CLAIMS

What is claimed is:

1. A solid dosage form comprising 300 mg or less of an API and at least one of croscarmellose sodium or microcrystalline cellulose, wherein the API is crystalline anhydrous (2S,3R,4R,5S,6R)-2-(4-chloro-3-(4-ethoxybenzyl)phenyl)-6-(methylthio)tetrahydro-2H-pyran-3,4,5-triol.

2. The solid dosage form of claim 1, wherein the API is crystalline anhydrous (2S,3R,4R,5S,6R)-2-(4-chloro-3-(4-ethoxybenzyl)phenyl)-6-(methylthio)tetrahydro-2H-pyran-3,4,5-triol Form 1.

3. The solid dosage form of claim 1, wherein the API is crystalline anhydrous (2S,3R,4R,5S,6R)-2-(4-chloro-3-(4-ethoxybenzyl)phenyl)-6-(methylthio)tetrahydro-2H-pyran-3,4,5-triol Form 2.

4. The solid dosage form of claim 1, which further comprises silicon dioxide.

5. The solid dosage form of claim 1, which is a tablet.

6. The solid dosage form of any of claims 1-5, which further comprises a second therapeutic agent, which second therapeutic agent is an anti-diabetic agent, anti-hyperglycemic agent, hypolipidemic/lipid lowering agent, anti-obesity agents, anti-hypertensive agent, or appetite suppressant.

7. The solid dose form of claim 6, wherein the second therapeutic agent is a DPP-4 inhibitor (e.g., sitagliptin, dutogliptin).

8. A granule comprising an API and at least one of croscarmellose sodium, collodial silicon dioxide, and microcrystalline cellulose, wherein the API is (2S,3R,4R,5S,6R)-2-(4-chloro-3-(4-ethoxybenzyl)phenyl)-6-(methylthio)tetrahydro-2H-pyran-3,4,5-triol.

9. The granule of claim 8, wherein the API is crystalline anhydrous (2S,3R,4R,5S,6R)-2-(4-chloro-3-(4-ethoxybenzyl)phenyl)-6-(methylthio)tetrahydro-2H-pyran-3,4,5-triol Form 1.

10. The granule of claim 8, wherein the API is crystalline anhydrous (2S,3R,4R,5S,6R)-2-(4-chloro-3-(4-ethoxybenzyl)phenyl)-6-(methylthio)tetrahydro-2H-pyran-3,4,5-triol Form 2.

~~11. The use of a dosage form of any of claims 1-7 in the manufacture of a medicament for improving the cardiovascular and metabolic health of a patient.~~

~~12.~~11. The use of a dosage form of any of claims 1-7 in the manufacture of a medicament for treating diabetes in a patient.

~~13.~~12. The use of claim ~~12~~11, wherein the patient has taken, or is currently taking, a second therapeutic agent, which second therapeutic agent is an anti-diabetic agent, anti-hyperglycemic agent, hypolipidemic/lipid lowering agent, anti-obesity agents, anti-hypertensive agent, or appetite suppressant.

~~14.~~13. The use of claim ~~13~~12, wherein the second medication is a biguanide (e.g., metformin, phenformin).