

**KEY COMPLIANCE REQUIREMENTS AND DEADLINES FOR ONC and CMS RULES REGARDING INTEROPERABILITY,
INFORMATION BLOCKING, PATIENT ACCESS, AND HEALTH IT CERTIFICATION**

| ONC FINAL RULE | | |
|---|--|--|
| Final Rule: 21st Century Cures Act: Interoperability, Information Blocking, and the ONC Health IT Certification Program | | |
| Regulatory Requirement Description | Relevant Regulatory Section(s) and Description | Compliance or Enforcement Deadline |
| General Effective Date of Rules | <p>Applies to <u>removal</u> of pre-2015 Edition certification criteria from:</p> <ul style="list-style-type: none"> • 45 C.F.R. § 170.315(a)(6) (problem list) • 45 C.F.R. § 170.315 (a)(7) (medication list) • 45 C.F.R. § 170.315 (a)(8) (medication allergy list) • 45 C.F.R. § 170.315 (a)(11) (smoking status) • 45 C.F.R. § 170.315 (b)(4)(CCDS –Create) • 45 C.F.R. § 170.315 (b)(5)(CCDS –Receive) <p>and <u>revisions</u> of the following:</p> <ul style="list-style-type: none"> ▪ 45 C.F.R. § 170.315(b)(3) ePrescribing ▪ 45 C.F.R. § 170.315(b)(7) Security Tags – Summary of Care (send) (formerly, DS4P – Send) ▪ 45 C.F.R. § 170.315(b)(8) Security Tags – Summary of Care (receive) (formerly, DS4P – Receive) ▪ 45 C.F.R. § 170.315(c)(3) CQMs – Report ▪ 45 C.F.R. § 170.315(d)(2) Adjustable Events and Tamper-Resistance ▪ 45 C.F.R. § 170.315(d)(3) Audit Report(s) ▪ 45 C.F.R. § 170.315(d)(10) Auditing Actions on Health Information | <p>60 days after publication date.</p> <p>For removed criteria, certified health IT products will no longer need to certify to including these elements. Any certification of health IT submitted or subject to maintenance of certification 60 days after publication date will need to comply with revised criteria definitions.</p> |
| General Information | <ul style="list-style-type: none"> • 45 C.F.R. § 170.401 (information blocking condition of certification) | 6 months after publication |

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| Blocking Compliance Date | requirement) <ul style="list-style-type: none"> • 45 C.F.R. § 170.402(a)(1) (information blocking assurances) • 45 C.F.R. part 171 (definitions and exceptions) | date. |
| Phase out of ONC-ACB certification issuances | <ul style="list-style-type: none"> • 45 C.F.R. § 170.315(a)(10)(drug/formulary preferred drug list checks), • 45 C.F.R. § 170.315 (a)(13) (patient-specific education resource), • 45 C.F.R. § 170.315 (b)(6) (data export), and • 45 C.F.R. § 170.315 (e)(2) (secure messaging) | Ends January 1, 2022 with sunset of Medicare Promoting Interoperability Program (except for § 170.315(b)(6), which ends 36 months after publication date). |
| Interim “Electronic Health Information” Definitions in effect (Compliance with USCDI Only) | 45 C.F.R. § 170.315 (see also Table 1 . 2015 Edition Cures Update in ONC Final Rule) <ul style="list-style-type: none"> • 45 C.F.R. § 170.315(b)(1) Transitions of Care • 45 C.F.R. § 170.315(b)(2) Clinical Information Reconciliation and Incorporation • 45 C.F.R. § 170.315(e)(1) View, Download, and Transmit to a 3rd Party • 45 C.F.R. § 170.315(f)(5) Transmission to Public Health Agencies – Electronic Case Reporting • 45 C.F.R. § 170.315(g)(6) Consolidated CDA Creation Performance • 45 C.F.R. § 170.315(g)(9) Application Access – All Data Request • 45 C.F.R. § 170.315(g)(10) Standardized API for Patient and Population Services | Months 6-24 after publication date. |

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| | (new certification criterion that also refers to the USCDI) | |
| Full EHI Definitions in Effect | 45 C.F.R. § 170.315 (2015 Edition certified Health IT certification criteria - see also Table 1 . 2015 Edition Cures Update in ONC Final Rule) | 24 months after publication date. |
| HL7 FHIR API Capability Rollout | 45 C.F.R. § 170.215 (standard) and 45 C.F.R. § 170.315(g)(10) (use) | No later than 24 months after publication date. |
| EHI Export Capability Rollout | 45 C.F.R. § 170.315(b)(10) | No later than 36 months after publication date. |
| Attestations for Conditions and Maintenance of Certification (CMC) | 45 C.F.R. § 170.406 (attestations for certified Health IT CMCs) Health IT developers will be able to submit their attestations within a designated 30-day window twice a year for purposes of compliance. | First 30-day attestation submission window begins April 1, 2021, for attestation periods between the effective date of the final rule and March 31, 2021. Subsequent semiannual submissions during 30-day windows (e.g., October 1, 2021-October 31, 2021 - window for attestation period |

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| | | of April 1 through September 30, 2021). |
| CMC Requirements Related to Information Blocking | 45 C.F.R. § 170.401 Prohibits any “health IT developer” who has at least one health IT product certified under the Program from taking any action that constitutes information blocking as defined by section 3022(a) of the Public Health Service Act and proposed in 45 C.F.R. § 171.103. | 6 months after publication date. |
| Conditions and Maintenance of Certification (CMC) Requirements Related to Information Blocking Assurances: | 45 C.F.R. § 170.402 45 C.F.R. § 170.315 Requires that a health IT developer, as a Condition and Maintenance of Certification requirement under the ONC Health IT Certification Program (“Program”), provide assurances to the Secretary, unless for legitimate purposes specified by the Secretary, that it will not take any action that constitutes information blocking as defined in section 3022(a) of the PHSA, or any other action that may inhibit the appropriate exchange, access, and use of electronic health information (EHI) | 6 months after publication date. |
| CMC Requirements Related to Protected | 45 C.F.R. § 170.403 Prohibits health IT developers from restricting “protected communications” | 6 months after publication date. |

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| Communications | regarding the following subjects: usability of the health information technology; interoperability of the health information technology; security of the health information technology; relevant information regarding users' experiences when using the health information technology; business practices of developers of health information technology related to exchanging electronic health information; and manner in which a user of the health information technology has used such technology. Certain narrow exceptions to the above are set out in the regulation. | |
| CMC Requirements Related to APIs | <ul style="list-style-type: none"> • 45 C.F.R. § 170.404(API technology supplier, API data provider, and API user definitions, among others) • 45 C.F.R. § 170.213 (U.S. Core Data for Interoperability (USCDI) definition) • 45 C.F.R. § 170.215 (API standards - FHIR and USCDI provisions) • 45 C.F.R. § 170.315(g)(10) (standardized API for Patient and Population Services (new certification criterion that also refers to the USCDI)) <p>Requires health IT developers to publish APIs that allow “health information from such technology to be accessed, exchanged, and used without special effort through the use of APIs or successor technology or standards, as provided for under applicable law.”</p> | 6 months after publication date. |
| First Real-World Testing | 45 C.F.R. § 170.405 | Dec. 15, 2020 (deadline for publishing the plan via publicly |

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| Plans Due | <p>Health IT developers must successfully test the real world use of the technology for interoperability in the type of setting in which such technology would be marketed.</p> <p>NOTE: CMS finalized that document-level tagging remains applicable for up to 24 months after the publication date of the final rule. For certification and compliance of Health IT Modules certified after 24 months after the publication date of this final rule, only the full scope of the HL7 Data Segmentation for Privacy (DS4P) standard is applicable. HL7 DS4P standard provides a means to express obligations and disclosure restrictions that may exist for sensitive data.</p> | accessible hyperlink on the Certified Health IT Products List) |
| Information Blocking Civil Monetary Penalties | <p>TBD – Proposed Rule Currently Under Review at OMB</p> <p>NOTE: This does not foreclose the possibility that the OIG will still rely on its statutory authority to accept complaints regarding information blocking allegations, investigate such claims, and/or engage in other monitoring or audit activities with respect to information blocking independently or in coordination with ONC, or from pursuing enforcement against actors for false attestations submitted to the ONC related to health IT certification criteria in coordination with the Department of Justice under the False Claims Act.</p> | Delayed until publication of final rules from HHS Office of the Inspector General |

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CMS PATIENT ACCESS RULE

[CMS Finalizes New Requirements on Health Plans to Release Health Data; Updates Medicare Conditions of Participation for Hospitals to Send Electronic Notifications](#)

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| Digital Contact Information to be reported on NPPES | <p>Cures Act section 4003</p> <p>Requires the HHS Secretary to create a provider digital contact information index that includes all individual health care providers and facilities. CMS plans to update the National Plan and Provider Enumeration System (NPPES) to include data fields to collect, for example, Direct Addresses, FHIR server URLs, query endpoints and/or other “electronic service information.”</p> <p>CMS will publicly report the names and NPIs of those providers who do not have digital contact information that can be used to facilitate the secure sharing of health information. CMS has not yet identified where this reporting will be publicized.</p> | <p>Second half of 2020</p> <p>CMS information collection request under the Paperwork Reduction Act of 1995 will be published for review and comment.</p> |
| Admission, Discharge and Transfer Event Notifications | <p>Hospitals - 42 C.F.R. § 482.24(d),</p> <p>Psychiatric Hospitals - 42 C.F.R. § 482.61(f),</p> <p>Critical Access Hospitals (CAHs) - 42 C.F.R. § 485.638(d).</p> <p>Revises the Conditions of Participation for Medicare- and Medicaid-participating hospitals, psychiatric hospitals, and CAHs by adding a</p> | <p>Fall 2020 (estimated in September)</p> |

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| | <p>new standard, “Electronic Notifications,” that will require hospitals, psychiatric hospitals, and CAHs to make electronic patient event notifications available to applicable post-acute care services providers and suppliers, and to community practitioners such as the patient’s established primary care practitioner, established primary care practice group or entity, or other practitioner or practice group or entity identified by the patient as primarily responsible for his or her care. This requirement is limited to only those hospitals, psychiatric hospitals, and CAHs that utilize electronic medical records systems, or other electronic administrative systems, which are conformant with the content exchange standard at 45 C.F.R. § 170.205(d)(2).</p> | |
| Patient Access API | <ul style="list-style-type: none"> ▪ 42 C.F.R. § 422.119 (Medicare Advantage (MA) organizations) ▪ 42 C.F.R. § 431.60 (Medicaid Fee-for-Service (FFS)) ▪ 42 C.F.R. § 438.242 (Medicaid Managed Care) ▪ 42 C.F.R. § 457.730 (Children’s Health Insurance Program (CHIP) FFS) ▪ 42 C.F.R. § 457.123 (CHIP Managed Care) ▪ 45 C.F.R. § 156.221 (QHP issues on Federally Facilitated Exchanges) | January 1, 2021 |
| Provider Directory API | <ul style="list-style-type: none"> ▪ 42 C.F.R. 422.120 for (MA organizations) | January 1, 2021 |

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| | <ul style="list-style-type: none"> ▪ 42 C.F.R. 431.70 (Medicaid state agencies) ▪ 42 C.F.R. 438.242(b)(6) (Medicaid managed care plans) ▪ 42 C.F.R. 457.760 (CHIP state agencies) ▪ 42 C.F.R. 457.1233(d)(3) (CHIP managed care entities) | |
| Payer-to-Payer Data Exchange | <ul style="list-style-type: none"> ▪ 42 C.F.R. 422.119(f) (MA organizations) ▪ 42 C.F.R. 438.62(b)(1)(vi) (Medicaid managed care plans) ▪ 42 C.F.R. § 457.1216 (CHIP managed care entities), ▪ 45 C.F.R. 156.221(f) (Qualified Health Plan (QHP) issuers on the Federally Facilitated Exchanges) ▪ 45 C.F.R. § 170.213 (exchange content standard – currently USCDI version 1) | <p>January 1, 2022</p> <p>CMS-regulated payers are required to exchange patient clinical data at the patient’s request, as patients move between CMS-regulated payers <u>for services dated on or after January 1, 2016.</u></p> |
| Frequency of Federal-State Data Exchanges for Dual-Eligible Patients | <p>42 C.F.R. §§ 406.26 and 407.40</p> <p>Requires all states to participate in daily exchange of buy-in data with CMS, with “daily” meaning every business day, but that if no new transactions are available to transmit, data would not need to be submitted on a given business day</p> | <p>April 1, 2022</p> |

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| Physician Compare Reporting | Indicator that eligible clinicians and groups submitted “no” responses to any of the three prevention of information blocking statements for Merit-based Incentive Payment System (MIPS) | Starting with the 2019 performance period data available for public reporting starting in late 2020. |
| Public Website Reporting | List of eligible hospitals or CAHs attesting under the Medicare FFS Promoting Interoperability Program that submitted a “no” response to any of the three attestation statements related to the prevention of information blocking | Starting with the 2019 performance period data available for public reporting starting in late 2020. |