Costs and Limitations

For Certified Healthcare IT EHR

EyeDoc Version 10.1

06/2020

Penn Medical Informatics Systems, Inc

A Division of Eye Care Leaders, LLC

Costs and Limitations for EyeDoc EMR Version 10.1

Capability and Description

2015 Edition criteria applicable to EyeDoc EMR Version 10.1: a1, a2, a3, a5, a9, a12, a13, a14, b1, b2, b9, d1, d2, d3, d4, d5, d6, d7, d8, g3, g4, g5, g6

EyeDoc EMR Version 10.1 is an EHR that supports healthcare professionals in ophthalmology and optometry in outpatient ambulatory environments. . It allows users to perform a wide range of functions such as to document, review, and edit patient health information including but not limited to problem lists, medication lists, medication allergy lists, family health history, and all aspects of the patient's eye exam. Perform CPOE (computerized provider order entry) for medications, laboratory orders and imaging procedures

EyeDoc EMR Version 10.1 requires integration of third-party components to perform the following tasks (fees may apply):

- electronically create prescriptions and prescription-related information for electronic transmission to pharmacies,
- measure CQMs (clinical quality measures) and to export these in standard file formats,
- be alerted to possible CDS (clinical decision support) interventions, and
- Report information to PHAs (public health agencies) and clinical data registries.
- Portal related activities

Types of Costs or Fees and Additional Types of Costs or Fees

EyeDoc EMR Version 10.1 solution includes one-time software license and implementation / setup fees and a monthly subscription fee (which includes support and upgrades). These fees are determined based on the number of providers and additional fees are required to increase the number of providers.

Onsite training is available for separate per day fees.

E-Prescribing, Secure Mail and Patient Portal are an additional cost in the EyeDoc EMR V10.1 for all clients

Additional software may be required dependent upon any other installed programs apart from The EyeDoc EMR Version 10.1 (i.e., Peripheral and/or Imaging device interfaces such as Visual Field, Automatic Refractors, Corneal Topography, etc). Separate fees may apply depending on the software or device manufacturer/vendor. The PennMedical Fee for device integration is based on whether the peripheral or imaging device is Screening (data only) or Imaging (Data + Images). Fee includes a one-time implementation charge and an annual maintenance fee

American Academy of Ophthalmology (AAO) patient education documents are available as an annual subscription. Subscription Fees apply and are subject to automatic annual renewal.

Clinical Decision Support documentation via InfoButton technology is available upon request. Subscription Fees apply and are subject to automatic annual renewal

Penn Medical does not charge any additional fees for electronic integration with third-party systems such as public health registries, clinical data registries, HIEs, ACOs or CINs, assuming such electronic integration is based on an established technical capability of EyeDoc EMR Version 10.1 or the Regulatory Compliance Platform

Limitations (Contractual/Business)

Base contractual obligation is for 1 years. Automatic annual renewal thereafter.

This Health IT Module is 2015 Edition compliant and has been certified by an ONC-ACB in accordance with the applicable certification criteria adopted by the Secretary of the U.S. Department of Health and Human Services. This certification does not represent an endorsement by the U.S. Department of Health and Human Services.

Vendor	Version	Date Certified	Certification Number
Penn Medical Informatics Systems, Inc	EyeDoc EMR Version 10.1	Dec 27, 2017	15.04.04.2150.EyeD.10.00.1.171227

Criteria Certified

- 170.315 (a)(1) CPOE - Medications
- 170.315 (a)(2) CPOE - Laboratory
- 170.315 (a)(3) CPOE-Diagnostic Imaging
- 170.315 (a)(5) Demographics
- 170.315 (a)(9) Clinical Decision Support
- 170.315 (a)(12) Family Health History
- 170.315 (a)(13) Patient-Specific Education Resources
- 170.315 (a)(14) Implantable device list
- 170.315 (b)(1) Transitions of Care
- 170.315 (b)(2) Clinical Information Reconciliation and Incorporation
- 170.315 (b)(6) Data export
- 170.315 (b)(9) Care Plan
- 170.315 (d)(1) Authentication, Access Control and Authorization
- 170.315 (d)(2) Auditable Events and Tamper-resistance
- 170.315 (d)(3) Audit Report(s)
- 170.315 (d)(4) Amendments
- 170.315 (d)(5) Automatic Log-off
- 170.315 (d)(6) Emergency Access
- 170.315 (d)(7) End-user Device Encryption
- 170.315 (d)(8) Integrity
- 170.315 (g)(3) Safety-enhanced Design
- 170.315 (g)(4) Quality Management System
- 170.315 (g)(5) Accessibility- Centered Design
- 170.315 (g)(6) Consolidated CDA Creation Performance

Additional Software for Demonstration

Windows Professional, Microsoft SQL 2005 or Newer, Windows Server, Regulatory Compliance Platform Version 1.2

END OF DOCUMENT