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CA 2363053 C 2011/01/25

(11)(21) **2 363 053**

(12) **BREVET CANADIEN
CANADIAN PATENT**

(13) **C**

(22) Date de dépôt/Filing Date: 2001/11/09

(41) Mise à la disp. pub./Open to Public Insp.: 2003/05/09

(45) Date de délivrance/Issue Date: 2011/01/25

(51) Cl.Int./Int.Cl. *A61K 31/4365* (2006.01),
A61K 9/20 (2006.01), *A61P 7/02* (2006.01)

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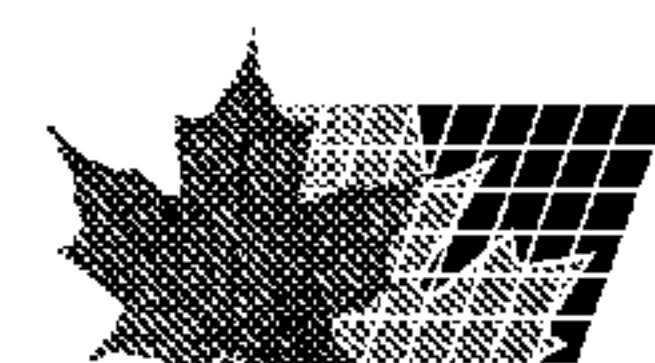
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(54) Titre : FORMULATION POUR PASTILLES DE BISULFATE DE CLOPIDOGREL

(54) Title: CLOPIDOGREL BISULFATE TABLET FORMULATION

(57) Abrégé/Abstract:

Pharmaceutical tablets comprising clopidogrel bisulfate and a lubricant selected from zinc stearate, stearic acid, and sodium stearyl fumarate.



ABSTRACT

5 Pharmaceutical tablets comprising clopidogrel bisulfate and a lubricant
selected from zinc stearate, stearic acid, and sodium stearyl fumarate.

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CLOPIDOGREL BISULFATE TABLET FORMULATION

Background of the Invention

5 Clopidogrel is a compound disclosed in U.S. patent 4847265 to be therapeutically useful as an inhibitor of platelet agglomeration.

Tablets containing clopidogrel as clopidogrel bisulfate are sold in the United States and elsewhere under the tradename PlavixTM.

10 According to the labelling of PlavixTM, each tablet contains 98 mg of clopidogrel bisulfate, which is the molar equivalent of 75 mg of clopidogrel. The tablets are made as cores which are then film coated. The cores comprise, in addition to the clopidogrel bisulfate, the following inactive
15 ingredients: lactose, hydrogenated castor oil, microcrystalline cellulose, polyethylene glycol 6000 and pregelatinized starch. The weight of each tablet is about 240 mg.

Lactose and microcrystalline cellulose are ingredients that are commonly
20 used as fillers and binders. Starch is commonly used as a disintegrant to cause the tablets to disintegrate in gastrointestinal fluid. Hydrogenated castor oil and polyethylene glycol are sometimes, although rarely, used as lubricants.

In the manufacture of tablets, it is generally necessary to include a lubricant in
25 the mix of ingredients to prevent sticking of the compressed tablets to the punches, and also to prevent binding between the punches and the dies.

The most commonly used lubricant is magnesium stearate. U.S. patent
30 4847265, in examples 1 and 2, shows the use of magnesium stearate as the lubricant.

TM - registered trademark

5 It is thus unusual that PlavixTM tablets do not contain magnesium stearate as the lubricant, but instead contain hydrogenated castor oil and polyethylene glycol as lubricants. Both of these ingredients have occasionally been used as lubricants, but they are not as effective as magnesium stearate, and their use in place of magnesium stearate must be expected to make it difficult to produce tablets at high speed without experiencing sticking to the punches or binding between the punches and dies. It appears that the reason that magnesium stearate is not used in PlavixTM tablets is that there is an interaction between clopidogrel bisulfate and magnesium stearate that causes degradation of the clopidogrel bisulfate, so that the stability of clopidogrel sulfate tablets which include magnesium stearate as lubricant is insufficient to enable sale of such tablets.

15 In light of this prior art, the object of the present invention is to provide stable clopidogrel hydrochloride tablets that contain a lubricant that is more effective than hydrogenated castor oil and polyethylene glycol 6000.

20 Brief Summary of the Invention

25 It has surprisingly been found that clopidogrel bisulfate tablets, comprising a lubricant selected from the group consisting of zinc stearate, stearic acid, and sodium stearyl fumarate, are much more stable than tablets comprising either magnesium stearate or calcium stearate as lubricant. Zinc stearate, stearic acid and sodium stearyl fumarate are also more effective lubricants than either hydrogenated castor oil or polyethylene glycol 6000. The invention is thus pharmaceutical tablets comprising clopidogrel bisulfate and a lubricant selected from zinc stearate, stearic acid and sodium stearyl fumarate.

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Illustrative embodiments of the present invention provide a pharmaceutical tablet which comprises clopidogrel bisulfate and a lubricant selected from the group consisting of zinc stearate, sodium stearyl fumarate and stearic acid.

Illustrative embodiments of the present invention provide a tablet described herein which comprises zinc stearate.

Illustrative embodiments of the present invention provide a tablet described herein which comprises sodium stearyl fumarate.

Illustrative embodiments of the present invention provide a tablet described herein which comprises stearic acid.

Illustrative embodiments of the present invention provide a tablet described herein which does not comprise microcrystalline cellulose.

Illustrative embodiments of the present invention provide a tablet described herein further comprising lactose.

Illustrative embodiments of the present invention provide a tablet described herein further comprising methylcellulose.

Illustrative embodiments of the present invention provide a tablet described herein further comprising crospovidone.

Illustrative embodiments of the present invention provide a tablet described herein further comprising croscarmellose.

Illustrative embodiments of the present invention provide a tablet described herein further comprising sodium starch glycolate.

Detailed Description of the Invention

5 Pharmaceutical tablets comprising clopidogrel bisulfate will generally contain,
in addition to the active ingredient, one or more ingredients that serve as
fillers and binders such as, for example, lactose, microcrystalline cellulose, or
methylcellulose. It has also surprisingly been found that microcrystalline
cellulose inhibits dissolution of clopidogrel bisulfate, so that the tablets will
10 preferably exclude microcrystalline cellulose. The tablets will preferably
comprise lactose or methylcellulose.

The tablets will further comprise at least one ingredient as a lubricant to avoid
sticking to the tooling in the tableting process. As aforesaid, it has been
found that improved stability is achieved by selecting the lubricant from zinc
15 stearate, stearic acid and sodium stearyl fumarate.

When the lubricant is zinc stearate or sodium stearyl fumarate, the amount
will preferably be from about 0.5% to about 3% of the weight of the tablet.
When the lubricant is stearic acid, the amount will preferably be from about
20 1% to about 6% of the weight of the tablet.

Clopidogrel bisulfate tablets will preferably further comprise a disintegrant to
cause the tablets to swell and disintegrate in gastrointestinal fluid after
ingestion. As aforesaid, PlavixTM tablets comprise starch which serves as a
25 disintegrant, but is desirable to use a more efficient disintegrant; i.e. one that
is effective in a smaller quantity. Preferred disintegrants are croscarmellose
sodium, sodium starch glycolate, and crospovidone.

30 The amount of disintegrant will preferably be from about 0.5% to about 10% of
the weight of the tablet.

The tablets of this invention will optionally also comprise other ingredients, such as, for example, colloidal silicon dioxide as a glidant.

5 The tablets will also optionally be coated with a film coat.

10 The clopidogrel bisulfate tablets of this invention can be prepared by conventional tablet forming techniques such as, for example, wet granulation and dry granulation. In the wet granulation process, the active ingredient is mixed with some or all of the filler. This blend is then wet granulated with water or an organic solvent, optionally containing a binder in solution. The resultant wet granulation is then dried and milled. The granules are then mixed with the remaining ingredients, which will include the lubricant, to produce the final mix, which is then compressed into tablets.

15 In the dry granulation process, the active ingredient is mixed with the other ingredients without addition of any solvent, and thus without the need for drying. Again the final mix is compressed into tablets. The dry granulation approach is preferred as it is simpler and thus less costly.

20 The invention will be further understood from the following examples:

Example No.:	1	2	3	4	5
25 Clopidogrel Bisulfate	98.0	98.0	98.0	98.0	98.0
Anhydrous Lactose	49.0	49.0	49.0	49.0	49.0
Methylcellulose 15CPS	17.0	17.0	17.0	17.0	17.0
Crospovidone	12.0	12.0	12.0	12.0	12.0
Magnesium Stearate	3.8	X	X	X	X
Calcium Stearate	X	3.8	X	X	X
30 Zinc Stearate	X	X	3.8	X	X
Sodium Stearyl Fumarate	X	X	X	3.8	X
Stearic Acid	X	X	X	X	8.8
Colloidal Silicon Dioxide	0.2	0.2	0.2	0.2	0.2
	180.	180.	180.	180.	185.

5 For each of the 5 examples, the ingredients in the proportions listed were mixed together. The powder mixture was then compressed into tablets of weight 180 mg for examples 1 to 4, and 185 mg for example 5. Because stearic acid is not as efficient a lubricant as the lubricants of the other examples, a larger quantity of lubricant was used in example 5 than in examples 1 to 4.

10 Sample tablets from each example were stored at 60°C for 2 weeks and then tested by an HPLC method to determine degradation products as a percentage of the initial clopidogrel bisulfate content.

The results were as follows:

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<u>Example No:</u>	<u>1</u>	<u>2</u>	<u>3</u>	<u>4</u>	<u>5</u>
Total Degradation Products	3.8%	3.8%	1.3%	1.0%	1.2%

20 It thus can be seen that, when any of zinc stearate, sodium stearyl fumarate or stearic acid is used as lubricant, the degradation rate is substantially less than when either magnesium stearate or calcium stearate is used.

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Claims

1. A pharmaceutical tablet which comprises clopidogrel bisulfate and a lubricant selected from the group consisting of zinc stearate, sodium stearyl fumarate and stearic acid.
2. The tablet of claim 1 which comprises zinc stearate.
3. The tablet of claim 1 which comprises sodium stearyl fumarate.
4. The tablet of claim 1 which comprises stearic acid.
5. The tablet of any one of claims 1 to 4 which does not comprise microcrystalline cellulose.
6. The tablet of any one of claims 1 to 5 further comprising lactose.
7. The tablet of any one of claims 1 to 6 further comprising methylcellulose.
8. The tablet of any one of claims 1 to 7 further comprising crospovidone.
9. The tablet of any one of claims 1 to 7 further comprising croscarmellose.
10. The tablet of any one of claims 1 to 7 further comprising sodium starch glycolate.