

the same terms and conditions in effect on that date.

In providing for the grandfathering of satellite facilities of hospitals and hospital units, we believed it was appropriate to require that the satellite facilities operate under the same terms and conditions that were in effect on September 30, 1999. There are similarities between the definition of the two types of satellite facilities and the definition of hospitals-within-hospitals (that is, hospitals-within-hospitals and satellite facilities are both physically located in acute care hospitals that are paid for their inpatient services on a prospective payment basis). Also, satellite facilities of both excluded hospitals and hospital units and hospitals-within-hospitals provide inpatient hospital services that are paid at a higher rate than would apply if the facilities were treated by Medicare as part of an acute care hospital.

We are proposing to revise § 412.22(f) to specify that, effective with cost reporting periods beginning on or after October 1, 2003, a hospital operating as a hospital-within-a-hospital on or before September 30, 1995, is exempt from the criteria in § 412.22(e)(1) through (e)(5) only if the hospital-within-a-hospital continues to operate under the same terms and conditions in effect as of September 30, 1995. The intent of the “grandfathering” provision was to ensure that hospitals that had been in existence prior to the effective date of our hospital-within-hospital requirements should not be adversely affected by those requirements. To the extent hospitals were already operating as hospitals-within-hospitals without meeting those requirements, we believe it is appropriate to limit the “grandfathering” provision to those hospitals that continue to operate in the same manner as they had operated prior to the effective date of those rules. However, if a hospital changes the way it operates (for example, adds more beds) subsequent to the effective date of the new rules, it should no longer receive the benefit of the “grandfathering” provision.

Under § 412.22(e), we specify the criteria that a hospital-within-a-hospital is required to meet in order to be excluded from the IPPS. One of these criteria, under § 412.22(e)(5)(i), requires that a hospital-within-a-hospital is able to perform basic hospital functions (for example, medical record services and nursing services) that are presently included in the Medicare hospital conditions of participation under Part 482 of the Medicare regulations. These requirements were first included in Part 412 in response to hospitals organizing

themselves as what is referred to as the hospital-within-a-hospital model. Thus, to avoid recognizing nominal hospitals, while allowing hospitals adequate flexibility and opportunity for legitimate networking and sharing of services, we included, by reference, certain hospital conditions of participation as additional criteria in part 412 for hospitals-within-hospitals that request exclusion from the IPPS. (Further discussion can be found in a final rule published in the **Federal Register** on September 1, 1994 (59 FR 45389).) Modifications to the conditions of participation have been made since the publication of that September 1, 1994 final rule. Thus, we need to update the references to the conditions of participation in § 412.22(e)(5)(i) to make them consistent with existing provisions under the basic hospital conditions of participation. Therefore, we are proposing to amend § 412.22(e)(5)(i) to add references to § 482.43 (discharge planning) and § 482.45 (organ, tissue, and eye procurement) as basic hospital functions that a hospital-within-a-hospital would also be required to meet.

C. Clarification of Classification Requirements for LTCHs

Under § 412.23(e)(2), to qualify to be excluded from the IPPS as a LTCH and to be paid under the LTCH PPS, a hospital must have an average Medicare length of stay of greater than 25 days (which includes all covered and noncovered days of stay for Medicare patients) as calculated under the criteria of § 412.23(e)(3). In calculating this average Medicare inpatient length of stay, data from the hospital's most recently filed cost report are used to make this determination. However, if the hospital has not yet filed a cost report or if there is an indication that the most recently filed cost report does not accurately reflect the hospital's current Medicare average length of stay, data from the most recent 6-month period are used.

Our interpretation of § 412.23(e)(3)(ii) and (e)(3)(iii) was to allow hospitals that submit data for purposes of exclusion from the IPPS to use a period of at least 5 months of the most recent data from the preceding 6-month period. This longstanding policy interpretation was necessary in order to comply with the time requirement in § 412.22(d) that specifies that, for purposes of the IPPS, status is determined at the beginning of each cost reporting period and is effective for the entire cost reporting period. Therefore, we are proposing to revise §§ 412.23(e)(3)(ii) and (iii) to reflect our longstanding interpretation of the regulations.

D. Criteria for Payment on a Reasonable Cost Basis for Clinical Diagnostic Laboratory Services Performed by CAHs

Section 1820 of the Act provides for the establishment of Medicare Rural Hospital Flexibility Programs, under which individual States may designate certain facilities as critical access hospitals (CAHs). Facilities that are so designated and meet the CAH conditions of participation in 42 CFR part 485, subpart F, will be certified as CAHs by CMS. Section 1834(g) of the Act states that the amount of payment for outpatient services furnished by a CAH will be the reasonable costs of the CAH in providing these services.

Regulations implementing section 1834(g) of the Act are set forth at § 413.70. These regulations state, in paragraph (b)(2)(iii), that payment to a CAH for outpatient clinical diagnostic laboratory tests will be made on a reasonable cost basis only if the individuals for whom the tests are performed are outpatients of the CAH, as defined in 42 CFR 410.2, at the time the specimens are collected. The regulations also state that clinical diagnostic laboratory tests for persons who are not patients of the CAH at the time the specimens are collected will be paid for in accordance with the provisions of sections 1833(a)(1)(D) and 1833(a)(2)(D) of the Act. These provisions, which also are the basis for payment for clinical diagnostic laboratory tests performed by independent laboratories and by hospitals on specimens drawn at other locations, set payment at the least of: (1) Charges determined under the fee schedule as set forth in section 1833(h)(1) or section 1834(d)(1) of the Act; (2) the limitation amount for that test determined under section 1833(h)(4)(B) of the Act; or (3) a negotiated rate established under section 1833(h)(6) of the Act. Payments determined under this methodology are typically referred to as “fee schedule payments,” and are so described here both for ease of reference and to differentiate them from payments determined on a reasonable cost basis.

The definition of an “outpatient” in 42 CFR 410.2 states that an outpatient means a person who has not been admitted as an inpatient but who is registered on hospital or CAH records as an outpatient and receives services (rather than supplies alone) directly from the hospital or CAH.

Recently, we have received numerous questions about how Medicare pays for laboratory services that a CAH may furnish to Medicare beneficiaries in various settings other than the CAH.