



US 20100273829A1

(19) **United States**

(12) **Patent Application Publication**  
**Wisler**

(10) **Pub. No.: US 2010/0273829 A1**  
(43) **Pub. Date: Oct. 28, 2010**

(54) **METHODS FOR TREATING DISORDERS ASSOCIATED WITH HYPERLIPIDEMIA IN A MAMMAL**

(76) Inventor: **Gerald L. Wisler**, Gladstone, NJ (US)

Correspondence Address:  
**GOODWIN PROCTER LLP**  
**PATENT ADMINISTRATOR**  
**53 STATE STREET, EXCHANGE PLACE**  
**BOSTON, MA 02109-2881 (US)**

(21) Appl. No.: **12/646,146**

(22) Filed: **Dec. 23, 2009**

**Related U.S. Application Data**

(63) Continuation of application No. 11/582,876, filed on Oct. 18, 2006, now abandoned.

(60) Provisional application No. 60/727,664, filed on Oct. 18, 2005, provisional application No. 60/788,616, filed on Apr. 3, 2006.

**Publication Classification**

(51) <b>Int. Cl.</b>	
<i>A61K 31/4468</i>	(2006.01)
<i>A61P 3/06</i>	(2006.01)
<i>A61P 7/00</i>	(2006.01)
<i>A61P 1/16</i>	(2006.01)
<i>A61P 9/10</i>	(2006.01)
(52) <b>U.S. Cl.</b> .....	<b>514/325</b>

(57) **ABSTRACT**

The invention is directed to methods for treating disorders associated with hyperlipidemia in a mammal. The methods involve combination therapies using a microsomal triglyceride transfer protein (MTP) inhibitor (for example, BMS-201038 and implitapide) and a fibrate (for example, fenofibrate). Co-administration of the MTP inhibitor with the fibrate produces a therapeutic benefit, for example, a reduction in the concentration of cholesterol and/or triglycerides in the blood stream, but with fewer or reduced side effects than when higher dosages of the MTP inhibitor are used during monotherapy to provide the same or similar therapeutic benefit.

## METHODS FOR TREATING DISORDERS ASSOCIATED WITH HYPERLIPIDEMIA IN A MAMMAL

### RELATED APPLICATIONS

[0001] This application is a continuation of U.S. application Ser. No. 11/582,876, filed Oct. 18, 2006, which claims the benefit of U.S. Provisional Patent Application Ser. No. 60/788,616, filed Apr. 3, 2006, and U.S. Provisional Patent Application Ser. No. 60/727,664, filed Oct. 18, 2005, the entire disclosures of which are incorporated by reference herein.

### FIELD OF THE INVENTION

[0002] This invention relates generally to methods of reducing the concentration of cholesterol and/or triglycerides in the blood of a mammal. More particularly, the invention relates to combination therapies using a microsomal triglyceride transfer protein (MTP) inhibitor and a fibrate for reducing the concentration of cholesterol and/or triglycerides in the blood but with a reduced adverse event profile relative to MTP inhibitor monotherapy.

### BACKGROUND OF THE INVENTION

[0003] There are several known risk factors for atherosclerotic cardiovascular disease (ASCVD), the major cause of mortality in the Western world. One key risk factor is hyperlipidemia, which is the presence of elevated levels of lipids in blood plasma. Various epidemiological studies have demonstrated that drug mediated lowering of total cholesterol (TC) and low density lipoprotein (LDL) cholesterol (LDL-C) is associated with a significant reduction in cardiovascular events. The National Cholesterol Education Program's (NCEP's) updated guidelines recommends that the overall goal for high-risk patients is to achieve less than 100 mg/dL of LDL, with a therapeutic option to set the goal for such patients to achieve a LDL level less than 70 mg/dL.

[0004] One form of hyperlipidemia is known as hypertriglyceridemia and results in the presence of elevated amounts of triglycerides in the blood. Although triglycerides are necessary for good health, higher-than-normal triglyceride levels, often are associated with known risk factors for heart disease.

[0005] Another form of hyperlipidemia, known as hypercholesterolemia, which is the presence of elevated amounts of cholesterol in the blood, is a polygenic disorder. Modifications in lifestyle and conventional drug treatment are usually successful in reducing cholesterol levels. However, in some cases, as in familial hypercholesterolemia (FH), the cause is a monogenic defect. Treatment of a patient with FH can be more challenging because the levels of LDL-C remain elevated despite aggressive use of conventional therapy.

[0006] For example, one type of FH, homozygous familial hypercholesterolemia (hoFH), is a serious life-threatening genetic disease caused by homozygosity or compound heterozygosity for mutations in the low density lipoprotein (LDL) receptor. Patients with hoFH typically have total plasma cholesterol levels over 400 mg/dL, resulting in premature atherosclerotic vascular disease. When left untreated, most patients develop atherosclerosis before age 20 and generally do not survive past age 30. However, patients diagnosed with hoFH are largely unresponsive to conventional

cholesterol synthesis and upregulating the hepatic LDL receptor, have negligible effect in patients whose LDL receptors are non-existent or defective. A mean LDL-C reduction of only less than about 20% has been recently reported in patients with genotype-confirmed hoFH treated with the maximal dose of statins (atorvastatin or simvastatin administered at 80 mg/day). The addition of ezetimibe 10 mg/day to this regimen resulted in a total reduction of LDL-C levels of 27%, which is still far from optimal. Non-pharmacological options have also been tested, including surgical interventions, such as portacaval shunt and ileal bypass, and orthotopic liver transplantation, but with clear disadvantages and risks. Therefore, there is a tremendous unmet medical need for new medical therapies for hoFH.

[0007] Microsomal triglyceride transfer protein (MTP) inhibitors have been developed as potent inhibitors of MTP-mediated neutral lipid transfer activity. MTP catalyzes the transport of triglyceride, cholesteryl ester, and phosphatidylcholine between small unilamellar vesicles. One exemplary MTP inhibitor is BMS-201038, developed by Bristol-Myers Squibb. See, U.S. Pat. Nos. 5,739,135; and 5,712,279. Studies using an animal model for homozygous FH indicated that BMS-201038 effectively reduced plasma cholesterol levels in a dose dependent manner, for example, at 25 mg/day, suggesting that this compound might be effective for treating patients with hoFH. It was noticed, however, that certain patients treated with 25 mg/day of BMS-201038 experienced certain adverse events, for example, gastrointestinal perturbations, abnormalities in liver function, and hepatic steatosis. Although a promising therapeutic agent, large scale clinical trials of BMS-201038 have been discontinued. Another potent MTP inhibitor known as implitapide has been developed. See, U.S. Pat. Nos. 6,265,431, 6,479,503, 5,952,498. During clinical studies, dosages of implitapide of 80 mg/day or greater, although therapeutically effective, were found to result in certain adverse events, for example, gastrointestinal disturbances, abnormalities in liver function, and hepatic steatosis. Large scale clinical studies using implitapide have also been discontinued.

[0008] Accordingly, there is still a need for methods for aggressively treating hyperlipidemias that effectively lower, for example, circulating cholesterol and triglycerides levels but with fewer or reduced adverse effects that typically result when higher dosages of the MTP inhibitor are used alone in monotherapy.

### SUMMARY OF THE INVENTION

[0009] The invention provides methods for lowering the concentration of cholesterol and/or triglycerides in the blood, and/or reducing the amount of one or more markers of atherosclerosis. The method includes administering an MTP inhibitor, such as, BMS-201038 or implitapide, in combination with a fibrate, such as fenofibrate. The MTP inhibitors can be administered at certain lower dosages that are still therapeutically effective when combined with a fibrate but yet create fewer or reduced adverse effects when compared to therapies using therapeutically effective dosages of the MTP inhibitors during monotherapy.

[0010] In one aspect, the invention provides a method of reducing the concentration of cholesterol and/or triglycerides in the blood of a mammal, and/or the amount of a marker of atherosclerosis in a mammal. The method comprises a combination therapy including a combination of a fibrate and

this protocol, BMS-201038 initially is administered at a first dosage in the range of 1 to 5 mg/day for at least 4 weeks, is then administered at a second dosage in the range of 3 to 7 mg/day for at least 4 weeks, and is then administered at a third dosage in the range of 6 to 9 mg/day for at least 4 weeks. Optionally, the method further comprises administering a fourth dosage of BMS-201038 in the range of 9 to 12 mg/day for at least 4 weeks. Optionally, the method further comprises administering a fifth dosage of BMS-201038 in the range of 12 to 17 mg/day for at least 4 weeks.

**[0011]** In one embodiment, the first dosage of BMS-201, 038 is 2.5 mg/day. In another embodiment, the second dosage is 5 mg/day. In another embodiment, the third dosage is 7.5 mg/day. In another embodiment, the optional fourth dosage is 10 mg/day. In another embodiment, the optional fifth dosage is 15 mg/day. Furthermore, the fibrate is administered at a dosage of 25 to 500 mg/day, optionally at a dosage of 25 to 250 mg/day, and optionally at a dosage of 100 to 200 mg/day. In certain embodiments, the fibrate is administered at a dosage of 160 mg/day.

**[0012]** The fibrate and BMS-201038 can be administered together in the same dosage form, or in different dosage forms. In the case of the separate dosage forms, the fibrate can be administered before, after, or simultaneously with BMS-201038.

**[0013]** The foregoing method may reduce the concentration of at least one of cholesterol and triglycerides in the blood but with a reduced incidence of an adverse event as compared to administration of a dosage of 25 mg/day of BMS-201038 in monotherapy. In addition, the method may reduce the number or amount of plaques on a wall of a blood vessel of the mammal but with a reduced incidence of an adverse event as compared to administration of a dosage of 25 mg/day of BMS-201038 in monotherapy. Contemplated adverse events include, for example, gastrointestinal disturbances, abnormalities in liver function, and hepatic steatosis.

**[0014]** In another aspect, the invention provides a method of reducing the concentration of cholesterol and/or triglycerides in the blood of a mammal, and/or the amount of a marker of atherosclerosis in a mammal. The method comprises administering each day to the mammal a combination of a fibrate and implitapide.

**[0015]** The implitapide can be administered at a dosage in the range of 0.01 to 60 mg/day. It is understood that the implitapide preferably is administered at a dosage in the range of 20-60 mg/day, for example, 20 mg/day, 25 mg/day, 30 mg/day, 35 mg/day, 40 mg/day, 45 mg/day, 50 mg/day, 55 mg/day or even 60 mg/day. The fibrate can be administered at a dosage of 25 to 250 mg/day, and optionally in the range of 100 to 200 mg/day. In one embodiment, the fibrate is administered at a dosage of 160 mg/day. The implitapide and fibrate can be administered together in the same dosage form or in different dosage forms. In the case of separate dosage forms, the fibrate can be administered before, after, or simultaneously with implitapide.

**[0016]** This method may reduce the concentration of at least one of cholesterol and triglycerides in the blood but with a reduced incidence of an adverse event as compared to administration of a dosage of 80 mg/day or greater of implitapide, for example, 80 mg/day and 160 mg/day, during monotherapy. Furthermore, this method may reduce the num-

ber of adverse events, for example, compared to administration of a dosage of 80 mg/day or greater of implitapide, for example, 80 mg/day or 160 mg/day, during monotherapy.

**[0017]** The foregoing methods can be used to treat (i) patients with hyperlipidemia, for example, hypercholesterolemia (for example, homozygous or heterozygous familial hypercholesterolemia) or hypertriglyceridemia, (ii) patients resistant to statin monotherapy, (iii) statin-intolerant patients, and/or (iv) patients having a combination of (i) and (ii), (i) and (iii), (ii) and (iii), and (i), (ii) and (iii).

## DETAILED DESCRIPTION

**[0018]** This invention relates, in part, to methods of reducing at least one of (i) the concentration of cholesterol and/or triglycerides in the blood of a mammal, and (ii) the amount of a marker of atherosclerosis in a mammal. The methods are based on combination therapies where an MTP inhibitor, for example, BMS-201038 or implitapide, is administered with a fibrate, for example, fenofibrate. The disclosed methods use lower dosages of the MTP inhibitor but, which in combination with the fibrate, can be effective at reducing the concentration of cholesterol and/or triglycerides in the blood but with fewer adverse events, less severe adverse events and/or reduced frequency of adverse events resulting from the use of higher dosages of the MTP inhibitor during monotherapy.

## 1. DEFINITIONS

**[0019]** For convenience, certain terms used in the specification, examples, and appended claims are collected in this section.

**[0020]** The phrase “combination therapy,” as used herein, refers to co-administering an MTP inhibitor, for example, BMS-201038 and implitapide, or a combination thereof, and a fibrate, for example, fenofibrate, as part of a specific treatment regimen intended to provide the beneficial effect from the co-action of these therapeutic agents. The beneficial effect of the combination includes, but is not limited to, pharmacokinetic or pharmacodynamic co-action resulting from the combination of therapeutic agents. Administration of these therapeutic agents in combination typically is carried out over a defined time period (usually weeks, months or years myasthenia depending upon the combination selected). Combination therapy is intended to embrace administration of multiple therapeutic agents in a sequential manner, that is, wherein each therapeutic agent is administered at a different time, as well as administration of these therapeutic agents, or at least two of the therapeutic agents, in a substantially simultaneous manner. Substantially simultaneous administration can be accomplished, for example, by administering to the subject a single tablet or capsule having a fixed ratio of each therapeutic agent or in multiple, single capsules for each of the therapeutic agents. Sequential or substantially simultaneous administration of each therapeutic agent can be effected by any appropriate route including, but not limited to, oral routes, intravenous routes, intramuscular routes, and direct absorption through mucous membrane tissues. The therapeutic agents can be administered by the same route or by different routes. For example, a first therapeutic agent of the combination selected may be administered by intravenous injection while the other therapeutic agents of the combination may be administered orally. Alternatively, for example, all therapeutic agents may be administered orally.

[0021] Combination therapy also can embrace the administration of the therapeutic agents as described above in further combination with other biologically active ingredients and non-drug therapies. Where the combination therapy further comprises a non-drug treatment, the non-drug treatment may be conducted at any suitable time so long as a beneficial effect from the co-action of the combination of the therapeutic agents and non-drug treatment is achieved. For example, in appropriate cases, the beneficial effect is still achieved when the non-drug treatment is temporally removed from the administration of the therapeutic agents, perhaps by days or even weeks.

[0022] The components of the combination may be administered to a patient simultaneously or sequentially. It will be appreciated that the components may be present in the same pharmaceutically acceptable carrier and, therefore, are administered simultaneously. Alternatively, the active ingredients may be present in separate pharmaceutical carriers, such as, conventional oral dosage forms, that can be administered either simultaneously or sequentially.

[0023] The terms, "individual," "patient," or "subject" are used interchangeably herein and include any mammal, including animals, for example, primates, for example, humans, and other animals, for example, dogs, cats, swine, cattle, sheep, and horses. The compounds of the invention can be administered to a mammal, such as a human, but can also be other mammals, for example, an animal in need of veterinary treatment, for example, domestic animals (for example, dogs, cats, and the like), farm animals (for example, cows, sheep, pigs, horses, and the like) and laboratory animals (for example, rats, mice, guinea pigs, and the like).

[0024] The term, "patient resistant to statin monotherapy," as used herein includes those patients for whom conventional statin monotherapy has been found ineffective or less effective than desired. A physician designing lipid reduction therapy for a patient will be able to determine via diagnosis and observation of periodic blood cholesterol and/or triglyceride levels whether such a patient is or has been resistant to statin monotherapy.

[0025] The term, "statin-intolerant patient," as used herein includes those patients for whom conventional statin therapy, for example, for serum lipid reduction, has been found to be ineffective and/or for whom an effective lipid-reducing dose of statins is too high to be tolerated or that there is an unacceptable adverse event associated with a particular dose. For example, statin therapy may be discontinued by the physician/patient due to concern over an adverse event such as Liver Function Test abnormality, muscle aches and pains or inflammation—myalgia or myositis, elevation in enzymes (CK) showing muscle adverse event. A physician designing lipid reduction therapy for a patient will be able to determine via diagnosis and observation of periodic blood cholesterol and/or triglyceride levels whether such a patient is statin-intolerant.

[0026] The phrase "minimizing adverse effects," "reducing adverse events," or "reduced adverse events," as used herein refer to an amelioration or elimination of one or more undesired side effects associated with the use of MTP inhibitors of the present invention. Side effects of traditional use of the MTP inhibitors include, without limitation, nausea, gastrointestinal disorders, steatorrhea, abdominal cramping, distention, elevated liver function tests, fatty liver (hepatic steatosis), hepatic fat build-up, and hypoglycemia.

rhinitis, dizziness, arthritis, peripheral edema, gastroenteritis, liver function tests abnormal, colitis, rectal hemorrhage, esophagitis, eructation, stomatitis, biliary pain, cheilitis, duodenal ulcer, dysphagia, enteritis, melena, gum hemorrhage, stomach ulcer, tenesmus, ulcerative stomatitis, hepatitis, pancreatitis, cholestatic jaundice, paresthesia, amnesia, libido decreased, emotional lability, incoordination, torticollis, facial paralysis, hyperkinesia, depression, hypesthesia, hypertonia, leg cramps, bursitis, tenosynovitis, myasthenia, tendinous contracture, myositis, hyperglycemia, creatine phosphokinase increased, gout, weight gain, hypoglycemia, anaphylaxis, angioneurotic edema, and bullous rashes (including erythema multiforme, Stevens-Johnson syndrome, and toxic epidermal necrolysis). Accordingly, the methods described herein provide an effective therapy while at the same time causing fewer or less significant adverse events.

[0027] In certain embodiments, side effects are partially eliminated. As used herein, the phrase "partially eliminated" refers to a reduction in the severity, extent, or duration of the particular side effect by at least 30%, 40%, 50%, 60%, 70%, 80%, 90%, 95% and 99% relative to that found by administering 25 mg/day of BMS-201038 during monotherapy or either 80 mg/day or 160 mg/day of implitapide during monotherapy. In certain embodiments, side effects are completely eliminated. Those skilled in the art are credited with the ability to detect and grade the severity, extent, or duration of side effects as well as the degree of amelioration of a side effect. In some embodiments, two or more side effects are ameliorated.

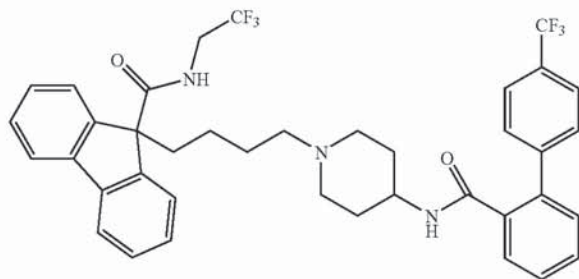
[0028] The term, "therapeutically effective" refers to the ability of an active ingredient, for example, BMS-201038 and implitapide, to elicit the biological or medical response that is being sought by a researcher, veterinarian, medical doctor or other clinician. Non-limiting examples include reduction of cholesterol (for example, LDL-C) and/or triglyceride levels in a patient, reduction of the amount of plaques, for example, arterial plaques, on the wall of a blood vessel, and the like.

[0029] The term, "therapeutically effective amount" includes the amount of an active ingredient, for example, BMS-201038 and implitapide, that will elicit the biological or medical response that is being sought by the researcher, veterinarian, medical doctor or other clinician. The compounds of the invention are administered in amounts effective at lowering the cholesterol concentration in the blood, and/or the triglyceride concentration in the blood and/or reducing the amount of plaques, for example, arterial plaques disposed upon the blood contacting wall of one or more blood vessels. Alternatively, a therapeutically effective amount of an active ingredient is the quantity of the compound required to achieve a desired therapeutic and/or prophylactic effect, such as the amount of the active ingredient that results in the prevention of or a decrease in the symptoms associated with the condition (for example, to meet an end-point).

[0030] The terms, "pharmaceutically acceptable" or "pharmacologically acceptable" refer to molecular entities and compositions that do not produce an adverse, allergic or other untoward reaction when administered to an animal, or to a human, as appropriate. The term, "pharmaceutically acceptable carrier" includes any and all solvents, dispersion media, coatings, antibacterial and antifungal agents, isotonic and absorption delaying agents and the like. The use of such media and agents for pharmaceutical active substances is well known in the art. Excipients for pharmaceutical use include

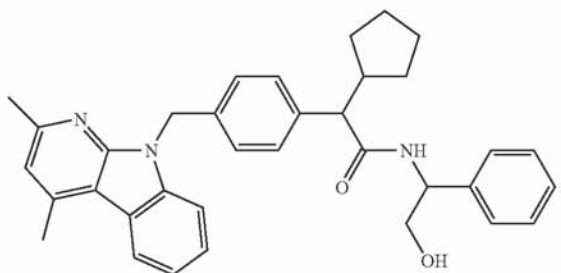
therapeutic compositions is contemplated. Supplementary active ingredients can also be incorporated into the compositions.

**[0031]** As used herein, the phrase, “BMS-201038” refers to a compound known as N-(2,2,2-Trifluorethyl)-9-[4-[4-[[[4’-(trifluoromethyl)[1,1’biphenyl]-2-Yl]carbonyl]amino]-1-piperidiny]butyl]9H-fluorene-9-carboxamide, having the formula:



stereoisomers thereof, and/or pharmaceutically acceptable salts or esters thereof.

**[0032]** As used herein, the phrase “implitapide” refers to a compound known as (2S)-2-cyclopentyl-2-[4-[(2,4-dimethyl-9H-pyrido[2,3-b]indol-9-yl)methyl]phenyl]-N-[(1S)-2-hydroxy-1-phenylethyl]ethanamide and having the structure shown below:



stereoisomers thereof, and/or pharmaceutically acceptable salts or esters thereof.

**[0033]** Pharmaceutically acceptable salts of the foregoing compounds can be synthesized, for example, from the parent compound, which contains a basic or acidic moiety, by conventional chemical methods. Generally, such salts can be prepared by reacting the free acid or base forms of these compounds with a stoichiometric amount of the appropriate base or acid in water or in an organic solvent, or in a mixture of the two; generally, non-aqueous media like ether, ethyl acetate, ethanol, isopropanol, or acetonitrile are preferred. Lists of suitable salts are found in Remington’s Pharmaceutical Sciences, 20th ed., Lippincott Williams & Wilkins, Baltimore, Md., 2000, p. 704.

**[0034]** As used herein, the term “stereoisomers” refers to compounds made up of the same atoms bonded by the same bonds but having different spatial structures which are not interchangeable. The three-dimensional structures are called configurations. As used herein, the term “enantiomers” refers

“racemic mixture” or “racemic modification” refer to a mixture of equal parts of enantiomers.

## 2. METHODS OF THE INVENTION

**[0035]** In general the invention provides methods for treating hyperlipidemia using one or more MTP inhibitors, for example, BMS-201038 or implitapide. The MTP inhibitors can be used at dosages lower than those already found to result in one or more adverse events, for example, gastrointestinal disorders, abnormalities in liver functional and/or hepatic steatosis (for example, 25 mg/day of BMS-201038, 80 mg/day of implitapide and 160 mg/day of implitapide have been found to cause gastrointestinal disorders, abnormalities in liver function and/or hepatic steatosis) but are still are therapeutically effective when combined with a fibrate, for example, fenofibrate.

**[0036]** (a) Combination Therapies Using BMS-201038 and Fibrate

**[0037]** In certain aspects, the invention provides a method of reducing at least one of (i) the concentration of cholesterol and/or triglycerides in the blood of a mammal, and (ii) the amount of a marker of atherosclerosis in the blood stream of a mammal. The method comprises a combination therapy, which can be achieved by co-administering to the mammal, each day, a fibrate and BMS-201038. In one protocol, BMS-201038 initially is administered at a first dosage in the range of 1 to 5 mg/day for at least 4 weeks, is then administered at a second dosage in the range of 3 to 7 mg/day for at least 4 weeks, and is then administered at a third dosage in the range of 6 to 9 mg/day for at least 4 weeks. The protocol may optionally include a fourth dosage in the range of 9 to 12 mg/kg for at least 4 weeks. The protocol may optionally include a fifth dosage in the range of 12 to 17 mg/kg for at least 4 weeks.

**[0038]** The first dosage of BMS-201038 can be for example 2.5 mg/day. The second dosage of BMS-201038 can be 5 mg/day. The third dosage of BMS-201038 can be 7.5 mg/day. The optional fourth dosage can be 10 mg/day. The optional fifth dosage can be 15 mg/day. In certain embodiments, the second dosage is administered immediately following the first dosage, i.e., the second dosage is administered starting at five weeks from the initial first dosage. Similarly, in certain other embodiments, the third dosage of BMS-201038 is administered immediately following the second dosage, e.g., the second dosage is administered at nine weeks from the initial first dosage. Similarly, in certain other embodiments the fourth dosage is administered immediately following the third dosage, e.g., the fourth dosage is administered at thirteen weeks from the initial first dosage. Similarly, in certain other embodiments the fifth dosage is administered immediately following the fourth dosage, e.g., the fifth dosage is administered at seventeen weeks from the initial first dosage.

**[0039]** In this approach, BMS-201038 is administered with a fibrate. Exemplary fibrates include fenofibrate (also known as Tricor), bezafibrate, ciprofibrate, clofibrate and gemfibrozil (also known as Lopid). The fibrate is administered at a dosage of 25 to 500 mg/day, optionally at a dosage of 25 to 250 mg/day, and optionally at a dosage of 100 to 200 mg/day. In certain embodiments, the fibrate is administered at a dosage of 160 mg/day. The fibrate and BMS-201038 can be administered together in the same dosage form, or in different dosage forms. In the case of the separate dosage forms, the fibrate can be administered before, after, simultaneously,

# Explore Litigation Insights

Docket Alarm provides insights to develop a more informed litigation strategy and the peace of mind of knowing you're on top of things.

## Real-Time Litigation Alerts



Keep your litigation team up-to-date with **real-time alerts** and advanced team management tools built for the enterprise, all while greatly reducing PACER spend.

Our comprehensive service means we can handle Federal, State, and Administrative courts across the country.

## Advanced Docket Research



With over 230 million records, Docket Alarm's cloud-native docket research platform finds what other services can't. Coverage includes Federal, State, plus PTAB, TTAB, ITC and NLRB decisions, all in one place.

Identify arguments that have been successful in the past with full text, pinpoint searching. Link to case law cited within any court document via Fastcase.

## Analytics At Your Fingertips



Learn what happened the last time a particular judge, opposing counsel or company faced cases similar to yours.

Advanced out-of-the-box PTAB and TTAB analytics are always at your fingertips.

## API

Docket Alarm offers a powerful API (application programming interface) to developers that want to integrate case filings into their apps.

## LAW FIRMS

Build custom dashboards for your attorneys and clients with live data direct from the court.

Automate many repetitive legal tasks like conflict checks, document management, and marketing.

## FINANCIAL INSTITUTIONS

Litigation and bankruptcy checks for companies and debtors.

## E-DISCOVERY AND LEGAL VENDORS

Sync your system to PACER to automate legal marketing.