



Eligible Professional Meaningful Use Menu Set Measures Measure 3 of 10

Stage 1

Date issued: November 7, 2010

| Patient Lists | |
|---------------|--|
| Objective | Generate lists of patients by specific conditions to use for quality improvement, reduction of disparities, research, or outreach. |
| Measure | Generate at least one report listing patients of the EP with a specific condition. |
| Exclusion | No exclusion. |

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Definition of Terms

Specific Conditions -- Those conditions listed in the active patient problem list.

Attestation Requirements

YES / NO

Eligible professionals (EPs) must attest YES to having generated at least one report listing patients of the EP with a specific condition to meet this measure.

Additional Information

- This objective does not dictate the report(s) which must be generated. An EP is best positioned to determine which reports are most useful to their care efforts.
- The report generated could cover every patient whose records are maintained using certified EHR technology or a subset of those patients at the discretion of the EP.
- The report generated is required to include only patients whose records are maintained using certified EHR technology.

Related Meaningful Use FAQs

To see the FAQs, click the New ID # hyperlinks below, or visit the CMS FAQ web page at <https://questions.cms.gov/> and enter the New ID # into the Search Box, clicking the “FAQ #” option to view the answer to the FAQ. (Or you can enter the OLD # into the Search Box and click the “Text” option.)

How should EPs select menu objectives? [New ID #2903](#), [Old ID #10162](#)

Certification and Standards Criteria

Below is the corresponding certification and standards criteria for electronic health record technology that supports achieving the meaningful use of this objective.

| Certification Criteria | |
|---------------------------------------|---|
| §170.302(i) Generate patient lists | Enable a user to electronically select, sort, retrieve, and generate lists of patients according to, at a minimum, the data elements included in: (1) Problem list; (2) Medication list; (3) Demographics; and (4) Laboratory test results. |
| Standards Criteria | |
| | N/A |

Related Certification FAQs

Click on the green numbers to view the answer to the FAQ.

- I'm in the process of implementing EHR technology developer XYZ's certified Complete EHR [or certified EHR Module] "E-HealthSystem2010."
Scenario 1: Can I reconfigure E-HealthSystem2010 without compromising the certified status of my implementation of E-HealthSystem2010?
Scenario 2: EHR technology developer XYZ communicated to my organization that they relied upon a 3rd party software program "PatientInfoTracker 2.0" for the purposes of demonstrating compliance with the "generate patient lists" certification criterion specified at 45 CFR 170.302(i) in achieving E-HealthSystem2010's certification. I have already implemented, use, and would like to continue using "SuperListGenerator 7.0." I have determined that I can reconfigure SuperListGenerator 7.0 to work with E-HealthSystem2010. Can I use SuperListGenerator 7.0 in lieu of PatientInfoTracker 2.0 without compromising the certified status of my implementation of E-HealthSystem2010? [9-10-016-1](#)



- Is an EP limited to demonstrating meaningful use in the exact way that EHR technology was tested and certified? Similarly, if the EHR technology was tested and certified with certain clinical decision support rules, are those the only clinical decision support rules an eligible health care provider is permitted to use when demonstrating meaningful use? [3-11-024-1](#)