



(86) Date de dépôt PCT/PCT Filing Date: 2001/06/15  
 (87) Date publication PCT/PCT Publication Date: 2001/12/27  
 (45) Date de délivrance/Issue Date: 2003/12/09  
 (85) Entrée phase nationale/National Entry: 2002/11/28  
 (86) N° demande PCT/PCT Application No.: US 2001/019100  
 (87) N° publication PCT/PCT Publication No.: 2001/097612  
 (30) Priorité/Priority: 2000/06/16 (60/211,967) US

(51) Cl.Int.<sup>7</sup>/Int.Cl.<sup>7</sup> A01N 37/12  
 (72) Inventeurs/Inventors:  
 SINGER, CLAUDE, IL;  
 PILARSKI, GIDEON, IL;  
 PESACHOVICH, MICHAEL, IL;  
 SCHWARTZ, EDUARD, IL  
 (73) Propriétaire/Owner:  
 TEVA PHARMACEUTICAL INDUSTRIES LTD., IL  
 (74) Agent: RIDOUT & MAYBEE LLP

(54) Titre : GABAPENTINE STABLE CONTENANT PLUS DE 20 PPM D'IONS CHLORE  
 (54) Title: STABLE GABAPENTIN CONTAINING MORE THAN 20 PPM OF CHLORINE ION

(57) Abrégé/Abstract:

Pharmaceutical compositions containing substantially pure and stable gabapentin are disclosed wherein gabapentin contains an anion of a mineral acid, such as chloride, in amounts greater than 20 ppm.

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

(19) World Intellectual Property Organization  
International Bureau



(43) International Publication Date  
27 December 2001 (27.12.2001)

PCT

(10) International Publication Number  
**WO 01/97612 A1**

- (51) International Patent Classification<sup>7</sup>: **A01N 37/12**
- (21) International Application Number: PCT/US01/19100
- (22) International Filing Date: 15 June 2001 (15.06.2001)
- (25) Filing Language: English
- (26) Publication Language: English
- (30) Priority Data:  
60/211,967 16 June 2000 (16.06.2000) US
- (71) Applicant (for all designated States except BB): **TEVA PHARMACEUTICAL INDUSTRIES LTD.** [IL/IL]; Basel Street 5, P.O. Box 3190, 49131 Petah Tiqva (IL).
- (71) Applicant (for BB only): **TEVA PHARMACEUTICALS USA, INC.** [US/US]; 1090 Horsham Road, P.O. Box 1090, North Wales, PA 19454-1090 (US).
- (72) Inventors; and
- (75) Inventors/Applicants (for US only): **SINGER, Claude** [IL/IL]; Haemek 17/9, Kfar Saba (IL). **PILARSKI, Gideon** [IL/IL]; Ataroth 12, Holon (IL). **PESACHOVICH, Michael** [IL/IL]; Zvulon Hammer 5/11, Givat Shmuel (IL).
- (74) Agents: **BIRDE, Patrick, J.** et al.; Kenyon & Kenyon, One Broadway, New York, NY 10004 (US).
- (81) Designated States (*national*): AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW.
- (84) Designated States (*regional*): ARIPO patent (GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).
- Published:**  
— with international search report
- For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.*



**WO 01/97612 A1**

(54) Title: STABLE GABAPENTIN CONTAINING MORE THAN 20 PPM OF CHLORINE ION

(57) Abstract: Pharmaceutical compositions containing substantially pure and stable gabapentin are disclosed wherein gabapentin contains an anion of a mineral acid, such as chloride, in amounts greater than 20 ppm.

WO 01/97612

PCT/US01/19100

**Stable Gabapentin Containing More than 20 ppm of Chlorine  
Ion**

**Cross-reference to Related Applications**

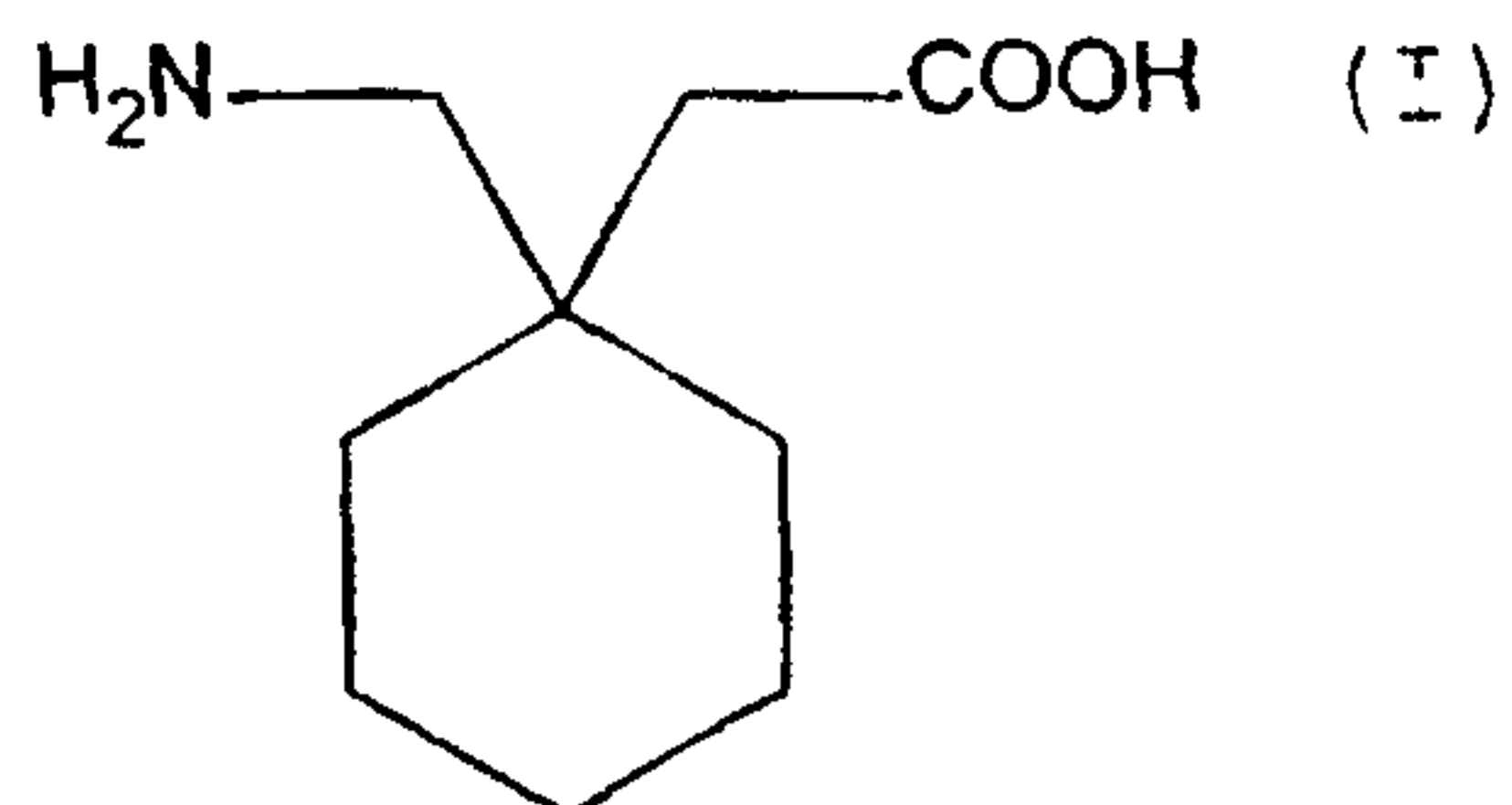
5           This invention relates to PCT Publication No. WO  
98/28255, filed July 2, 1998, also assigned to the assignee  
of the present invention.

10          **Field of the Invention**

          The present invention relates to a pharmaceutical  
composition containing therapeutically effective amount of  
gabapentin and its derivatives in combination with effective  
carriers. More particularly, the present invention relates  
15       to a composition and a process for manufacturing pure and  
stable gabapentin having greater than 20 ppm of chloride  
ion.

**Background of the Invention**

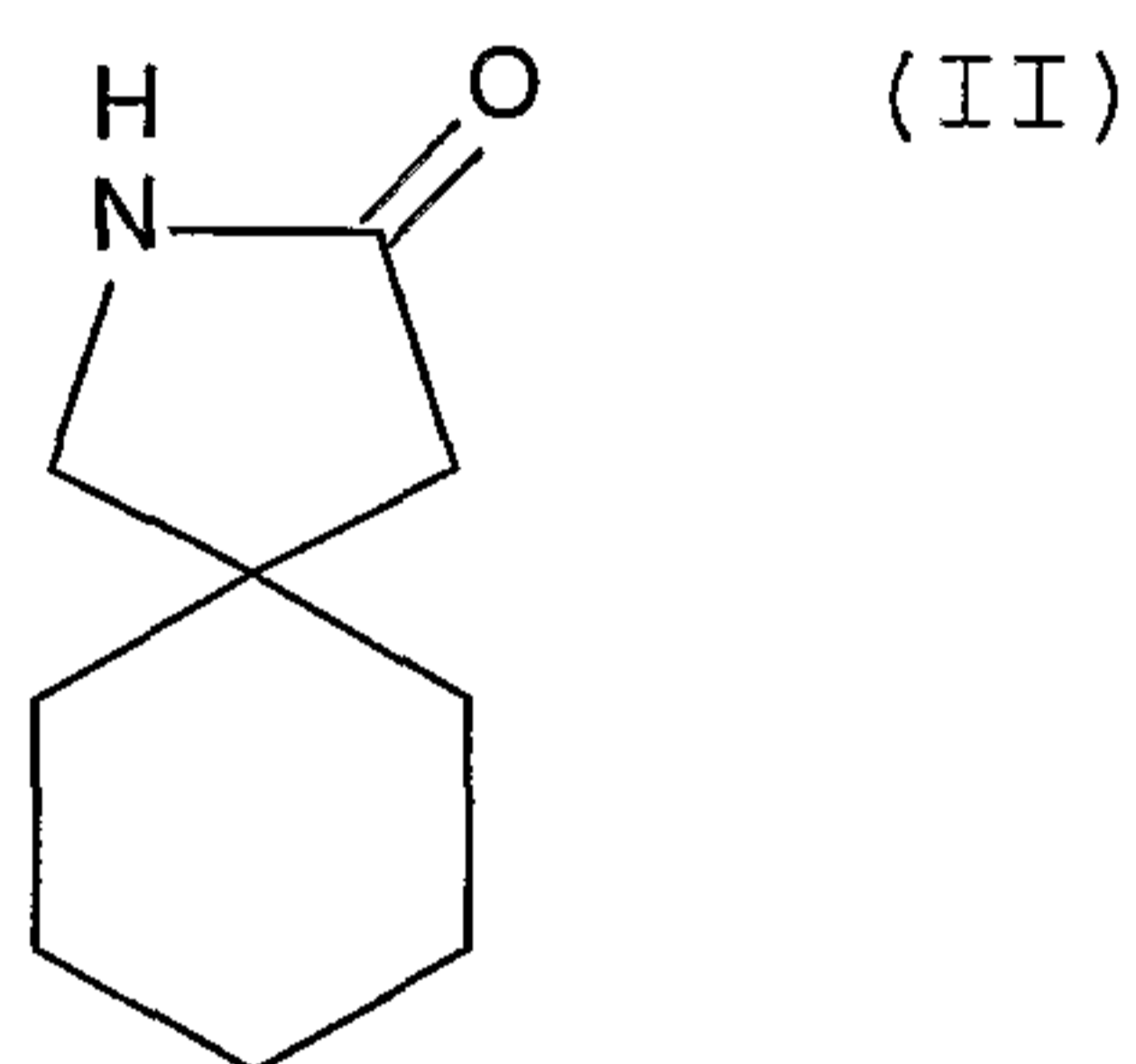
          Gabapentin is 1-(aminomethyl)-1-cyclohexaneacetic acid,  
20       having the chemical structure of formula I:



          Gabapentin is used for treating cerebral diseases such  
as epilepsy, faintness attacks, hypokinesia and cranial  
traumas. United States Patent No. 4,024,175 to Satzinger et  
25       al., incorporated herein by reference, discloses that  
gabapentin of formula (I) shows hypothermic and, in some

cases, narcosis-potentiating or sedating properties as well as protective effect against cardiozole cramp in animals. Finally, gabapentin has been found especially useful in treating geriatric patients. As such, there has been a need  
5 for producing pure and stable gabapentin.

United States Patent No. 6,054,482 to Augart et al. discloses that preparation and long-term storage of gabapentin presents several problems since (i) during the preparation the compounds shows considerable variations  
10 without apparent reason, and (ii) the long-term storage of even very pure gabapentin showed differing stabilities with progressively long storage times. Augart further discloses that the toxic lactam compound of formula (II)



15 forms during the preparation and storage of gabapentin. According to Augart, because the lactam has a higher toxicity than gabapentin, its presence in gabapentin should be limited if not eliminated. To combat lactam formation  
20 and provide product stability, Augart stresses the importance of (i) starting with gabapentin raw material that contains 0.5% or less of corresponding lactam, (ii) not allowing the anion of a mineral acid in the composition to exceed 20 ppm, and (iii) using a specifically selected  
25 adjuvant that is not adverse to gabapentin stability.

According to Augart, the following adjuvants (or excipients) had no noticeable influence on the stability of gabapentin, and as such, they were taught to be acceptable

adjuvants for use with gabapentin:

hydroxypropylmethylcellulose, polyvinylpyrrolidone,  
crospovidon, poloxamer 407™, poloxamer 188™, sodium starch  
glycolate, copolyvidone, maize starch, cyclodextrin,  
5 lactose, talc, as well as co-polymers of dimethylamino-  
methacrylic acid and neutral methacrylic acid ester.

Conversely, Augart discloses that the following  
adjuvants reduce the stability of gabapentin and should be  
avoided: modified maize starch, sodium croscarmellose,  
10 glycerol behenic acid ester, methacrylic acid co-polymers  
(types A and C), anion exchangers titanium dioxide and  
silica gels such as Aerosil 200™.

The composition and method disclosed in Augart are  
industrially impractical and technically unnecessary. It  
15 has now been found that Augart's reliance on maintaining  
the anion of a mineral acid as not exceeding 20 ppm is  
misplaced. Thus, gabapentin and pharmaceutical  
formulations of gabapentin can be prepared and stored such  
that initially they do not contain more than 0.5% of the  
20 lactam, and even after one year of storage at 25 °C and 60%  
atmospheric humidity, the conversion of gabapentin to its  
corresponding lactam does not exceed 0.2% by weight of  
gabapentin. That is, gabapentin and pharmaceutical  
formulations of gabapentin have been found to be stable  
25 even though such formulations do not meet Augart's  
requirements (ii) and (iii).

The specific mineral acid disclosed by Augart is  
hydrochloric acid (column 3, lines 61-63; column 5, lines  
24-29; examples 1 and 2). The specification states, in  
30 particular

The active materials of formula (I) [including gabapentin] must be prepared as highly purified, nonderivatized free amino acids, for example, from the corresponding hydrochloride by ion exchange. The  
5 proportion of remaining hydrochloride admixtures should

thereby not exceed 20 ppm.

(Column 5, lines 24-29)

20 ppm of gabapentin hydrochloride corresponds to roughly 3 ppm of chloride ion, due to the higher molecular weight of gabapentin.

Augert's claims require gabapentin with "less than 20 ppm of the anion of a mineral acid", e.g. chloride.

#### Summary of the Invention

Accordingly, the present invention relates to a pharmaceutical composition containing a pharmaceutically effective amount of gabapentin containing more than 20 ppm of an anion of a mineral acid and which initially contains less than 0.5% of a corresponding lactam and after one year of storage at 25 °C and 60% atmospheric humidity the conversion of gabapentin to its corresponding lactam does not exceed 0.2% by weight of gabapentin. The anion of the mineral acid may comprise a halide.

The present invention also relates to a process for preparing a stable pharmaceutical formulation containing gabapentin with more than 20 ppm of the anion of a mineral acid and which initially contains less than 0.5% of a corresponding lactam and after storage for one year at 55 °C and 60% atmospheric humidity the conversion of gabapentin to its corresponding lactam does not exceed 0.2% by weight of gabapentin.

#### Detailed Description of the Preferred Embodiments

The subject invention will now be described in greater detail for preferred embodiments of the invention, it being understood that these embodiments are intended only as illustrative examples and the invention is not to be limited thereto.

As will be illustrated through exemplary embodiments 1-16, gabapentin may be prepared from the hydrochloride salt of gabapentin (gabapentin hydrochloride) and that in purified form gabapentin may contain more than 20 ppm of chloride ion in the composition as measured by the amount of chloride ion in the composition.

Exemplary embodiments 17-19 illustrate formulations of gabapentin containing varying amounts of chloride ion, some of which are greater than 20 ppm and some less, and all of which initially contain less than 0.5% of lactam and after one year of storage at 25 °C and 60% humidity, the conversion of gabapentin to its corresponding lactam is measured not to exceed 0.2% by weight of gabapentin.

Commonly known adjuvants (also referred to as excipients) which can be utilized in a gabapentin formulation of the present invention may include for example, modified maize starch, sodium croscarmellose, titanium dioxide, and silica gels such as Aerosil 200. Hydroxypropylmethylcellulose, polyvinylpyrrolidone, crospovidon, poloxamer 407, poloxamer 188, sodium starch glycolate, copolyvidone, maize starch, cyclodextrin, lactose, talc, co-polymers of dimethylamino-methacrylic acid and neutral methacrylic acid ester may also be used. The list of adjuvants is not an exhaustive list and it would be within the scope of the claimed invention to use any known adjuvant that would behave similar to those enumerated herein.

Certain specific representative embodiments of the invention are described in detail below, the materials, apparatus and process steps being understood as examples that are intended for illustrative purposes only. Consequently, it will be noted that the invention is not intended to be limited to the methods, materials,

conditions, process parameters, apparatus and the like specifically recited herein.

In the examples below chloride ion concentration is measured by any commonly known method, such as for example, by titration with  $\text{AgNO}_3$ , or pH electrode or chromatography.

### EXAMPLE 1

The following raw material were used:

Gabapentin hydrochloride	18.2 g
Isopropanol for dissolution	160 ml
Active carbon SX1	1.1 g
Ethylacetate	268 ml
Tributylamine	19.5 g
Methanol for washing	23 ml

#### A) Preparation of Crude gabapentin

Gabapentin hydrochloride was dissolved in 130 ml of dry isopropanol at 25°C by mixing. Next, 1.1 grams of active carbon was added and the suspension was heated to 40°C and maintained at this temperature for 2 hours. The suspension was then filtered at 40°C and the filter cake was washed twice with additional 15 ml of isopropanol each time. The washings were added to the already separated solution of gabapentin hydrochloride in isopropanol. The solution was concentrated to dryness in vacuum (Approximately 10 mm Hg) to a constant weight. The temperature of the heating bath was maintained (maximally) at 35°C during this operation. Thereafter, 245 ml of ethylacetate was added to the dry residue of gabapentin hydrochloride and the solution was mixed. After half an hour of mixing at 25°C, an amount of 19.5 grams of tributylamine was added during the subsequent 30 minutes. The mixing continued for an additional two hours at the same temperature.

The gabapentin base which was formed during this operation was separated from the suspension through filtration. The filter cake was washed with 23 ml of ethylacetate and 23 ml of methanol to give crude gabapentin.

5 B) Gabapentin Purification

The following raw material were used:

Methanol for suspending	52.5 ml
Methanol for washing	2x15 ml

10 Wet crude gabapentin prepared according to Step A was suspended in 52.5 ml of methanol for 14 hours at approximately 25°C and stirred. Thereafter, the solid gabapentin was separated from the suspension by filtration. The filter cake was washed twice with 15 ml of methanol and than dried under vacuum giving pure gabapentin. The yield  
15 was 72%.

The following data regarding the chlorine anion content of the above-prepared gabapentin were obtained:

**TABLE 1 - Anion content and pH values  
after the reslurry in methanol**

20

Run	Cl <sup>-</sup> (ppm)	pH
A	4	6.94
B	20	7.01
C	< 5	7.04
D	40	6.97
E	35	6.92
F	15	6.84

25

30 Gabapentin purified according to these procedures contains less than 0.5% lactam as measured by HPLC vs. standard. After a year of storage at 25°C and 60% relative humidity, the conversion of gabapentin to its corresponding lactam does not exceed 0.2% by weight of gabapentin.

For a better control of the pH of pure gabapentin several basic agents were added. Some examples of added basic agents are given in the following Examples.

**EXAMPLE 2**

5 The following raw material were used:

Methanol for suspending	52.5 ml
Methanol for washings	2x15 ml
Tributylamine	~0.3 equivalents

10 The wet crude gabapentin(as in Step 1A) was suspended in 52.5 ml of methanol for 14 hours and at 25°C and stirred. Tributylamine was added to the suspension. After 14 hours of stirring the solid gabapentin was separated from the suspension by filtration. The filter cake was then washed twice, each time with 15 ml of methanol and than dried under  
15 vacuum resulting in pure gabapentin with a yield of 87%, pH of 7.15 and chlorine anion content of 50 ppm. Gabapentin so prepared contains less than 0.5% by weight of lactam, and, after a year of storage at 25°C and 60% relative humidity the conversion of gabapentin to its corresponding lactam is  
20 found not to exceed 0.2% by weight of gabapentin.

**EXAMPLE 3**

The following raw material were used:

Methanol for suspending	52.5 ml
Methanol for washing	2x15 ml
25 Sodium methoxide	~0.001 equivalents

The wet crude gabapentin (as in Example 1, step A) was suspended in 52.5 ml of methanol for 14 hours and kept at 25°C. Sodium methoxide was added to the suspension. After

14 hours of stirring, the solid gabapentin was separated from the suspension by filtration. The filter cake was then washed twice with 15 ml of methanol, then dried under vacuum, resulting in pure gabapentin having a yield of 85%, pH of 6.8, and chlorine anion content of 50 ppm. Gabapentin so prepared contained less than 0.5% by weight of lactam, and, after a year of storage at 25°C and 60% relative humidity, the conversion of gabapentin to its corresponding lactam is found not to exceed 0.2% by weight of gabapentin.

It should be noted that the solvents and the base used in Example 1A were not unique. In addition, it should be noted that in Examples 4-9 gabapentin pure was always prepared as in Example 1B and the results (Cl<sup>-</sup> content and yield) refer to gabapentin pure.

#### EXAMPLE 4

The following raw material were used:

Gabapentin hydrochloride (100%)	18.2 g
Isopropanol for dissolution	160 ml
Active carbon SX1	1.1 g
Tributylamine	19.5 g
Methanol for washing	23 ml

In this Example, gabapentin hydrochloride was dissolved in 130 ml of dry isopropanol at 25°C. Then 1.1 grams of active carbon was added and the suspension was heated to 40° C and maintained at this temperature for 2 hours. The suspension was filtered at 40° C and the filter cake was then washed twice, each time with an additional 15 ml of isopropanol. The washings were added to the already separated solution of gabapentin hydrochloride in isopropanol. After half an hour of mixing at 25° C, 19.5

grams of tributylamine was added during half an hour and the mixing was continued for two hours at the same temperature. The formed gabapentin base was separated from the suspension by filtration and washed with 23 ml of methanol to give  
 5 gabapentin crude. After reslurry as in Example 1B gabapentin pure was obtained at a yield of 58.8% and chloride anion content of 7 ppm Cl<sup>-</sup>.

Gabapentin so prepared contained less than 0.5% by weight of lactam, and, after a year of storage at 25°C and  
 10 60% relative humidity, the conversion of gabapentin to its corresponding lactam is found not to exceed 0.2% by weight of gabapentin.

#### **EXAMPLE 5**

The following raw material were used:

15	Gabapentin hydrochloride (100%)	18.2 g
	Isopropanol for dissolution	160 ml
	Active carbon SX1	1.1 g
	Ethylacetate	268 ml
	Trihexylamine	28.3 g
20	Methanol for washing	23 ml

Gabapentin hydrochloride was dissolved in 130 ml dry isopropanol at 25° C by mixing, then 1.1 g of active carbon was added and the suspension was heated to 40° C and maintained for two hours at 40° C. The suspension was  
 25 filtered at 40°C and the filter cake was washed twice with additional 15 ml of isopropanol each time. The washings were added to the already separated solution of gabapentin hydrochloride in isopropanol. The solution was concentrated to dryness in vacuum (approximately 10 mm Hg) to constant

weight. The temperature of the heating bath was maintained at maximum 35° C during this operation. Next, 245 ml of ethylacetate was added to the dry residue of gabapentin hydrochloride and the mixing was started. After half an hour of mixing at 25°C, an amount of 28.3 grams of trihexylamine was added during half an hour and the mixing was continued for an additional two hours at the same temperature. The formed gabapentin base was separated from the suspension by filtration. The filter cake was washed with 23 ml of ethylacetate and 23 ml of methanol to give gabapentin crude. After reslurry as in Example 1B, gabapentin pure was obtained having a yield of 75.0% and chloride anion content of 213 ppm.

#### EXAMPLE 6

The following raw material were used:

Gabapentin hydrochloride (100%)	18.2 g
Isopropanol for dissolution	160 ml
Active carbon SX1	1.1 g
Ethylacetate	268 ml
Tripropylamine	15 g
Methanol for washing	23 ml

Gabapentin hydrochloride was dissolved in 130 ml dry isopropanol at 25°C by mixing, then 1.1 g of active carbon was added and the suspension was heated to 40°C and maintained during two hours at 40° C. The suspension was filtered at 40° C and the filter cake was washed twice with additional 15 ml of isopropanol each time. The washings were added to the already separated solution of gabapentin hydrochloride in isopropanol. The solution was concentrated to dryness in vacuum (~10 mm Hg) to constant weight. The

temperature of the heating bath was maintained at maximum 35° C during this operation. Next, 245 ml of ethylacetate was added to the dry residue of gabapentin hydrochloride and mixing commenced. After half an hour of mixing at 25° C, 5 fifteen grams of tripropylamine was added during half an hour and the mixing was continued for two hours at the same temperature. The formed gabapentin base was separated from the suspension by filtration. The filter cake was washed with 23 ml of ethylacetate and 23 ml of methanol to give 10 gabapentin crude. After a reslurry process, as in Example 1B, gabapentin pure was obtained having a yield of 68.0% and chloride anion content of 142 ppm.

#### EXAMPLE 7

The following raw material were used:

15	Gabapentin hydrochloride (100%)	18.2 g
	Isopropanol for dissolution	160 ml
	Active carbon SX1	1.1 g
	Acetonitrile	268 ml
	Tributylamine	19.5 g
20	Methanol for washing	23 ml

Gabapentin hydrochloride was dissolved in 130 ml of dry isopropanol at 25° C by mixing, then 1.1 g of active carbon was added and the suspension was heated to 40° C and maintained for two hours at 40° C. The suspension was 25 filtered at 40° C and the filter cake was washed twice with additional 15 ml of isopropanol. The washings were added to the already separated solution of gabapentin hydrochloride in isopropanol. The solution was concentrated to dryness in vacuum (~10 mm Hg) to constant weight. The temperature of 30 the heating bath was maintained at a maximum temperature of

35°C during this operation. Next, 245 ml of acetonitrile was added to the dry residue of gabapentin hydrochloride and mixing commenced. After half an hour of mixing at 25° C, an amount of 19.5 g of tributylamine was added during 30  
5 minutes and the mixing was continued for two hours at the same temperature. The formed gabapentin base was separated from the suspension by filtration. The filter cake was washed with 23 ml of acetonitrile and 23 ml of methanol to give gabapentin crude. After reslurry as in Example 1B,  
10 gabapentin pure was obtained having a yield of 67.8%, and anion content of 142 ppm.

#### EXAMPLE 8

The following raw material were used:

	Gabapentin hydrochloride (100%)	18.2 g
15	Isopropanol for dissolution	160 ml
	Active carbon SX1	1.1 g
	Dimethylcarbonate	268 ml
	Tributylamine	19.5 g
	Methanol for washing	23 ml

20 Gabapentin hydrochloride was dissolved in 130 ml of dry isopropanol at 25° C by mixing, then 1.1 g of active carbon was added and the suspension was heated to 40°C and maintained at 40° C for two hours. The suspension was filtered at 40° C and the filter cake was washed twice with  
25 additional 15 ml of isopropanol. The washings were added to the already separated solution of gabapentin hydrochloride in isopropanol. The solution was concentrated to dryness in vacuum (~10 mm Hg) to constant weight. The temperature of the heating bath was maintained at maximum of 35° C during  
30 this operation. Next, 245 ml of dimethylcarbonate was added

to the dry residue of gabapentin hydrochloride and the mixing was started. After half an hour of mixing at 25° C, an amount of 19.5 g of tributylamine was added during half an hour and the mixing was continued for two hours at the same temperature. The formed gabapentin base was separated from the suspension by filtration. The filter cake was washed with 23 ml of dimethylcarbonate and 23 ml of methanol to give gabapentin crude. After reslurry as in Example 1B, gabapentin pure was obtained, having a yield of 57.9%, and anion content of 142 ppm.

#### EXAMPLE 9

The following raw material were used:

	Gabapentin hydrochloride (100%)	18.2 g
	Isopropanol for dissolution	160 ml
15	Active carbon SX1	1.1 g
	Isopropylacetate	268 ml
	Tributylamine	19.5 g
	Methanol for washing	23 ml

Gabapentin hydrochloride is dissolved in 130 ml of dry isopropanol at 25° C by mixing, then 1.1 g of active carbon was added and the suspension was heated to 40°C and maintained for two hours at 40°C. The suspension was filtered at 40°C and the filter cake was washed twice with additional 15 ml of isopropanol each time. The washings were added to the already separated solution of gabapentin hydrochloride in isopropanol. The solution was concentrated to dryness in vacuum (~10 mm Hg) to constant weight. The temperature of the heating bath was maintained at maximum 35° C during this operation. Next, 245 ml of isopropylacetate was added to the dry residue of gabapentin

hydrochloride and mixing commenced. After half an hour of mixing at 25°C, an amount of 19.5 g of tributylamine was added during half an hour and the mixing was continued for two hours at the same temperature. The formed gabapentin base was separated from the suspension by filtration. The filter cake was washed with 23 ml of isopropylacetate and 23 ml of methanol to give gabapentin crude. After reslurry as in Example 1B, gabapentin pure was obtained having a yield of 57.9% and an anion content of 142 ppm.

#### **EXAMPLE 10**

(The neutralization reaction as in Example 1, however, the reslurry in methanol is replaced by a crystallization in methanol.)

The following raw material were used:

15	Methanol for dissolution	180 ml
	Methanol for washing	2x12 ml

The gabapentin crude (Step 1A) was suspended in 180 ml of methanol at 25° C. The suspension was heated while mixing to 55°C when gabapentin was dissolved. The solution was then cooled slowly for an hour to 25°C. At 25° C the solution was concentrated to a volume of 50 ml. The suspension was stirred for twelve hours at 25°C. After 12 hours, the solid gabapentin was separated from the suspension by filtration. The filter cake was washed twice with 12 ml of methanol then dried under vacuum to give gabapentin pure (yield: 72%). Following Cl<sup>-</sup> contents of gabapentin and pH values were obtained and tabulated in TABLE 2 as follows:

**TABLE 2 - Anion content and pH values for crystallization in methanol**

Run	Cl <sup>-</sup> (ppm)	pH
A	4	6.94
B	< 5	7.2
C	150-200	6.9

5

For a better control of the pH of gabapentin pure several basic agents were added. Some examples of added basic agents are given in the following examples

**EXAMPLE 11**

10 The following raw material were used:

Methanol for suspending                    52.5 ml  
Methanol for washings                    2x15 ml  
Tributylamine                                ~0.34 equivalents

15 The gabapentin crude was suspended in 180 ml of methanol at 25°C. The suspension was then heated, while mixing, to 55°C when gabapentin was dissolved. Tributylamine was added to the solution and the solution was cooled slowly during an hour to a temperature of 25° C. At 25° C the solution was concentrated to a volume of 50 ml. The  
20 suspension was stirred for twelve hours at 25° C. After 12 hours the solid gabapentin was separated from the suspension by filtration. The filter cake was washed twice with 12 ml of methanol and then dried under vacuum to give gabapentin pure having a yield of 81.4%, pH of 7.25 and chlorine anion  
25 content of 35 ppm.

Gabapentin so prepared contained less than 0.5% by weight of lactam, and, after a year of storage at 25°C and 60% relative humidity, the conversion of gabapentin to its corresponding lactam does not exceed 0.2% by weight of  
30 gabapentin.

**EXAMPLE 12**

The following material were used:

Methanol for suspending 52.5 ml

Methanol for washings 2x15 ml

5 Sodium methoxide ~0.001 equivalents

Crude gabapentin was suspended in 180 ml of methanol at 25°C. The suspension was heated under mixing to 55° C when gabapentin was dissolved. Sodium methoxide was added to the solution and the solution was cooled slowly during one hour  
10 to 25° C. At 25°C the solution was concentrated to a volume of 50 ml. The suspension was stirred for twelve hours at 25°C. After 12 hours the solid gabapentin was separated from the suspension by filtration. The filter cake was washed twice with 12 ml of methanol, then dried under vacuum  
15 to give gabapentin pure at a yield of 81.4%, pH of 7.08 and anion content of Cl<sup>-</sup> 20 ppm.

Gabapentin so prepared contained less than 0.5% by weight of lactam, and, after a year of storage at 25°C and 50% relative humidity, the amount of lactam remained less  
20 than 0.5% by weight. After one year of storage at 25°C and 60% relative humidity, the conversion of gabapentin to its corresponding lactam is found not to exceed 0.2% by weight of gabapentin.

**EXAMPLE 13**

25 The following material were used:

Methanol for suspending 52.5 ml

Methanol for washings 2x15 ml

Sodium bicarbonate ~0.05 equivalents

Crude gabapentin was suspended in 180 ml of methanol at 25° C. The suspension was heated under mixing to 55°C when gabapentin was dissolved. Sodium bicarbonate was added to the solution and the solution was cooled slowly for one hour  
5 to 25°C. At 25°C the solution was concentrated to a volume of 50 ml. The suspension was stirred for twelve hours at 25°C. After 12 hours the solid gabapentin was separated from the suspension by filtration. The filter cake was washed twice with 12 ml of methanol, then dried under vacuum  
10 to give gabapentin pure having a yield of 72.4%, pH of 7.28 and anion (Cl<sup>-</sup>) content of 20 ppm.

Gabapentin so prepared contained less than 0.5% by weight of lactam, and, after a year of storage at 55°C and 50% relative humidity, the amount of lactam remained less  
15 than 0.5% by weight. After a year of storage at 25°C and 60% relative humidity, the conversion of gabapentin to its corresponding lactam is found not to exceed 0.2% by weight of gabapentin.

#### **EXAMPLE 14**

20 The following raw material were used:

Methanol for suspending	52.5 ml
Methanol for washing	2x15 ml
Tetramethylammoniumhydroxide	~0.002 equivalents

Crude gabapentin was suspended in 180 ml of methanol at 25° C. The suspension was heated under mixing to 55° C when gabapentin was dissolved. Tetramethylammoniumhydroxide was added to the solution and the solution was cooled slowly for one hour to 25°C. At 25° C the solution was concentrated to a volume of 50 ml. The suspension was stirred for 12 hours  
25 at 25°C. After 12 hours the solid gabapentin was separated from the suspension by filtration. The filter cake was  
30

washed twice with 12 ml of methanol than dried under vacuum to give gabapentin pure having a yield of 75.8%, pH of 7.03 and anion content of (Cl<sup>-</sup>) 20 ppm.

5 Gabapentin so prepared contained less than 0.5% by weight of lactam.

#### **EXAMPLE 15**

The following raw material were used:

	Methanol for suspending	52.5 ml
	Methanol for washing	2x15 ml
10	Tetrabutylammoniumhydroxide	~0.002 equivalents

Crude gabapentin was suspended in 180 ml of methanol at 25° C. The suspension was heated under mixing to 55°C when gabapentin was dissolved. Tetrabutylammoniumhydroxide was added to the solution and the solution was cooled slowly during one hour to 25°C. At 25°C the solution was concentrated to a volume of 50 ml. The suspension was stirred for 12 hours at 25°C. After 12 hours the solid gabapentin was separated from the suspension by filtration. The filter cake was washed twice with 12 ml of methanol, then dried under vacuum to give gabapentin pure having a yield of 77.6%, pH of 7.22 and anion (Cl<sup>-</sup>) content of 20 ppm.

#### **EXAMPLE 16**

The following raw material were used:

	Methanol for suspending	52.5 ml
25	Methanol for washing	2x15 ml
	Sodiumtetraborate	~0.05 equivalents

Crude gabapentin was suspended in 180 ml of methanol at 25° C. The suspension was heated under mixing to 55°C when

gabapentin was dissolved. Sodiumtetraborate was added to the solution and the solution was cooled slowly for one hour to 25°C. At 25°C the solution was concentrated to a volume of 50ml. The suspension was stirred for 12 hours at 25°C. After 12 hours the solid gabapentin was separated from the suspension by filtration. The filter cake was washed twice with 12 ml of methanol and then dried under vacuum to give gabapentin pure having a yield of 75%, pH of 7.17 and anion content (Cl<sup>-</sup>) of 10 ppm.

#### EXAMPLE 17

The following gabapentin tablet formulation is prepared using gabapentin containing chloride ion ranging from 5 to 40 ppm and pH in the range of 6.84 - 7.04 according to Example 1. The following material is used:

Ingredients	Amounts
gabapentin	125 g
Corn Starch NF	200 g
Cellulose, Microcrystalline	46 g
Sterotex Powder HM	4 g
Purified Water	q.s. or 300 ml

Combine corn starch, cellulose, and gabapentin together in a mixer and mix for 2-4 minutes. Add water to this combination and mix for an addition 1-3 minutes. The resulting mix is spread on trays and dried in convection oven at 45-55 °C until a moisture level of 1 to 2% is obtained. The dried mix is then milled and added back to the mill mixture and the total is blended for additional 4-5 minutes. Compressed tables of 150 mg, 375 mg and 750 mg are formed using appropriate punches from the total mix.

The formulation is measured to contain less than 0.5%

lactam and after one year of storage at 25 °C and 60% atmospheric humidity, the conversion of gabapentin to its corresponding lactam is found not to exceed 0.2% by weight of gabapentin.

5 **EXAMPLE 18**

Gabapentin of Example 2 (having chloride ion content of 50 ppm and pH of 7.15) is used to formulate tablets as in EXAMPLE 17, except that corn starch is replaced in each sample by one of the following adjuvants: pregelatinized  
10 starch, croscarmellose sodium, silica gel, titanium dioxide, talc, modified maize starch and maize starch.

The resulting gabapentin tablet of each sample is initially measured to have 0.5% by weight of a corresponding lactam, more than 50 ppm of chloride anion, and pH exceeding  
15 6.8. The tablet is stored for one year at 25 °C and 60% atmospheric humidity and the increase in the lactam concentration is found not to exceed 0.2% by weight.

**EXAMPLE 19**

EXAMPLE 18 is repeated except that gabapentin of  
20 Example 4, having chloride ion of 7 ppm is used for formulating tablets. The resulting gabapentin tablet of each sample is initially measured to have 0.5 % by weight of lactam and approximately 7 ppm of chloride anion. The tablet is stored for one year at 55 °C and 50% atmospheric  
25 humidity and the increase in the lactam concentration is found not to exceed 0.2% by weight.

Examples 17-19 show that, contrary to Augart's disclosure, the presence of anion of a mineral acid in an amount greater than 20 ppm does not adversely affect the  
30 stability of gabapentin when stored for one year at 25 °C and 60% humidity. The examples also show that the

gabapentin formulations prepared in accordance with the invention showed adequate stability regardless of the type of adjuvant that were used.

WE CLAIM:

1. A pharmaceutical composition comprising gabapentin and initially containing less than 0.5% by weight of a corresponding lactam with respect to the weight of gabapentin and having greater than 20 ppm of an anion of a mineral acid with respect to the weight of gabapentin, which, after one year of storage at 25 °C and 60% humidity the conversion of gabapentin to its corresponding lactam does not exceed 0.2% by weight of gabapentin.
2. The pharmaceutical composition of claim 1, further comprising at least one adjuvant.
3. The pharmaceutical composition of claim 2, wherein at least one adjuvant is selected from the group consisting of modified maize starch, glycerol behenic acid ester, sodium croscarmellose, methacrylic acid co-polymers (types A and C), anion exchangers, titanium dioxide, silica gels, hydroxypropylmethylcellulose, polyvinylpyrrolidone, crospovidon, poloxamer 407™, poloxamer 188™, sodium starch glycolate, copolyvidone, maize starch, cyclodextrin, lactose, talc, co-polymers of dimethylamino-methacrylic acid and neutral methacrylic acid ester.
4. The pharmaceutical composition of claim 3, wherein said silica gel is Aerosil 200™.
5. The pharmaceutical composition of claim 1, wherein said anion of a mineral acid is a halide.
6. The pharmaceutical composition of claim 5, wherein said halide is chloride.

7. The pharmaceutical composition of claim 1, wherein the amount of said anion of a mineral acid does not exceed 100 ppm.

8. Gabapentin which contains less than 0.5% of a  
5 corresponding lactam with respect to the weight of  
gabapentin and between 20 and 100 ppm of an anion of a  
mineral acid with respect to the weight of gabapentin, and  
which, after one year of storage at 25 °C and 60% humidity  
the conversion of gabapentin to the corresponding lactam  
10 does not exceed 0.2% by weight of gabapentin.

9. A pharmaceutical composition comprising gabapentin  
and at least one adjuvant, and initially containing less  
than 0.5% by weight of a corresponding lactam with respect  
to the weight of gabapentin and having greater than 20 ppm  
15 of chloride with respect to the weight of gabapentin,  
which, after one year of storage at 25 °C and 60% humidity  
the conversion of gabapentin to the corresponding lactam  
does not exceed 0.2% by weight of gabapentin.