



## Consumer Product Safety Improvement Act (CPSIA) General Conformity Certificate

As required by section 102 of the Consumer Product Safety Improvement Act of 2008, codified at 15 U.S.C. §2063(a), the following certifies the product referenced meets all applicable testing standards

Amgen NDC Number: 55513-0073-07  
Amgen Drug Product: 038 FDP US 30mg 7Tbl Sample  
Dosage: 30mg  
Finished Drug Product Lot Number: 1010002

Consumer Product Safety Commission (CPSC) safety requirements to which the above-referenced product is being certified: Child-resistant packaging requirements under the Poison Prevention Packaging Act. The applicable CPSC regulations are codified at 15 CFR Parts 1700 and 1701.

Date of (month/year) and place of Manufacture of Finished Drug Product:  
03/23/2009, Mississauga, Ontario Canada.

Drug Product Package Packaging Testing [Per Amgen Technical Assessment # TA-004171](#)

Date of testing: [5/21/2001](#)

Location of testing (incl. city/county): [Perritt Laboratories, Hightstown, NJ 08520](#)

U.S Importer of Record/ Issuer of Certificate	Amgen U.S.A 12000 Plantside Drive Louisville, KY 40299 502-266-2706
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For additional information, please contact the following party responsible for maintaining the test results:

Amgen Inc  
Attn: Amgen Medical Information  
One Amgen Center Drive  
Thousand Oaks, CA 91320  
Phone Number: 800-77-AMGEN  
Contact fax Number: 866-29-AMGEN  
Email: [www.amgenmedinfo.com](http://www.amgenmedinfo.com)