

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: 1042667

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 16 CFR Parts 1700 and 1701.

Date of (month/year) and place of Manufacture of Finished Drug Product:

Date of Manufacture: February 2013

Place of Manufacture: Patheon, Inc., Toronto Region Operations
2100 Syntex Court
Mississauga, Ontario L5N7K9
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	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
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