



(12) UK Patent (19) GB (11) 2 119 246 B

(54) Title of invention

Solid shaped articles

(51) INT CL⁴; A61K 9/00

(21) Application No
8232189

(22) Date of filing
11 Nov 1982

(30) Priority data

(31) **8136360**

(32) **2 Dec 1981**

(33) **United Kingdom (GB)**

(43) Application published
16 Nov 1983

(45) Patent published
22 May 1985

(52) Domestic classification
A5B 832 833 M N
C1B 3F1 3FX 4

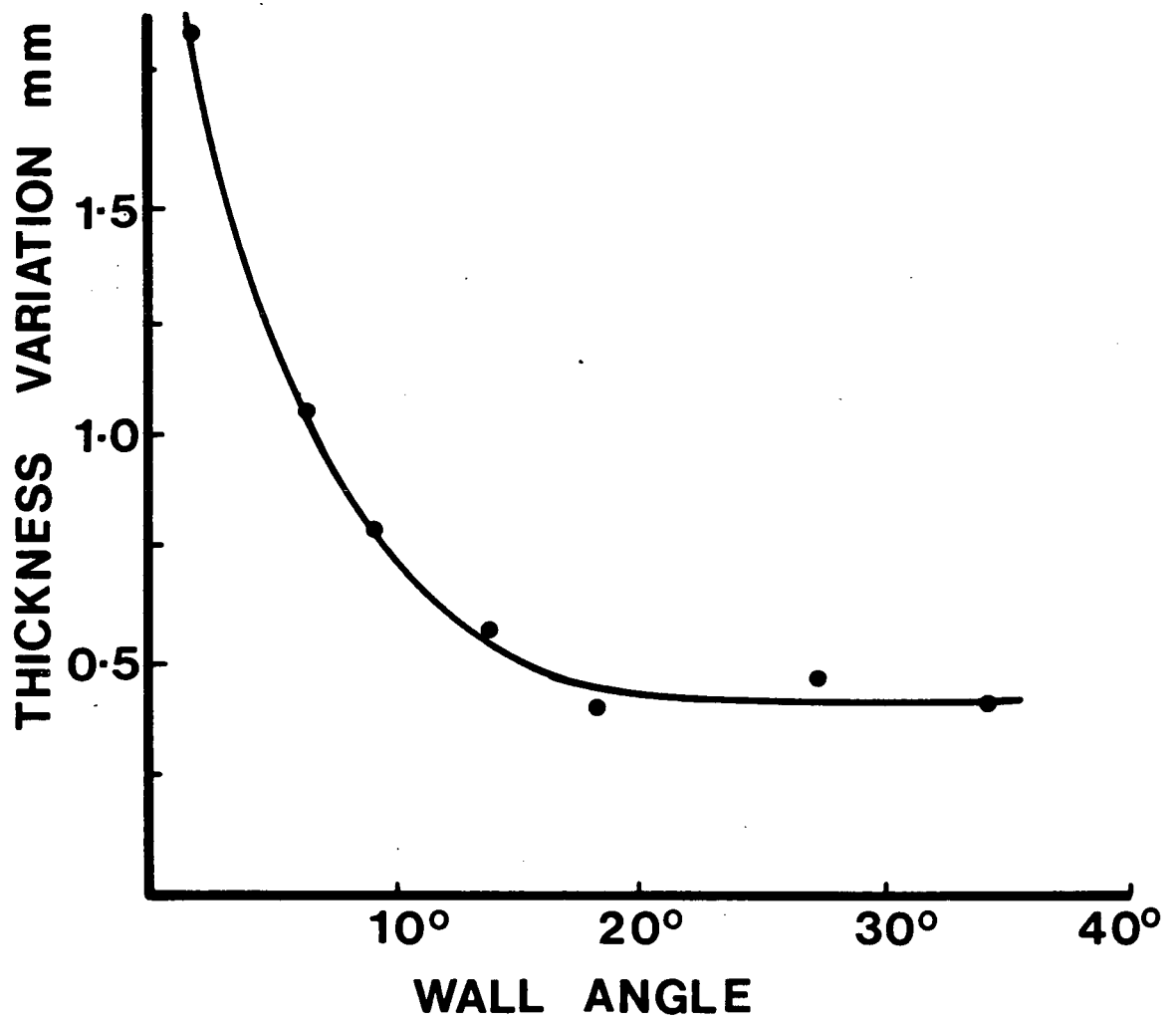
(56) Documents cited
None

(58) Field of search
A5B

(73) Proprietor
John Wyeth & Brother Limited
(United Kingdom),
Huntercombe Lane South
Taplow
Maidenhead
Berkshire SL6 0PH

(72) Inventor
George Keith Emerson Gregory

(74) Agent and/or
Address for Service
K. J. S. Brown,
c/o Wyeth Laboratories
Huntercombe Lane South
Taplow
Berks SL6 0PH



- 1 -

2119246

SOLID SHAPED ARTICLES

This invention relates to a solid shaped article carrying a predetermined unit quantity of chemical and to a novel process for preparing such an article.

It is known to produce shaped articles carrying
5 chemicals by a process involving sublimation of a solvent from a composition in the solid state comprising the chemical and a solution in a solvent of a carrier material. Such a process is described, for example, in our French Specification No. 7729663 (publication 2366835) and in our
10 UK Patent Specification 1,548,022. The process generally involves freezing the composition in a mould and then freeze drying the frozen composition. The mould can be, for example, a depression in a metal plate but it is preferably a depression in a sheet of filmic material.
15 The filmic material preferably contains more than one depression. The filmic material may be similar to that employed in conventional blister packs which are used for packaging oral contraceptive tablets and like medicament forms. For example the filmic material may be made of
20 thermoplastic material with the depressions formed by thermoforming. The preferred filmic material is a polyvinyl chloride film. Laminates such as polyvinyl chloride/polyvinylidene chloride, polyvinyl chloride/polytetrafluoroethylene or polyvinyl chloride/polyvinylidene chloride/polyethylene may also be used.
25 Our investigations of the process described in the above mentioned specifications have shown that one of the rate limiting steps in the production of the shaped article, especially on a commercial scale, is the time required for sublimation of the solvent in the freeze
30 drying apparatus. We have found that the drying time can be reduced substantially if the depth of the composition in the mould (prior to freezing) is kept to a minimum. In general, drying times tend to be proportional to the square of the thickness. We have found that if the mean
35 depth of composition in the mould is about 4.0 mm or less

(preferably about 3.5 mm or less) reasonable drying times can be attained. However, when shaped articles of this mean thickness were produced in conventional moulds with substantially vertical side walls, there was considerable variation in thickness in each sample if the moulds were not completely filled. The shaped articles in general had a minimum thickness near their centre and a maximum thickness near their edges, this variation apparently being derived from the meniscus effect of the composition at the side walls of the mould. The variation in thickness had disadvantages, for example, in handling the formed article and in obtaining consistent and minimal drying times in their preparation. It is an object of the present invention to alleviate these disadvantages.

Accordingly the present invention provides a process for preparing a solid shaped article carrying a predetermined unit quantity of a chemical which process comprises adding to a mould a composition comprising the predetermined amount of chemical and a solution in a solvent of a carrier material inert towards the chemical, the side wall or walls of the mould diverging outwards from the base and making an angle with the vertical of at least 5° at the surface of the composition and the mean depth of the composition in the mould being about 4.0mm or less, freezing the composition in the mould and then subliming solvent from the frozen composition so as to produce a network of carrier material carrying the chemical.

Preferably the side wall or walls slope outwardly from a substantially flat base and make a constant angle with the vertical, the angle being at least 5° . However, the side walls need not make a constant angle provided that the angle is at least 5° at the surface of the composition. For example the angle below the surface may be less than that at the surface. In addition, or alternatively, the

angle above the surface could be greater than the angle at the surface.

The mould is preferably a depression in a sheet of filmic material as hereinabove described. The chemical
5 may be any chemical (including, for example, agricultural or horticultural chemicals or chemical reagents) which it is desired to administer, dispense or utilise in predetermined unit quantities. Preferably the chemical is a pharmaceutical substance and the solid shaped article
10 carrying the predetermined unit quantity of pharmaceutical substance is a pharmaceutical dosage form, particularly a pharmaceutical dosage form suitable for oral administration. By choice of a carrier material that is water soluble or water dispersible it is possible to prepare shaped articles, including
15 pharmaceutical dosage forms, that are capable of being rapidly disintegrated by water, for example, within ten seconds (or preferably 5 seconds) when tested by, for example, the test procedure described in the above mentioned specifications. Examples of suitable carrier materials are
20 given in the above mentioned specifications. For example, the carrier may be formed from polypeptides such as gelatin, particularly gelatin which is partially hydrolysed, e.g. by heating in water. For example, the gelatin may be partially hydrolysed by heating a solution of the gelatin in water,
25 e.g. in an autoclave at about 120°C for up to 2 hours, e.g. from about 5 minutes to about 1 hour, preferably from about 30 minutes to about 1 hour. The hydrolysed gelatin is preferably used at concentrations of about 1 to 6% weight/vol., most preferably at 2 to 4% e.g. about 3%. Other carrier
30 materials may be used in place of partially hydrolysed gelatin for example polysaccharides such as hydrolysed dextran, dextrin and alginates (e.g. sodium alginate) or mixtures of above mentioned carriers with each other or with other carrier materials such as polyvinyl alcohol,
35 polyvinylpyrrolidone or acacia.

Besides the chemical and the carrier material the composition may contain other additional ingredients.

For example, when preparing pharmaceutical dosage forms the composition may include pharmaceutically acceptable adjuvants such as colouring agents, flavouring agents, preservatives and the like. In addition the composition
5 may contain ingredients which aid the preparation of the shaped articles. For example, the composition may include a surfactant, e.g. Tween 80 [polyoxyethylene (20) sorbitan mono-oleate; Tween is a registered Trade Mark], to aid in the dispersion of the chemical. The composition
10 may also include ingredients such as fillers (e.g. mannitol, sorbitol) which improve the physical properties of the shaped article.

The solvent for the composition is preferably water but it may contain a co-solvent (such as an alcohol) if it is
15 desired to improve the solubility of the chemical.

The ingredients for the composition may be filled into the mould individually or in admixture, preferably as a mixture comprising the complete composition.

The volume of the composition in the mould is
20 preferably about 3 ml or less (for example, about 0.25 to 1 ml) especially when the shaped article is a pharmaceutical dosage form for oral administration. The depth of the composition in the mould, the cross sectional area of the mould and the angle of the side wall or walls will
25 all affect the volume of the composition. The mould can be, for example, a regular shape such as a circle or a polygon (e.g. a rectangle particularly a square) in horizontal cross section with the area of the section at the base of the mould being less than the area nearer
30 the open top, in view of the sloping sides. Preferably the mould is circular in cross section and the formed shaped article resembles a flat disc having sloping side walls.

In order to keep drying times in the freeze dryer
35 to a minimum it is preferable that the depth of composition in the mould should be less than 3.5 mm, for

example, 1.5 (or more preferably 2) to 2.5 mm. If the depth is below about 1.5 mm the shaped articles tend to be difficult to handle.

5 The invention further provides a shaped article produced by the process of the invention. The invention particularly provides a solid shaped article produced by freezing a composition comprising a predetermined amount of chemical and a solution in a solvent of a carrier material inert towards the chemical and subliming solvent
10 from the frozen composition so as to produce a network of carrier material carrying the chemical characterised in that the shaped article has a mean thickness of about 4.0 mm or less and has side wall or walls diverging outwards from a bottom surface, said side walls making
15 an angle with the vertical of at least 5° at the upper surface.

In order to investigate the effect of the angle of the side wall or walls had on the variations in thickness of the formed shaped article, a number of blister trays
20 were thermoformed from 200 μ UPVC having depressions in them of varying profiles. The depressions were circular in cross section and the profiles, in general, were of inverted truncated cone shape with the side walls making various angles with the vertical. The dimensions of
25 the different profiles together with that of a conventionally formed flat base blister pack (control) are given below:-

TABLE I

	Profile	Diameter at base (mm)	Depth of depression (mm)	Upper diameter (mm)	Capacity (ml)	Angle of side wall to the vertical
5						
	1	12.0	3	13.0	0.36	9.5°
	2	11.5	3	13.0	0.35	14.0°
	3	11.0	3	13.0	0.34	18.5°
	4	10.0	3.5	14.6	0.41	33.0°
10	5	9.0	3.5	14.8	0.39	39.5°
	6	14.2	3.5	15.0	0.59	6.5°
	control	14.5	6.0	15.0	1.0	2°

An aqueous composition having the following constitution was prepared:

15 3% hydrolysed gelatin
 3% mannitol B.P.

(The 3% hydrolysed gelatin is prepared by suspending 30 g. of powdered gelatin in 800 ml of water, autoclaving it at 121°C for 60 minutes and adjusting the final volume to 1 litre).

Varying amounts of the composition with and without the addition of a surfactant (1% Tween 80) were added to the depressions (i.e. moulds) of Table I. The compositions were then frozen and the solvent sublimed in a freeze dryer to give solid shaped articles, samples of which, when tested by the procedure in the aforementioned specifications, disintegrated rapidly in water. The dimensions of the shaped articles were determined using a micrometer system and the variation in thickness of the individual samples was calculated. The mean maximum thickness

variation for the different samples is shown in the following Table II and is illustrated graphically in Figure 1 of the accompanying drawings.

TABLE II		
5 Profile	Angle of side wall to the vertical	Mean maximum thickness variation (mm)
1	9.5°	0.8
2	14.0°	0.57
10 3	18.5°	0.4
4	33.0°	0.48
5	39.5°	0.42
6	6.5°	1.07
Control	2°	1.88

15 The results show that there was considerable variation in thickness of the samples when the side walls of the mould approached the vertical as, for example in the control. The maximum thickness was at the edge and the minimum thickness was at the centre in these samples,

20 this variation being apparently derived from the meniscus effect although it was not significantly affected by the presence of surfactant. The articles made in depressions in which the side walls made angles with the vertical of considerably more than 5° had much flatter top surfaces.

25 In many of these latter cases the small variation in thickness that did occur was the result of an increased thickness in the centre (a "blip") apparently produced during the freezing procedure. The variations in thickness due to the meniscus effect could be avoided by filling the moulds

30 completely to their upper surface, but, using such full moulds can give rise to further difficulties in that there can be spillage during filling or when the moulds are subsequently handled while the contents are still

liquid. In addition, further difficulties can arise if a cover layer is applied over the freeze dried articles in the moulds; the articles may stick to the cover layer if their upper surfaces are substantially level
5 with the upper surfaces of the moulds.

In the present invention a mould is used in which the side wall or walls make an angle with the vertical of at least 5°. The angle with the vertical may be, for example, from 5° to about 40° and the advantage of minimal
10 thickness variation is retained. Especially preferred angles for this advantage as shown in the above profiles, are from about 9° to 40°. However, if the side walls make too big an angle with the vertical the moulds waste unnecessary space in the freeze drier and also the
15 products produced have sharp edges which make them difficult to handle. Accordingly it is preferred that the angle made with the vertical is within the range 9° to 20°, for example 10° to 15°. A particularly preferred angle is about 12°.

20 The following Examples illustrate the invention:

EXAMPLE 1

Pharmaceutical Dosage Forms containing 15 mg oxazepam

Formulation:

25	Oxazepam	15 mg
	Tween 80 BPC	0.25 mg
	Mannitol BP	15 mg
	3% hydrolysed gelatin to	0.5 ml

The 3% hydrolysed gelatin is prepared by suspending 30 g of powdered gelatin in 800 ml of cold distilled water in a 1 litre flask and autoclaving it at 121°C for 60 minutes. When cool, the final volume is adjusted to 1 litre.

Oxazepam (30 g) is suspended in 3% hydrolysed gelatin solution containing dissolved mannitol (30 g) and Tween 80 (0.5 g) using ultrasonics for 5 minutes and the suspension made up to 1 litre with 3% gelatin solution. 0.5 ml portions of the suspension are dosed, using an automatic filling machine, into pockets in polyvinyl chloride blister trays. Each of the pockets in the blister tray is of inverted truncated conical shape of 3.5 mm depth, a bottom diameter of 14.5 mm and a top diameter of 16 mm (i.e. the side walls make an angle with the vertical of about 12°). The contents of the pockets are then frozen by passing the trays through a freeze tunnel into which liquid nitrogen is injected.

The blister trays containing the frozen compositions are then transferred to a freeze drier. The pressure is adjusted to 0.5 mm Hg. The temperature of the shelves in the freeze drier is set at 60°C for 1 hour and then lowered to 40°C. After 2 hours the trays are removed from the freeze drier. A peelable aluminium foil is then sealed to the blister pack around the depressions containing the pharmaceutical dosage forms. The pharmaceutical dosage forms are of substantially uniform thickness and disintegrate rapidly, for example, in two seconds or less, when taken orally.

30

EXAMPLES 2 TO 11

The procedure of Example 1 is followed to prepare pharmaceutical dosage forms containing the following active ingredients:

Example

	2	Oxazepam 30 mg and 50 mg
	3	Lorazepam 1, 2, 2.5 and 4 mg
	4	Temazepam 10 and 20 mg
	5	Lormetazepam 1 mg
5	6	Frusemide 40 mg
	7	Bendrofluazide 5 mg
	8	Cyclopenthiazide 0.5 mg
	9	Isosobide dinitrate 2.5, 5 and 10 mg
	10	Indomethacin 25 and 50 mg
10	11	Prochlorperazine maleate 50 mg

EXAMPLE 12

Formulation:

	Indole acetic acid	1 mg
	Mannitol BP	15 mg
15	3% hydrolysed gelatin	to 0.5 ml

The procedure of Example 1 is followed substituting 2g of indole acetic acid for the 30g of oxazepam and omitting the Tween 80. Each of the resulting freeze dried products can be added to 1 litre of water to
20 give a composition useful as a plant growth promoter.

EXAMPLE 13

The procedure of Example 12 is repeated replacing the indole acetic acid by indole butyric acid. Each of the freeze dried products containing 1 mg
5 of indole butyric acid may be added to 1 litre of water to give a composition useful as a rooting growth promoter for plant cuttings.

CLAIMS

1. A process for preparing a solid shaped article carrying a predetermined unit quantity of a chemical which process comprises adding to a mould a composition comprising the predetermined amount of chemical and a solution in a solvent of a carrier material inert towards the chemical, the side wall or walls of the mould diverging outwards from the base and making an angle with the vertical of at least 5° at the surface of the composition and the mean depth of the composition in the mould being about 4.0mm or less, freezing the composition in the mould and then subliming solvent from the frozen composition so as to produce a network of carrier material carrying the chemical.

2. A process as claimed in Claim 1 wherein the volume of the composition in the mould is about 3 ml or less.

3. A process as claimed in claim 1 wherein the volume of the composition in the mould is about 0.25 to 1 ml.

4. A process as claimed in any one of the preceding claims wherein the depth of the composition in the mould is less than 3.5 mm.

5. A process as claimed in any one of the preceding claims wherein the depth of the composition in the mould is 1.5 to 2.5 mm.

6. A process as claimed in any one of the preceding claims wherein the angle with the vertical is from 9° to 20°.

7. A process as claimed in any one of the preceding claims wherein the angle with the vertical is from 10° to 15° .

5 8. A process as claimed in any one of the preceding claims wherein the mould is a depression in a sheet of filmic material.

9. A process as claimed in any one of the preceding claims wherein the carrier material is hydrolysed gelatin.

10. A process as claimed in any one of the preceding claims wherein the chemical is a pharmaceutical substance and the solid shaped article is a pharmaceutical dosage form.

5 11. A solid shaped article produced by freezing a composition comprising a predetermined amount of chemical and a solution in a solvent of a carrier material inert towards the chemical and subliming solvent from the frozen composition so as to produce
10 a network of carrier material carrying the chemical characterised in that the shaped article has a mean thickness of about 4.0 mm or less and has side wall or walls diverging outwards from a bottom surface, said side walls making an angle with the vertical
15 of at least 5° at the upper surface.

12. A solid shaped article as claimed in claim 11 wherein the carrier material is hydrolysed gelatin.

20 13. A solid shaped article as claimed in claim 11 or 12 wherein the chemical is a pharmaceutical substance.

14. A solid shaped article whenever prepared by the process claimed in any one of claims 1 to 10.

5 15. A solid shaped article substantially as hereinbefore described with reference to any one of the Examples.

16. A process for preparing a solid shaped article substantially as hereinbefore described with reference to any one of the Examples.

Publication No.
2119246 A dated 16 November 1983

Patent Granted:

WITH ENTRY FROM
SECTION 2(1)(1)

22 MAY 1985

Application No.
8232189 filed on 11 November 1982

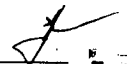
Priority claimed:
2 December 1981 in United Kingdom doc: 8136360

Title:
Solid shaped articles

Applicant:
John Wyeth and Brother Limited (United Kingdom) Huntercombe Lane South Taplow
Maidenhead Berkshire SL6 0PH

Inventor:
George Keith Emerson Gregory 12 Hazel Close Marlow Bottom Marlow Buckinghamshire

Classified to:
A5B C1B

Examination requested  APR 1984

Address for Service:
K J S Brown c/o Wyeth Laboratories Huntercombe Lane South Taplow Berks SL6 0PH