

Attachment A

Dear Healthcare Provider:

As you may be aware Cephalon, Inc. recently entered into a global civil, criminal, and administrative settlement with the United States and individual states in connection with its promotion of three Cephalon products. This letter provides you with additional information about the settlement, explains Cephalon's commitments going forward, and provides you with access to information about those commitments. In general terms, the Government alleged that Cephalon unlawfully promoted three drugs (Actiq, Gabitril, and Provigil) for uses not approved by the Food & Drug Administration (FDA). To resolve these matters, Cephalon pled guilty to a misdemeanor criminal violation and agreed to pay a total of \$425 million to the Federal Government and state Medicaid programs. In addition, Cephalon paid \$6,150,000 in a companion settlement with the Connecticut Attorney General related to unfair trade practice laws. Additional information about the settlements may be found at the following websites [include a link to the USAO, Cephalon (www.cephalon.com), and Attorney General of Connecticut's websites.]

As part of the federal settlement, Cephalon also entered a five-year corporate integrity agreement with the Office of Inspector General of the U.S. Department of Health and Human Services. The corporate integrity agreement is available at <http://oig.hhs.gov/fraud/cia/index.html>. Under this agreement, Cephalon agreed to undertake certain obligations designed to promote compliance with Federal health care program and FDA requirements. We also agreed to notify healthcare providers about the settlement and inform them that they can report any questionable practices by Cephalon's representatives to Cephalon's Compliance Department or the FDA.

Please call or email Cephalon at 1-866-900-7167 or questions@cephalon.com if you have questions about the settlement referenced above or to report any instances in which you believe that a Cephalon representative inappropriately promoted a product or engaged in other questionable conduct. Alternatively, you may report any such instances to the FDA's Division of Drug Marketing, Advertising, and Communications at 301-796-1200. You should direct medical questions or concerns about the products to Cephalon's Medical Information department.

We appreciate your time and attention. We are dedicated to ensuring that we bring you the scientific and medical information you need to make well-informed decisions about whether Cephalon products are right for your patients.

Sincerely,

Chief Executive Officer
Cephalon, Inc.