Costs and Limitations

For Certified Healthcare IT EHR

ManagementPlus Version

07/21/2017

Costs and Limitations for ManagementPlus 6.0

Capability and Description

2015 Edition criteria applicable to ManagementPlus 6.0: a1, a2, a3, a5, a6, a9, a11, a12, a13, a14, b1, b2, b6, b9, d1, d2, d3, d4, d5, d6, d7, d8, g3, g4, g5, g6

ManagementPlus 6.0 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,
- 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.

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

In order to achieve MU, the client will need to purchase eRX and Secure mail solutions.
eRx is provided by DrFirst, a one-time credential fee of $75/doctor as well as a $510/doctor annual fee.
Secure E-mail is provided by Secure Exchange for a fee of $200/doctor annually.

Limitations (Contractual/Business)

We do not impose any limitations on the implementation or use of our certified software or those 3rd party software integrated to achieve Meaningful Use
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Limitations (Technical/Practical)

The Regulatory Compliance Platform Version 1.2 is used as part of an EHR solution package. It allows users to perform a wide range of functions such as monitor MIPS Performance, CQM calculation, PHA reporting, Bulk Export of CCD files, Send/receive messages and attached documents to/from the HISP via Direct Edge Protocol, create gold standard documents required for referrals and transitions of care

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
ManagementPlus	7.0	Dec 31, 2017	15.04.04.1864.Mana.07.00.1.171231

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)(6) Problem List
- 170.315 (a)(9) Clinical Decision Support
- 170.315 (a)(11) Smoking Status
- 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

Meinberg NTP, Dr First, MedLine Plus, LDM, Secure Exchange Solutions, CQM Solution, Regulatory Compliance Platform Version 1.2

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