

3101	How will eligible professionals (EPs) be required to show that they are meeting the Medicaid or needy individual patient volume thresholds of 30% for the Medicaid EHR Incentive Program?	To show that EPs are meeting the Medicaid or needy individual patient volume thresholds of 30% for the Medicaid EHR Incentive Program, States will need to propose one or more methods of calculating patient volume to CMS in their State Medicaid Health Information Technology Plans and would need to identify verifiable data sources available to the provider and/or the State. Please contact your State Medicaid Agency for more information on how your state is calculating patient volume. For more information about the Medicare and Medicaid EHR Incentive Program, please visit http://www.cms.gov/EHRIncentivePrograms Keywords: FAQ10523
3107	Are the criteria for needy patient volumes under the Medicaid EHR Incentive Program only applied to eligible professionals (EPs) practicing predominantly in Federally Qualified Health Centers (FQHCs) and/or Rural Health Clinics (RHCs), or can they also apply to hospital patient volumes?	Criteria for minimum patient volumes attributable to needy individuals apply only to EPs practicing predominantly in an FQHC or RHC. These criteria do not apply to hospital patient volumes. For more information about the Medicare and Medicaid EHR Incentive Program, please visit http://www.cms.gov/EHRIncentivePrograms Keywords: FAQ10526
3121	If a State proposes a new definition for meaningful use under its Medicaid EHR Incentive Program, will it need to include the new definition of meaningful use in its State Medicaid Health Information Technology Plan (SMHP)? When are the SMHPs due?	Yes, if a State wishes to request flexibility with the definition of meaningful use, to the extent permissible under the Medicare and Medicaid EHR Incentive Programs final rule, it would do so via its SMHP. There is no due date for SMHPs. States are implementing their Medicaid EHR Incentive Programs on a rolling basis. The SMHPs are therefore expected to be iterative, as States implement their programs incrementally, especially in the early years. For more information about the Medicare and Medicaid EHR Incentive Program, please visit http://www.cms.gov/EHRIncentiveProgram words: FAQ10533
3123	If a State has a team of staff members who will be administering the Medicaid EHR Incentive Program from 2011-2021 (answering provider questions, engaging in reporting and analysis, assisting providers with eligibility and verifying provider eligibility, appeals, etc.), would there be 90% Federal Financial Participation for this team on an ongoing basis once approval is received from CMS on State Medicaid Health Information Technology Plan and the Health Information Technology Implementation Advance Planning Document?	Yes. However, if state staff members are not working full-time on the Medicaid EHR Incentive Program, their salaries need to be cost-allocated appropriately. For more information about the Medicare and Medicaid EHR Incentive Program, please visit http://www.cms.gov/EHRIncentivePrograms Keywords: FAQ10534
3373	How does CMS define pediatrician for purposes of the Medicaid EHR Incentive Program?	CMS does not define pediatrician for this program. Pediatricians have special eligibility and payment flexibilities offered under the program and it is up to States to define pediatrician, consistent with other areas of their Medicaid programs. You can find your State's contact information http://www.cms.gov/apps/files/statecontacts.pdf here For more information about the Medicare and Medicaid EHR Incentive Program, please visit http://www.cms.gov/EHRIncentivePrograms Keywords: FAQ10715
5995	For the Medicaid EHR Incentive Program, how do we determine Medicaid patient volume for procedures that are billed globally, such as obstetrician (OB) visits or some surgeries? Such procedures are billed to Medicaid at a global rate where one global rate might cover several visits.	CMS leaves it up to the states how to operationalize the patient volume considerations of global payments with the following guidance: the numerator and denominator must be incorporated consistently. The total encounters can be kept global, or broken down into individual visits. If a global payment is broken down into separate visits in the numerator, then for purposes of the denominator, the state must break down any other global payments received from other payers. We recognize this could be administratively challenging and are open to reviewing strategies for doing this that may involve sampling (e.g., if the Medicaid global payment for OB averages 12 visits, we would expect to see the numerator expanded to 12 visits for Medicaid encounters, and a denominator constructed using sample data from a random file review that similarly breaks down any global payments into separate visits for Medicaid and non-Medicaid payers). Additionally, if the state's approach to global payments excludes providers from the Medicaid EHR Incentive Program who would otherwise be eligible, the state must create a mechanism to re-review their eligibility. Keywords: FAQ10957
2817	The billing provider on a claim is an eligible professional (EP) but the performing provider type is not an EP. If we use claims to validate patient volume or meaningful use for the Medicaid Electronic Health Record (EHR) Incentive Program, should we count performing providers (person rendering the service) or the billing provider?	In establishing an encounter for purposes of patient volume, please see the regulations at 495.306(e)(2)(i)-(ii) at 75 FR 44579. Furthermore, in estimating patient volume for any EP or hospital, we do not specify any requirements around billing, but rather we discuss patients. For example, if a physician's assistant (PA) provides services, but they are billed through the supervising physician, it seems reasonable that a State has the discretion to consider the patient as part of the patient volume for both professionals. However, this policy would need to be applied consistently. In this scenario, using services provided by the PA but billed under the physician in the physician's numerator (e.g., Medicaid encounters) also would increase the physician's denominator (all encounters), because the State would need to adequately reflect the total universe of patients (both Medicaid and non-Medicaid) who the PA saw, but for whom the physician billed. In terms of meaningful use, because each eligible professional must demonstrate meaningful use of certified EHR technology him or herself, if the State cannot not distinguish between the physician's claims and the PA's individual claims, then this would not be an adequate audit methodology. To view the final rule, please visit: http://edocket.access.gpo.gov/2010/pdf/2010-17207.pdf For more information about the Medicare and Medicaid EHR Incentive Program, please visit http://www.cms.gov/EHRIncentivePrograms Keywords: FAQ10098
3015	When eligible professionals work at more than one clinical site of practice, are they required to use data from all sites of practice to support their demonstration of meaningful use and the minimum patient volume thresholds for the Medicaid EHR Incentive Program?	CMS considers these two separate, but related issues. Meaningful use: Any eligible professional demonstrating meaningful use must have at least 50% of their patient encounters during the EHR reporting period at a practice/location or practices/locations equipped with certified EHR technology capable of meeting all of the meaningful use objectives. Therefore, States should collect information on meaningful users' practice locations in order to validate this requirement in an audit. Patient volume: Eligible professionals may choose one (or more) clinical sites of practice in order to calculate their patient volume. This calculation does not need to be across all of an eligible professional's sites of practice. However, at least one of the locations where the eligible professional is adopting or meaningfully using certified EHR technology should be included in the patient volume. In other words, if an eligible professional practices in two locations, one with certified EHR technology and one without, the eligible professional should include the patient volume at least at the site that includes the certified EHR technology. When making an individual patient volume calculation (i.e., not using the group/clinic proxy option), a professional may calculate across all practice sites, or just at the one site. For more information on applying the group/clinic proxy option, see FAQ #10362 or here For more information about the Medicare and Medicaid EHR Incentive Program, please visit http://www.cms.gov/EHRIncentivePrograms Keywords: FAQ10416
3017	Can tribal clinics be treated as Federally Qualified Health Centers (FQHCs) for the Medicaid Electronic Health Record (EHR) Incentive Program?	CMS previously issued guidance stating that health care facilities owned and operated by American Indian and Alaska Native tribes and tribal organizations ("tribal clinics") with funding authorized by the Indian Self-Determination and Education Assistance Act (Public Law 93-638, as amended) must be reimbursed as FQHCs in order to be considered FQHCs in the Medicaid EHR Incentive Program. CMS revised this policy and will allow any such tribal clinics, as well as urban clinics that are funded by urban Indian organizations receiving funds under title V of the Indian Health Care Improvement Act (Public Law 94-437, as amended) for the provision of primary health services http://www.ihms.gov/ihm/index.cfm?module=dsp_ihm_pc_p3c19#3-19.2D to be considered as FQHCs for the Medicaid EHR Incentive Program, regardless of their reimbursement arrangements. For more information on how FQHCs are defined, please see https://questions.cms.gov/faq.php?id=5005&faqId=3017 For more information about the Medicare and Medicaid EHR Incentive Program, please visit http://www.cms.gov/EHRIncentivePrograms Keywords: FAQ10417
3099	Do Federally Qualified Health Center (FQHC) sites have to meet the 30% minimum Medicaid patient volume threshold to receive payment under the Medicaid EHR Incentive Program?	Eligible professionals may participate in the Medicaid EHR Incentive Program if: 1) They meet Medicaid patient volume thresholds; or 2) They practice predominantly in an FQHC or Rural Health Clinic (RHC) and have 30% needy individual patient volume. FQHCs and RHCs are not eligible to receive payment under the program. Please contact your State Medicaid agency for more information on which types of encounters qualify as Medicaid/needy individual patient volume. For more information about the Medicare and Medicaid EHR Incentive Program, please visit http://www.cms.gov/EHRIncentivePrograms Keywords: FAQ10522
2845	How does CMS define Federally Qualified Health Center (FQHC) and Rural Health Center (RHC) for the purposes of the Medicaid EHR Incentive Program?	The Social Security Act at section 1905(j)(2) defines an FQHC as an entity which, "(i) is receiving a grant under section 330 of the Public Health Service Act, or (ii)(I) is receiving funding from such a grant under a contract with the recipient of such a grant and (II) meets the requirements to receive a grant under section 330 of the Public Health Service Act, (iii) based on the recommendation of the Health Resources and Services Administration within the Public Health Service, and is determined by the Secretary to meet the requirements for receiving such a grant including requirements of the Secretary that an entity may not be owned, controlled, or operated by another entity; or (iv) was treated by the Secretary, for purposes of Part B of title XVIII, as a comprehensive Federally-funded health center as of January 1, 1990, and includes an outpatient health program or facility operated by a tribe or tribal organization under the Indian Self-Determination Act or by an urban Indian organization receiving funds under Title V of the Indian Health Care Improvement Act for the provision of primary health services. RHCs are defined as clinics that are certified under section 1861(aa)(2) of the Social Security Act to provide care in underserved areas, and therefore, to receive cost-based Medicare and Medicaid reimbursements. In considering these definitions, it should be noted that programs meeting the FQHC requirements commonly include the following (but must be certified and meet all requirements stated above): Community Health Centers, Migrant Health Centers, Healthcare for the Homeless Programs, Public Housing Primary Care Programs, Federally Qualified Health Center Look-Alikes, and Tribal Health Centers. For more information about the Medicare and Medicaid EHR Incentive Program, please visit http://www.cms.gov/EHRIncentivePrograms Keywords: FAQ10127

7649	May a hospital include zero pay Medicaid eligible days in the Medicaid hospital EHR Incentive Program payment calculation?	No, zero pay Medicaid eligible days must be excluded from the Medicaid hospital incentive calculation. Section 1903(t)(5)(C) of the Act requires the Medicaid share to be calculated "in the same manner as the Medicare share." In all ways possible, the Medicaid hospital incentive calculation is similar to Medicare, based on this language. Medicare retrieves data for the calculation exclusively from the Medicare cost report. Although Medicaid offers additional flexibility in data sources, the data parameters for Medicaid are the same as Medicare. This is cited in the Stage 1 final rule where CMS said: "The statute requires us to calculate the Medicaid share 'in the same manner' as the Medicare share under section 1886(n)(2)(D) of the Act and such substitute service days would not be considered 'in the same manner.' Thus, we proposed that for purposes of the Medicaid formula, we would count only those days that would count as inpatient-bed-days for Medicare purposes under section 1886(n)(2)(D) of the Act." In the CMS Stage 1 final rule, CMS also made clear: "[T]he EHR hospital incentive payment calculation requires the inclusion of only paid inpatient-bed-days." 75 Fed. Reg. at 44500. Given this, a joint FAQ was published (FAQ # 3471) that mirrored cost report data sources for the calculation. Per the cost report instructions, all acute inpatient days must be paid. While CMS uses line 2 of worksheet S-3 part 1, which contains HMO data as well as other data used to calculate the Disproportionate Share Hospital (DSH) calculation (including zero pay days), Medicare is removing all of the DSH data from line 2 and using only the paid managed care days. Medicare does not include unpaid days as acute inpatient days, so following the same manner for Medicaid means using only paid days as well. Additionally, 1903(t)(5)(C) states that the Secretary establishes how the "inpatient bed-days" used in the Medicaid numerator are counted. The statute specifically says that the Medicaid share has as its numerator "the amount that is equal to the number of inpatient-bed-days (as established by the Secretary) which are attributable to individuals who are receiving medical assistance under this title." By using only paid inpatient Medicaid days, the Secretary has "established" how she counts the number of inpatient bed days per statutory authority.
9822	For Eligible Professionals (EP) in the Medicaid Electronic Health Records (EHR) Incentive Program using the group proxy method of calculating patient volume, how should the EPs calculate patient volume using the "12 months preceding the EP's attestation" approach, as not all of the EPs in the group practice may use the same 90-day period.	In the Stage 2 final rule, CMS adopted a final policy that allows states the option for their providers to calculate patient volume in any representative, continuous 90-day period in the 12 months preceding the eligible professional's (EP) attestation (see 77 FR 54121, 42 CFR 495.306(b)). This option is in addition to the method of calculating patient volume in any representative, continuous 90-day period in the calendar year preceding the payment year for which the EP is attesting. For EPs who calculate patient volume at the group practice or clinic level under 42 CFR 495.306(h), although we expect the same 90-day period to be used for all EPs in the group practice or clinic, we understand this may not be feasible in scenarios where EPs attest on different dates. For example, for the 2013 payment year, if one EP in the group attested on April 1, 2013 and another EP in the group attested on February 1, 2014, there would not be a continuous 90-day period that occurred within the 12 months preceding the first EP's attestation and also within the 12 months preceding the second EP's attestation. In such scenarios where it would be impossible to use the same representative, continuous 90-day period for EPs in the group practice or clinic, we would allow different representative, continuous 90-day periods to be used, as long as all of the provisions of 42 CFR 495.306(h) are satisfied. For more information, please visit the Stage 2 Final Rule: http://www.gpo.gov/fdsys/pkg/FR-2012-09-04/pdf/2012-21050 .
2819	Do States need to verify the "installation" or "a signed contract" for adopt, implement, or upgrade (AIU) in the Medicaid EHR Incentive Program?	States should make clear to providers when they attest for AIU what documentation they must maintain, and for how long, in case of audit. If States determine that certain provider types are a high risk for potential fraud/abuse for AIU, then they can ask for some verification of adopting, implementation or upgrading but CMS encourages that this be done in a targeted manner, with the most electronic and simple means possible and not in such a way that would be burdensome to providers. For AIU, a provider does not have to have installed certified EHR technology. The definition of AIU in 42 CFR 495.302 allows the provider to demonstrate AIU through any of the following: (a) acquiring, purchasing or securing access to certified EHR technology capable of meeting meaningful use; (b) installing or commencing utilization of certified EHR technology capable of meeting meaningful use requirements; or (c) expanding the available functionality of certified EHR technology capable of meeting meaningful use requirements at the practice site, including staffing, maintenance, and training, or upgrade from existing EHR technology to certified EHR technology per the EHR certification criteria published by the Office of the National Coordinator of Health Information Technology (ONC). Thus, a signed contract indicating that the provider has adopted or upgraded would generally be sufficient. However, if a provider has been identified as high risk, states could further investigate the provider's intention (or lack thereof) to AIU. While a signed contract indicating that the provider has adopted or upgraded might generally be sufficient to establish an intent to AIU, the state could still determine that other, contradictory evidence demonstrated that the provider in fact had no intent to AIU. In such cases, that contradictory evidence might outweigh the presence of a signed contract. For more information about the Medicare and Medicaid EHR Incentive Program, please visit " http://www.cms.gov/EHRIncentivePrograms " Keywords: FAQ10100 Updated on 4/2/2014
2831	For calculation of a Medicaid hospital's electronic health record (EHR) incentive payment, is the estimated growth rate for hospitals most recent three years based on growth in total days or growth in discharges?	The average annual growth rate should be for discharges (see 1903(t)(5)(B), referring to the annual rate of growth of the most recent 3 years for "discharge data.") We agree that the sources are different. Hospitals would probably have to use MMIS or auditable hospital records to get accurate discharge data rate of growth. To view the Stage 1 final rule, please visit: http://edocket.access.gpo.gov/2010/pdf/2010-17207.pdf For more information about the Medicare and Medicaid EHR Incentive Program, please visit http://www.cms.gov/EHRIncentivePrograms Keywords: FAQ10108
2839	Under the Medicaid Electronic Health Record (EHR) Incentive Program, how long is the EHR reporting period for each participation year demonstrating meaningful use?	Under the Medicaid Electronic Health Record (EHR) Incentive Program, if a provider adopts, implements or upgrades (AIU) certified EHR technology in their first year, the provider will not have to demonstrate meaningful use (MU) in order to receive payment. In the second year they will have to demonstrate MU for a 90 day period only. A provider that is already a meaningful user would have to demonstrate for a 90 day period the first year and subsequent years they would have to demonstrate it for the full year. The exception to this rule is for the year 2014. 24 CFR 495.4 establishes a one-time exception for providers attesting to meaningful use in 2014 during which the reporting period for Medicaid providers is any continuous 90-day period within the reporting year. For more information about the Medicare and Medicaid EHR Incentive Program, please visit http://www.cms.gov/EHRIncentivePrograms Keywords: FAQ10112
3113	Can a State access enhanced matching funds for the Medicaid EHR Incentive Program to participate in the creation of a Health Information Exchange (HIE) that is not directly administered by the State Medicaid Agency?	The enhanced match rate depends upon whether the Health Information Exchange solution is using Medicaid Management Information System (MMIS) funding or Health Information Technology for Economic and Clinical Health (HITECH) funding. Governance only is relevant under the MMIS regulations, as it pertains to the matching rate determination. States should talk to CMS about their ideas in draft, informally, so that CMS can give a more State-specific response around appropriate funding, matching rates, etc. For more information about the Medicare and Medicaid EHR Incentive Program, please visit http://www.cms.gov/EHRIncentivePrograms Keywords: FAQ10529
7809	What funding sources may States use to fund the 10% non-federal share of HITECH administrative expenditures?	States must fund the 10 percent non-federal share of the Health Information Technology (HITECH) Act administrative expenditures consistent with the law and regulations applicable to the non-federal share for all Medicaid expenditures. Consistent with that authority, which includes Social Security Act sections 1902(a)(2), 1903(a), 1903(w), and 42 CFR part 433, subpart B, states may fund the non-federal share of Medicaid expenditures through legislative appropriations to the Medicaid agency, intergovernmental transfers (IGTs), certified public expenditures (CPES), permissible health-care related taxes, and bona-fide donations. States must submit their proposed strategies for funding the non-federal share of HITECH administrative payments to CMS for review as part of the HIT plan approval process. CMS will review each individual State's proposal to ensure that each proposed non-federal share funding source meets federal requirements. During this process, CMS can address specific questions about funding the non-federal share of Medicaid expenditures. CMS strongly urges States to work on their funding proposals with their CMS HIT Coordinators as early as possible before claiming for HITECH administrative expenditures, to ensure funding structures are appropriate. HITECH administrative expenditures, like other title XIX expenditures, are subject to audit, and federal funds may be at risk if funding sources are found not to be in compliance with federal requirements. Below are some statutory and regulatory citations pertaining to non-federal share financing requirements. Please note this is not an all-inclusive list of funding requirements. Use of Federal Funds Social Security Act §1903 42 CFR 433.51(c) State Appropriations Social Security Act §1902(a)(2) 42 CFR 433.51 Intergovernmental Transfers Social Security Act §1903(w)(6)(A) 42 CFR 433.51 Certified Public Expenditures Social Security Act §1903(w)(6)(A) 42 CFR 433.51 Healthcare-Related Taxes and Provider-Related Donations 42 CFR part 433, subpart B Created 2/7/2013
8902	Can a state capture electronic Clinical Quality Measures, or eCQMs, for the Medicaid EHR Incentive Program through a Health Information Exchange (HIE)	Yes, a state can capture clinical quality data for eCQMs using an HIE, and states should consider the health data landscape of their state when designing a system to collect eCQMs for the Medicaid EHR Incentive Program. Utilizing an HIE can allow the state to collect more sophisticated patient-level data, to encourage provider adoption, and to facilitate alignment between various programs, such as those authorized under the HITECH Act, Accountable Care Organizations, and Medical Homes. In order to use an HIE for quality data collection, a state would need to develop infrastructure to capture electronic clinical quality measures through the Quality Reporting Data Architecture (QRDA) format. In addition, eligible professionals and hospitals would either have to generate the QRDA files using the provider's Certified EHR Technology and/or the HIE itself would have to be certified as an EHR module for eCQMs. The Office of the National Coordinator for Health Information Technology and CMS published a state-focused electronic clinical quality reporting toolkit to provide support for states developing the policy and IT infrastructure for electronic clinical quality measurement. Further, states can request Federal Financial Participation at the 90/10 HITECH rate to assist in building the infrastructure to submit eCQMs electronically for the Medicaid EHR Incentive program. For more information, please see the State Medicaid Directors Letter #11-004: http://downloads.cms.gov/cmsgov/archived-downloads/SMDL/ Created on 7/24/2013
8037	Can eligible professionals (EPs) or eligible hospitals round their patient volume percentage when calculating patient volume in the Medicaid Electronic Health Records (EHR) incentive program?	To participate in the Medicaid EHR incentive program, EPs are required to demonstrate a patient volume of at least 30% Medicaid patients over a 90-day period in the prior calendar year or in the 12 months before attestation. The Centers for Medicare and Medicaid Services allow rounding 29.5% and higher to 30% for purposes of determining patient volume. Similarly, pediatric patient volume may be rounded from 19.5% and higher to 20%. Finally, acute care hospitals are required to demonstrate a patient volume of at least 10% Medicaid patients over a 90-day period in the prior fiscal year preceding the hospital's payment year or in the 12 months before attestation. Hospitals' patient volume may be rounded from 9.5% and higher to 10%. Created on 3/13/2013
2767	Are eligible professionals (EPs) who practice in State Mental Health and Long Term Care Facilities eligible for Medicaid electronic health record (EHR) incentive payments if they meet the eligibility criteria (e.g., patient volume, non-hospital based, certified EHR)?	The setting in which a physician, nurse practitioner, certified nurse-midwife, or dentist practices is not relevant in determining eligibility for the Medicaid EHR Incentive Program (except for purposes of determining whether an EP can qualify through "needy individual" patient volume). Setting is relevant for physician assistants (PA), as they are eligible only when they are practicing at a Federally Qualified Health Center (FQHC) that is led by a PA or a Rural Health Center (RHC) that is so led. All providers must meet all program requirements prior to receiving an incentive payment (e.g. adopt, implement or meaningfully use certified EHR technology, patient volume, etc.). For more information about the Medicare and Medicaid EHR Incentive Program, please visit " http://www.cms.gov/EHRIncentivePrograms " Keywords: FAQ10069 Updated 5/12/2016

3471	If the State chooses to use the cost report in the Medicaid EHR incentive hospital payment calculation, what data elements should be used in the Medicare cost report, Form CMS 2552-96 and the Form CMS 2552-10?	<p>Based on the Medicare cost report guidance, Form CMS 2552-96 will be used until the implementation of the new Medicare cost report, Form CMS 2552-10. Although the State may choose to use the following data elements, it is the States' and hospitals' responsibility to ensure the integrity and regulatory compliance of the data. The CMS 2552-96 data elements are as follows: Total Discharges - Worksheet S-3 Part 1, Column 15, Line 12-Medicaid Days - Worksheet S-3, Part I, Column 5, Line 1 + Lines 6-10-Medicaid HMO Days - Worksheet S-3, Part I, Column 5, Line 2-Total Inpatient Days - Worksheet S-3 Part 1, Column 6, Line 1, 2 + Lines 6 -10-Total Hospital Charges - Worksheet C Part 1, Column 8, Line 101-Charity Care Charges - Worksheet S-10, Column 1, Line 30 The CMS 2552-10 data elements are as follows:-Total Discharges - Worksheet S-3 Part 1, Column 15, Line 14-Medicaid Days - Worksheet S-3, Part I, Column 7, Line 1 + Lines 8-12-Medicaid HMO Days - Worksheet S-3, Part I, Column 7, Line 2-Total Inpatient Days - Worksheet S-3 Part 1, Column 8, Line 1, 2 + Lines 8 - 12- Total Hospital Charges - Worksheet C Part 1, Column 8, Line 200-Charity Care Charges - Worksheet S-10, Column 3, Line 20 For information about the cost report data elements that are used in the Medicare hospital incentive calculation, please see http://questions.cms.hhs.gov/app/answers/detail/a_id/10717/ FAQ #10717 For more information about the Medicare and Medicaid EHR Incentive Program, please visit http://www.cms.gov/EHRIncentivePrograms"</p> <p>Keywords: FAQ10771</p>
		<p>To meet the "Protect Electronic Health Information" core objective for Stage 1, eligible professionals (EP), eligible hospitals or critical access hospitals (CAH) must conduct or review a security risk analysis in accordance with the requirements under 45 CFR 164.308(a)(1) and implement security updates as necessary and correct identified security deficiencies as part of the provider's risk management process. In Stage 2, in addition to meeting the same security risk analysis requirements as Stage 1, EPs and hospitals will also need to address the encryption and security of data stored in the certified EHR technology (CEHRT). These steps may be completed outside or the EHR reporting period timeframe but must take place within the same calendar year as the EHR reporting period, and if the provider attests prior to the end of the calendar year, it must be conducted prior to the date of attestation. For example, a EP who is reporting Meaningful Use for a 90-day EHR reporting period may complete the appropriate security risk analysis requirements outside of this 90-day period as long as it is completed no earlier than January 1st of the EHR reporting year and no later than the date the provider submits their attestation for that EHR reporting period. This meaningful use objective complements but does not impose new or expanded requirements on the HIPAA Security Rule. In accordance with the requirements under (45 CFR 164.308(a)(1)(ii)), providers are required to conduct an accurate and thorough analysis of the potential risks and vulnerabilities to the confidentiality, integrity, and availability of electronic protected health information (ePHI). Once the risk analysis is completed, providers must take any additional "reasonable and appropriate" steps to reduce identified risks to reasonable and appropriate levels. Please note that a security risk analysis or review needs to be conducted during each EHR reporting year for Stage 1 and Stage 2 of meaningful use to ensure the privacy and security of their patients' protected health information. For more information about completing a security risk analysis, please see the following resources: Security Risk Assessment Tip Sheet: https://www.cms.gov/Regulationsand-Guidance/Legislation/EHRIncentivePrograms/Downloads/SecurityRiskAssessment_FactSheet_Updated20131122.pdf">https://www.cms.gov/Regulations-and-Health Information Privacy and Security: A 10 Step Plan: a http://www.healthit.gov/providers-professionals/ehr-privacy-security/Created 10/6/2014 Updated 11/5/2014 Archived 12/15/15</p>