

**(12) STANDARD PATENT
(19) AUSTRALIAN PATENT OFFICE**

(11) Application No. AU 2007327594 B2

(54) Title
Stable lactide particles

(51) International Patent Classification(s)
C08G 63/00 (2006.01)

(21) Application No: **2007327594** (22) Date of Filing: **2007.11.28**

(87) WIPO No: **WO08/065132**

(30) Priority Data

(31) Number
07113211.2 (32) Date
2007.07.26
06124934.8 (33) Date
2006.11.28 (33) Country
EP
EP

(43) Publication Date: **2008.06.05**
(44) Accepted Journal Date: **2013.08.22**

(71) Applicant(s)
PURAC Biochem BV

(72) Inventor(s)
De Vos, Sicco

(74) Agent / Attorney
Griffith Hack, GPO Box 1285, Melbourne, VIC, 3001

(56) Related Art
US 5264592 A

(12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(19) World Intellectual Property Organization
International Bureau



(43) International Publication Date
5 June 2008 (05.06.2008)

PCT

(10) International Publication Number
WO 2008/065132 A1

(51) International Patent Classification:
C08G 63/00 (2006.01)

(81) Designated States (unless otherwise indicated, for every kind of national protection available): AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BH, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DO, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LY, MA, MD, ME, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RS, RU, SC, SD, SE, SG, SK, SL, SM, SV, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW.

(21) International Application Number:
PCT/EP2007/062919

(22) International Filing Date:
28 November 2007 (28.11.2007)

(25) Filing Language: English

(26) Publication Language: English

(30) Priority Data:
06124934.8 28 November 2006 (28.11.2006) EP
07113211.2 26 July 2007 (26.07.2007) EP

(71) Applicant (for all designated States except US): PURAC Biochem BV [—/NL]; Arkelsedijk 46, NL-4206 AC Gorinchem (NL).

(84) Designated States (unless otherwise indicated, for every kind of regional protection available): ARIPO (BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW), Eurasian (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European (AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, MT, NL, PL, PT, RO, SE, SI, SK, TR), OAPI (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG).

(72) Inventor; and

Published:

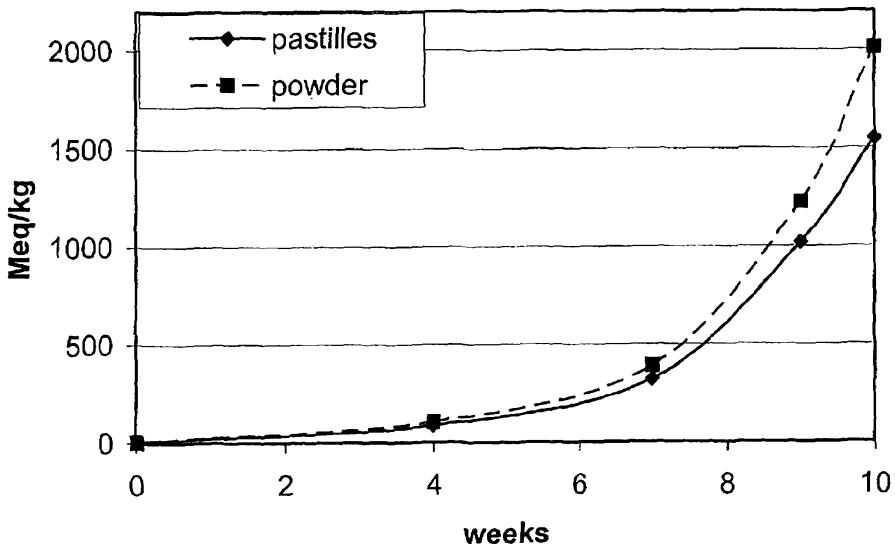
(75) Inventor/Applicant (for US only): DE VOS, Sicco [NL/NL]; Top Naeffstraat 2, NL-6836 PW Arnhem (NL).

— with international search report
— before the expiration of the time limit for amending the claims and to be republished in the event of receipt of amendments

(74) Agents: HESSELINK, Dinah Elisabeth et al.; Overschiestraat 180, NL-1062 XK Amsterdam (NL).

(54) Title: STABLE LACTIDE PARTICLES

Free acid content after storage at 20 degrees
Celcius in air



(57) **Abstract:** The present invention is directed to stable lactide particles, more specifically lactide particles which are stable enough to be stored and transported at room temperature and have a quality high enough for use as starting material for polylactic acid. The lactide particle has a surface/volume ratio of lower than 3000 m^{-1} . Preferably the lactide in the particle has an optical purity of at least 95%. The lactide particles are prepared by subjecting lactide to a shaping step comprising extrusion, pastillation, prilling, tabletting, or flaking.

WO 2008/065132 A1

STABLE LACTIDE PARTICLES

The present invention relates to lactide particles, more specifically to lactide particles which are stable enough to be stored and transported at room temperature and which have a quality high enough for use as starting material for 5 polylactic acid.

The continued depletion of landfill space and the problems associated with incineration of waste have led to the need for development of truly biodegradable polymers to be utilized as substitutes for non-biodegradable or partially 10 biodegradable, petrochemical-based polymers in packaging, paper coating and other non-medical industry applications, hereinafter referred to as bulk applications. The use of lactic acid and lactide to manufacture a biodegradable polymer is well known in the medical industry. As disclosed by 15 Nieuwenhuis et al. (US 5,053,485), such polymers have been used for making biodegradable sutures, clamps, bone plates and biologically active controlled release devices. It will be appreciated that processes developed for the manufacture of polymers to be utilized in the medical industry have 20 incorporated techniques that respond to the need for high purity and biocompatibility in the final polymer product. Furthermore, the processes were designed to produce small volumes of high dollar-value products, with less emphasis on manufacturing cost and yield.

25 It is known that lactic acid undergoes a condensation reaction to form polylactic acid upon dehydration. Dorough recognized and disclosed in US 1,995,970, that the resulting polylactic acid is limited to a low

molecular weight polymer of limited value, based on physical properties, due to a competing depolymerization reaction in which the cyclic dimer of lactic acid, lactide, is generated. As the polylactic acid chain lengthens, the polymerization 5 reaction rate decelerates until it reaches the rate of the depolymerization reaction, which effectively, limits the molecular weight of the resulting polymers.

Therefore, in most publications, processes for the production for polylactic acid are described wherein from 10 lactic acid first a prepolymer is prepared, said prepolymer is depolymerised by means of a catalyst to form crude lactide (i.e. the ring-closure reaction), said crude lactide is subsequently purified and lactide is used as starting material for the preparation of polylactic acid by ring-opening 15 polymerization. For the purpose of this description the term polylactic acid and polylactide are used interchangeably. It is well known that lactic acid exists in two forms which 20 are optical enantiomers, designated as D-lactic acid and L-lactic acid. Either D-lactic acid, L-lactic acid, or mixtures thereof may be polymerized to form an intermediate molecular weight polylactic acid which, after the ring-closure reaction, generates lactide as earlier disclosed. The lactide (sometimes 25 also referred to as dilactide), or the cyclic dimer of lactic acid, may have one of three types of optical activity depending on whether it consists of two L-lactic acid molecules, two D-lactic acid molecules or an L-lactic acid molecule and a D-lactic acid molecule combined to form the dimer. These three dimers are designated L-lactide, D-lactide, and meso-lactide, respectively. In addition, a 50/50 mixture 30 of L-lactide and D-lactide with a melting point of about 126 °C is often referred to in the literature as D,L-lactide. The optical activity of either lactic acid or lactide is known to alter under certain conditions, with a tendency toward

equilibrium at optical inactivity, where equal amounts of the D and L enantiomers are present. Relative concentrations of D and L enantiomers in the starting materials, the presence of impurities or catalysts and time at varying temperatures, and 5 pressures are known to affect the rate of such racemization. The optical purity of the lactic acid or the lactide is decisive for the stereochemistry of the polylactid acid obtained upon ring-opening polymerization of the lactide. With respect to polylactic acid, stereochemistry, and molecular 10 weight are the key parameters for polymer quality.

When preparing polylactic acid for the medical industry often crystalline powdery lactide is used as the starting material. This application is for instance described in EP-A1-1 310 517. These crystals, which are commercially 15 available for over 30 years now, are highly hygroscopic and are packed under inert atmosphere in damp- and air tight packages and stored in freezers (temperature below 12 °C). It will be clear that these precautions cannot be taken when 20 polylactic acid is used for bulk applications because it would render the product too expensive. Lactide powder or crystals usually have particles sizes ranging from 0.05 to approximately 0.05 mm.

In publications describing processes for the preparation of polylactic acid for bulk applications, the 25 lactide formed and purified is directly fed in its molten, liquid form to a polymerization reactor to form polylactide. See for instance EP 0,623,153 and US 6,875,839. By the direct conversion of the lactide prepared to polylactic acid, the negative effects of the relative instability of lactide can be 30 decreased by controlling the residence time of the lactide in the reactor. However, this process requires that the lactide production and polylactic acid production are combined. This makes the process rather inflexible and creates an entrance

21 Jun 2013
2007327594

barrier for new polylactic acid producers, because it requires large investments in equipment. Secondly, as the quality of the lactide is decisive for the molecular weight and stereo-chemistry that can be obtained in the polylactic acid, and the ring-closure process and purification require strict control of the temperature, pressure and residence time, it is also the most delicate part of the polylactic acid production process. The risk of failure in this part of the process enlarges the entrance barrier even more. If new polylactic acid producers for bulk applications could simply be provided with stable high quality lactide, this burden would be taken from them and substitution of petrochemical-based polymers with polylactic acid could actually take place. It has been suggested to transport lactide in its melted form (melting point of D-lactide and L-lactide is 97 °C) . Beside the fact that this type of transport is expensive, the transport and storage of melted lactide is also detrimental to the quality of the lactide because racemization, hydrolysis, and oxidation reactions are accelerated at these temperatures. The same problem occurs in the direct conversion process when the residence time of the lactide is not precisely controlled.

To this end the present invention is directed to stable lactide particles wherein the surface/volume ratio of the particle is lower than 3000 m^{-1} . According to the present invention there is provided a process for the preparation of lactide particles wherein lactide is subjected to a shaping process to form particles having a surface/volume ratio lower than 3000 m^{-1} , wherein the shaping process comprises extrusion, pastillation, prilling, tabletting, or flaking. The shaping process is carried out under inert gas or dry atmosphere.

We have found that lactide particles that fulfill this requirement are stable enough for storage and transport at room temperature and can readily be used as starting material for the production of lactic acid for bulk applications. With stable lactide particles is meant that when storing the lactide particles having an initial free acid content of at most 5 meq/kg at 20 degrees Celcius in air, the free acid content will

still be below 2000 after 10 weeks of storage. Crystalline powdery lactides used for the medical industry appeared not stable over time.

As mentioned-above, the optical purity of the lactide is
5 very important for the stereochemistry of the polylactic acid
that is obtained. Therefore, it is preferred that the lactide
present in the particles according to the invention contains
more than 95% by weight D- or L-lactide, preferably more than
98.5% by weight D- or L-lactide, most preferably more than 99.5%
10 D- or L-lactide by weight.

The lactide particles according to the invention can be
prepared by subjecting lactide (for instance in the melted or
crystalline powdery form) to a shaping process comprising
extrusion, pastillation, prilling, tabletting, or flaking. The
15 particles formed in the shaping process can be considered
pellets, pastilles, granules and/or agglomerates. These terms
are used throughout the description dependent from the term
commonly used in the shaping process concerned.

By melted is meant that at least part of the lactide is at
20 a temperature at or above the melting point of the lactide.

The apparatus used for the shaping process, or at least
those parts that will be in contact with the lactide, preferably
are prepared from corrosive-resistant material such as stainless
steel. Further, to avoid water uptake of the lactide particles,
25 the shaping process is preferably conducted under inert gas or
dry atmosphere such as under nitrogen or dry air.

By means of extrusion through one or more dies cylindrical
or rod-like particles can be obtained. When looking at the
surface/volume ratio of the lactide particles, these cylindrical
30 or rod-shaped particles are preferred. This shaping process is
further preferred because processing equipment for the
preparation of polylactide from lactide readily can handle
particles of this shape because of the

relatively uniform particle size and shape. The extruder is optionally cooled to avoid local overheating of the lactide. Any extruder conventionally used in the plastics, metal powder, food and ceramics industry such as screw extruders, 5 such as single- and twin-screw extruders and radial screen extruders etcetera is suitable.

Suitable pastillation machines are for instance the disc pastillator, ex GMF® or a rotoformer ® ex Sandvik. Herein the lactide is melted and droplets are placed on a disk or 10 belt with controlled temperature. We have found that by means of pastillation robust, uniformly shaped pellets can be made of lactide. Even though the surface/volume ratio of the resulting substantially hemi-spherical lactide particles is somewhat higher than for cylindrical or rod-shaped particles, 15 hemispherical lactide particles are preferred because processing equipment for the preparation of polylactide from lactide readily can handle particles of this shape because of the relatively uniform particle size and shape. Moreover, with this shaping process virtually no dusting takes place and the 20 resulting pastilles are hardly susceptible to abrasion during transport or any other mechanical handling. Compared to extruder-made particles, pastilles usually can easier be dosed in polylactic acid reactors especially when reactive extrusion polymerization is used. The term "relatively uniform" means 25 that at least 90 percent by weight of the pastilles are within plus/minus 30 percent of the mean diameter. Preferably, at least 95 percent by weight of the particles are within plus/minus 10 percent of the mean diameter. The term 30 "substantially hemi-spherical" means that the form of the particle is basically hemi-spherical, but can be flattened somewhat, i.e. the height of the particle is between 50 and 30% of its diameter.

When using flaking for the shaping process, optionally a sieving step is performed after the shaping to avoid dusting during transport and further processing to form polylactide.

5 With prilling lactide droplets fall in a liquid bath and thus spherical particles can be obtained. If water is used for the bath, extensive drying of the lactide particles is necessary.

10 Irrespective of the shape, particles with an average diameter of at least 3 millimeters are preferred, because then an optimum surface/volume ratio is ensured. More preferably the particles have an average diameter between 3 and 10 millimeters.

15 The water content of the lactide is an important factor for the stability of the lactide particles.

Contamination by water vapor leads to ring-cleavage causing the lactide to convert to lactoyl lactic acid and lactic acid. It was found that if the water content is below 200 ppm the stability of the lactide particles when stored at room 20 temperature in air-tight and vapor-tight packages is ensured for several months. Preferably, the water content is below 100 ppm because it further increases the stability of the lactide. The water content of the lactide can be measured by means of a Karl-Fisher titration as will be known by the artisan.

25 Also the acid content of the lactide (either lactic acid or lactoyl lactic acid) is important for the stability and quality of the lactide. The presence of lactic acid and or lactoyl lactic acid in the feed to the final polymerization step will result in polymers of limited molecular weight. If 30 the free acid content is below 50 milli-equivalents per Kg lactide (meq.Kg⁻¹) the stability of the lactide particles when stored at room temperature in air-tight and vapor-tight packages is ensured for several months. Preferably, the acid

content is below 20 meq.Kg⁻¹ because it further increases the stability of the lactide. Most preferably the acid content is between 0 and 10 meq.Kg⁻¹. The acid content can be measured by means of titration using for instance sodium methanoate or 5 potassium methanoate, as will be clear for the artisan.

The lactide used as starting material for the shaping process may have been prepared by any conventional lactide process such as water removal from a lactic acid solution or condensation reaction of lactate esters, followed 10 by a ring-closure reaction in a lactide reactor with the help of a catalyst. Optionally the crude lactide is further purified by for instance distillation and/or crystallization prior to the shaping process.

The lactide reactor can be of any suitable type that 15 is designed for heat sensitive materials. A reactor that can maintain a uniform film thickness, such as a falling film or agitated thin-film evaporator is most preferred, because film formation increases the rate of mass transfer. When the rate of mass transfer is increased, lactide can quickly form and 20 vaporize, and as lactide vaporizes, more lactide is produced as dictated by the polylactic acid/lactide equilibrium reaction. Optionally these lactide reactors are operated under reduced pressure such as between about 1 mmHg and 100 mmHg. The temperature of the lactide formation is kept between 150 25 °C and 250 °C. Many suitable catalysts are known, such as metal oxides, metal halides, metal dusts, and organic metal compounds derived from carboxylic acids or the like. Normally a tin(II) oxide or tin(Oct)₂ catalyst is used for lactide formation.

30 Stabilizers may also be added to the lactide reactor in order to facilitate lactide formation and discourage degenerative lactic acid and lactide reactions. Stabilizers, such as antioxidants, can be used to reduce the number of

degradation reactions that occur during the process of polylactic acid and lactide production. Stabilizers may also reduce the rate of lactide formation during this process. Therefore, efficient production of lactide requires proper 5 reactor design for minimal thermal severity and a proper balance between the catalyst and any use of process stabilizers.

A variety of stabilizers may be used. The stabilizing agent may include primary antioxidants and/or 10 secondary antioxidants. Primary antioxidants are those which inhibit free radical propagation reactions, such as alkylidene bisphenols, alkyl phenols, aromatic amines, aromatic nitro and nitroso compounds, and quinones. To prevent formation of free radicals secondary (or preventive) antioxidants break down 15 hydroperoxides. Some examples of secondary antioxidants include: phosphites, organic sulfides, thioethers, dithiocarbamates, and dithiophosphates. Antioxidants, when added to the lactide reactor can reduce the extent of racemization during lactide production. This reduction 20 indicates that the addition of antioxidants is an additional means to control optical purity. Antioxidants include such compounds as trialkyl phosphites, mixed alkyl/aryl phosphites, alkylated aryl phosphites, sterically hindered aryl phosphites, aliphatic spirocyclic phosphites, sterically 25 hindered phenyl spirocyclics, sterically hindered bisphosphonites, hydroxyphenyl propionates, hydroxy benzyls, alkylidene bisphenols, alkyl phenols, aromatic amines, thioethers, hindered amines, hydroquinones, and mixtures thereof. Preferably, phosphite-containing compounds, hindered 30 phenolic compounds, or other phenolic compounds are used as process stabilizing antioxidants. Most preferably, phosphite-containing compounds are used. The amount of process stabilizer used can vary depending upon the optical purity

desired of the resulting lactide, the amount and type of catalyst used, and the conditions inside of the lactide reactor. Normally amounts varying from 0.01 to 0.3 wt.% process stabilizer can be used. Next to stabilizers also 5 dehydration or anti-hydrolysis agents may be used. These dehydration agents favor the formation of lactide. Further, they may be used in a later stage of the manufacturing process for polylactic acid as well as for preventing chain scission by water. Compounds based on peroxide may be used for this 10 purpose but preferred are compounds containing the carbodiimide functionality. The carbodiimide compound is a compound having one or more carbodiimide groups in a molecule and also includes a polycarbodiimide compound. As a monocarbodiimide compound included in the carbodiimide 15 compounds, dicyclohexyl carbodiimide, diisopropyl carbodiimide, dimethyl carbodiimide, diisobutyl carbodiimide, dioctyl carbodiimide, diphenyl carbodiimide, naphthyl carbodiimide, etc. may be exemplified. In particular industrially easily available compounds such as dicyclohexyl 20 carbodiimide, diisopropyl carbodiimide or products like Stabaxol® by Rheinchemie are used.

It is also possible to add above-mentioned process stabilizers and dehydration agents to the lactide at a later stage, such as for instance prior to the shaping and/or after 25 the shaping step. If the stabilizers are added to the lactide after shaping, the stabilizers may be sprayed or coated onto the lactide particles.

It is of course desired to have as little as possible material such as process stabilizers present in the 30 lactide particles other than lactide. Therefore, the lactide particle usually comprises more than 95% by weight lactide, preferably more than 98.5% by weight lactide, most preferably more than 99.5% by weight.

Depending on the lactide preparation and/or purification method and the type of shaping process, the shaping process can either be combined with the preparation and or purification, or not. For instance, if the lactide is 5 obtained form distillation, it makes sense to directly couple a pastillation machine to the distillation column because the lactide is already in its melted form. If the final purification step of the lactide comprises crystallization, the use of an extruder is more opportune. Said extrusion can 10 also take place at a later point in time.

We have found that the presence of the above-mentioned process stabilizers also increases the stability of the lactide particles during storage.

15 The invention is further illustrated by means of the following non-limiting examples

EXAMPLE 1

Pastillation of L-lactide using lab-scale disc pastillator.

20 Fresh L-lactide ex. Purac ® (<1 meq/Kg free lactic acid) was molten using a microwave and subsequently poured into a double-walled metal container that was continuously heated by means of a hot air current. The lactide was thus kept in the molten state, and covered with a metal plunger. At 25 the bottom of the heated container, a nozzle with a cylindrical die (D=1 mm) was mounted. A slight pressure was applied to the lactide melt resulting in droplets falling onto a RVS disc that was mounted 6-7 mm below the nozzle. The RVS disc (D=400 mm) was slowly rotating (1-2 rpm), did not have active cooling and had a temperature of 15-20 °C (RT). The 30 clear lactide melt discharged from the nozzle solidified and crystallized on the RVS disc producing white pastilles. The droplet falling rate and the disc rotation speed were matched

in order to get circular arrays of pastilles on the disc. As soon as a circular array of pastilles was full, the position of the nozzle over the disc was adapted to start a new array, thus producing a cooling disc ultimately covered with

5 concentric arrays of pastilles. Pastilles did not stick to the metal disc and could be collected easily. Solidified lactide pastilles of uniform dimensions could thus be produced (Average particle diameter 5.5- 6 mm with a thickness of between 1.6-1.8 mm).

10

EXAMPLE 2

Cylindrical L-lactide pellets produced by extrusion

Fresh L-lactide ex. Purac ® (<1 meq/Kg free lactic acid) was extruded through a single capillary die of a Prism 15 Pharmalab 16 Series co-rotating twin-screw extruder of Thermo Fisher Scientific Corporation. The screw diameter was 16 mm and the processing length L/D was 40. The temperatures (°C) of the electrically heated zones (#1-11) of the extruder barrel were:

20

#	die		mixing section				mixing section				feed
	11	10	9	8	7	6	5	4	3	2	
°C	92	95	90	85	80	75	70	70	60	50	10

The extruder was operated with a screw speed of 150 rpm and L-lactide powder was metered in water-cooled zone 1 at 25 a solids rate of 1.8-2.4 Kg/h by means of a volumetric feeder. The temperature of the white paste discharged from the die was 88-92 °C. The resulting strands broke spontaneously when they fell down some 20-40 cm upon discharge from the extruder onto an RVS tray. As a result, cylindrical pellets with a randomly 30 distributed length of several millimeters are obtained (the

particle diameter was about 3 mm while the length varied from 5 to 15 mm).

The white lactide pellets initially exhibited a free lactic acid content of 4 meq/Kg.

5

COMPARATIVE EXAMPLE 3

The stability of powdery lactide particles was tested. The surface/volume ratio of powdery lactide is given in the TABLE below:

10

Shape	Average particle diameter (mm)	Surface/volume ratio (m^2/m^3)
Powder (spherical)	0.001	6,000,000
	0.005	1,200,000
	0.01	600,000
	0.02	300,000
	0.1	60,000
	0.2	30,000
	0.5	12,000

The stability of powdery material having a diameter of about 1 mm (surface/volume ratio of $6000 m^{-1}$) was measured after storage for 1 year in air-tight and vapor-tight bags (comprising a polyethylene inner bag and an aluminum outer bag) with a hole in it. The initial free acid content was 0.080 meq/Kg. After 1 year at 4 °C the free acid content was increased to 0.09 meq/Kg and after 1 year at 25 °C the free acid content was increased to 1131 meq/Kg. This means shows that powdery material is not stable enough for storage at room temperature for several months.

COMPARATIVE EXAMPLE 4

The stability of powdery material having a diameter of about 1 mm was measured after storage for 1 year in a single polyethylene bag (vapor-tight but not air-tight). The 5 initial free acid content was 0.09 meq/Kg. After 6 months at 25 °C the free acid content was increased to 405 meq/Kg, and thus not suitable anymore as a starting material for the preparation of polylactid acid.

10 EXAMPLE 5

In the TABLE below the surface/volume ratio is given for cylindrical and hemi-spherical shaped lactide particles.

Shape	Average particle length X diameter (mm X mm)	Surface/volume ratio (m ² /m ³)
Cylindrical	2 X 1.5	2000
	3 X 1.5	1333.4
	4 X 1.5	1000
	5 X 1.5	800
	6 X 1.5	666.7
	7 X 1.5	571.4
	8 X 1.5	500
	9 X 1.5	444.4
	10 X 1.5	400

15

Shape	Average particle diameter (mm)	Surface/volume ratio (m ² /m ³)
Hemi-spherical	2	4500
	3	3000
	4	2250
	5	1800
	6	1500
	7	1286
	8	1125
	9	1000
	10	900

Example 6

The stability of lactide pastilles as prepared in Example 1 were compared with powdered lactide with an average particle size of 100 micrometers. The surface/volume ratio of 5 the pastilles was 1600 m^{-1} , while the surface/volume ratio of the powdered lactide was 6000 m^{-1} . To this end both lactid pastilles and lactid powder with an initial free acid content of 5 meq/kg were subjected to stability tests at 20 and 40 degrees Celcius. The lactide samples were kept in a polyethylene bag 10 (vapour-tight but not air tight). The free acid content of the samples were measured after various periods of storage. The results are compiled in figures 1 and 2. Figure 1 gives the results of storage at 20 degrees Celcius. These results show that the free acid content of powdered lactide increases much 15 faster over time than pastillated lactide. In fact, the free acid content of powdered lactide had increased to over 2000 after storage for 10 weeks, which had rendered the powdered lactide unsuitable for polylactic acid production.

Figure 2 gives the results of storage at 40 degrees 20 Celcius. These results show that at higher temperatures the free acid content increases faster than with storage at 20 degrees Celcius. Here also the free acid content of powdered lactide increases much faster over time than pastillated lactide.

25 It is to be understood that, if any prior art publication is referred to herein, such reference does not constitute an admission that the publication forms a part of the common general knowledge in the art, in Australia or any other country.

In the claims which follow and in the preceding description of the invention, except where the context requires otherwise due to express language or necessary implication, the word "comprise" or variations such as "comprises" or 5 "comprising" is used in an inclusive sense, i.e. to specify the presence of the stated features but not to preclude the presence or addition of further features in various embodiments of the invention.

THE CLAIMS DEFINING THE INVENTION ARE AS FOLLOWS:

1. Process for the preparation of lactide particles wherein lactide is subjected to a shaping process to form particles having a surface/volume ratio lower than 3000 m^{-1} , wherein the shaping process comprises extrusion, pastillation, prilling, tabletting, or flaking, and wherein the shaping process is carried out under inert gas or dry atmosphere.
2. Process according to claim 1 wherein a lactide is extruded or compressed to form cylindrical, cubical or rod-shaped particles.
3. Process according to claim 1 or 2 wherein a lactide melt is pastillated to form substantially hemi-spherical particles.
4. Process according to claim 3, wherein the shaping step comprises pastillation and the lactide is obtained from a distillation process in melted form and is directly fed to a pastillation machine.
5. Process according to claim 1 or 2, wherein the shaping step is an extrusion step and the lactide originates from a process wherein the final purification step is a crystallization step.
6. Process according to claim 1, wherein the shaping process is a flaking process, and the flakes are subjected to a sieving step after the shaping.
7. Process according to any one of the preceding claims, wherein the apparatus used for the shaping process, or at least those parts of which that will be in contact with the lactide are prepared from corrosion-resistant material.
8. Process according to any one of the preceding claims, which is carried out in such a manner that particles with a diameter of at least 3 mm are obtained.

9. Process according to any one of the preceding claims, wherein the lactide particles have a surface/volume ratio lower than 3000 m^{-1} , and are stable lactide particles having an initial free acid content of at most 5 meq/kg at 20 degrees Celcius in air, and after 10 weeks of storage the free acid content will still be below 2000 meq/kg.
10. Process according to any one of the preceding claims, wherein the lactide particles comprise more than 95% by weight lactide, more than 98.5% by weight lactide or more than 99.5% by weight lactide.
11. Process according to any one of the preceding claims, wherein the lactide present in the particles contains more than 95% by weight D-lactide, more than 98.5% by weight D-lactide or more than 99.5% by weight D-lactide, or wherein the lactide in the particle contains more than 95% by weight L-lactide, more than 98.5% by weight L-lactide or more than 99.5% by weight L-lactide.
12. Process according to any one of the preceding claims, wherein the lactide particles have a water content below 200 ppm, below 100 ppm or below 50 ppm.
13. Process according to any one of the preceding claims, wherein the particles have a free lactic acid content below 50 milli-equivalents per Kg lactide (meq.Kg^{-1}), below 20 meq.Kg^{-1} or between 0 and 10 meq.Kg^{-1} .
14. Lactide particles obtained by the process according to any one of claims 1 to 13.
15. Use of lactide particles obtained by the process according to any one of claims 1 to 13 which have a surface/volume ratio lower than 3000 m^{-1} in the manufacture of polylactic acid for bulk applications.

19 Jul 2013

2007327594

16. Process for the preparation of lactide particles according to any one of claims 1 to 14, or lactide particles obtained by the process, substantially as herein described with reference to the accompanying examples (excluding the comparative examples) and/or drawings.

Figure 1: Free acid content after storage at 20 degrees Celcius in air

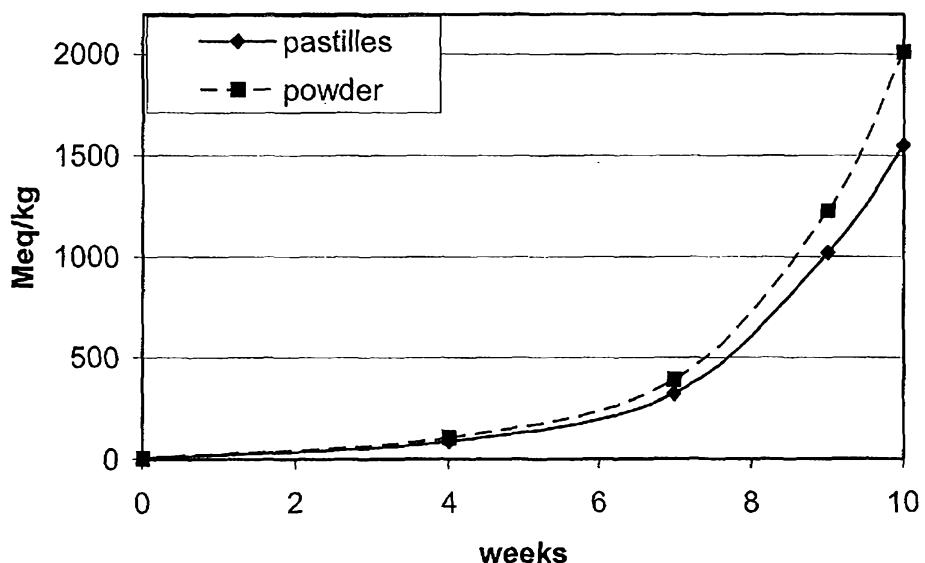


Figure 2: Free acid content after storage at 40 degrees Celcius in air

