OFICIO 600-7840-16

Bogotá D.C., Septiembre 15 de 2016

ASUNTO: Acuso Recibo de Eventos Adversos Serios

Dando cumplimiento a lo estipulado en la Resolución No. 2011020764 del 10 de junio de 2011 “Por la cual se establece el reglamento relativo al contenido y periodicidad de los reportes de eventos adversos en la fase clínica con medicamentos en humanos de que trata el artículo 146 del decreto 677 de 1995”, la Dirección de Medicamentos y Productos Biológicos del INVIMA, da acuso de recibo de la información allegada con los siguientes radicados, correspondientes al reporte de eventos adversos serios nacionales e internacionales y otros documentos relacionados. El INVIMA en cualquier momento y como parte del proceso de análisis de la información, podrá requerir datos adicionales, para efectos de Inspección Vigilancia y Control.

**Eventos Adversos Serios Internacionales**

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Atentamente,

LUZ HELENA FRANCO CHAPARRO
Directora de Medicamentos y Productos Biológicos

Proyectó: Profesionales del Grupo de Investigación Clínica