§ 414.806 Penalties associated with the failure to submit timely and accurate ASP data.

Section 1847A(d)(4) specifies the penalties associated with misrepresentations associated with ASP data. If the Secretary determines that a manufacturer has made a misrepresentation in the reporting of ASP data, a civil money penalty in an amount of up to $10,000 may be applied for each price misrepresentation and for each day in which the price misrepresentation was applied. Section 1927(b)(3)(C) of the Act, as amended by section 303(i)(4) of the MMA, specifies the penalties associated with a manufacturer’s failure to submit timely information or the submission of false information.

Subpart K—Payment for Drugs and Biologicals Under Part B

Source: 69 FR 66424, Nov. 15, 2004, unless otherwise noted.

§ 414.900 Basis and scope.

(a) This subpart implements sections 1842(o), 1847A, and 1847B of the Act and outlines two payment methodologies applicable to drugs and biologicals covered under Medicare Part B that are not paid on a cost or prospective payment system basis. (b) Examples of drugs that are subject to the requirements specified in this subpart are:

(1) Drugs furnished incident to a physician’s service; durable medical equipment (DME) drugs.
(2) Separately billable drugs at independent dialysis facilities not under the ESRD composite rate.
(3) Statutorily covered drugs, for example—
   (i) Influenza.
   (ii) Pneumococcal and Hepatitis B vaccines.
   (iii) Antigens.
   (iv) Hemophilia blood clotting factor.
   (v) Immunosuppressive drugs.
   (vi) Certain oral anti-cancer drugs.

(69 FR 66424, Nov. 15, 2004, as amended at 70 FR 36093, July 6, 2005)