluation and management service may be billed. Therefore, a comprehensive eye examination and a medical examination, such as a diabetic eye evaluation, may not be billed on the same date of service. Instead of billing two examinations separately, providers should select a level of service representative of the combined visits and submit the appropriate code for that level. The less extensive procedure is bundled into the more extensive procedure.
- The services furnished and associated medical record documentation must meet the definition of the CPT code billed. This is especially important when providers bill the highest levels of visit and consultation codes. For example, to bill a comprehensive eye exam - new patient, the patient may not have been examined by a provider in the practice within the past three years, the history must meet the CPT code's definition of a comprehensive history, and all components of an examination need to be recorded, including dilation or equivalent. The provider may use professional discretion whether to dilate at subsequent visits for existing patients, but dilation is expected at the initial visit and at least every 3 years.
- Medical necessity of a service is the overarching criterion for payment, in addition to the individual requirements of a CPT code. It would not be medically necessary or appropriate to bill a higher level of evaluation and management service when a lower level of service is warranted or performed. In a similar vein, it would not be warranted to bill for services if medical necessity is not established by standards of medical or optometric practice.
- The date of service on the claim should always match the date of service on the medical record and the medical record should include complete documentation related to all billed services.
- The comprehensive nature of the examination codes includes a number of tests and evaluations. Some of these procedures have their own CPT code. When these procedures are broken out and billed in conjunction with a comprehensive examination is referred to as "unbundling" and is an inappropriate billing practice. This type of billing practice will be subject to action from a health plan or insurance carrier.

The most common errors include:

- Billing for a dilated fundus examination with the indirect ophthalmoscope and using the codes 92225, 92226, or separately billing visual fields using 92081.
- Billing color vision testing using 92283.
- Billing sensory motor testing using 92060
- Billing gonioscopy using 92020.

The appropriate and correct use of the CPT (procedure) and diagnosis code is the responsibility of every health care provider.

In all instances, the medical record should reflect the intensity of examination that is being billed. MARCH® will audit claim submissions to ensure compliance. Audits will include the review of medical records. These are some of the criteria that are used when MARCH® performs retrospective random chart audits based on claims submitted. Claims submitted that deviate from this standard may trigger a medical record audit. Audits that reveal chronic billing problems, or trends, or quality of care issues will require a Corrective Action Plan ("CAP"). Failure to execute the CAP may lead to termination as a MARCH® provider.

In an effort to improve HEDIS and Star Ratings performance, MARCH® Vision Care requires providers to submit CPT II and ICD-10 codes, on claims, to demonstrate performance and diagnosis of the following for diabetic members:

- Retinal of dilated eye exams;
- Negative retinal or dilated eye exams;
- Diabetes;
- Diabetic retinopathy

Please see Exhibit F: Performance Measurement & Reporting for more information.

3.3 Billing for Replacements and Repairs

Replacements and repairs are generally only covered under certain circumstances. For this reason, confirmation numbers are required for replacements and repairs. Replacement and repair services must be billed with the applicable modifier. The valid modifiers are provided below:

RA (Replacements)

RB (Repairs)

Reimbursement for materials billed with the RB (Repairs) modifier will be reimbursed at 50% of the contracted rate.

3.4 Billing for Glaucoma Screenings

The screening examination for glaucoma must include the following two (2) components:

1. Dilated exam with intraocular pressure (IOP) measurement;
2. Either direct ophthalmoscopy or slit lamp biomicroscopy.

CMS mandates payment for a glaucoma screening examination that is performed on an eligible beneficiary after at least eleven (11) months have passed following the month in which the last glaucoma screening examination was performed.

3.5 Frame Warranty

Frames from the MARCH[®] frame kit are fully guaranteed against manufacturing defects for a period of one (1) year from the date the frame was dispensed.

If the provider determines that the defective frame is covered under the warranty, please contact MARCH[®] at (844) 506-2724. Please do not send broken glasses to MARCH[®] or the contracted lab.

3.6 Order Cancellations

Orders placed with the MARCH[®] contracted lab for frames and lenses are final.

- Providers are responsible for the cost of frames and/or lenses if the order is incorrect due to provider error.
- In the event of an error, do not resubmit a corrected order. Please contact MARCH[®] at (844) 506-2724.

3.7 Claim Filing Limits

MARCH[®] imposes claim filing limits in accordance with the applicable provider services agreement and governing entity regulations. Claim filing limits for Medicaid are 180 days and begin on the date services are rendered.

Proof of Timely Filing

In cases where there is documentation proving “good cause” for a filing delay and a claim has not been submitted to MARCH[®] or a claim has been denied by MARCH[®] for exceeding the filing limit, MARCH[®] will consider issuing payment following a review of the “good cause” documentation.

The following are examples of acceptable forms of documentation to show “good cause” for delayed filing:

- Explanation of payment/denial from the primary payor dated within the timely filing period.
- Explanation of payment/denial from the believed payor dated within the timely filing period.

IMPORTANT: Please attach delayed filing “good cause” documentation to late filed claims. Submit late filed claims on a red CMS-1500 form and clearly indicate on the claim that the submission is a **late file claim with good cause documentation attached**. This ensures the information will be considered during claims processing and will help prevent payment delays.

3.8 Prompt Claim Processing

Claim payments are issued in accordance with the applicable provider services agreement and governing entity regulations. Prompt payment processing time for Medicaid is 30 calendar days for paper and electronic data interchange (EDI) claims. The processing time limit generally begins on the date the claim is received by MARCH[®].

3.9 Corrected Claims

A corrected claim may be submitted through the ^{eye}Synergy[®] web portal, under the Claims Details page. Providers will only have the option to submit a corrected claim after the claim has been paid. When using the “correct claim” function in ^{eye}Synergy[®], providers are to indicate the reason for the correction in the note section field. If attachments are required to

process the claim, please do not submit the corrected claim through ^{eye}Synergy[®]. Instead, please submit your corrected claim on a red CMS-1500 form along with the proof of timely filing or coordination of benefits attachment(s).

All other corrected claims, not submitted via ^{eye}Synergy[®] during the initial claim submission, must also be submitted on a red CMS-1500 form. Clearly indicate on the claim that the submission is a **“corrected claim.”** This ensures the corrected information will be considered during claims processing and will help prevent payment delays.

Please mail corrected claims to:

MARCH[®] Vision Care
6701 Center Drive West, Suite 790
Los Angeles, CA 90045

3.10 Claim Reconsideration, Appeal, State Fair Hearing and Provider Grievance Process

MARCH[®] is committed to ensuring provider satisfaction. Our Customer Service department can be reached at (844) 506-2724. In addition to contacting our customer service department, MARCH[®] provides a mechanism for you to communicate disputes in writing.

Claim Reconsideration

- Reconsideration is defined as a request by a provider for an MCO to review a claim decision.
- Reconsideration is an optional process available to providers prior to submitting an appeal.
- Requests must be submitted within one hundred twenty (120) calendar days from the remittance date, plus three (3) calendar days from the date the notice is sent.
- Reconsideration requests can be submitted through various means:
 - By phone: (844) 506-2724
 - Email: marchproviderappeal@marchvisioncare.com
 - By mail: MARCH[®] Vision Care
Attention: PDR Unit
6701 Center Drive West, Suite 790
Los Angeles, CA 90045
- Providers may terminate the reconsideration process and submit a formal appeal request within sixty (60) calendar days of the original remittance notice of action, plus three (3) calendar days from the date the notice is sent.
- If you disagree with a claim reconsideration decision, you have the right to file a formal claim appeal within 60 calendar days of the reconsideration notice of action.
- Providers have the right to represent him/herself or be represented by legal counsel or another spokesperson when requesting reconsideration or an appeal.

Appeal

- Appeal must be filed in writing within sixty (60) calendar days of the date of the provider remittance or notice of action, plus three (3) calendar days from the date the notice is sent.
- Request must state in the document this is a “formal appeal”.
- State the specific reason for denial as stated on the remittance or notice of action.
- Enclose all relevant documentation with the appeal request.
- Providers cannot submit a reconsideration following the appeal decision.
- Appeals can be submitted:
 - By mail: MARCH[®] Vision Care
Attention: PDR Unit
6701 Center Drive West, Suite 790
Los Angeles, CA 90045
- If you disagree with the appeal outcome, you can file a State Fair Hearing.

State Fair Hearing

- Providers must exhaust UHC appeal process prior to submitting a State Fair Hearing.
- Request must specifically request a “fair hearing” and should describe the decision appealed and the specific reasons for the appeal.
- Requests must be submitted within one hundred twenty (120) calendar days from the date of notice, plus an additional three (3) calendar days from the date the notice is sent.
- State Fair Hearing requests can be submitted through various means:

- By mail: Office of Administrative Hearings
1020 S. Kansas Avenue
Topeka, KS 66612
- By fax: (785) 296-4848

Provider Grievance Process

A grievance is any expression of dissatisfaction about any matter. Provider has one hundred eighty (180) calendar days from the date of service or incident to notify MARCH[®] of a grievance.

Grievances can be submitted by a provider or provider's authorized representative to MARCH[®] Vision Care either verbally or in writing as follows:

- Verbally: (844) 506-2724
 - In writing: You may send a written request with supporting documentation to providers@marchvisioncare.com or via mail or fax to:
MARCH[®] Vision Care
Attention: Provider Services
6701 Center Drive West, Suite 790
Los Angeles, CA 90045
Fax (877) 627-2488
- MARCH[®] will resolve each grievance fairly and consistent with our policies and covered benefits within thirty (30) calendar days from the date MARCH[®] receives the grievance. The grievance resolution will be communicated in writing to the Provider.
 - MARCH will acknowledge each grievance within ten (10) calendar days of receipt. If a grievance is resolved the same day of receipt, MARCH[®] is not required to issue a separate acknowledgement but will acknowledgement receipt in our resolution response.

This grievance process is the same for both participating and non participating providers.

3.11 Overpayment of Claims

If MARCH[®] determines a claim was overpaid or was paid incorrectly, MARCH[®] will notify the provider in writing. Overpayment refund requests are issued in accordance with the applicable provider services agreement and governing entity regulations. MARCH[®] does not issue overpayment refund requests more than three hundred and sixty five (365) days following the payment date, even when permitted by governing entity regulations.

Once an overpayment refund request is issued, if MARCH[®] does not receive an overpayment dispute request or refund of the overpaid amount within sixty (60) calendar days, with three (3) additional days for mailing, MARCH[®] may offset the overpayment against future claim payments if not prohibited by governing entity regulations.

3.12 Balance Billing

"Balance Billing" means charging or collecting an amount in excess of the Medicaid or contracted reimbursement rate for services covered under a Medicaid beneficiary's plan. "Balance Billing" does not include charging or collecting deductibles or copayments and coinsurance required by the beneficiary's plan.

Providers are prohibited from balance billing MARCH[®] members. The explanation codes MARCH[®] provides in the explanation of payment remittance advice clearly indicate when balance billing for a service is not permissible.

3.13 Coordination of Benefits

Coordination of Benefits (COB) is a method of integrating health benefits payable under more than one health insurance plan, allowing patients to receive up to 100% coverage for services rendered. Patients that have health benefits under more than one health insurance plan are said to have "dual coverage". In some cases patients may have primary, secondary, and tertiary coverage. When a patient has multiple plans or "dual coverage", it is necessary to know what plan is primary and what plan is secondary or tertiary. The primary plan must be billed first and the claim is billed just like any other claim would be billed. The secondary plan is billed once an explanation of payment (EOP) and possibly a payment is received from the primary plan. The claims submitted to a secondary or tertiary plan are considered "COB claims". When billing a secondary plan, the bill must have the primary insurance plans' EOP attached. The payments received from the primary plan should be

indicated in field twenty-nine (29) of the CMS 1500 form. If the secondary plan is billed without an attached primary insurance EOP, the claim will be contested and the primary insurance EOP will be requested. Medicaid will not make an additional payment if the amount received from the primary insurance company is equal to or greater than the Medicaid reimbursement amount.

MARCH[®] processes COB claims in accordance with the applicable provider services agreement and governing entity regulations. When MARCH[®] is the secondary payor, the primary payor information is required for calculation of the secondary payment. MARCH[®] subtracts the primary insurance payment from the lesser of the MARCH[®] allowable or the primary insurance plan allowable, not to exceed the applicable MARCH[®] reimbursement rates and benefit allowance.

The timeframe for filing a claim in situations involving third party benefits (COB and subrogation) shall begin on the date that the third party documented resolution of the claim. COB claims must be submitted as paper claims on a red CMS 1500 form.

Please mail COB claims to:

MARCH[®] Vision Care
6701 Center Drive West, Suite 790
Los Angeles, CA 90045

4.1 Access Standards

MARCH[®] optometrists and ophthalmologists are required to meet minimum standards of accessibility for members at all times as a condition of maintaining participating provider status.

In connection with the foregoing, MARCH[®] has established the following accessibility standards, when otherwise not specified by regulation or by client performance standards:

- Appointments for routine, non-urgent eye examinations and eyeglass or contact lens fittings and dispensing are available within thirty (30) calendar days.
- Rescheduling an appointment in a manner that is appropriate for the enrollee's health care needs and ensures continuity of care consistent with good professional practice.
- When MARCH[®] is contractually responsible for more than routine eye examinations, appointments for urgent/emergent eye care services, within the optometrist's or ophthalmologist's scope of practice, are available within twenty-four (24) hours.
- Providers are required to employ an answering service or a voice mail system during non-business hours, which provide instructions to members on how they may obtain urgent or emergency care. The message may include:
 - An emergency contact number (i.e. cell number, auto forwarding call system, pager);
 - Information on how to contact another provider who has agreed to be on-call to triage or screen by phone, or if needed, deliver urgent or emergency care; and/or
 - Instructions to call 911 or go to the local emergency room.
- Members with scheduled appointments will wait no more than thirty (30) minutes from their appointment time before being seen by a provider. Wait time is defined as the time spent in the lobby and in the examination room prior to being seen by a provider.

Additional state-specific requirements are provided below. In the event of a conflict between any standard above and those of a particular state, the more stringent standard shall apply.

4.2 Access Monitoring

MARCH[®] is responsible for monitoring compliance with accessibility standards. MARCH[®] will bear responsibility for reviewing and exercising oversight regarding matters such as member wait times, both for appointments and in the office, as well as other barriers to accessibility that may be reflected in member grievances, informal comments received by MARCH[®] employees or otherwise noted.

The following are some of the mechanisms that will be employed by MARCH[®] to verify access and compliance with its accessibility standards:

- Blast Fax requests may be used to gather information from providers to determine demographic, access and language information.
- Telephone access surveys will be conducted by MARCH[®] through random calls to optometrist and ophthalmologist offices to verify capacity to ensure that appointments are scheduled on a timely basis, with appropriate office wait time, and that appropriate after hours answering systems are being utilized.
- MARCH[®]'s grievance system also serves to identify access-related concerns. The tracking of grievances and an investigation of grievance patterns may result in the implementation of new policies and procedures and/or the education of participating optometrists, ophthalmologists, and staff members.
- Members may be provided with a Member Satisfaction Survey to comment on the service and products received from MARCH[®] and its providers.
- The appointment books of participating optometrists and ophthalmologists may be periodically reviewed during on-site inspections to validate the availability of appointments for services within reasonable time frames. Waiting rooms may also be periodically monitored to determine how long members wait for scheduled appointments.

The coordination of access monitoring is facilitated by MARCH[®]'s Department of Health Care Services. Reports of the results of these initiatives are prepared and presented to the Quality Improvement Committee and the Board of Directors which is responsible to ensure compliance with such standards.

5.1 Quality Improvement Program

Provider participation is of key importance to a successful Quality Improvement Program.

Provider participation in Quality Management (“QM”) activities includes:

- Participate in MARCH[®] Quality Committees including the Quality Improvement Committee, Peer Review Sub-Committee, Utilization Management Sub-Committee, and the Professional Review Committee.
- Participate in disease management programs.
- Adhere to adopted clinical care guidelines.
- Timely and appropriate response to member appeals and grievances including the provision of Medical Records when requested.
- Meet member access requirements.
- Participate in clinical reviews.
- Maintain medical record standards.
- Maintain the confidentiality of member information and records.

If you are interested in an active role in one of the committees noted above, please contact MARCH[®]'s Director, Health Care Services, Reva Sober at (310) 216-2300.

Please refer to the links below for additional information regarding the 2017 Quality Improvement Program.

[2018 Quality Improvement Program Non-Commercial Group](#)

5.2 Coordination with Primary Care Providers

Providers are encouraged to contact a member's Primary Care Provider (PCP) should they notice any additional medical needs while providing vision services. The assigned PCP is noted on the front of the member's ID card. Additionally, you may contact the Health Plan directly for assistance in coordinating any other needs for the member.

5.3 Medical Charting for Eye Care Services

In an effort to ensure quality of services and to combat fraud, waste and abuse, MARCH[®]'s Health Care Services Department perform audits of medical records used as supporting documentation to substantiate post-payment claims submissions. MARCH[®]'s PEER Review Sub-Committee has identified over seventeen (17) elements necessary in a comprehensive eye examination and, using a proprietary scoring system; records are evaluated and assigned a point value for each element based on their hierarchy of significance. The cumulative total point value is then used to determine the adequacy of the supporting documentation. When a comprehensive examination is billed, if any of the critical elements are skipped 10 out of 10 times, the audit score automatically defaults to the failing Severity Level score 4. These critical elements include: biomicroscopy/slit lamp exam, intraocular pressure, optic nerve head evaluation, and dilated fundus exam.

If any of these elements are missing or inadequately documented in the medical chart, MARCH[®] may send a request for a corrective action plan (“CAP”), asking you to address the documentation issue(s) identified during the audit.

Below are items to keep in mind for ensuring your medical chart supporting documentation is sufficient to pass an audit:

Paper Charts

- The encounter must record critical general health care information as well as the traditional refractive data. Details of a patient's medicine list and a formal review of systems are critical elements of the eye exam.
- Notes on pulse, blood pressure and body mass index.
- Providers must query about tobacco use and alcohol use, assess patient orientation to time and place, and rate the patient's emotional state during the exam.

Traditional paper charts may need to be updated to meet these standards. In addition to the requirements noted above, the form must include adequate space for a detailed slit lamp exam, notations for drugs that are administered during the exam, and a detailed posterior pole exam. A sample form that meets these requirements can be found in Exhibit E.

Electronic Medical Records

For providers using Electronic Medical Records (“EMR”), the following issues may be problematic. It is important to take them into consideration to ensure supporting documentation is sufficient:

- The templates for each encounter type, including the eye exam are customizable. Many providers have customized their office system in a way that has deleted key elements of the eye exam. Deleting some elements may make your charts non-compliant.
- EMR’s have “defaults” for normal findings that often fill in descriptive, detailed language for normal structures/findings. Caution should be used with defaults so that the clinical data and test results correlate with the diagnosis, assessment and management plan.
- When documentation is worded exactly like or similar to previous entries, the documentation is referred to as “cloned”. Cloning of documentation from a previous visit lacks the encounter-specific information necessary to support services rendered to patients.
- A review of the EMR for consistency, logical assessment, and treatment plans should be completed before signing the chart. The chart should not be manipulated or corrected once it is signed by the provider.

Critical Elements of an Eye Exam

The goal in medical chart review is to assist the providers in the improvement of the eye care encounter to meet today’s standards. For both paper charts and EMRs, the following elements are required for all comprehensive eye examinations:

Element 1 Reason for Visit

- **Why important:** This element should trigger the encounter type and then direct the examination to meet the needs of the encounter. The “reason for visit” should be addressed in the diagnosis/impression section at the end of the exam as well as in the treatment/management plan. The reason may be related to the time since the last examination and the patient may not have symptoms or abnormal signs.
- **What is expected:** The patient should be directly questioned as to why they presented for the encounter. The patient should also be asked about issues with their eyes and vision or other problems that may be related to the visual system. The answers to these queries should be documented in the medical record.
- **Who can collect data:** Doctor or Technician but findings must be attested to by doctor stating data has been reviewed.
- **How to document findings:** The information should be entered in free text or with bullet points.
- **When should optional testing be performed?** The responses to the reason for visit may redirect the exam to a problem focused visit rather than a routine eye examination. Testing and examination should follow the reason for the visit.
- **Quality point value:** 3 points
- **Critical element if box checked:**

Element 2 Review of Systems

- **Why important:** In addition to this review being a requirement for billing comprehensive eye examination codes and the Medical Vision evaluation and management codes, a review of systems documents all reported health issues and allows the doctor to discuss compliance with recommendations and follow-up with any necessary medical treatment with providers of the health care team. Historical information can assist in providing guidance as to required testing during the eye examination.
- **What is expected:** Each of the following systems should be queried and the patients’ response recorded. For all positive responses, additional questioning may be indicated.
 - Cardiovascular
 - Constitutional
 - Endocrine
 - Gastrointestinal
 - Head
 - Hematologic/Lymphatic
 - Immunologic
 - Integumentary
 - Musculoskeletal

- Neurological
- Psychiatric
- Respiratory
- **Who can collect data:** Doctor or Technician but findings must be attested to by doctor stating data has been reviewed.
- **How to document findings:** Findings are recorded as positive or negative. All positive findings should be questioned further and responses recorded in the patient's record.
- **When should optional testing be performed?** If history reveals a condition that may have manifestations in the eye, adnexa or visual pathways, additional testing may be warranted.
- **Quality point value:** 3 points
- **Critical element if box checked:**

Element 3 Medications and Allergies

- **Why important:** A patient's current medication list is an indicator of the overall health of the patient. Patients taking a number of medications have chronic health issues that can affect ocular health and the ultimate visual outcome. A patient's current list of medications also directs the eye examination so that the provider focuses more closely on certain components of the exam. For example, patients on several medicines for heart and circulation may develop optic nerve damage at a lower IOP and are at risk to develop macular degeneration. Some patient will report "no medical problems" because they assume that the use of medicines eliminates the problems. For example, in some cases, only a review of the medication list will reveal that the patient is a diabetic. The list of the patient's allergies is also critical as a patient may be allergic to some of the medications used in the eye examination. The patient may also call at some point after the exam and need a prescription for conjunctivitis or other medical eye conditions. Most providers review the last examination notes to assess the clinical situation and prescribe medications.
- **What is expected:**
 - Medications: Medication name and dosage for all drugs or supplements the patient is taking. If taking no medication, this should be indicated on the chart as none and not left blank.
 - Allergies: For allergies related to medications, the name should be listed as well as the adverse effect the member experienced. If the patient experiences environmental or food allergies, these should be noted as well. If no allergies are reported, the chart should indicate this result.
- **Who can collect data:** Data is collected from the patient intake form and verified by the doctor / technician during the history. It may also be collected during the history. It is required for each exam or patient encounter.
- **How to document findings:** Document as a list in the history section of the chart.
- **When should optional testing be performed?** If history reveals a condition that may have manifestations in the eye, adnexa or visual pathways, additional testing may be warranted
- **Quality point value:** 3 points
- **Critical element if box checked:**

Element 4 Ocular, Family History

- **Why important:**
 - **Ocular:** A patient's ocular history is one of the most important elements of the eye examination. It is impossible to provide a meaningful eye examination without the knowledge of previous problems, procedures and conditions
 - **Family History:** The modern understanding of genetics has opened new considerations for the treatment and management of ocular disease. From the routine problems of cataracts and glaucoma to the full spectrum of macular degeneration, the family history is critical in the treatment and management plan for each patient.
- **What is expected:** A detailed list of the patient's previous eye problems and procedures should be listed. The family history should query medical problems including diabetes, hypertension, thyroid problems and cancer in addition to eye problems such as cataracts, glaucoma, and macular degeneration.
- **Who can collect data:** This is collected from the patient intake form and verified by the doctor / technician during the history but findings must be attested to by doctor stating data has been reviewed.
- **How to document findings:** Document as a list or in free text in the history section of the chart.
- **When should optional testing be performed?** If history reveals a condition that may have manifestations in the eye, adnexa or visual pathways, additional testing may be warranted
- **Quality point value:** 3 points
- **Critical element if box checked:**

Element 5
Entering Visual Acuity at Distance and Near

- **Why important:** For medico-legal reasons entering visual acuity must be measured. In addition, the patient's acuity level establishes both a baseline for and a guide to further testing. All additional refractive findings should relate back to and be consistent with the entering visual acuity.
- **What is expected:** A measurement of visual acuity both uncorrected and with the patient's habitual correction should be performed at both distance and near.
- **Who can collect data:** Doctor or Technician
- **How to document findings:** Distance Visual Acuity is recorded as a Snellen fraction with the numerator the testing distance and the denominator the level of visual acuity the patient read. The most appropriate measure of near visual acuity is a fraction with the target size seen in meters as the numerator and the testing distance in meters as the denominator. Alternatively, near vision may be recorded using reduced Snellen acuity followed by the testing distance.
- **When should optional testing be performed:** If vision is abnormal or inadequate, the examination should be geared to finding the cause. If this is related to a stable pathology that is not resolvable, a low vision evaluation may be indicated.
- **Quality point value:** 10 points
- **Critical element if box checked:**

Element 6
Entering Tests
Vital Signs and
External Examination (Pupil testing/Extra Ocular Muscle testing
The Cover test/Screening Visual Field)

- **Why important:**
 - **Vital signs** are mandated by the Stage 2 Meaningful Standards for the appropriate use of an electronic health record. MARCH[®] requires vital signs to be recorded in both electronic and hand written medical records.
 - **External Examination** includes a battery of entering tests to assess a significant portion of the physical examination of the patient. Each test not only reveals clues to visual function, but also provides important screening of the neurological system.
- **What is expected: Measurement of-**
 - Height
 - Weight
 - Body Mass Index
 - Blood Pressure – for patients age 13 and older
 - Pulse
 - Testing of pupil response
 - Direct
 - Consensual
 - Swinging flashlight
 - Extra Ocular Muscle testing
 - Cover test
 - Visual field
 - Confrontation or
 - Automated test
- **Who can collect data:** Doctor or technician but findings must be attested to by doctor stating data has been reviewed.
- **How to document findings:** The information should be documented as each test is completed with the appropriate results listed for each test.
- **When should optional testing be performed?** If testing reveals a condition that is abnormal, additional testing may be warranted.
- **Quality point value:** 10 points
- **Critical element if box checked:**

Element 7 Refraction

- **Why important:** The subjective refraction is used to establish the final prescription.
- **What is expected:** The refraction required by the MARCH[®] standards is the subjective test that allows for the patients visual perception of the physical refractive error. Auto-refraction, by itself, is not an acceptable measurement
- **Who can collect data:** Doctor
- **How to document findings:** Sphere power, Cylinder power and axis for each eye as well as prism and bifocal power as indicated.
- **When should optional testing be performed?** If history reveals a condition that may have manifestations in the eye, adnexa or visual pathways, additional testing may be warranted
- **Quality point value:** 10 pts.
- **Critical element if box checked:**

Element 8 Near Point Testing

- **Why important:** If the patient is presbyopic or demonstrates signs or symptoms of near point problems, testing is indicated.
- **What is expected:** Testing may include measurements of accommodation and/ or convergence as well as additional testing as determined by the provider (e.g. evaluation of saccadic eye movements)
- **Who can collect data:** Doctor
- **How to document findings:** Responses to testing documented in medical record
- **When should optional testing be performed:** If reading issues are elicited during case history or if additional testing is indicated based upon clinical examination
- **Quality point value:** 3 points
- **Critical element if box checked:**

Element 9 Current Optical Prescriptions

- **Why important:** To prescribe eyeglasses, the provider must compare the patient's perception of their current vision with and without their current glasses, the entering visual acuity, the refractive testing and the patient's vision demands. The measurement and recording of the current prescription is a very necessary part of this decision making.
- **What is expected:** The current glasses prescription should be recorded in the refractive testing area.
- **Who can collect data:** This may be recorded by a technician or the doctor.
- **How to document findings:** Readings documented in medical record
- **When should optional testing be performed:** N/A
- **Quality point value:** 3 points
- **Critical element if box checked:**

Element 10 Corneal Curvature

- **Why important:** This test can be an important part of the refractive evaluation. It is a required element for contact lens fittings but it is an optional test for the eye examination. The clarity, regularity and quality of the mires can point to the cornea or the tear film as a cause of reduced vision or other symptoms. In cases where the final acuity is less than expected, a Keratometry reading would be expected as part of the refractive work up. In lieu of keratometry, corneal topography is an acceptable procedure.
- **What is expected:** The measurement should be recorded in the refractive testing area when indicated.
- **Who can collect data:** This may be recorded by a technician or the doctor.
- **How to document findings:** Readings documented in medical record
- **When should optional testing be performed:** N/A
- **Quality point value:** 3 points
- **Critical element if box checked:**

Element 11 Biomicroscopy

- **Why important:** Biomicroscopy is useful in diagnosing infections, allergies, inflammations and other disease entities affecting structures in the anterior segment of the eye.
- **What is expected:** Use of the biomicroscope to inspect all anterior segment eye structures including the lids and lashes, tear film, cornea, anterior chamber, angle grade, iris and lens. The documentation must be individualized based on the findings of the examination. Cloned language in Electronic Health Records should be carefully reviewed and revised to be consistent with the rest of the documentation in the record.
- **Who can collect data:** Doctor
- **How to document findings:** Positive and negative findings are recorded by structure (e.g. cornea, lids, etc.). Using “WNL” or a line through a list of anatomical parts is not adequate documentation.
- **When should optional testing be performed:** Findings may indicate the need to perform gonioscopy if narrow anterior chamber angles are suspected as well as retinal examination with a hand-held fundus lens or 4 mirror lens.
- **Quality point value:** 10 points
- **Critical element if box checked:**

Element 12 Intraocular Pressure

- **Why important:** Intraocular pressure must be measured at each comprehensive eye examination. This is a critical evaluation of eye health. The pressure result in each eye is important as well as its relationship to the fellow eye. The trend over time is significant for each patient.
- **What is expected:** The type of instrument used as well as the time of measurement should be included with the numerical finding.
- **Who can collect data:** Doctor /Technician (subject to applicable laws)
- **How to document findings:** The test may be performed by any of the accepted methods such as Goldman Applanation, non-contact, or Tonopen, including digital or finger tension depending on the clinical situation. The numerical value should be charted along with the time of measurement.
- **When should optional testing be performed:** Appropriate treatment or additional testing should be done depending on the results and correlation with other clinical findings.
- **Quality point value:** 10 points
- **Critical element if box checked:**

Element 13 Optic Nerve Head Evaluation

- **Why important:** The health and status of the optic nerve head is critical to vision and ocular health. The relationship to the fellow eye and any change over time is clinically significant.
- **What is expected:** The optic nerve must be visualized and details recorded at each visit. The details of the evaluation of the Optic nerve should include all aspects of the nerve itself including: -Cup to disc ratio, - disc margin, - disc size, - color, - thickness, - vessel caliber. The exam may be performed with a fundus lens, the direct ophthalmoscope, indirect ophthalmoscope, or photographically. At a minimum a fundus lens should be utilized.
- **Who can collect data:** Doctor
- **When should optional testing be performed:** Appropriate treatment or additional testing should be done depending on the results and correlation with other clinical findings.
- **Quality point value:** 10 points
- **Critical element if box checked:**

For additional information, you may refer to the following link:

http://eyewiki.aao.org/Examination_of_the_optic_nerve_at_the_slit-lamp_biomicroscope_with_a_handheld_lens.

Element 14 Dilated Fundus Examination

- **Why important:** A dilated retinal examination is performed to detect any abnormal findings for baseline documentation. Retinal abnormalities may indicate the presence of local and/or systemic disease and indicate the need for further diagnosis and/or treatment. MARCH[®] has approved criteria which requires the entire ocular fundus be examined on the initial visit and periodically thereafter depending on each patient’s risk factors, but at least every three (3) years. Patients with diabetes must be dilated every year.

- **What is expected:** A thorough inspection of the optic nerve, macula, vascular tree and retinal surface with a fundus lens and biomicroscope, a binocular indirect ophthalmoscope and/or a wide angle retinal camera. Document the method of examination.
- **Who can collect data:** Doctor
- **How to document findings:** Positive and negative findings are recorded by structure (e.g. optic nerve, macula etc.). Using “WNL”, a line through a list of anatomical parts, or noting clear is not adequate documentation. The name of the dilating drops as well as the time of instillation should be documented. Confirm drops have not expired prior to instilling them.
- **When should optional testing be performed:** Suspected lesions, macular abnormalities, retinal holes or tears, etc. may require photographic imaging, ocular coherence tomography, ultra-sound, etc. to determine a diagnosis and treatment plan.
- **Quality point value:** 10 points
- **Critical element if box checked:**

Element 15 Diagnosis

- **Why important:** Following each clinical encounter, the provider must list each relevant diagnosis.
- **What is expected:** These can be a refractive diagnosis such as Myopia, Astigmatism, Emmetropia, Hyperopia, or Presbyopia or medical eye diagnoses such as Cataract, Corneal Dystrophy, Choroidal Nevus or Glaucoma. Pertinent medical diagnoses such as diabetes should also be listed.
- **Who can collect data:** Doctor
- **How to document findings:** The provider should list these at the end of the exam chart. The diagnosis of each eye examination is often forwarded to the Primary Care Provider as part of the MARCH[®] quality initiative to coordinate care.
- **Quality point value:** 3 points
- **Critical element if box checked:**

Element 16 Assessment/Management/Treatment Plan

- **Why important:** The review, summation and recommended treatment plan requires the critical thinking by the doctor to summarize the eye examination.
- **What is expected:** In this section, the provider should summarize the overall examination, and clarify the points that need to be managed. The treatment/management plan should spell out the steps to be taken to address the chief concerns identified in the clinical findings. In healthy patients, this can be as simple as, “Normal Exam, return in 1 year for re-examination.” For a patient with refractive error, the verbiage can include the diagnosis and be stated as “Myopia, order glasses to be used for distance only, return in 1 year. For patients with pathology, this section should be more specific and address patient education, glasses, contact lenses, low-vision aids, medications prescribed with directions for use, referrals, recommended testing, time frames and follow-up schedules. Other clinicians, reviewers, and any party evaluating this clinical encounter will look to this section to determine the important clinical points of the case and identify the plan of action/recommended follow-up.
- **Who can collect data:** Doctor
- **How to document findings:** The provider should list these at the end of the exam chart.
- **Quality point value:** 3 points
- **Critical element if box checked:**

Element 17 Legible Records

- **Why important:** Charts must be readable and legible so they may be interpreted by anyone authorized to review them, including MARCH[®]. This is important for review of historical patient data, continuity of patient care, auditing purposes, quality measures and for medico-legal reasons.
- **What is expected:** Records that are easily deciphered, following a consistent examination sequence, that are complete and document all findings, clinical decisions and any continuity of care recommendations. If using electronic medical records, it is important to review any “pre-populated” and/or “cloned” default data for accuracy, attest to the doctor personally reviewing history and medications and review all recorded data to ensure it reflects the examination findings and recommendations. A signature is required on all charts.
- **Who can collect data:** Doctor or staff
- **How to document findings:** Examination results are recorded on a “paper” chart or entered into an electronic medical record.

- **When optional testing is performed:** Any additional testing that generates data (e.g. retinal photography, visual fields, etc.) should be printed and included in the patient's medical record, scanned and attached to the electronic medical record or the location of the information documented within the patient's chart. For testing that requires it, an "Interpretation and Report" analysis must also be included in the record.
- **Quality point value:** 3 points
- **Critical element if box checked:**

6.1 Anti-Fraud Plan

Pursuant to Health and Safety Code Section 1348, MARCH[®]'s anti-fraud plan includes, but is not limited to, the following requirements:

1. The designation of an organization with specific investigative expertise in the management of fraud investigations.
2. Training of personnel and contractors concerning the detection of health care fraud.
3. Procedures for managing incidents of suspected fraud.
4. Procedures for referring suspected fraud to the appropriate government agency.

Designation of an Organization with Specific Investigative Expertise in the Management of Fraud Investigations

MARCH[®] has designated the law firm of Katten Muchin & Rosenman, LLP ("KMR") as its fraud investigator. KMR has substantial experience in the management of fraud investigations.

Training of Personnel and Contractors Concerning the Detection of Health Care Fraud

MARCH[®] recognizes the importance of properly educating and training its personnel and contractors to detect fraud by MARCH[®], MARCH[®]'s providers, and MARCH[®]'s members. As part of its anti-fraud plan, MARCH[®] requires its personnel and contractors to receive the following training in the detection of health care fraud:

Training of MARCH[®] Personnel

All MARCH[®] personnel will be annually trained in the detection of fraud and all new personnel will be trained in the detection of fraud upon hire.

The training of MARCH[®] personnel will include a general training session for all MARCH[®] personnel regarding the most common types of health care fraud that impact managed care organizations and may include specialized training for MARCH[®] personnel who work in the enrollment, credentialing, claims, and marketing areas regarding the identification and detection of fraud that is likely to specifically impact their jobs. In addition, the Chief Executive Officer shall establish such other training and dissemination of information to all employees concerning the necessity of complying with all applicable laws and regulations and shall keep MARCH[®] personnel abreast of current trends and issues relating to fraud on an ongoing basis through informational bulletins and discussions.

MARCH[®] personnel shall sign an Employee Statement of Understanding regarding the anti-fraud plan both at the time of their initial anti-fraud training, and thereafter on a yearly basis. The initial signed Statements of Understanding shall be kept in each employee's personnel file. The annual attestation is collected through an electronic form and signature once the Compliance Department completes the annual training.

Training of MARCH[®]'s Participating Providers

All of MARCH[®]'s participating providers will receive a copy of MARCH[®]'s anti-fraud plan. They will be required to either adopt and comply with MARCH[®]'s anti-fraud plan, or to have their own anti-fraud plan/compliance program in place that meets or exceeds the standards of MARCH[®]'s anti-fraud plan. MARCH[®] will also issue provider communications from time to time concerning fraud detection and related issues.

Areas of Training

Training includes an overview of health care fraud, a summary of the applicable fraud and abuse laws, training on how to identify potentially fraudulent claims (including indicators of fraud), examples of fraudulent activity that has been uncovered and the procedure for referring suspected fraudulent activity to the Chief Executive Officer.

Training topics will include, but not be limited to, methods of detecting the following types of fraud:

1. Detection of Fraud by the Plan
 - a) Marketing - Using marketing techniques that coerce, mislead or confuse potential members and engaging in marketing that discriminates among potential members based on their health status.

- b) Underutilized/Quality of Care - Failing to employ or contract with sufficient providers to accommodate all members; failing to provide geographically reachable services to members; and categorically denying payment of claims.
- c) Enrollment Fraud - Using unnecessarily complex disenrollment procedures and materials.
- d) Licensure/Credentialing - Not adequately credentialing providers; contracting with unlicensed providers.
- e) Kickbacks - Accepting kickbacks in order to refer certain members to a particular provider.

2. Detection of Fraud by Providers

- a) Marketing - Failing to comply with the applicable licensing board's advertising guidelines.
- b) Kickbacks - Providers paying kickbacks to MARCH[®] employees in order to be referred members.
- c) False Claims - Billing for services that were never performed or were not medically necessary; and waivers of copayments or deductibles.
- d) Licensure/Credentialing - Misrepresenting licensure status to MARCH[®].

3. Detection of Fraud by Members

- a) Enrollment Fraud - Members claiming to be eligible for MARCH[®] health coverage when they are, in fact, ineligible.

4. Identification of Possible Indicators of Fraud

The training will emphasize that certain circumstances may be indicative of fraudulent activity, and should be reviewed further. Such circumstances include, but are not limited to, the following:

- a) Inconsistency between the services billed and the services rendered.
- b) A provider's advertisement of "free" services.
- c) An unusually high number of members/member visits in a given time frame.
- d) A provider's lack of supporting documentation for a claim selected for audit.
- e) A high-dollar claim for services dated soon after the effective date of coverage or just before the termination of coverage.

Procedures for Managing Incidents of Suspected Fraud

Upon reports or reasonable indications of fraud, the Chief Executive Officer will promptly initiate steps to investigate the conduct in question to determine whether fraudulent activity has occurred. As needed, the fraud investigator will be requested to conduct the investigation. If the Chief Executive Officer and/or fraud investigator determines that fraudulent activity has occurred, the Chief Executive Officer will develop an appropriate response, as described below.

Discovery of Fraudulent Activities

1. Reporting Incidents of Suspected Fraud

All MARCH[®] personnel are responsible for preventing, detecting and reporting suspected fraud. If an employee detects any suspicious activity, he/she is required to notify the Chief Executive Officer. The person reporting fraud may make himself/herself known by reporting the suspected fraud in person, or may report the suspected fraud anonymously via inter-office mail or U.S. Mail.

The manager of each department will be responsible for the early detection of fraud within his/her department. If fraud is suspected within a department, that department's manager is required to immediately notify the Chief Executive Officer. Each manager's performance evaluation will be based in part on his/her efforts to detect fraud.

2. Implementation of a Monitoring and Audit Program

The Chief Executive Officer will implement a monitoring and audit program, as necessary. Through the use of ongoing auditing and monitoring, the Chief Executive Officer will investigate any changes from the baseline audit that may be indicative of fraud. Ongoing auditing and monitoring will enable MARCH[®] to gather some of the information MARCH[®] will need to make annual reports to the Department of Managed Health Care as required by Health and Safety Code Section 1348(c).

3. As determined to be necessary by the Chief Executive Officer, the implementation of the monitoring and audit program may involve the following steps:

- a) Interviewing personnel involved in enrollment, credentialing, claims, marketing, and related areas to detect potential improper conduct.
- b) Reviewing medical and financial records and other source documents that support claims for reimbursement.
- c) Reviewing written materials and documentation prepared by the different departments within MARCH®.

Investigate the Incident to Determine Whether there is a Violation of Law/Regulation/MARCH® Policy

The Chief Executive Officer or his/her designee will investigate all credible incidents of suspected fraud that are reported and all credible incidents that are uncovered pursuant to the auditing and monitoring program. The investigation will involve interviews and document review. In the case where employee fraud is suspected, the Chief Executive Officer will determine whether the employee should be removed from his/her duties until the investigation is completed and whether or not immediate steps should be taken to prevent the destruction of documents or other evidence relevant to the investigation. The Chief Executive Officer shall record the progress of the investigation, including the results of interview and document reviews.

Take Appropriate Remedial Measures

If fraudulent activity has occurred, the Chief Executive Officer will consult with the manager of the department in which the fraudulent activity has occurred to determine the appropriate action necessary to correct the matter. The following remedial measures will be taken, as applicable:

1. Deny/Recoup Payment - If the fraudulent activity involves payment to a provider or to a member, the payment will be denied if not yet made, and will be recouped if already made.
2. Terminate Contract/Discipline Employee Appropriately - If appropriate, contracts with providers will be terminated, and employees will be disciplined. Corrective action will be based upon the individual circumstances and the severity of the incident. All personnel will be disciplined similarly, regardless of their position within MARCH®.
3. File Appropriate Reports - If fraudulent behavior constitutes a reportable offense, a report will be made to the appropriate entity. Examples include reports required by California Business & Professions Code Section 805, and reports required by the National Practitioner Data Bank.
4. Notify Appropriate Government Agencies - See below.
5. Take Further Remedial Measures - In order to decrease the possibility that fraud will reoccur, the Chief Executive Officer will educate MARCH® personnel and participating providers regarding how to avoid the recurrence of any fraudulent activities that are discovered. In addition, the Chief Executive Officer will undertake additional investigations or other actions if it appears there may be a continuing pattern of fraud.

Procedures for Referring Suspected Fraud to the Appropriate Government Agency

MARCH® is committed to aggressively investigate suspected fraud and is committed to referring fraud for prosecution as appropriate. At least annually, MARCH® shall submit a report to the Department of Managed Health Care regarding MARCH®'s adherence to its anti-fraud plan generally and the results of investigations conducted by MARCH® regarding suspected fraud.

The Chief Executive Officer will discuss the findings of fraud investigations with legal counsel to determine whether or not a violation of federal or state law or health care program requirements has occurred, whether or not the conduct should be disclosed to a governmental agency, and, if so, to which agency. Such disclosure will observe the following guidelines:

1. Providers that are found to be in violation of state licensing requirements will be reported to the appropriate state licensing board.
2. Plan employees, providers or members who are found to be in violation of other state laws will be reported to the District Attorney's Office.
3. Providers that are found to be in violation of a federal, criminal, civil or administrative law related to a federal health care program will be reported to the Office of Inspector General, Department of Justice or the Centers for Medicare and Medicaid Services, as appropriate.
4. Plan employees, providers or members who are found to be in violation of other federal laws will be reported to the Department of Justice/U.S. Attorney's Office.

Anti-Fraud Plan Oversight

MARCH[®]'s Board of Directors is responsible for overseeing MARCH[®]'s anti-fraud plan. The Chief Executive Officer is responsible for implementing MARCH[®]'s anti-fraud plan and will make quarterly reports to Board of Directors regarding anti-fraud activities to enable the Board of Directors to monitor the anti-fraud plan and recommend any necessary changes.

7.1 Credentialing and Re-Credentialing

All potential providers are required to submit their CAQH number for credentialing.

CAQH ProView

MARCH[®] accepts CAQH numbers for the purpose of credentialing which will expedite the credentialing process as well as decrease the amount of paperwork for you and your staff. To expedite credentialing, please provide us with your CAQH number as soon as possible. CAQH ProView does not accept paper applications. To further avoid delays in processing; please be sure to give MARCH[®] permission on the CAQH ProView site to access the provider's record.

Please ensure the following documents are up-to-date:

- Completed W-9 form.
- State license.
- Current malpractice face sheet showing expiration dates, limits and provider's name.
- Curricula vita/resume to include work history if application does not cover last five (5) years.
- Board certificate (if applicable).
- CDS, CSR certificate, and/or DEA certificate (if applicable).

Credentialing Process

Upon receipt of the CAQH number, credentialing information is reviewed by the Credentialing Coordinator for completeness. All data, licenses and certificates are electronically confirmed by the applicable regulatory agencies, and any provider not in good standing with his/her respective regulatory agency is pended. The confirmed CAQH number is forwarded to the Professional Review Committee Chairperson for review and consideration. If consideration is favorable, the provider is approved. If the consideration is not favorable, the information is sent back to the Credentialing Coordinator with recommendations for further review.

Re-Credentialing Process

All providers are re-credentialed every three (3) years. The Provider Services Agreement stipulates automatic yearly renewal. The provider must forward to MARCH[®] on an annual basis a current photocopy of his or her yearly state license renewal and malpractice insurance. Failure to provide updated information may affect claims payments. Membership in good standing is re-confirmed.

Health Plan Credentialing Process

Health plans may perform Primary Source Verification on their own or in parallel. In order to comply with any state and/or health plan specific policies, you may be required to provide all pertinent credentialing documents on more than one occasion.

7.2 National Provider Identifier

The National Provider Identifier ("NPI") is a Health Insurance Portability and Accountability Act ("HIPAA") Administrative Simplification Standard. The NPI is a unique identification number for covered health care providers. Covered health care providers, all health plans and health care clearinghouses must use NPIs in the administrative and financial transactions adopted under HIPAA.

In accordance with 45 CFR § 162.410, MARCH[®] shall require each provider rendering services to members to have a National Provider Identifier.

7.3 Disclosure of Criminal Conviction, Ownership and Control Interest

In accordance with 42 CFR, Part 455, Subpart B and as required by CMS, individual physicians and other healthcare professionals must disclose criminal convictions, while facilities and businesses must additionally disclose ownership and control interest, **prior to payment for any services rendered to Medicaid enrollees.**

Prior to participation, all potential providers must accurately complete and sign the Disclosure of Ownership and Control Interest Statement Form. The State of Kansas Disclosure of Ownership and Control Interest Statement form is available on the Kansas Medical Assistance Program website at <https://www.kmap-state-ks.us/Documents/Content/Forms/DNO.pdf>.

The Disclosure of Ownership and Control Interest Statement is to be submitted with the provider's initial credentialing and recredentialing application (every three (3) years), or at initial and renewal of a contract or agreement and any time there is a revision to the information. This form must also be provided within thirty-five (35) days of a request for this information. If a provider or health care professional is a member of a group practice, **both** the individual member and group practice must submit a signed Statement attesting to the requirements under these regulations.

In order to comply with these Federal Regulations MARCH[®] Vision Care has suspended payments to providers who have failed to comply and have not submitted a valid and completed disclosure form to MARCH[®] Vision Care.

8.1 Interpreter Services and Language Assistance

Access to Interpreters

When a provider office identifies a member as being Limited English Proficient (LEP) and the member is present in the office, telephone interpretation should be used immediately to avoid any delay in services. There are new federal requirements for language services. The federal guidance, published as Section 1557 of the Affordable Care Act (ACA), provides specific limitations on the use of Bilingual Staff and minors as interpreters. These requirements are not limited to federal programs.

Providers are at risk if they use in house bilingual staff if they are not qualified interpreters. Qualified interpreters:

- Adhere to generally accepted interpreter ethics principles, including client confidentiality;
- Have demonstrated proficiency in speaking and understanding both spoken English and at least one other spoken language;
- Is able to interpret effectively, accurately, and impartially, both receptively and expressly, to and from such language(s) and English, using any necessary specialized vocabulary, terminology and phraseology.

Minors may not be used as interpreters except in emergency situations involving an imminent threat to the safety or welfare of an individual or the public where there is no qualified interpreter for the individual with limited English proficiency immediately available; **No one** can give permission to use a minor in a non-emergency.

The Provider shall not:

- (1) Require an individual with limited English proficiency to provide his or her own interpreter;
- (2) Rely on an adult accompanying an individual with limited English proficiency to interpret or facilitate communication, except:
 - a. In an emergency involving an imminent threat to the safety or welfare of an individual or the public where there is no qualified interpreter for the individual with limited English proficiency immediately available;
 - b. Where the individual with limited English proficiency specifically requests that the accompanying adult interpret or facilitate communication, the accompanying adult agrees to provide such assistance, and reliance on that adult for such assistance is appropriate under the circumstances;

Non-compliance may expose providers to the risk of violating a consumer's civil rights resulting in civil rights law suits and also be subject to law suits filed by the Office of Civil Rights. Enforcement and law suits may occur up to a year after the Date of Service.

Providers should document all actions taken to comply with this law – this documentation must be accessible and complete.

Suggestions:

- Appoint an employee to oversee compliance;
- Make sure aids and services comply with the law;
- Draft the required nondiscrimination notice and, if the entity has 15 or more employees, grievance policy;
- Review covered services to identify if any changes are needed
- Conduct training.

Health care providers are responsible for ensuring that patients have a full understanding of their diagnosis and treatment guidelines, regardless of their preferred language. To ensure that all limited English proficient members receive appropriate access to vision care, all providers are expected to comply with federal and state requirements regarding cultural and linguistic services. It is not permissible to turn a member away; to limit the member's participation or access to services because of language barriers; to subject a member to unreasonable delays due to language barriers; or to provide services to LEP members that are lower in quality than those offered in English.

Interpreting Services

Access to free language assistance services for members with LEP is required by various regulations. Interpreters must be professionally trained and versed in medical terminology and health care benefits. Providers can contact UnitedHealthcare's Provider Services Center at (877) 542-9235 to arrange language assistance services.

A Customer Service Representative will require the following information:

Member Information:

- Member name and identification number.
- Language requested.

Provider information:

- Provider name and telephone number.
- Office address.

Medical Record Documentation for LAP

For all LEP members, it is best practice to document the member's preferred language in paper and/or electronic medical records in the manner that best fits your practice flow. In addition, when possible, provider should attempt to collect and document member's race, ethnicity, and preferred written language in member's medical record.

If a member refuses or declines interpretive services, as a participating provider with MARCH® you should document the refusal/declination of services in the medical record. This documentation not only protects you and your practice, it also ensures consistency if your medical records are monitored through site reviews or audits.

Documentation of Provider/Staff Language Capabilities

The provider directory lists fluent languages spoken by providers. This information is received via self-reported Provider Demographic Forms updated on a quarterly basis, or whenever there is a demographic change. The information you provide will be used to update MARCH®'s provider database, which is used to generate our provider directories and to provide members with online and automated information to assist them in identifying provider offices that may meet their language needs.

Translation of Written Material

Translations of written informational material such as applications, consent forms, denial notices and explanation of payments are available through the member's Health Plan (the number on the back of their Identification Card).

9.1 Cultural Competency

MARCH[®] shall ensure that all health plan members receive equitable and effective treatment in a culturally and linguistically appropriate manner. As a health care provider, MARCH[®] expects you to be culturally sensitive to the diverse population you serve by effectively and appropriately providing services to people of all races, cultures, religions, ethnic backgrounds, education, and medical status in a manner that recognizes values, affirms and respects the worth of each individual member, and protects and preserves the dignity of each.

What is cultural competency?

Culture refers to integrated patterns of human behavior that include the language, thoughts, actions, customs, beliefs, values, and institutions that unite a group of people. It impacts the care given to members because it describes:

- Concepts of health, healing
- How illness, disease, and their causes are perceived
- The behaviors of patients who are seeking health care
- Attitudes toward health care providers

It also defines health care expectations such as:

- Who provides treatment
- What is considered a health problem
- What type of treatment
- Where care is sought
- How symptoms are expressed
- How rights and protections are understood

And why is it important?

Cultural competency is one the main ingredients in closing the disparities gap in health care. It's the way patients and doctors can come together and talk about health concerns without cultural differences hindering the conversation, but enhancing it. Quite simply, health care services that are respectful of and responsive to the health beliefs, practices and cultural and linguistic needs of diverse patients can help bring about positive health outcomes.

There are many cultural influences that impact the office visit. Some cultural preferences to remember include:

- Do members feel their privacy is respected?
- Are they the health care decision maker?
- Does their belief in botanical treatments and healers contradict standard medical practices and does it impact their decisions?
- What type of language skills and preferences do they use in their interactions?

Because health care is a cultural construct based in beliefs about the nature of disease and the human body, cultural issues are actually central in the delivery of health services.

Culture impacts every health care encounter. By understanding these influences and by communicating clearly at each visit you fulfill the opportunity to build rapport, help improve adherence and safety.

10.1 Secure Transmission of Protected Health Information (PHI)

To ensure that all communications (email, phone, or fax) containing Protected Health Information (PHI) (i.e. member number, name, address, etc.) from provider organizations meet HIPAA privacy guidelines, we are asking providers to follow the recommended guiding principles when exchanging PHI with MARCH[®] Vision Care.

- First, please determine if it is business necessary to exchange PHI with MARCH[®], the MARCH[®] recipient of PHI is appropriate, and include only the "minimum necessary" information.
- If you have a business need to exchange PHI with MARCH[®] personnel via email, please check with your IT personnel to make sure they have a secure transmission setup with MARCH[®] email systems. For more details, follow steps described in Exhibit D: "Sending a Secure Email to MARCH[®] Vision Care for PHI related data" to ensure that HIPAA guidelines are being met and PHI is secured. This will prevent MARCH[®] from receiving unencrypted or unsecured emails with PHI.
- While sending PHI securely via encrypted emails, please be aware that the HIPAA Privacy Rule still requires that PHI only be shared with those who are permitted to have the information and share only the minimum amount of PHI necessary to accomplish the business purpose.
- Please be aware that when contacting MARCH[®] by phone, email, or fax that we are required to confirm your name, associated provider/physician organization, and contact information before exchanging or confirming PHI.
- If you receive PHI or Personally Identifiable Information ("PII") directed to, or meant for, another provider or someone other than you, you agree to promptly destroy all such PHI or PII and not further use or disclose it. In addition, if such an event occurs, you agree to cooperate with any remediation efforts undertaken by MARCH[®].

Thank you in advance for following these recommended steps as we improve our business processes.

Exhibits

- Exhibit A Non-Covered Service Fee Acceptance Form
- Exhibit B MARCH[®] Lab Order Form
- Exhibit C Clinical Practice Guidelines
- Exhibit D Sending a Secure E-mail to MARCH[®] Vision Care for PHI Related Data
- Exhibit E Examination Record Template
- Exhibit F Performance Measurement & Reporting

Non-Covered Service Fee Acceptance Form

I _____, a member of _____ wish to obtain and pay for _____, a service which is not covered as a covered benefit under the Medicaid/Medicare Program under which I have coverage.

Dr. _____ has explained to me that I will be solely responsible for the cost of _____, which is \$_____. I agree to accept responsibility for payment of \$_____. I understand that I am not obligated to pay for the above service if it is later found that the service was covered under the Medicaid/Medicare Program under which I have coverage at the time it was provided, even if Medicaid/Medicare did not pay Dr. _____ for the service because he or she did not satisfy Medicaid/Medicare billing requirements.

I acknowledge that I have been given a copy of this agreement.

Member's Signature

Printed Name

Date

Formulario de aceptación del cargo por servicios no cubiertos

Yo _____, miembro de _____ deseo obtener y pagar el costo de _____, un servicio que no tiene cobertura como beneficio cubierto en el programa de Medicaid/Medicare bajo el cual tengo cobertura.

El/la Dr(a). _____ me explicó que yo seré el único responsable del costo total de _____, que es \$ _____. Acepto responsabilizarme del pago de \$ _____. Entiendo que no tengo la obligación de pagar por el servicio indicado arriba si posteriormente se determina que cuando se me brindó el servicio sí tenía cobertura en el programa de Medicaid/Medicare bajo el cual tengo cobertura, aunque Medicaid/Medicare no le haya pagado al/a la Dr(a). _____ el servicio porque él o ella no cumplió con los requisitos de facturación de Medicaid/Medicare.

Confirmando que recibí una copia de este acuerdo.

Firma del miembro

Nombre en letra de imprenta

Fecha

MARCH[®] Lab Order Form Instructions

Lab order forms may be submitted online through eyeSynergy[®].

- Complete the Lab Order Form on the following page. Please print clearly.
- Use of this Lab Order Form for non-plan members is prohibited.

IMPORTANT: If you choose not to submit lab orders through eyeSynergy[®], you **must** fax your order to our Customer Service Center at (855) 640-6737.

If you need to contact one of our contracted labs, please refer to the table below to determine the appropriate lab assigned to your state:

State	Lab Information	Contact Information
Kansas	Classic Optical Laboratories, Inc.	3710 Belmont Avenue Youngstown, OH 44505 Phone (888) 522-2020 MARCH [®] Vision Care fax: (855) 640-6737

MARCH Lab Order Form

Please fax completed form to (855) 640-6737

MEMBER INFORMATION

Member's name: _____ Today's date: _____
 Member's ID number: _____ Date of eye exam (if known): _____

PROVIDER INFORMATION

TIN: _____ Confirmation Number: _____
 Provider name: _____ Phone Number: _____

Address: _____

Material to Order Check all that apply.

- Frame Right Lens Left Lens Uncut Lenses

Is this a replacement? Yes No

	Sphere	Cylinder	Axis	Prism In / Out	Prism Up/ Down	Add Power	Seg Height
Right							
Left							

	Distant PD	Near PD	Requested Base Curve
Right			
Left			

Materials:

- Plastic Hi-Index 1.60
 Glass Hi-Index 1.67
 Polycarbonate Photochromic: Grey or Brown
 Trivex Edge Polish

Segment Style

- SV PAL Standard Trifocal 7x28
 FT28 PAL Standard Short Round 22 or 24
 FT35 PAL Premium Slab Off
 FT45 PAL Premium Short

Coating Options:

- Solid Mirror Coating Mirror Type: _____
 Gradient Scratch Coating AR Standard
 Double Gradient UV AR Premium
 Color: _____% or Density #1 #2 #3

Frame Selection:

- Patient Supplied Frame/Non-Formulary Frame Rimless Drill 2 Hole Rimless Drill 4 Hole
 (Please include copy of order form with shipment of PSF/NFF. Please ship frame to lab within 48 hours of submitting order to MARCH[®]).

Frame Manufacturer: _____ Lens Size: _____ Bridge Size: _____

Frame Model: _____ B Measurement _____ ED Measurement: _____

Frame Color: _____ Temple Size _____ Edge Type _____

Other Instructions/Special Notes

I certify that the prescription information supplied above is medically indicated and necessary to the health of this patient and was personally furnished by me or my employee under my personal direction. This is to certify that the foregoing information is true, accurate and complete. I understand that payment and satisfaction of this order will be from Federal and State funds, and that any false claims, statements, or documents or concealment of a material fact may be prosecuted under applicable Federal and State laws.

Provider Signature: _____

Clinical Practice Guidelines

Clinical practice guidelines describe the expected standard of practice for participating providers that is specific to the membership demographics and service needs and serves as the basis for a health management programs benefit interpretation and quality/performance measurements.

MARCH[®] is committed to providing high quality services to its members. MARCH[®] does not pressure health care providers or institutions to render care beyond the scope of their training or experience. The Quality Improvement Committee has adopted the following guidelines for its providers:

Standard of Care for Eyeglass Dispensing/Fitting and Contact Lens Fitting

EYEGLASS DISPENSING/FITTING

- Assist with frame fashion selection.
- Evaluate frame for appropriate eye size, bridge, and A, B, and ED for required lenses.
- Take physical measurements including PD, Seg Height.
- Order materials via eyeSynergy[®] or fax order to MARCH[®] (see Exhibit B).
- Monitor laboratory for appropriate turnaround time and follow up with MARCH[®] and the member as necessary.
- When materials have been received, measure lens power, PD, and Seg Height and physically inspect frame and lenses for manufacturer defects.
- Promptly contact the member when the eyewear has passed inspection.
- Adjust frame as needed to assure proper fit and alignment of lenses.
- Discuss proper use.

CONTACT LENSES FITTING

- Assess the health of the eyes in relationship to wearing contact lenses (age/anatomy etc.).
- Assess the anatomical appropriateness of the eyelids.
- Assess the quality and volume of tear film.
- Perform refractive tests and calculations related to contact lenses.
- Examine for issues and physical findings related to contact lenses.
- Measure cornea by keratometry and/or topography.
- Conduct diagnostic contact lens evaluation.
- Train patient on safe and effective lens care, and insertion and removal of lenses.
- Dispense final lenses or provide final prescription.
- Follow up visits for one month as indicated.

Standards for Dilation

All new members require a dilated fundus exam or equivalent (if acceptable per state/federal regulation). Diabetics require dilation every year at a minimum, more often if they have retinopathy. Members with other certain pathology such as lattice degeneration, choroidal nevi, or retinoschisis for example, may also need a dilated exam every year or as medically indicated. Members with no risk factors should be dilated thereafter based on the professional judgment of the provider or every three (3) years, whichever occurs first.

MARCH[®] Vision Care's Quality Improvement Committee has developed the following standards regarding the examination of members with diabetes:

- The history should include the name and, if available, contact information of the Primary Care Physician ("PCP"), or the provider managing the diabetes.
- List of all diabetes medications.
- The HA1c should be documented in the chart - this may come from the patient, a lab report, or the PCP.
- Dilation is required. All common eye changes that result from diabetes should be documented in the medical record. These include, but are not limited to: retinopathy, dry eye, blepharitis, cataract, and low tension glaucoma.
- The retina examination must be detailed and subtle background changes should be noted.
- Education and counseling about blood sugar control and the required numbers to prevent vision loss should be emphasized.
- **It is highly recommended to communicate and coordinate with the PCP** - Send a full report of the dilated eye examination results for all patients with diabetes to the PCP and/or diabetes provider. The assigned PCP is noted on the front of the member's ID card. Additionally, you may contact the Health Plan directly or the PCP for assistance in coordinating additional medical needs for the member, as identified while providing vision services.

- Include the correct coding for retinopathy on your claim; the appropriate ICD-10 code if positive or CPT II code 3072F if negative.

Medical record documentation will be evaluated based on these standards, when they are reviewed as part of a quality audit.

Clinical Criteria*

The state specific criteria in the Provider Reference Guide (PRG) outline the benefits according to the member's plan. This chart is not an indication that the member has a specific benefit. Rather this chart is used to define the medically necessary indications when the PRG indicates that the benefit is available to a member and when no regulatory/client criteria is available.

Benefit	Available When	Clinical Criteria
Eyewear After Eye Surgery	Determined to be medically necessary.	The stable refractive prescription changes are more than +/-0.75 diopters in any meridian or more than 20 degrees of axis shift or a change in add power greater than 0.50 diopters.
Oversize Lens	Needed for physiological reasons.	The pupillary distance is 70mm or greater or other facial or ocular anomalies requiring a large lens.
Trifocal Lens	Member has a special need due to a job training program or extenuating circumstances.	The base prescription is greater than +/- 1.00 and a bifocal greater than or equal to 2.00
Necessary Contact Lens	Such lenses provide better management of a visual or ocular condition than can be achieved with spectacle lenses, including, but not limited to the diagnosis of:	Irregular astigmatism; unilateral aphakia; keratoconus when vision with glasses is less than 20/40; corneal transplant when vision with glasses is less than 20/40 or anisometropia that is greater than or equal to 4.00 diopter
Color Tinting	Light sensitivity which will hinder driving or seriously handicap the outdoor activity of such member is evident.	The member has photophobia, aniridia, uveitis, corneal dystrophy, cataracts, albinism, or use a medication that has a side effect of photophobia.
Single Vision Eyeglasses In Lieu Of Bifocals	Need is substantiated in member's medical record by clinical data.	The need for distance correction > +/- 1.50 diopter AND Net combination of distance RX and bifocal > +1.00 or -2.00 AND you are unable to tolerate a multifocal lens.
Progressive Lenses	Need is substantiated in member's medical record by clinical data.	Epilepsy, childhood disorders with multiple impairments.
Transitions Lenses	Need is substantiated in member's medical record by clinical data.	Chronic iritis or uveitis, albinism.
Polycarbonate Lenses	Need is substantiated in member's medical record by clinical data.	<ul style="list-style-type: none"> ▪ The member has a prescription of +/-8.00; or ▪ Permanently reduced vision in one eye to less than 20/60; or ▪ A facial deformity or disease that interferes with eye glass fit; or ▪ A documented occupational hazard.
Ultra Violet Coating	Need is substantiated in member's medical record by clinical data	<ul style="list-style-type: none"> ▪ Provided to members with aphakia, albinism, members that have clinical evidence of macular degeneration, or are taking medicine that makes them more sensitive to ultra violet light.
Replacement Due To Outgrown Glasses	Need is substantiated in member's medical record by clinical data	<ul style="list-style-type: none"> ▪ Available for children under 18 when the member's pupil distance is wider than the frame's mechanical optical center by greater than 5mm. ▪ Available when the new frame size is at least 3mm larger than the existing frames.
Second Opinion Examination	Need is substantiated in member's medical record by clinical data	<ul style="list-style-type: none"> ▪ Available when medical chart review of the first examination shows inadequate examination, documentation, or when clinical issues are not adequately addressed.

High Index lenses (Higher than Polycarbonate)	Need is substantiated in member's medical record by clinical data	<ul style="list-style-type: none"> ▪ Available when weight of a standard prescription could cause facial development issues (primarily for children). ▪ Available when lab cannot practically produce lenses with a lower index lens.
Allergy To Certain Frames	Need is substantiated in member's medical record by clinical data	<ul style="list-style-type: none"> ▪ Alternative frame to be provided when a provider documents a rash or other adverse reaction to all MARCH frame kit materials.
SLAB Off/Prism	Need is substantiated in member's medical record by clinical data	<ul style="list-style-type: none"> ▪ Available for bifocal or trifocal prescriptions that generate greater than 2 prism diopters of imbalance at the reading plane.
Safety Frames	Need is substantiated in member's medical record by clinical data	<ul style="list-style-type: none"> ▪ Used with polycarbonate lenses based on polycarbonate criteria noted above; and ▪ Member is in and around a hazardous environment where, in the discretion of the patient, (parent) and the provider, extra ocular safety measures are required ▪ These would be considered "deluxe frames" and covered by MARCH. ▪ These must meet ANSI standards.
Non-Standard Frames	Need is substantiated in member's medical record by clinical data	<ul style="list-style-type: none"> ▪ Used when member has facial parameters where standard frames do not fit correctly. ▪ Used when optical correction will not fit practically in a standard frame.
Low Vision Rehabilitation	Need is substantiated in member's medical record by clinical data.	<ul style="list-style-type: none"> ▪ Visual loss with best corrected visual acuity of 20/50 or worse in the better eye. ▪ Constriction of visual fields to be less than 20 degrees or hemianopia. ▪ Limited contrast sensitivity due to underlying pathology. ▪ Initial consult codes of 97241 – 97245 or 99244. ▪ Maximized medical treatment of conditions such as, but not limited to, diabetic retinopathy, macular degeneration, optic atrophy, and glaucoma. ▪ Diagnosis codes consistent with low vision pathology. Under certain circumstances, medical records may be requested. If requested, they need to demonstrate that medical, surgical, and other treatments that have been tried and failed. They must have a diagnosis as noted below AND reduced vision. The appropriate diagnosis codes are necessary, including, but not limited to: <ul style="list-style-type: none"> ▪ D49.81 ▪ G.35 ▪ H47.099 ▪ H33.08-H33-303 ▪ E11.319, E10,319 ; H35.00-H35.443 ▪ H40.001-H40-2234 ▪ H53.40-H53-483 ▪ H54.2-H54.60 ▪ H46.00-H47.333 ▪ H55.00-H55.01 ▪ Or others by pre-approval ▪ A Low Vision Rehabilitation request form must be completed and submitted. ▪ Before proceeding, prior approval is required.

Dilation of Eyes	Initial examination required. Subsequent examinations as follows:	<ul style="list-style-type: none"> All new members require a dilated fundus exam or equivalent (if acceptable per state/federal regulation). Diabetics require dilation every year at a minimum, more often if they have retinopathy. Members with other certain pathology such as lattice degeneration, choroidal nevi, or retinoschisis for example, may also need a dilated exam every year or as medically indicated. Members with no risk factors should be dilated thereafter based on the professional judgment of the provider or every three (3) years, whichever occurs first.
Polarized Lenses	Need is substantiated in member's medical record by clinical data.	<ul style="list-style-type: none"> Chronic iritis, uveitis, or other active inflammatory eye disease with fixed and dilated pupils or aniridia.
Necessary Contact Lens Replacement	Such lenses provide better management of a visual or ocular condition than can be achieved with spectacle lenses (see criteria above).	<ul style="list-style-type: none"> The member meets criteria as noted above for necessary contact lens and there is: <ul style="list-style-type: none"> -Change of +/- 1.00 diopter in power -Change of 0.50 mm in base curve -Change of 0.30 mm in optic zone -Change of 0.75 mm in peripheral curve radius -Change of 0.30 mm in peripheral curve width
Replacement Glasses When A Member Can Not Adapt To Bifocals	Member has presbyopia and unable to adapt to bifocals.	<ul style="list-style-type: none"> Members should attempt to make the adjustment to bifocal lenses for a minimum of two (2) weeks. When lens manufacturers and/or the laboratory provides a warranty for "non-adapts", this should be used. When two pairs of glasses is the solution, each pair must have a sphere power of at least +/- 1.00 or a cylinder power of greater than +/-0.75 in at least one eye. In cases where one of the final single vision Rx calculation yield lower powers, the member will just be entitled to distance only or near distance only glasses. The frame that was used for the bifocals will be reused for one of the new single vision glasses.
Medically Necessary Contact Lenses And Glasses For Aphakia In Children Aged 2 Weeks To 12 Years.	Post surgically, for children born with a visually significant Cataract(s), or other medical eye problems that result in pediatric aphakia.	<p>Coverage for either medically necessary contact lenses or glasses in a given benefit period, but not both except for the following circumstances:</p> <ul style="list-style-type: none"> The patient has greater than three (3) diopters of astigmatism in one or both eyes and requires this correction over the contact lens or lenses. The patient has vision less than 20/200 in the poorer eye, or pathology where 20/200 or less is expected but can not be measured (ie. PHPV, RD, macula scarring, coloboma involving the posterior pole) and a spectacle lens is needed for protection of the good eye.
Prescription/ Fitting check	Glasses are dispensed, including when a member has ongoing vision issues using new materials	<ul style="list-style-type: none"> Included in the fitting fee/payment for materials for up to 45 days after member has received materials.
Eye Care of Patient with Diabetes Mellitus	Person has Diabetes Mellitus	<ul style="list-style-type: none"> MARCH adopted the American Optometric Association ("AOA") Guidelines for treating Diabetes Mellitus".

* QIC approval 2/24/2017

Sending a Secure Email to MARCH[®] Vision Care for PHI Related Data

NOTE:

This document is technical in nature and will require expertise in understanding the workings of the Microsoft Exchange Server Infrastructure. The information provided in this document can be used by your IT administrator to implement secure email transmission with MARCH[®] Vision Care. For any support questions please call Microsoft Support for more details.

The below details are from Microsoft TechNet Article on Secure Your E-mail Traffic

Secure Your E-Mail Traffic

As part of establishing e-mail coexistence between your local Microsoft Exchange Server environments, we recommend that you implement Transport Layer Security (TLS) send and receive capability in your local Exchange Server environment. This is necessary because, during coexistence with Exchange Online, e-mail that was previously sent and received within your organization will now be sent over the Internet. The instructions in this section describe how to secure e-mail traffic on Microsoft Exchange 2000 Server and Exchange Server 2003 and Exchange Server 2007.

To secure your e-mail traffic with TLS, you will require a certificate that is granted by a recognized certification authority (CA). To implement TLS in your local Exchange Server environment, you are required to:

1. Identify the Exchange Server on which to install the certificate.
2. Generate a certificate request.
3. Acquire the certificate.
4. Install the certificate.
5. Create a Simple Mail Transfer Protocol (SMTP) connector.
6. Enable TLS.

Step 1: Identify the Exchange Server on Which to Install the Certificate

TLS should be enabled on the bridgehead server of your local Exchange Server environment. That is the computer that directs your organization's e-mail to and from the Internet. For more information about bridgehead servers and Exchange Server message routing, see [Exchange Server 2003 Message Routing Topology](#).

If you have separate bridgehead servers for sending and receiving e-mail from the Internet, you will need to acquire and install a certificate on the SMTP server of each bridgehead server computer running Exchange Server; however, you will need to set up a connector and enable TLS only on the server that is used for sending e-mail to the Internet.

NOTE:

- If your Exchange Server environment relies on an external relay server to send and receive e-mail to and from the Internet, you will need to contact the administrator of the external service about their TLS support. When TLS has been enabled on the external service, secure e-mail will flow between their relay server and Microsoft Online Services.
- If you have third-party bridgehead software or service, refer to that documentation to see how you can configure TLS.

If you have a local Exchange Server bridgehead server running the standard SMTP virtual server, continue reading this topic.

Step 2: Generate a Certificate Request

Use the Exchange System Manager in Exchange Server to generate a certificate request on your bridgehead server. You must provide the fully qualified domain name (FQDN) of the bridgehead server. For more information, see [Creating a Certificate or Certificate Request for TLS](#).

Step 3: Acquire the Certificate

Locate a recognized certification authority (CA), such as VeriSign, Comodo, or GoDaddy. Submit the certificate request file that you generated in the previous section. The CA will provide you with a certificate (CER) file that contains the certificate for your server.

Step 4: Install the Certificate

Use the Exchange System Manager to install the certificate file. You must provide the path to the certificate file that you received from the CA.

Step 5: Create an SMTP Connector

Based on your current e-mail environment, use one of the following procedures to create an SMTP connector or Send connector.

To create an SMTP connector in Exchange 2000 or Exchange 2003

1. In Exchange System Manager, right-click **Connectors**, and then click **New SMTP Connector**.
2. Type a name for the connector (for example, MicrosoftOnline).
3. On the **General** tab, select **Forward all e-mail through this connector to the following smart host**, and then type **mail.global.frontbridge.com**.

IMPORTANT: When you use the URL **mail.global.frontbridge.com**, e-mail messages are routed through servers to follow a path that balances the network load efficiently. If you want e-mail messages to be routed through servers in the United States instead of being routed through servers that might be located in other countries, type the following URL: **mail.us.messaging.microsoft.com**.

4. Under **Local Bridgeheads**, click **Add**, and then select your bridgehead server computer running Exchange Server.
5. On the **Address Space** tab, click **Add**, and then type your organization's Microsoft Online Services e-mail routing domain (for example, contoso1.microsoftonline.com).

For more information about creating SMTP connectors, see [How to configure the SMTP connector in Exchange 200x](#).

To create a Send connector in Exchange 2007

1. Open the Exchange Management Console, and then do one of the following:
 - On the computer that has the Edge Transport server role installed, select **Edge Transport**, and then, in the work pane, click the **Send Connectors** tab.
 - On the computer with the Hub Transport server role installed, in the console tree, expand **Organization Configuration**, select **Hub Transport**, and then, in the work pane, click the **Send Connectors** tab.
2. In the action pane, click **New Send connector**. The new SMTP Send Connector wizard starts.
3. On the **Introduction** page, do the following:
 - In the **Name** field, type a meaningful name for the connector (for example, type MicrosoftOnlineServices)
 - In the **Select the intended use for this Send connector** field, select **Internet**, and then click **Next**.
4. On the **Address Space** page, click **Add**.
5. In the **Add Address Space** dialog box, in the **Address** field, type your organization's Microsoft Online Services e-mail routing domain (for example, contoso1.microsoftonline.com), and then click **OK**.
6. On the **Address Space** page, click **Next**.
7. On the **Network Settings** page, select **Route all mail through the following smart hosts**, and then click **Add**.
8. In the **Add Smart Host** dialog box, select **Fully qualified domain name (FQDN)**, type **mail.global.frontbridge.com**, and then click **OK**.

IMPORTANT: When you provide the URL **mail.global.frontbridge.com**, e-mail messages are routed through servers to follow a path that balances the network load efficiently. If you want e-mail messages to be routed through servers in the United States instead of being routed through servers that might be located in other countries, type the following URL: **mail.us.messaging.microsoft.com**.

9. On the **Network Settings** page, click **Next**.
10. On the **Configure Smart host authentication settings** page, select **None**, and then click **Next**.

The Source Server page appears only on a computer with the Hub Transport server role installed. By default, the Hub Transport server that you are currently working on is listed as a source server.

11. To add a source server, click **Add**.
12. In the **Select Hub Transport and subscribed Edge Transport servers** dialog box, select one or more Hub Transport servers in your organization, and then click **OK**.

Step 6: Enable TLS

After you install the certificate, your server will be able to receive TLS e-mail. However, it cannot send TLS e-mail until you enable TLS.

To enable TLS

1. In Exchange System Manager, expand **Connectors** and locate the MicrosoftOnline connector that you created in the previous procedure.
2. Right-click the connector and then click **Properties**.
3. On the **Advanced** tab, click **Outbound Security**, and then select **TLS Encryption**.

Eye Examination Record MARCH[®] Vision Care

Patient Name:

DOS:

	Last name	First Name	Middle Initial
Date of Birth:			Patient ID:
Reason for Visit <small>(Chief Complaint/ Concern)</small>			
Medical History			
Eye History			Date of last DFE
Family Medical and Eye History			
Allergies:			
Current Medicines:			
Social History:	Tobacco:		Alcohol:
Orientation /Mood	Oriented to time and place:	Normal	Abnormal
	Mood or Affect:	Normal	Abnormal
Comments:			
Physical Findings:	BP:	Pulse:	Height: Weight: BMI:

Review of Systems

System	Findings	Problem:
Constitution	Neg	
Ear/Nose/Throat	Neg	
Neurological	Neg	
Psychological	Neg	
Cardiovascular	Neg	
Respiratory	Neg	
Gastrointestinal	Neg	
Genital urinary	Neg	
Muscular-Skeletal	Neg	
Integument	Neg	
Endocrine	Neg	
Hematology/Lymphatic	Neg	
Allergy/Immunology	Neg	

Vision:

Vcc: Distance	R: 20/	L: 20/	Both: 20/
Vcc: Near	R: 20/	L: 20/	Both: 20/
Vsc: Distance	R: 20/	L: 20/	Both: 20/
Vsc: Near	R: 20/	L: 20/	Both: 20/

Current RX:	OD	OS
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External Exam:

Pupils:		
Cover:	Distance	Near
Motility:		
Confrontation Fields:	OD	OS
Keratometry/Topo:	OD	OS
Color Vision:	OD	OS
Depth Perception:		

Refractions:

Auto: OD	20/	OS	20/
Static: OD	20/	OS	20/
Dry: OD	20/	OS	20/
Wet: OD	20/	OS	20/

Patient Name: _____ **DOS:** _____
Last name First Name Middle Initial

Near Testing:	Add:
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Slit Lamp Examination:

Lids/ Lashes/Adnexa:	OD	OS
Cornea:	OD	OS
Conjunctiva:	OD	OS
AC:	OD	OS
Iris:	OD	OS
Lens:	OD	OS

Intra Ocular Pressure

OD	OS	Time:
Method: AP	Puff	Tono FT

Gonioscopy:	OD	OS
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Medicines: Prop	Tetra	Fluress	NaF	Myd	Paradryn	Cyclo	Other:
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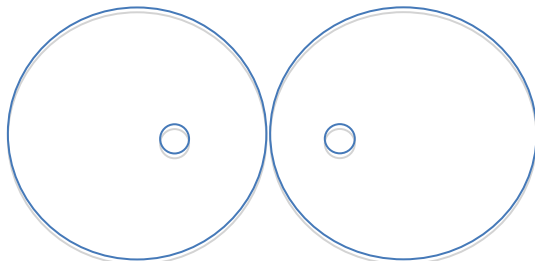
Fundus:

Direct	Indirect	Slit Lamp Lens	Photo
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Nerve:

C/D:	OD	OS
Rim:	OD	OS
Color:	OD	OS
Comments:		

Macula:	OD	OS
Post Pole:	OD	OS
Vessels:	OD	OS
Vitreous:	OD	OS
Rim:	OD	OS
Periphery:	OD	OS



Diagnosis Impression:		
Assessment:		
Management Plan:		
I have personally reviewed this medical record including the patient's health history.	Signature:	Date:
		Return:

Performance Measurement & Reporting

In an effort to improve HEDIS and Star Ratings performance, MARCH[®] Vision Care requires providers to submit CPT II* and ICD-10 codes, on claims, to demonstrate performance and diagnosis of the following for diabetic members:

- Retinal or dilated eye exams
- Negative retinal or dilated eye exams
- Diabetes
- Diabetic retinopathy

CPT II codes - Retinal or dilated eye exam

Code	Description
2022F	Dilated retinal eye exam with interpretation by an ophthalmologist or optometrist documented and reviewed.
2024F	7 standard field stereoscopic photos with interpretation by an ophthalmologist or optometrist documented and reviewed.
2026F	Eye imaging validated to match diagnosis from 7 standard field stereoscopic photos results documented and reviewed.
3072F	Low risk for retinopathy (no evidence of retinopathy in the prior year)

NEGATIVE or low risk for retinopathy:

Please always include two CPT II codes when the patient is negative or low risk for retinopathy, one from the 2022F - 2026F range PLUS 3072F.

ICD-10 codes - Diabetic retinopathy

Diagnosis	Type 1	Type 2
Unspecified DR with DME	E10.311	E11.311
Unspecified DR without DME	E10.319	E11.319
Mild NPDR without DME	E10.3299**	E11.3299**
Moderate NPDR without DME	E10.3399**	E11.3399**
Severe NPDR without DME	E10.3499**	E11.3499**
PDR without DME	E10.3599**	E11.3599**

POSITIVE for retinopathy:

Please always include one CPT II code from the 2022F-2026F range PLUS the appropriate ICD-10 code.

*CPT II codes are tracking codes used for performance measurement. They should be billed in the CPT/HCPCS field of CMS-1500 forms and submitted on the same claim as the CPT I code(s). CPT II codes do not have relative value and can be billed with a \$0.00 charge amount.

**Please note that the 7th character (9) indicates unspecified eye. Providers should use the correct digit to indicate which eye the condition applies to, or bilateral, if known.

IMPORTANT:

- Always bill the appropriate ICD-10 diagnosis code when submitting your claim. In particular, please include any medical diagnosis codes including, but not limited to, diabetes at the highest level of specificity.
- A patient's medical record should always support the CPT I, CPT II and ICD-10 codes billed.

Diabetes ICD-10 codes commonly billed by optometrists and ophthalmologists

Diagnosis	Type 1	Type 2
Unspecified DR with DME	E10.311	E11.311
Unspecified DR without DME	E10.319	E11.319
With diabetic cataract	E10.36	E11.36
With hyperglycemia	E10.65	E11.65
With other diabetic arthropathy	E10.618	E11.618
With other diabetic ophthalmic complication	E10.39	E11.39
With other specified complication	E10.69	E11.69
With unspecified complications	E10.8	E11.8
Without complications		E11.9

Normal billing rules still apply. The requirements listed in this document should be included in your billing process.