

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

(19) World Intellectual Property Organization

International Bureau



(10) International Publication Number

WO 2015/035212 A1

(43) International Publication Date

12 March 2015 (12.03.2015)

(51) International Patent Classification:

C07D 311/72 (2006.01) A61P 31/00 (2006.01)
A61K 31/355 (2006.01)

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(21) International Application Number:

PCT/US2014/054354

(22) International Filing Date:

5 September 2014 (05.09.2014)

(25) Filing Language:

English

(26) Publication Language:

English

(30) Priority Data:

61/874,823 6 September 2013 (06.09.2013) US

(81) Designated States (unless otherwise indicated, for every kind of national protection available): AE, AG, AL, AM, AO, AT, AU, AZ, BA, BB, BG, BH, BN, BR, BW, BY, BZ, CA, CH, CL, 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, IR, IS, JP, KE, KG, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LU, LY, MA, MD, ME, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PA, PE, PG, PH, PL, PT, QA, RO, RS, RU, RW, SA, SC, SD, SE, SG, SK, SL, SM, ST, SV, SY, TH, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW.

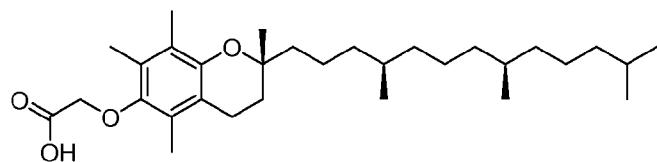
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(84) Designated States (unless otherwise indicated, for every kind of regional protection available): ARIPO (BW, GH, GM, KE, LR, LS, MW, MZ, NA, RW, SD, SL, ST, SZ, TZ, UG, ZM, ZW), Eurasian (AM, AZ, BY, KG, KZ, RU, TJ, TM), European (AL, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HR, HU, IE, IS, IT, LT, LU, LV, MC, MK, MT, NL, NO, PL, PT, RO, RS, SE, SI, SK, SM, TR), OAPI (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, KM, ML, MR, NE, SN, TD, TG).

Published:

— with international search report (Art. 21(3))

(54) Title: ALPHA-TEA SALT FORMS: COMPOSITIONS AND USES FOR TREATING DISEASE



(I)

(57) Abstract: The present disclosure relates to salts of the compound (I), polymorphic forms thereof, methods for preparation and use thereof, and pharmaceutical compositions thereof.

DESCRIPTION

ALPHA-TEA SALT FORMS: COMPOSITIONS AND USES FOR TREATING DISEASE

This application claims the benefit of US 61/874,823 filed on September 6, 2013, the entirety
5 of which is incorporated herein by reference.

BACKGROUND OF THE INVENTION

I. Field of the Invention

The present disclosure relates generally to salts of α -tocopheryloxyacetic acid (α -TEA), polymorphic forms thereof, and pharmaceutical compositions thereof. In other aspects, the present
10 disclosure also relates methods for preparation and use of these substances.

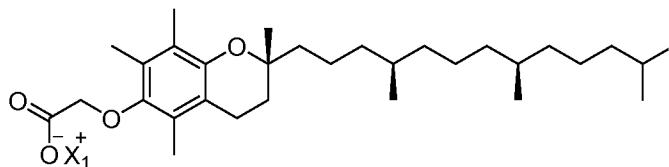
II. Description of Related Art

Vitamin E analogs such as α -tocopheryloxyacetic acid (α -TEA) have been previously reported (US 6,417,223, US 6,703,384 B2, US 6,770, 672 B1, and US 7,312,232 B2). The derivative α -TEA has been shown to suppress the growth of tumors through a mechanism which includes
15 mitochondrial depolarization and the generation of reactive oxygen species which trigger cell death (Yu, *et. al.*, 2010; Neuzil, *et. al.*, 2007; Kline, *et. al.*, 2007). Studies have shown that the compound inhibits the growth of a variety of different cancer cell lines without significant negative effects on
normal tissues (Lawson, *et al.*, 2003; Anderson, *et al.*, 2004; Hahn, *et al.*, 2009; Hahn, *et al.*, 2006).

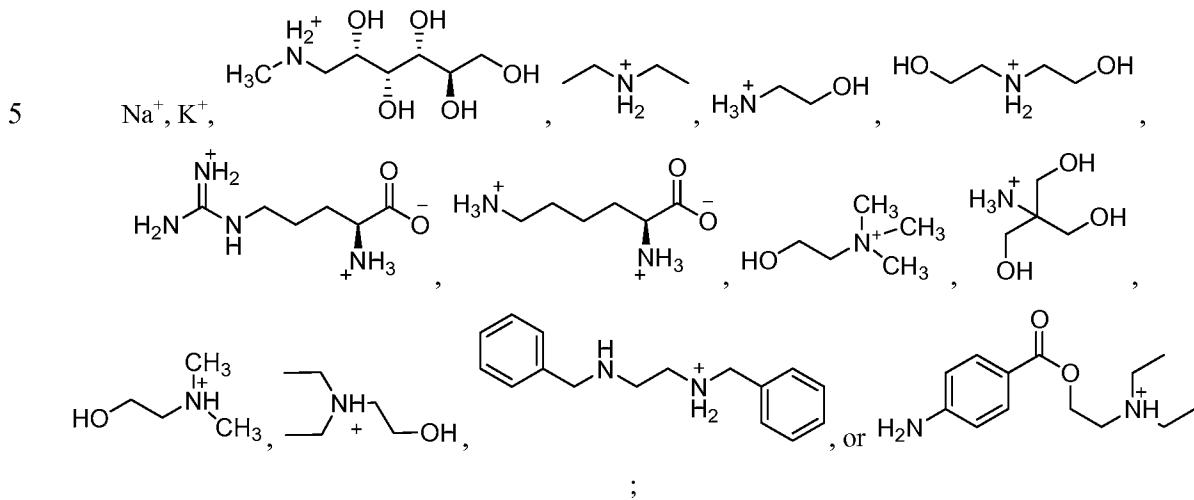
In view of these promising biological properties, and in consideration that biological activity
20 profiles and physical properties can be expected vary across different salt forms of a compound, it remains desirable to obtain novel salts forms with advantageous properties, including one that may ease process scale-up and formulation issues and/or possess biological activity or pharmacokinetic profiles that is well suited for the treatment or prevention of a given disease or indication. In view of the high degree of unmet medical need represented by the diseases and indications discussed herein, it
25 is therefore desirable to synthesize new solid forms, including salt forms, of α -TEA.

SUMMARY OF THE INVENTION

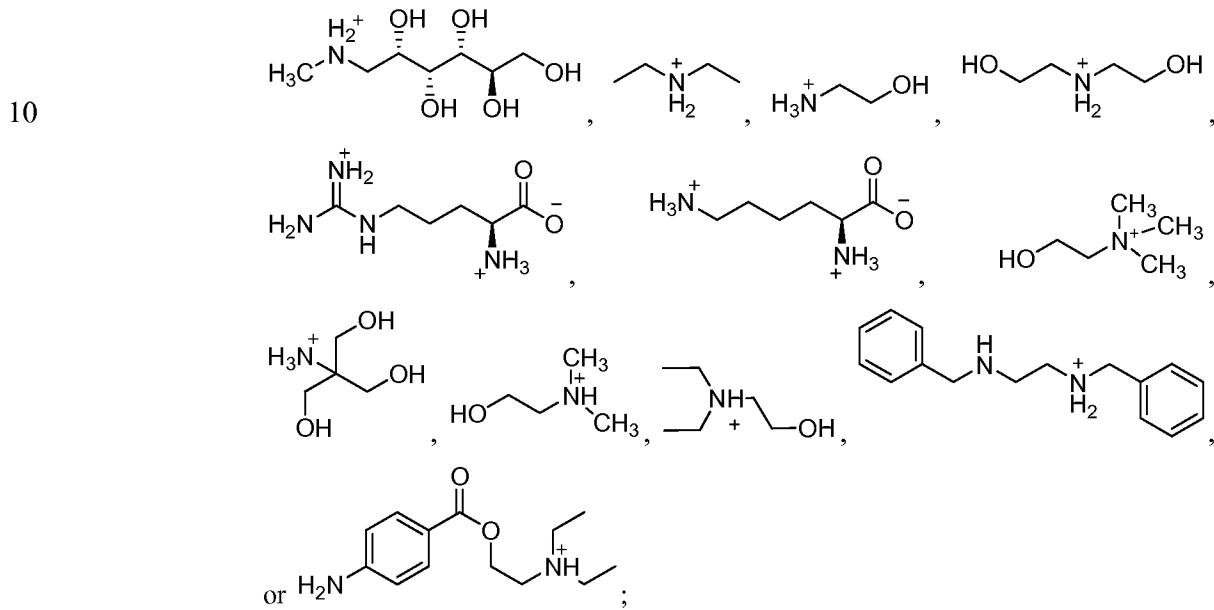
In one aspect, the present invention provides compounds of the formula:



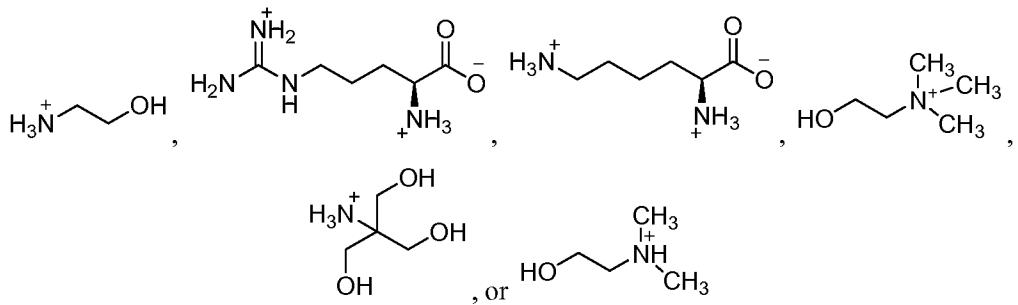
wherein X_1^+ is:



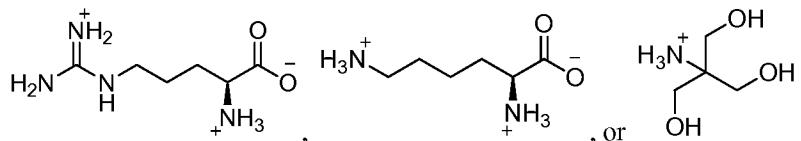
or a hydrate thereof. In some embodiments, X_1^+ is:



or a hydrate thereof. In some embodiments, X_1^+ is:

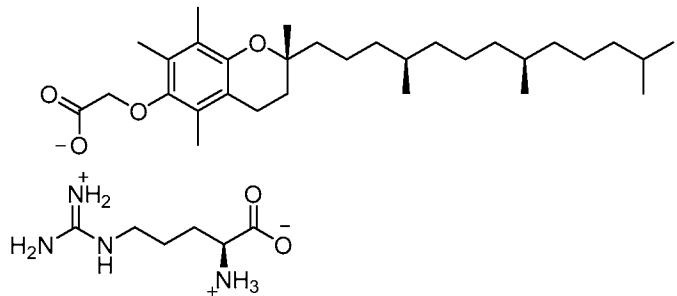


or a hydrate, thereof. In some embodiments, X_1^+ is



5 or a hydrate, thereof.

In some embodiments, the present disclosure provides a compound of the formula:

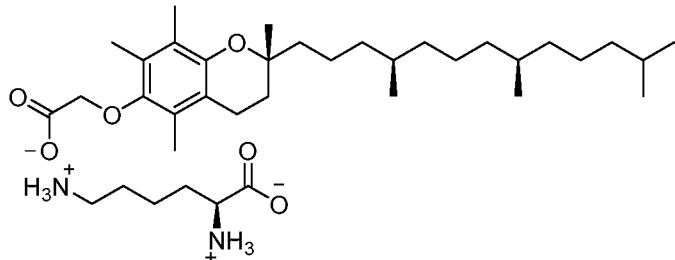


or a hydrate, thereof. In some embodiments, the compound is present as a polymorphic form having an X-ray powder diffraction pattern (CuK α) comprising two peaks at about 11.84 $^{\circ}$ 2 θ and 19.74 $^{\circ}$ 2 θ or

10 a hydrate, thereof. In some embodiments, the polymorphic form has an X-ray powder diffraction pattern (CuK α) further comprising a peak at about 20.73 $^{\circ}$ 2 θ . In some embodiments, the polymorphic form has an X-ray powder diffraction pattern (CuK α) further comprising 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, or all of peaks at about 6.24, 9.35, 14.92, 15.51, 17.03, 18.49, 21.70, 22.81, 23.51, 24.85, 27.69, 28.31, 30.93, 32.30, 33.21, 35.02, and 39.30 $^{\circ}$ 2 θ . In some embodiments, the polymorphic form has an X-ray powder diffraction pattern (CuK α) that is substantially as shown in FIG. 4. In some embodiments, the polymorphic form has a differential scanning calorimetry (DSC) isotherm comprising two endotherms from about 125 $^{\circ}$ C to about 140 $^{\circ}$ C and from about 170 $^{\circ}$ C to about 195 $^{\circ}$ C. In some embodiments, the polymorphic form has a differential scanning calorimetry (DSC) isotherm comprising two endotherms centered at about 131 $^{\circ}$ C and at around 184 $^{\circ}$ C, 15 respectively. In some embodiments, the polymorphic form has a differential scanning calorimetry isotherm that is substantially as shown in FIG. 5. In some embodiments, the polymorphic form has a thermogravimetric analysis (TGA) indicating a total volatile content of about 0.3 wt% over the temperature range of about 25-194 $^{\circ}$ C. In some embodiments, the polymorphic form has a dynamic vapor sorption (DVS) kinetic data and isotherm indicating a hemihydrate formation at about 40%

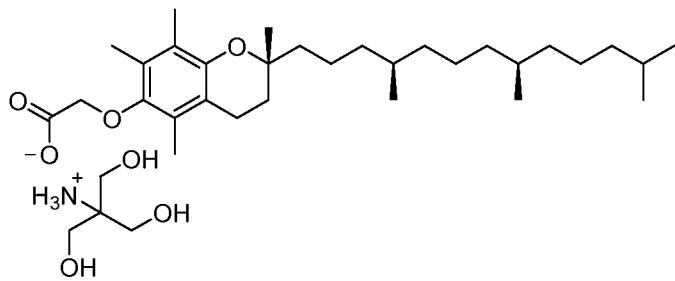
relative humidity. In some embodiments, the polymorphic form is anhydrous. In other embodiments, the polymorphic form is a hydrate. In some embodiments, the polymorphic form is a hemihydrate.

In yet another embodiment, the present disclosure provides a compound of the formula:



5 or a hydrate, thereof. In some embodiments, the compound is present as a polymorphic form having an X-ray powder diffraction pattern ($\text{CuK}\alpha$) comprising a peak at about $9.69^\circ 2\theta$ or a hydrate, thereof. In some embodiments, the X-ray powder diffraction pattern ($\text{CuK}\alpha$) further comprises a peak at about $20.10^\circ 2\theta$. In some embodiments, the X-ray powder diffraction pattern ($\text{CuK}\alpha$) further comprises a peak at about $19.78^\circ 2\theta$. In some embodiments, the X-ray powder diffraction pattern ($\text{CuK}\alpha$) further comprises 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, or all of peaks at about 5.90, 7.77, 14.08, 17.58, 17.91, 18.31, 21.26, 22.17, 23.16, 24.77, 28.57, 29.71, 31.10, 31.69, 34.45, 34.99, and 36.24 $^\circ 2\theta$. In some embodiments, the X-ray powder diffraction pattern ($\text{CuK}\alpha$) that is substantially as shown in FIG. 6. In some embodiments, the polymorphic form has a differential scanning calorimetry (DSC) isotherm comprising two endotherms from about 53°C to about 60°C and from about 180°C to about 235°C . In some embodiments, the polymorphic form has a differential scanning calorimetry (DSC) isotherm comprising two endotherms centered at about 55.8°C and at about 205°C , respectively. In some embodiments, the polymorphic form has a differential scanning calorimetry isotherm that is substantially as shown in FIG. 7. In some embodiments, the polymorphic form has a thermogravimetric curve from thermogravimetric analysis (TGA) indicating a total volatile content of about 0.4 wt% over the temperature range of about 25 - 168.8°C . In some embodiments, the polymorphic form has a dynamic vapor sorption (DVS) kinetic data and isotherm indicating a hydrate formation above about 75% relative humidity. In some embodiments, the polymorphic form is anhydrous. In other embodiments, the polymorphic form is a hydrate.

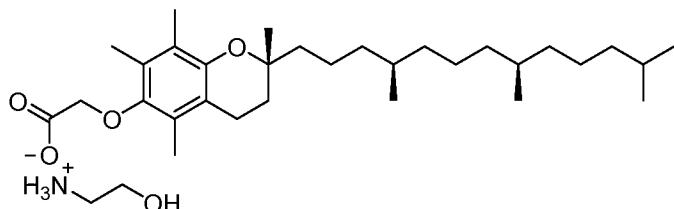
In another embodiment, the present disclosure provides a compound of the formula:



25 or a hydrate, thereof. In some embodiments, the compound is present as a polymorphic form having an X-ray powder diffraction pattern ($\text{CuK}\alpha$) comprising a peak at about $20.14^\circ 2\theta$ or a hydrate,

thereof. In some embodiments, the X-ray powder diffraction pattern (CuK α) further comprises peaks at about 13.63, 14.40, 19.31, and 22.34 $^{\circ}$ 20. In some embodiments, the X-ray powder diffraction pattern (CuK α) further comprises two peaks at about 18.25 and 18.49 $^{\circ}$ 20. In some embodiments, the X-ray powder diffraction pattern (CuK α) further comprises 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, or all of 5 peaks at about 9.67, 15.60, 21.47, 23.57, 27.07, 28.39, 30.49, 32.21, 33.30, 34.29, 34.96, 35.89, and 36.39 $^{\circ}$ 20. In some embodiments, the polymorphic form has an X-ray powder diffraction pattern (CuK α) that is substantially as shown in FIG. 10. In some embodiments, the polymorphic form has a differential scanning calorimetry (DSC) isotherm comprising two endotherms from about 97.5 $^{\circ}$ C to about 120 $^{\circ}$ C and from about 185 $^{\circ}$ C to about 225 $^{\circ}$ C. In some embodiments, the polymorphic form 10 has a differential scanning calorimetry (DSC) isotherm comprising two endotherms at about 106 $^{\circ}$ C and at about 192 $^{\circ}$ C, respectively. In some embodiments, the polymorphic form has a differential scanning calorimetry isotherm that is substantially as shown in FIG. 11. In some embodiments, the polymorphic form has a thermogravimetric curve from thermogravimetric analysis (TGA) indicating a 15 total volatile content of about 0.6 wt% over the temperature range of about 25-144 $^{\circ}$ C. In some embodiments, the polymorphic form has a dynamic vapor sorption (DVS) kinetic data and isotherm indicating a hemihydrate formation at about 55% relative humidity. In some embodiments, the polymorphic form is anhydrous. In other embodiments, the polymorphic form is a hydrate. In some embodiments, the polymorphic form is a hemihydrate.

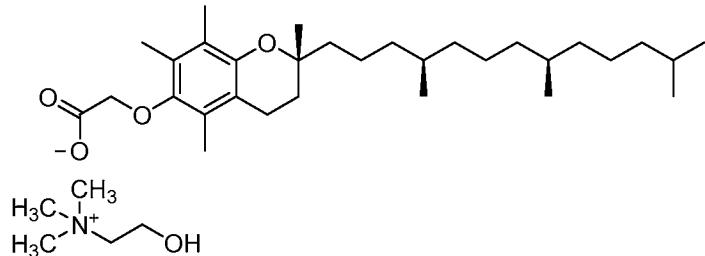
In yet another embodiment, the present disclosure provides a compound of the formula:



20 or a hydrate, thereof. In some embodiments, the compound is present as a polymorphic form having an X-ray powder diffraction pattern (CuK α) comprising a peak at about 19.4 $^{\circ}$ 20 or a hydrate, thereof. In some embodiments, the X-ray powder diffraction pattern (CuK α) further comprises peaks at about 10.4, 14.1, and 17.8 $^{\circ}$ 20. In some embodiments, the X-ray powder diffraction pattern (CuK α) further 25 comprises two peaks at about 20.8 $^{\circ}$ 20. In some embodiments, the polymorphic form has an X-ray powder diffraction pattern (CuK α) that is substantially as shown in FIG. 2. In some embodiments, the polymorphic form has a differential scanning calorimetry (DSC) isotherm comprising five endotherms from about 85 $^{\circ}$ C to about 97 $^{\circ}$ C, from about 97 $^{\circ}$ C to about 105 $^{\circ}$ C, from about 110 $^{\circ}$ C to about 125 $^{\circ}$ C, from about 145 $^{\circ}$ C to about 155 $^{\circ}$ C and from about 160 $^{\circ}$ C to about 210 $^{\circ}$ C. In some 30 embodiments, the polymorphic form has a differential scanning calorimetry (DSC) isotherm comprising five endotherms at about 90.1 $^{\circ}$ C, at about 100.7 $^{\circ}$ C, at about 117.9 $^{\circ}$ C, at about 149.7 $^{\circ}$ C, and at about 181.6 $^{\circ}$ C, respectively. In some embodiments, the polymorphic form has a differential scanning calorimetry isotherm that is substantially as shown in FIG. 3. In some embodiments, the

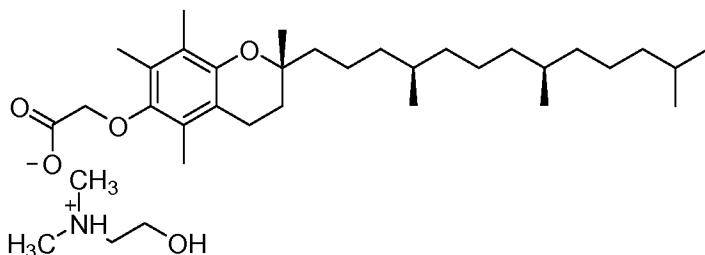
polymorphic form has a thermogravimetric curve from thermogravimetric analysis (TGA) indicating a total volatile content of about 0.4 wt% over the temperature range of about 25-95.5 °C.

In another embodiment, the present disclosure provides a compound of the formula:



5 or a hydrate, thereof. In some embodiments, the compound is present as a polymorphic form having an X-ray powder diffraction pattern (CuK α) comprising two peaks at about 18.40 and 19.77 °20 or hydrate, thereof. In some embodiments, the polymorphic form has an X-ray powder diffraction pattern (CuK α) further comprising peaks at about 4.28 and 10.75 °20. In some embodiments, the polymorphic form has an X-ray powder diffraction pattern (CuK α) further comprising two peaks at about 19.00 °20. In some embodiments, the polymorphic form has an X-ray powder diffraction pattern (CuK α) further comprising 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, or all of peaks at about 6.45, 12.90, 16.17, 16.73, 17.31, 17.72, 20.96, 21.72, 28.08, 28.71, 29.70, 31.05, 31.63, 32.25, 34.60, 35.18, 36.60, and 39.02 °20. In some embodiments, the polymorphic form has an X-ray powder diffraction pattern (CuK α) that is substantially as shown in FIG. 8. In some embodiments, the polymorphic form has a differential scanning calorimetry (DSC) isotherm comprising three endotherms from about 50 °C to about 60 °C, from about 75 °C to about 85 °C, and from about 150 °C to about 250 °C, respectively. In some embodiments, the polymorphic form has a differential scanning calorimetry (DSC) isotherm comprising three endotherms at about 50 °C, at about 80 °C, and at about 150 °C, respectively. In some embodiments, the polymorphic form has a differential scanning calorimetry isotherm that is substantially as shown in FIG. 9. In some embodiments, the polymorphic form has a thermogravimetric curve from thermogravimetric analysis (TGA) indicating a total volatile content of about 2.5 wt% over the temperature range of about 25-188.2 °C.

In another embodiment, the present disclosure provides a compound of the formula:

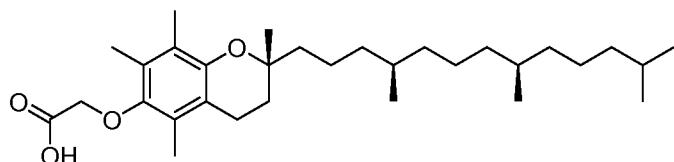


25 or a hydrate, thereof. In some embodiments, the compound is present as a polymorphic form having an X-ray powder diffraction pattern (CuK α) comprising a peak at about 21.73 °20 or a hydrate, thereof. In some embodiments, the polymorphic form has an X-ray powder diffraction pattern

(CuK α) further comprising peaks at about 9.34, 14.90, 16.06, 17.33, 18.51, and 22.94 $^{\circ}$ 20. In some embodiments, the polymorphic form has an X-ray powder diffraction pattern (CuK α) further comprising two peaks centered at about 20.22 $^{\circ}$ 20. In some embodiments, the polymorphic form has an X-ray powder diffraction pattern (CuK α) further comprising 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, or all of 5 peaks at about 10.05, 11.19, 14.13, 24.04, 27.65, 28.14, 29.73, 30.30, 31.06, 32.02, 33.45, and 34.49 $^{\circ}$ 20. In some embodiments, the polymorphic form has an X-ray powder diffraction pattern (CuK α) that is substantially as shown in FIG. 12. In some embodiments, the polymorphic form has a differential scanning calorimetry (DSC) isotherm comprising three endotherms from about 45 $^{\circ}$ C to about 55 $^{\circ}$ C, from about 65 $^{\circ}$ C to about 80 $^{\circ}$ C, and from about 80 $^{\circ}$ C to about 95 $^{\circ}$ C. In some 10 embodiments, the polymorphic form has a differential scanning calorimetry (DSC) isotherm comprising three endotherms at about 47 $^{\circ}$ C, at about 73 $^{\circ}$ C, and at about 87 $^{\circ}$ C, respectively. In some embodiments, the polymorphic form has a differential scanning calorimetry isotherm that is substantially as shown in FIG. 13. In some embodiments, the polymorphic form has a 15 thermogravimetric curve from thermogravimetric analysis (TGA) indicating a total volatile content of about 7.7 wt% over the temperature range of about 25-207 $^{\circ}$ C.

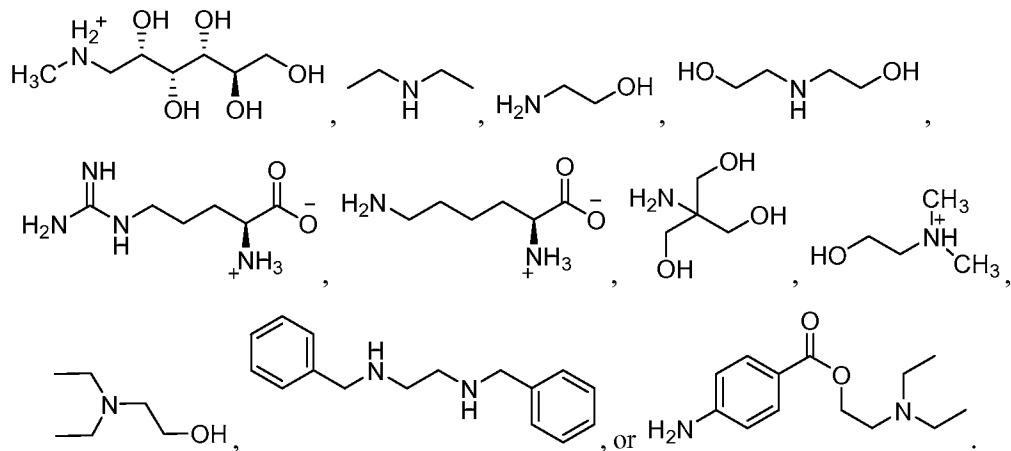
In another aspect, the present disclosure provides a pharmaceutical composition comprising a compound, salt, or polymorphic form of the present disclosure and an excipient. In some embodiments, the pharmaceutical composition is formulated for administration: orally, intraadiposally, intraarterially, intraarticularly, intracranially, intradermally, intralesionally, 20 intramuscularly, intranasally, intraocularly, intrapericardially, intraperitoneally, intrapleurally, intraprostatically, intrarectally, intrathecally, intratracheally, intratumorally, intraumbilically, intravaginally, intravenously, intravesicularily, intravitrealy, liposomally, locally, mucosally, parenterally, rectally, subconjunctival, subcutaneously, sublingually, topically, transbuccally, 25 transdermally, vaginally, in crèmes, in lipid compositions, via a catheter, via a lavage, via continuous infusion, via infusion, via inhalation, via injection, via local delivery, or via localized perfusion. In some embodiments, the pharmaceutical composition is formulated for oral, intraarterial, intravenous, intraperitoneal, topical, or inhalational administration.

In yet another aspect, the present disclosure provides a method of reacting a compound of the formula:

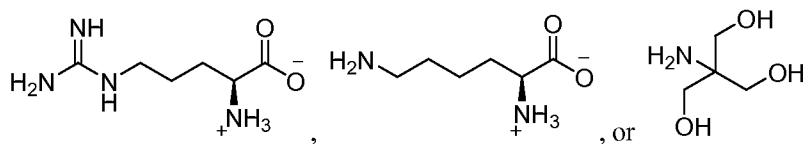


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to obtain a polymorphic form comprising combining an about equal molar amount of a base and the compound in an alcoholic solvent and mixing for a time period from about 1 to about 24 hours. In some embodiments, the base is NaOH, KOH,



In some embodiments, the base is:

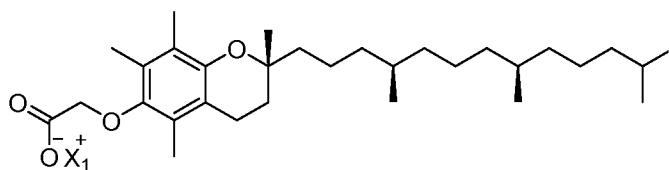


In some embodiments, the alcoholic solvent is of the formula: $\text{HO}-\text{R}_1$, wherein R_1 is alkyl_(C≤6). In some embodiments, the alcoholic solvent is methanol, 1-butanol, or isopropyl alcohol. In some embodiments, the alcoholic solvent is isopropyl alcohol. In some embodiments, the method further comprises allowing the solution to slowly evaporate at room temperature. In some embodiments, the method further comprises allowing the evaporation to occur under a constant flow of nitrogen gas. In some embodiments, the constant flow of nitrogen has a pressure of about 2 psi.

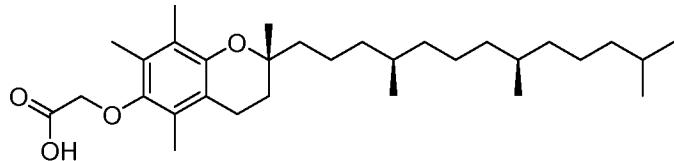
In still another aspect, the present invention provides a method of treating or preventing a hyperproliferative disease, comprising administering to the patient a therapeutically effective amount of the pharmaceutical composition, salt, polymorphic form, or compound of the present disclosure. In some embodiments, the hyperproliferative disease is a cancer. In some embodiments, the cancer is breast cancer, lung cancer, colon cancer, an endometrial tumor, cervical cancer, ovarian cancer, mammary cancer, prostate cancer, liver cancer, melanoma, lymphoma, or myeloma. In some embodiments, the cancer is breast cancer. In some embodiments, the method further comprises administering to the patient a second therapy. In some embodiments, the second therapy is an immunotherapy or surgery. In some embodiments, the second therapy is an immunotherapy. In some embodiments, the immunotherapy comprises administering a pharmaceutically effective amount of trastuzumab, adoptively transferred T lymphocytes, or a therapeutic antibody. In some embodiments, the therapeutic antibody selectively binds HER2/neu. In some embodiments, the immunotherapy is trastuzumab. In some embodiments, the immunotherapy comprises administering adoptively transferred T lymphocytes to the patient. In some embodiments, the adoptively transferred T lymphocytes are engineered to express a chimeric antigen receptor (CAR) to the patient. In some

embodiments, the immunotherapy comprises administering an immune modulating antibody to the patient. In some embodiments, the immune modulating antibody is an anti-PD-1, anti-4-1-BB, anti-GITR, anti-TIM3, anti-LAG3, anti-TIGIT, anti-CTLA-4 or an anti-LIGHT antibody. In other embodiments, the second therapy is radiotherapy or a chemotherapeutic agent. In other embodiments, 5 the second therapy is surgery. In other embodiments, the second therapy is a chemotherapeutic agent. In some embodiments, the chemotherapeutic agent is anthracyclines, taxanes, methotrexate, mitoxantrone, estramustine, doxorubicin, etoposide, vinblastine, carboplatin, vinorelbine, 5-fluorouracil, cisplatin, topotecan, ifosfamide, cyclophosphamide, epirubicin, gemcitabine, vinorelbine, 10 irinotecan, etoposide, vinblastine, pemetrexed, melphalan, capecitabine, oxaliplatin, BRAF inhibitors, and TGF-beta inhibitors.

In another aspect, the present disclosure provides a method of making a salt of the formula:



wherein: X_1^+ is a protonated base; comprising reacting a compound of the formula:



15 with the base sufficient to form a salt; or a hydrate thereof. In some embodiments, the base is: sodium hydroxide, potassium hydroxide, magnesium hydroxide, meglumine, diethylamine, ethanolamine, diethanolamine, L-arginine, L-lysine, choline, tris-hydroxymethylamine, N,N-dimethylethanolamine, 2-diethylethanolamine, calcium hydroxide, dibenzylethylenediamine, or procaine; or a hydrate thereof. In some embodiments, the base is ethanolamine, L-arginine, L-lysine, choline, tris-20 hydroxymethylamine, or N,N-dimethylethanolamine. In some embodiments, the base is L-arginine, L-lysine, or tris-hydroxymethylamine.

Other objects, features and advantages of the present disclosure will become apparent from the following detailed description. It should be understood, however, that the detailed description and the specific examples, while indicating specific embodiments of the invention, are given by way of 25 illustration only, since various changes and modifications within the spirit and scope of the invention will become apparent to those skilled in the art from this detailed description. Note that simply because a particular compound is ascribed to one particular generic formula does not mean that it cannot also belong to another generic formula.

BRIEF DESCRIPTION OF THE DRAWINGS

The following drawings form part of the present specification and are included to further demonstrate certain aspects of the present disclosure. The invention may be better understood by reference to one of these drawings in combination with the detailed description of specific 5 embodiments presented herein.

FIG. 1 – The CuK α powder x-ray diffractogram of the α -TEA free acid. The diffractogram shows the dominate peaks of the free acid to be about 17.78 $^{\circ}$ 20.

FIG. 2 – A CuK α powder x-ray diffractogram of the ethanolamine salt candidates from panels 1-3 compared to the spectra of the α -TEA free acid. The spectra for the free acid is shown at the bottom, 10 the ethanolamine salt from panel 2 is shown second from the bottom, the ethanolamine salt from panel 1 is shown in the second from the top spectra, and the ethanolamine salt from panel 3 is shown at the top.

FIG. 3 – A differential scanning calorimetry (DSC) isotherm of the ethanolamine salt candidate of the α -TEA free acid showing five endotherms with the three largest endotherms at about 90, 118, and 182 15 $^{\circ}$ C.

FIG. 4 – A CuK α powder x-ray diffractogram of a representative arginine salt candidate from panels 1-3. The spectra shows a dominate peak at about 19.74 $^{\circ}$ 20.

FIG. 5 – A differential scanning calorimetry (DSC) isotherm of the arginine salt candidate of the α -TEA free acid showing two endotherms with the endotherms at about 131 and 184 $^{\circ}$ C.

20 **FIG. 6** – A CuK α powder x-ray diffractogram of a representative lysine salt candidate. The spectra shows a dominate peak at about 9.69 $^{\circ}$ 20.

FIG. 7 – A differential scanning calorimetry (DSC) isotherm of the lysine salt candidate of the α -TEA free acid showing two endotherms with the endotherms at about 56 and 205 $^{\circ}$ C.

25 **FIG. 8** – A CuK α powder x-ray diffractogram of a representative choline salt candidate from panels 1-3. The spectra shows a pair of dominate peaks at about 18.4 and 19.77 $^{\circ}$ 20.

FIG. 9 – A differential scanning calorimetry (DSC) isotherm of the choline salt candidate of the α -TEA free acid showing three endotherms with the endotherms at about 50, 80 and 150 $^{\circ}$ C.

FIG. 10 – A CuK α powder x-ray diffractogram of a representative tris salt candidate. The spectra shows a dominate peak at about 20.14 $^{\circ}$ 20.

FIG. 11 – A differential scanning calorimetry (DSC) isotherm of the tris salt candidate of the α -TEA free acid showing two endotherms with the endotherms at about 106 and 192 °C.

FIG. 12 – A CuK α powder x-ray diffractogram of a representative N,N-dimethyl-ethanolamine salt candidate from panels 1-3. The spectra shows a dominate peak at about 21.73 °20.

5 **FIG. 13** – A differential scanning calorimetry (DSC) isotherm of the N,N-dimethyl-ethanolamine salt candidate of the α -TEA free acid showing three endotherms with the endotherms at about 47, 73, and 87 °C.

FIG. 14 – A graph showing HPLC traces of different α -TEA free acid samples that have been subjected to different stresses to test the stability of the compound in that form.

10 **FIG. 15** – A graph showing HPLC traces of different α -TEA lysine salt samples that have been subjected to different stresses to test the stability of the compound in that form.

FIG. 16 – A graph showing HPLC traces of different α -TEA arginine salt samples that have been subjected to different stresses to test the stability of the compound in that form.

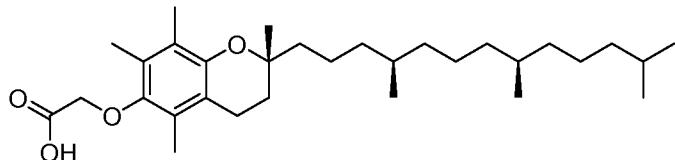
15 **FIG. 17** – A graph showing HPLC traces of different α -TEA tris salt samples that have been subjected to different stresses to test the stability of the compound in that form.

FIG. 18 – A graph of the average tumor size of BALB/c mice with a 4T1 mammary tumor in mammary fat pad for control group and mice feed a diet which contained the α -TEA lysine salt.

FIG. 19 – A graph of the Kaplan-Meier analysis of survival of the BALB/c mice on a diet of α -TEA lysine salt compared to mice fed a nutrient analogous control diet.

DESCRIPTION OF ILLUSTRATIVE EMBODIMENTS

The present disclosure provides in one aspect, salts of the compound:



and polymorphic forms of the compound, thereof. In another non-limiting example, the present
5 disclosure relates to pharmaceutical compositions of the polymorphic forms and the method of preparation of these compounds and the method of use of these compounds to treat diseases.

I. Definitions

When used in the context of a chemical group: “hydrogen” means $-H$; “hydroxy” means $-OH$; “oxo” means $=O$; “carboxylic acid”, “carboxy”, or “carboxyl” means $-C(=O)OH$ (also written as $-COOH$ or $-CO_2H$); “carboxylate” means $-C(=O)O^-$ (also written as $-COO^-$ or $-CO_2^-$); “halo” means independently $-F$, $-Cl$, $-Br$ or $-I$. Any undefined valency on an atom of a structure shown in this application implicitly represents a hydrogen atom bonded to the atom.

In the context of chemical formulas, the symbol “ $-$ ” means a single bond, “ $=$ ” means a double bond; and “ \equiv ” means triple bond. The symbol “ \nearrow ” means a single bond where the group attached to the thick end of the wedge is “out of the page.” The symbol “ \nwarrow ” means a single bond where the group attached to the thick end of the wedge is “into the page”. The symbol “ $\sim\sim$ ” means a single bond where the conformation (e.g., either *R* or *S*) or the geometry is undefined (e.g., either *E* or *Z*).

For the groups and classes below, the following parenthetical subscripts further define the group/class as follows: “(C_n)” defines the exact number (n) of carbon atoms in the group/class. 20 “(C_{≤n})” defines the maximum number (n) of carbon atoms that can be in the group/class, with the minimum number as small as possible for the group in question, e.g., “alkyl_(C≤8)” For example, “alkyl_(C≤10)” designates those alkyl groups having from 1 to 10 carbon atoms (e.g., 1, 2, 3, 4, 5, 6, 7, 8, 9, or 10, or any range derivable therein (e.g., 3 to 10 carbon atoms). (C_{n-n'}) defines both the minimum (n) and maximum number (n') of carbon atoms in the group. Similarly, “alkyl_(C2-10)” 25 designates those alkyl groups having from 2 to 10 carbon atoms (e.g., 2, 3, 4, 5, 6, 7, 8, 9, or 10, or any range derivable therein (e.g., 3 to 10 carbon atoms)).

The term “saturated” as used herein means the compound or group so modified has no carbon-carbon double and no carbon-carbon triple bonds, except as noted below. The term does not preclude carbon-heteroatom multiple bonds, for example a carbon oxygen double bond or a carbon nitrogen double bond. Moreover, it does not preclude a carbon-carbon double bond that may occur as part of keto-enol tautomerism or imine/enamine tautomerism.

The term “aliphatic” when used without the “substituted” modifier signifies that the compound/group so modified is an acyclic or cyclic, but non-aromatic hydrocarbon compound or group. In aliphatic compounds/groups, the carbon atoms can be joined together in straight chains, branched chains, or non-aromatic rings (alicyclic). Aliphatic compounds/groups can be saturated, that 5 is joined by single bonds (alkanes/alkyl), or unsaturated, with one or more double bonds (alkenes/alkenyl) or with one or more triple bonds (alkynes/alkynyl). When the term “aliphatic” is used without the “substituted” modifier only carbon and hydrogen atoms are present. When the term is used with the “substituted” modifier one or more hydrogen atom has been independently replaced by $-\text{OH}$, $-\text{F}$, $-\text{Cl}$, $-\text{Br}$, $-\text{I}$, $-\text{NH}_2$, $-\text{NO}_2$, $-\text{CO}_2\text{H}$, $-\text{CO}_2\text{CH}_3$, $-\text{CN}$, $-\text{SH}$, $-\text{OCH}_3$, $-\text{OCH}_2\text{CH}_3$, $-\text{C(O)CH}_3$, $-\text{N(CH}_3)_2$, $-\text{C(O)NH}_2$, $-\text{OC(O)CH}_3$, or $-\text{S(O)}_2\text{NH}_2$.
10

The term “alkyl” when used without the “substituted” modifier refers to a monovalent saturated aliphatic group with a carbon atom as the point of attachment, a linear or branched, cyclo, cyclic or acyclic structure, and no atoms other than carbon and hydrogen. Thus, as used herein cycloalkyl is a subset of alkyl. The groups $-\text{CH}_3$ (Me), $-\text{CH}_2\text{CH}_3$ (Et), $-\text{CH}_2\text{CH}_2\text{CH}_3$ (*n*-Pr), 15 $-\text{CH}(\text{CH}_3)_2$ (*iso*-Pr), $-\text{CH}(\text{CH}_2)_2$ (cyclopropyl), $-\text{CH}_2\text{CH}_2\text{CH}_2\text{CH}_3$ (*n*-Bu), $-\text{CH}(\text{CH}_3)\text{CH}_2\text{CH}_3$ (*sec*-butyl), $-\text{CH}_2\text{CH}(\text{CH}_3)_2$ (*iso*-butyl), $-\text{C}(\text{CH}_3)_3$ (*tert*-butyl), $-\text{CH}_2\text{C}(\text{CH}_3)_3$ (*neo*-pentyl), cyclobutyl, cyclopentyl, cyclohexyl, and cyclohexylmethyl are non-limiting examples of alkyl groups. When any of these terms is used with the “substituted” modifier one or more hydrogen atom has been independently replaced by $-\text{OH}$, $-\text{F}$, $-\text{Cl}$, $-\text{Br}$, $-\text{I}$, $-\text{NH}_2$, $-\text{NO}_2$, $-\text{CO}_2\text{H}$, $-\text{CO}_2\text{CH}_3$, $-\text{CN}$, $-\text{SH}$, 20 $-\text{OCH}_3$, $-\text{OCH}_2\text{CH}_3$, $-\text{C(O)CH}_3$, $-\text{N(CH}_3)_2$, $-\text{C(O)NH}_2$, $-\text{OC(O)CH}_3$, or $-\text{S(O)}_2\text{NH}_2$. The following groups are non-limiting examples of substituted alkyl groups: $-\text{CH}_2\text{OH}$, $-\text{CH}_2\text{Cl}$, $-\text{CF}_3$, $-\text{CH}_2\text{CN}$, $-\text{CH}_2\text{C(O)OH}$, $-\text{CH}_2\text{C(O)OCH}_3$, $-\text{CH}_2\text{C(O)NH}_2$, $-\text{CH}_2\text{C(O)CH}_3$, $-\text{CH}_2\text{OCH}_3$, $-\text{CH}_2\text{OC(O)CH}_3$, 25 $-\text{CH}_2\text{NH}_2$, $-\text{CH}_2\text{N(CH}_3)_2$, and $-\text{CH}_2\text{CH}_2\text{Cl}$. An “alkane” refers to the compound H–R, wherein R is alkyl. The term “alcohol” is an alkane as that term is defined herein wherein one or more of the hydrogen atoms have been replaced with a hydroxyl group.

The term “heteroaryl” when used without the “substituted” modifier refers to a monovalent aromatic group with an aromatic carbon atom or nitrogen atom as the point of attachment, said carbon atom or nitrogen atom forming part of one or more aromatic ring structures wherein at least one of the ring atoms is nitrogen, oxygen or sulfur, and wherein the heteroaryl group consists of no atoms other 30 than carbon, hydrogen, aromatic nitrogen, aromatic oxygen and aromatic sulfur. If more than one ring is present, the rings may be fused or unfused. As used herein, the term does not preclude the presence of one or more alkyl, aryl, and/or aralkyl groups (carbon number limitation permitting) attached to the aromatic ring or aromatic ring system. Non-limiting examples of heteroaryl groups include furanyl, imidazolyl, indolyl, indazolyl (Im), isoxazolyl, methylpyridinyl, oxazolyl, phenylpyridinyl, pyridinyl, 35 pyrrolyl, pyrimidinyl, pyrazinyl, quinolyl, quinazolyl, quinoxalinyl, triazinyl, tetrazolyl, thiazolyl, thienyl, and triazolyl. The term “*N*-heteroaryl” refers to a heteroaryl group with a nitrogen atom as the point of attachment. A “heteroarene” refers to the compound H–R, wherein R is heteroaryl.

Pyridine and quinoline are non-limiting examples of heteroarenes. When these terms are used with the “substituted” modifier one or more hydrogen atom has been independently replaced by –OH, –F, –Cl, –Br, –I, –NH₂, –NO₂, –CO₂H, –CO₂CH₃, –CN, –SH, –OCH₃, –OCH₂CH₃, –C(O)CH₃, –NHCH₃, –NHCH₂CH₃, –N(CH₃)₂, –C(O)NH₂, –OC(O)CH₃, or –S(O)₂NH₂.

5 The term “heterocycloalkyl” when used without the “substituted” modifier refers to a monovalent non-aromatic group with a carbon atom or nitrogen atom as the point of attachment, said carbon atom or nitrogen atom forming part of one or more non-aromatic ring structures wherein at least one of the ring atoms is nitrogen, oxygen or sulfur, and wherein the heterocycloalkyl group consists of no atoms other than carbon, hydrogen, nitrogen, oxygen and sulfur. If more than one ring
10 is present, the rings may be fused or unfused. As used herein, the term does not preclude the presence of one or more alkyl groups (carbon number limitation permitting) attached to the ring or ring system. Also, the term does not preclude the presence of one or more double bonds in the ring or ring system, provided that the resulting group remains non-aromatic. Non-limiting examples of heterocycloalkyl
15 groups include aziridinyl, azetidinyl, pyrrolidinyl, piperidinyl, piperazinyl, morpholinyl, thiomorpholinyl, tetrahydrofuranyl, tetrahydrothiofuranyl, tetrahydropyranyl, pyranyl, oxiranyl, and oxetanyl. The term “N-heterocycloalkyl” refers to a heterocycloalkyl group with a nitrogen atom as the point of attachment. When these terms are used with the “substituted” modifier one or more
20 hydrogen atom has been independently replaced by –OH, –F, –Cl, –Br, –I, –NH₂, –NO₂, –CO₂H, –CO₂CH₃, –CN, –SH, –OCH₃, –OCH₂CH₃, –C(O)CH₃, –NHCH₃, –NHCH₂CH₃, –N(CH₃)₂, –C(O)NH₂, –OC(O)CH₃, –S(O)₂NH₂, or –C(O)OC(CH₃)₃ (*tert*-butyloxycarbonyl, BOC).

The term “alkylamino” when used without the “substituted” modifier refers to the group –NHR, in which R is an alkyl, as that term is defined above. Non-limiting examples include: –NHCH₃ and –NHCH₂CH₃. The term “dialkylamino” when used without the “substituted” modifier refers to the group –NRR', in which R and R' can be the same or different alkyl groups. Non-limiting examples of dialkylamino groups include: –N(CH₃)₂ and –N(CH₃)(CH₂CH₃). Similarly, a trialkylamino is the group –RR'R'' in which R, R', and R'' can be the same or different alkyl groups. Non-limiting examples of dialkylamino groups include: –N(CH₃)₃⁺ and –N(CH₃)₂(CH₂CH₃)⁺. Furthermore, a trialkylamino group forms a positive charge when bonded to another compound. When any of these terms is used with the “substituted” modifier one or more hydrogen atom attached
25 to a carbon atom has been independently replaced by –OH, –F, –Cl, –Br, –I, –NH₂, –NO₂, –CO₂H, –CO₂CH₃, –CN, –SH, –OCH₃, –OCH₂CH₃, –C(O)CH₃, –NHCH₃, –NHCH₂CH₃, –N(CH₃)₂, –C(O)NH₂, –OC(O)CH₃, or –S(O)₂NH₂.

The use of the word “a” or “an,” when used in conjunction with the term “comprising” in the claims and/or the specification may mean “one,” but it is also consistent with the meaning of “one or
35 more,” “at least one,” and “one or more than one.”

Throughout this application, the term “about” is used to indicate that a value includes the inherent variation of error for the device, the method being employed to determine the value, or the

variation that exists among the study subjects. When used in the context of X-ray powder diffraction, the terms “about” and “substantially” are used interchangeably to indicate a value of ± 0.5 $^{\circ}$ 20 from the reported value, preferably a value of ± 0.2 $^{\circ}$ 20 from the reported value. When used in the context of differential scanning calorimetry, thermogravimetric analysis, or glass transition temperatures, the 5 terms “about” and “substantially” are used interchangeably to indicate a value of ± 10 $^{\circ}$ C relative to the maximum of the peak, preferably a value of ± 2 $^{\circ}$ C relative to the maximum of the peak.

The term “base” when used in the context of this application represents a compound which contains a group which contains a group which has a lone pair of electrons or a highly polarized and labile bond such as a metal alkane like butyllithium or a Gringard reagent. Such bases can form a 10 positively charged compound which interacts with the negatively charged carboxylate to form a “salt”. In some embodiments, the base is a nitrogenous base which contains a nitrogen atom with a lone pair of electrons which can act as a traditional Lewis base. Such nitrogenous bases is a compound which contains at least one amino, alkylamino, dialkylamino, trialkylamino, or a heteroaryl or heterocycloalkyl group which contains at least one nitrogen atom. In some embodiments, the base 15 can be a substituted version of these compounds wherein the substituted version comprises a compound wherein at least one hydrogen atom has been replaced with $-\text{OH}$, $-\text{F}$, $-\text{Cl}$, $-\text{Br}$, $-\text{I}$, $-\text{NH}_2$, $-\text{NO}_2$, $-\text{CO}_2\text{H}$, $-\text{CO}_2\text{CH}_3$, $-\text{CN}$, $-\text{SH}$, $-\text{OCH}_3$, $-\text{OCH}_2\text{CH}_3$, $-\text{C}(\text{O})\text{CH}_3$, $-\text{N}(\text{CH}_3)_2$, $-\text{C}(\text{O})\text{NH}_2$, $-\text{OC}(\text{O})\text{CH}_3$, or $-\text{S}(\text{O})_2\text{NH}_2$.

The terms “comprise,” “have” and “include” are open-ended linking verbs. Any forms or 20 tenses of one or more of these verbs, such as “comprises,” “comprising,” “has,” “having,” “includes” and “including,” are also open-ended. For example, any method that “comprises,” “has” or “includes” one or more steps is not limited to possessing only those one or more steps and also covers other unlisted steps.

The term “effective,” as that term is used in the specification and/or claims, means adequate 25 to accomplish a desired, expected, or intended result. “Effective amount,” “Therapeutically effective amount” or “pharmaceutically effective amount” when used in the context of treating a patient or subject with a compound means that amount of the compound which, when administered to a subject or patient for treating a disease, is sufficient to effect such treatment for the disease.

The term “hydrate” when used as a modifier to a compound means that the compound has 30 less than one (e.g., hemihydrate), one (e.g., monohydrate), or more than one (e.g., dihydrate) water molecules associated with each compound molecule, such as in solid forms of the compound.

As used herein, the term “ IC_{50} ” refers to an inhibitory dose which is 50% of the maximum response obtained. This quantitative measure indicates how much of a particular drug or other substance (inhibitor) is needed to inhibit a given biological, biochemical or chemical process (or 35 component of a process, i.e. an enzyme, cell, cell receptor or microorganism) by half.

An “isomer” of a first compound is a separate compound in which each molecule contains the same constituent atoms as the first compound, but where the configuration of those atoms in three dimensions differs.

As used herein, the term “patient” or “subject” refers to a living mammalian organism, such 5 as a human, monkey, cow, horse, sheep, goat, dog, cat, mouse, rat, guinea pig, or transgenic species thereof. In certain embodiments, the patient or subject is a primate. Non-limiting examples of human subjects are adults, juveniles, infants and fetuses.

As generally used herein “pharmaceutically acceptable” refers to those compounds, materials, 10 compositions, and/or dosage forms which are, within the scope of sound medical judgment, suitable for use in contact with the tissues, organs, and/or bodily fluids of human beings and animals without excessive toxicity, irritation, allergic response, or other problems or complications commensurate with a reasonable benefit/risk ratio.

“Prevention” or “preventing” includes: (1) inhibiting the onset of a disease in a subject or patient which may be at risk and/or predisposed to the disease but does not yet experience or display 15 any or all of the pathology or symptomatology of the disease, and/or (2) slowing the onset of the pathology or symptomatology of a disease in a subject or patient which may be at risk and/or predisposed to the disease but does not yet experience or display any or all of the pathology or symptomatology of the disease.

A “stereoisomer” or “optical isomer” is an isomer of a given compound in which the same 20 atoms are bonded to the same other atoms, but where the configuration of those atoms in three dimensions differs. “Enantiomers” are stereoisomers of a given compound that are mirror images of each other, like left and right hands. “Diastereomers” are stereoisomers of a given compound that are not enantiomers. Chiral molecules contain a chiral center, also referred to as a stereocenter or stereogenic center, which is any point, though not necessarily an atom, in a molecule bearing groups 25 such that an interchanging of any two groups leads to a stereoisomer. In organic compounds, the chiral center is typically a carbon, phosphorus or sulfur atom, though it is also possible for other atoms to be stereocenters in organic and inorganic compounds. A molecule can have multiple stereocenters, giving it many stereoisomers. In compounds whose stereoisomerism is due to tetrahedral stereogenic centers (e.g., tetrahedral carbon), the total number of hypothetically possible 30 stereoisomers will not exceed 2^n , where n is the number of tetrahedral stereocenters. Molecules with symmetry frequently have fewer than the maximum possible number of stereoisomers. A 50:50 mixture of enantiomers is referred to as a racemic mixture. Alternatively, a mixture of enantiomers can be enantiomerically enriched so that one enantiomer is present in an amount greater than 50%. Typically, enantiomers and/or diastereomers can be resolved or separated using techniques known in 35 the art. It is contemplated that for any stereocenter or axis of chirality for which stereochemistry has not been defined, that stereocenter or axis of chirality can be present in its R form, S form, or as a mixture of the R and S forms, including racemic and non-racemic mixtures. As used herein, the

phrase “substantially free from other stereoisomers” means that the composition contains $\leq 15\%$, more preferably $\leq 10\%$, even more preferably $\leq 5\%$, or most preferably $\leq 1\%$ of another stereoisomer(s).

“Treatment” or “treating” includes (1) inhibiting a disease in a subject or patient experiencing or displaying the pathology or symptomatology of the disease (e.g., arresting further development of the pathology and/or symptomatology), (2) ameliorating a disease in a subject or patient that is experiencing or displaying the pathology or symptomatology of the disease (e.g., reversing the pathology and/or symptomatology), and/or (3) effecting any measurable decrease in a disease in a subject or patient that is experiencing or displaying the pathology or symptomatology of the disease.

The above definitions supersede any conflicting definition in any of the reference that is incorporated by reference herein. The fact that certain terms are defined, however, should not be considered as indicative that any term that is undefined is indefinite. Rather, all terms used are believed to describe the invention in terms such that one of ordinary skill can appreciate the scope and practice the present invention.

II. Salts of α -TEA and Synthetic Methods

While the compound has shown improved efficacy in treating such tumors, α -TEA has proven difficult to formulate into a commercially successful drug given its liquid crystal like nature and ability to deform easily when manipulated. In order to overcome this challenge, in some embodiments, the present disclosure describes attempts to synthesize a salt of α -TEA using a variety of different bases which exhibit improved properties than α -TEA. During the development of the present disclosure, 16 potential bases were examined. Of the bases, 6 formed crystalline salts which showed improved characteristics over the free acid of α -TEA. The three α -TEA salts, the salts of lysine, arginine, and tris, were isolated and studied in detail. These three salts exhibited improved physical properties over the free acid without any loss of activity.

Salts of α -TEA can be prepared according to the methods described in the Examples section below. These methods can be further modified and optimized using the principles and techniques of organic chemistry as applied by a person skilled in the art. Such principles and techniques are taught, for example, in *March's Advanced Organic Chemistry: Reactions, Mechanisms, and Structure* (2007), which is incorporated by reference herein.

The salts of α -TEA may contain one or more asymmetrically-substituted carbon or nitrogen atoms, and may be isolated in optically active or racemic form. Thus, all chiral, diastereomeric, racemic form, epimeric form, and all geometric isomeric forms of a structure are intended, unless the specific stereochemistry or isomeric form is specifically indicated. The salts of α -TEA may occur as racemates and racemic mixtures, single enantiomers, diastereomeric mixtures and individual diastereomers. In some embodiments, a single diastereomer is obtained. The chiral centers of the present invention can have the *S* or the *R* configuration.

In addition, atoms making up the salts of α -TEA of the present invention are intended to include all isotopic forms of such atoms. Isotopes, as used herein, include those atoms having the same atomic number but different mass numbers. By way of general example and without limitation, isotopes of hydrogen include tritium and deuterium, and isotopes of carbon include ^{13}C and ^{14}C .

5 Similarly, it is contemplated that one or more carbon atom(s) of a compound of the present invention may be replaced by a silicon atom(s). Furthermore, it is contemplated that one or more oxygen atom(s) of the salts of α -TEA may be replaced by a sulfur or selenium atom(s).

The salts of α -TEA and polymorphic form thereof may also have the advantage that they may be more efficacious than, be less toxic than, be longer acting than, be more potent than, produce fewer 10 side effects than, be more easily absorbed than, and/or have a better pharmacokinetic profile (e.g., higher oral bioavailability and/or lower clearance) than, and/or have other useful pharmacological, physical, or chemical advantages over, compounds known in the prior art for use in the indications stated herein.

III. Hyperproliferative Diseases

15 Hyperproliferative diseases generally include any disease which causes a cell to begin to reproduce uncontrollably such as, e.g., cancer. In cancer, the cell's normal apoptotic cycle is interrupted and thus agents that can induce apoptosis in the cell can be important therapeutic agents for treating these diseases. α -TEA compounds have been shown to lead to cellular apoptosis and as such can potentially be used to treat a variety of types of cancer lines. As such, the salts of α -TEA 20 and the polymorphic forms thereof may be used as effective treatments for cancers such as an endometrial tumor, mammary cancer, lung cancer, ovarian cancer, prostate cancer, breast cancer, cervical cancer, liver cancer, colon cancer, lymphoma, or melanoma. In various aspects, it is anticipated that salts and polymorphic forms of the present invention may be used to treat virtually any malignancy.

25 The compounds and polymorphic forms may be used to treat cancer cells including but are not limited to cells from the bladder, blood, bone, bone marrow, brain, breast, colon, esophagus, gastrointestinal, gum, head, kidney, liver, lung, nasopharynx, neck, ovary, prostate, skin, stomach, pancreas, testis, tongue, cervix, or uterus. In addition, the cancer may specifically be used with the following histological type, though it is not limited to these: neoplasm, malignant; carcinoma; 30 carcinoma, undifferentiated; giant and spindle cell carcinoma; small cell carcinoma; papillary carcinoma; squamous cell carcinoma; lymphoepithelial carcinoma; basal cell carcinoma; pilomatrix carcinoma; transitional cell carcinoma; papillary transitional cell carcinoma; adenocarcinoma; gastrinoma, malignant; cholangiocarcinoma; hepatocellular carcinoma; combined hepatocellular carcinoma and cholangiocarcinoma; trabecular adenocarcinoma; adenoid cystic carcinoma; 35 adenocarcinoma in adenomatous polyp; adenocarcinoma, familial polyposis coli; solid carcinoma; carcinoid tumor, malignant; bronchiolo-alveolar adenocarcinoma; papillary adenocarcinoma;

chromophobe carcinoma; acidophil carcinoma; oxyphilic adenocarcinoma; basophil carcinoma; clear cell adenocarcinoma; granular cell carcinoma; follicular adenocarcinoma; papillary and follicular adenocarcinoma; nonencapsulating sclerosing carcinoma; adrenal cortical carcinoma; endometroid carcinoma; skin appendage carcinoma; apocrine adenocarcinoma; sebaceous adenocarcinoma; 5 ceruminous adenocarcinoma; mucoepidermoid carcinoma; cystadenocarcinoma; papillary cystadenocarcinoma; papillary serous cystadenocarcinoma; mucinous cystadenocarcinoma; mucinous adenocarcinoma; signet ring cell carcinoma; infiltrating duct carcinoma; medullary carcinoma; lobular carcinoma; inflammatory carcinoma; paget's disease, mammary; acinar cell carcinoma; adenosquamous carcinoma; adenocarcinoma w/squamous metaplasia; thymoma, malignant; ovarian 10 stromal tumor, malignant; thecoma, malignant; granulosa cell tumor, malignant; androblastoma, malignant; sertoli cell carcinoma; leydig cell tumor, malignant; lipid cell tumor, malignant; paraganglioma, malignant; extra-mammary paraganglioma, malignant; pheochromocytoma; glomangiosarcoma; malignant melanoma; amelanotic melanoma; superficial spreading melanoma; malig melanoma in giant pigmented nevus; epithelioid cell melanoma; blue nevus, malignant; 15 sarcoma; fibrosarcoma; fibrous histiocytoma, malignant; myxosarcoma; liposarcoma; leiomyosarcoma; rhabdomyosarcoma; embryonal rhabdomyosarcoma; alveolar rhabdomyosarcoma; stromal sarcoma; mixed tumor, malignant; mullerian mixed tumor; nephroblastoma; hepatoblastoma; carcinosarcoma; mesenchymoma, malignant; brenner tumor, malignant; phyllodes tumor, malignant; synovial sarcoma; mesothelioma, malignant; dysgerminoma; embryonal carcinoma; teratoma, 20 malignant; struma ovarii, malignant; choriocarcinoma; mesonephroma, malignant; hemangiosarcoma; hemangioendothelioma, malignant; kaposi's sarcoma; hemangiopericytoma, malignant; lymphangiosarcoma; osteosarcoma; juxtacortical osteosarcoma; chondrosarcoma; chondroblastoma, malignant; mesenchymal chondrosarcoma; giant cell tumor of bone; ewing's sarcoma; odontogenic tumor, malignant; ameloblastic odontosarcoma; ameloblastoma, malignant; ameloblastic 25 fibrosarcoma; pinealoma, malignant; chordoma; glioma, malignant; ependymoma; astrocytoma; protoplasmic astrocytoma; fibrillary astrocytoma; astroblastoma; glioblastoma; oligodendrogioma; oligodendroblastoma; primitive neuroectodermal; cerebellar sarcoma; ganglioneuroblastoma; neuroblastoma; retinoblastoma; olfactory neurogenic tumor; meningioma, malignant; neurofibrosarcoma; neurilemmoma, malignant; granular cell tumor, malignant; malignant lymphoma; 30 hodgkin's disease; hodgkin's; paragranuloma; malignant lymphoma, small lymphocytic; malignant lymphoma, large cell, diffuse; malignant lymphoma, follicular; mycosis fungoides; other specified non-hodgkin's lymphomas; malignant histiocytosis; multiple myeloma; mast cell sarcoma; immunoproliferative small intestinal disease; leukemia; lymphoid leukemia; plasma cell leukemia; erythroleukemia; lymphosarcoma cell leukemia; myeloid leukemia; basophilic leukemia; eosinophilic 35 leukemia; monocytic leukemia; mast cell leukemia; megakaryoblastic leukemia; myeloid sarcoma; and hairy cell leukemia. In certain aspects, the tumor comprises an osteosarcoma, angiosarcoma, rhabdosarcoma, leiomyosarcoma, Ewing sarcoma, glioblastoma, neuroblastoma, or leukemia.

IV. Pharmaceutical Formulations and Routes of Administration

For administration to a mammal in need of such treatment, the salts of α -TEA in a therapeutically effective amount may be used in combination with one or more excipients appropriate to the indicated route of administration. The salts of α -TEA may be used when admixed with lactose,

5 sucrose, starch powder, cellulose esters of alkanoic acids, cellulose alkyl esters, talc, stearic acid, magnesium stearate, magnesium oxide, sodium and calcium salts of phosphoric and sulfuric acids, gelatin, acacia, sodium alginate, polyvinylpyrrolidone, and/or polyvinyl alcohol, and tableted or encapsulated for convenient administration. Alternatively, the salts of α -TEA may be dissolved in water, polyethylene glycol, propylene glycol, ethanol, corn oil, cottonseed oil, peanut oil, sesame oil,

10 benzyl alcohol, sodium chloride, and/or various buffers. Other excipients and modes of administration are well and widely known in the pharmaceutical art.

In some embodiments, the pharmaceutical compositions useful in the present disclosure can be subjected to conventional pharmaceutical operations such as sterilization and/or contain conventional pharmaceutical carriers and excipients such as preservatives, stabilizers, wetting agents, 15 emulsifiers, buffers, *etc.*

In some embodiments, the salts of α -TEA are administered by a variety of methods, *e.g.*, orally or by injection (*e.g.* subcutaneous, intravenous, intraperitoneal, *etc.*). Depending on the route of administration, the active salts of α -TEA may be coated in a material to protect the compound from the action of acids and other natural conditions which may inactivate the compound. They may also 20 be administered by continuous perfusion/infusion of a disease or wound site.

To administer the therapeutic compound by other than parenteral administration, in some embodiments, it can be necessary to coat the salts of α -TEA with, or co-administer the salts of α -TEA with, a material to prevent its inactivation. For example, the therapeutic compound may be administered to a patient in an appropriate carrier, for example, liposomes, or a diluent. 25 Pharmaceutically acceptable diluents include saline and aqueous buffer solutions. Liposomes include water-in-oil-in-water CGF emulsions as well as conventional liposomes.

In some embodiments, the therapeutic salts of α -TEA are also administered parenterally, intraperitoneally, intraspinally, or intracerebrally. Dispersions can be prepared in glycerol, liquid 30 polyethylene glycols, and mixtures thereof and in oils. Under ordinary conditions of storage and use, these preparations can contain, in some embodiments, a preservative to prevent the growth of microorganisms.

Pharmaceutical compositions suitable for injectable use include sterile aqueous solutions (where water soluble) or dispersions and sterile powders for the extemporaneous preparation of sterile injectable solutions or dispersion are also envisioned. In all cases, the composition must be sterile and 35 must be fluid to the extent that easy syringability exists. It must be stable under the conditions of manufacture and storage and must be preserved against the contaminating action of microorganisms

such as bacteria and fungi. The carrier can be a solvent or dispersion medium containing, for example, water, ethanol, polyol (such as, glycerol, propylene glycol, and liquid polyethylene glycol, and the like), suitable mixtures thereof, and vegetable oils. The proper fluidity can be maintained, for example, by the use of a coating such as lecithin, by the maintenance of the required particle size in 5 the case of dispersion and by the use of surfactants. Prevention of the action of microorganisms can be achieved by various antibacterial and antifungal agents, for example, parabens, chlorobutanol, phenol, ascorbic acid, thimerosal, and the like. In many cases, it will be preferable to include isotonic agents, for example, sugars, sodium chloride, or polyalcohols such as mannitol and sorbitol, in the 10 composition. Prolonged absorption of the injectable compositions can be brought about by including in the composition an agent which delays absorption, for example, aluminum monostearate or gelatin.

Sterile injectable solutions can be prepared by incorporating the therapeutic salts of α -TEA in the required amount in an appropriate solvent with one or a combination of ingredients enumerated above, as required, followed by filtered sterilization. Generally, dispersions are prepared by incorporating the therapeutic compound into a sterile carrier which contains a basic dispersion 15 medium and the required other ingredients from those enumerated above. In the case of sterile powders for the preparation of sterile injectable solutions, the preferred methods of preparation are vacuum drying and freeze-drying which yields a powder of the active ingredient (*i.e.*, the therapeutic compound) plus any additional desired ingredient from a previously sterile-filtered solution thereof.

The therapeutic salts of α -TEA can be orally administered, for example, with an inert diluent 20 or an assimilable edible carrier. The therapeutic compound and other ingredients may also be enclosed in a hard or soft shell gelatin capsule, compressed into tablets, or incorporated directly into the subject's diet. For oral therapeutic administration, the therapeutic salts of α -TEA may be used with excipients and used in the form of ingestible tablets, buccal tablets, troches, capsules, elixirs, suspensions, syrups, wafers, and the like. The percentage of the therapeutic compound in the 25 compositions and preparations may, of course, be varied. The amount of the therapeutic salts of α -TEA in such therapeutically useful compositions is such that a suitable dosage will be obtained.

It is especially advantageous to formulate parenteral compositions in dosage unit form for ease of administration and uniformity of dosage. Dosage unit form as used herein refers to physically 30 discrete units suited as unitary dosages for the subjects to be treated; each unit containing a predetermined quantity of therapeutic salts of α -TEA calculated to produce the desired therapeutic effect in association with the required pharmaceutical carrier. The specification for the dosage unit forms of the invention are dictated by and directly dependent on (a) the unique characteristics of the therapeutic salts of α -TEA and the particular therapeutic effect to be achieved, and (b) the limitations inherent in the art of compounding such a therapeutic compound for the treatment of a selected 35 condition in a patient.

The therapeutic compound may also be administered topically to the skin, eye, or mucosa. Alternatively, if local delivery to the lungs is desired the therapeutic compound may be administered by inhalation in a dry-powder or aerosol formulation.

Active salts of α -TEA are administered at a therapeutically effective dosage sufficient to treat a condition associated with a condition in a patient. For example, the efficacy of the salts of α -TEA can be evaluated in an animal model system that may be predictive of efficacy in treating the disease in humans, such as the model systems shown in the examples and drawings.

The actual dosage amount of the salts of α -TEA of the present disclosure or composition comprising the salts of α -TEA of the present disclosure administered to a subject may be determined by physical and physiological factors such as age, sex, body weight, severity of condition, the type of disease being treated, previous or concurrent therapeutic interventions, idiopathy of the subject and on the route of administration. These factors may be determined by a skilled artisan. The practitioner responsible for administration will typically determine the concentration of active ingredient(s) in a composition and appropriate dose(s) for the individual subject. The dosage may be adjusted by the individual physician in the event of any complication.

An effective amount typically will vary from about 2 mg/kg to about 50 mg/kg, , in one or more dose administrations daily, for one or several days (depending of course of the mode of administration and the factors discussed above). In some particular embodiments, the amount is less than 5,000 mg per day with a range of 100 mg to 4500 mg per day.

The effective amount may be less than 10 mg/kg/day, less than 100 mg/kg/day, less than 250 mg/kg/day, less than 100 mg/kg/day, less than 50 mg/kg/day, less than 25 mg/kg/day or less than 10 mg/kg/day. It may alternatively be in the range of 1 mg/kg/day to 200 mg/kg/day.

In other non-limiting examples, a dose also comprises from about 10 mg/kg/body weight, about 100 mg/kg/body weight, about 10 g/kg/body weight, about 5 g/kg/body weight, or more per administration, and any range derivable therein. In non-limiting examples of a derivable range from the numbers listed herein, a range of about 1 mg/kg/body weight to about 100 mg/kg/body weight, about 5 g/kg/body weight to about 10 g/kg/body weight, *etc.*, can be administered, based on the numbers described above.

In certain embodiments, a pharmaceutical composition of the present disclosure comprises, for example, at least about 0.1% of a salt of α -TEA described in the present disclosure. In other embodiments, the compound of the present disclosure comprises between about 0.25% to about 75% of the weight of the unit, or between about 25% to about 60%, or between about 1% to about 10%, for example, and any range derivable therein.

Single or multiple doses of the agents are contemplated. Desired time intervals for delivery of multiple doses can be determined by one of ordinary skill in the art employing no more than routine experimentation. As an example, subjects may be administered two doses daily at approximately 12 hour intervals. In some embodiments, the agent is administered once a day.

The salts of α -TEA may be administered on a routine schedule. As used herein a routine schedule refers to a predetermined designated period of time. The routine schedule may encompass periods of time which are identical or which differ in length, as long as the schedule is predetermined. For instance, the routine schedule may involve administration twice a day, every day, every two days, 5 every three days, every four days, every five days, every six days, a weekly basis, a monthly basis or any set number of days or weeks there-between. Alternatively, the predetermined routine schedule may involve administration on a twice daily basis for the first week, followed by a daily basis for several months, *etc.* In other embodiments, the disclosure provides that the agent(s) can be taken orally and that the timing of which is or is not dependent upon food intake. Thus, for example, the 10 agent can be taken every morning and/or every evening, regardless of when the subject has eaten or will eat. In other embodiments, the disclosure is taken as a dietary supplement. In some embodiments, the α -TEA salts are taken before the onset of the tumor as a prophylaxis measure. In other embodiments, the α -TEA salts are taken as a treatment option for use as an antiproliferative agent.

15 V. Combination Therapy

In addition to being used as a monotherapy, the salts of α -TEA described in the present invention may also find use in combination therapies. Effective combination therapy may be achieved with a single composition or pharmacological formulation that includes both agents, or with two distinct compositions or formulations, administered at the same time, wherein one composition 20 includes a salt of α -TEA, and the other includes the second agent(s). The other therapeutic modality may be administered before, concurrently with, or following administration of the salts or polymorphic forms of α -TEA. The therapy using the salts or polymorphic forms of α -TEA may precede or follow administration of the other agent(s) by intervals ranging from minutes to weeks. In 25 embodiments where the other agent and the salts or polymorphic forms of α -TEA are administered separately, one would generally ensure that a significant period of time did not expire between the time of each delivery, such that each agent would still be able to exert an advantageously combined effect. In such instances, it is contemplated that one would typically administer the salts or polymorphic forms of α -TEA and the other therapeutic agent within about 12-24 hours of each other and, more preferably, within about 6-12 hours of each other, with a delay time of only about 12 hours 30 being most preferred. In some situations, it may be desirable to extend the time period for treatment significantly, however, where several days (2, 3, 4, 5, 6 or 7) to several weeks (1, 2, 3, 4, 5, 6, 7 or 8) lapse between the respective administrations.

It also is conceivable that more than one administration of a salt or polymorphic form of α -TEA, or the other agent will be desired. In this regard, various combinations may be employed. By 35 way of illustration, where the salts or polymorphic forms of α -TEA is "A" and the other agent is "B", the following permutations based on 3 and 4 total administrations are exemplary:

A/B/A B/A/B B/B/A A/A/B B/A/A A/B/B B/B/B/A B/B/A/B
A/A/B/B A/B/A/B A/B/B/A B/B/A/A B/A/B/A B/A/A/B B/B/B/A
A/A/A/B B/A/A/A A/B/A/A A/A/B/A A/B/B/B B/A/B/B B/B/A/B

Other combinations are likewise contemplated. Non-limiting examples of pharmacological

5 agents that may be used in the present invention include any pharmacological agent known to be of benefit in the treatment of a cancer or hyperproliferative disorder or disease. In some embodiments, combinations of the salts or polymorphic forms of α -TEA with a cancer targeting immunotherapy, radiotherapy, chemotherapy, or surgery are contemplated. Also contemplated is a combination of a salt or polymorphic form of α -TEA with more than one of the above mentioned methods including
10 more than one type of a specific therapy. In some embodiments, it is contemplated that the immunotherapy is a monoclonal antibody which targets HER2/neu such as trastuzumab (Herceptin[®]) or a similar antibody. In other embodiments, the immunotherapy can be other cancer targeting antibodies such as alemtuzumab (Campath[®]), bevacizumab (Avastin[®]), cetuximab (Eribitux[®]), and panitumumab (Vectibix[®]) or conjugated antibodies such as ibritumomab tiuxetan (Zevalin[®]),
15 tositumomab (Bexxar[®]), brentuximab vedotin (Adcetris[®]), ado-trastuzumab emtansine (KadcylaTM), or denileukin ditox (Ontak[®]) as well as immune cell targeting antibodies such as ipilimumab (Yervoy[®]), tremelimumab, anti-PD-1, anti-4-1-BB, anti-GITR, anti-TIM3, anti-LAG-3, anti-TIGIT, anti-CTLA-4, or anti-LIGHT. Furthermore, in some embodiments, the salts or polymorphic forms of
20 α -TEA are envisioned to be used in combination therapies with dendritic cell-based immunotherapies such as Sipuleucel-T (Provenge[®]) or adoptive T-cell immunotherapies.

Furthermore, it is contemplated that the salts or polymorphic forms of α -TEA are used in combination with a chemotherapeutic agent such as anthracyclines, taxanes, methotrexate, mitoxantrone, estramustine, doxorubicin, etoposide, vinblastine, carboplatin, vinorelbine, 5-fluorouracil, cisplatin, topotecan, ifosfamide, cyclophosphamide, epirubicin, gemcitabine, vinorelbine,
25 irinotecan, etoposide, vinblastine, pemetrexed, melphalan, capecitabine, oxaliplatin, BRAF inhibitors, and TGF-beta inhibitors. In some embodiments, the combination therapy is designed to target a cancer such as those listed above. In the preferred embodiments, the cancer the combination therapy is designed to treat is an endometrial tumor, mammary cancer, lung cancer, ovarian cancer, prostate cancer, breast cancer, cervical cancer or colon cancer.

30 **VI. Examples**

The following examples are included to demonstrate preferred embodiments of the invention. It should be appreciated by those of skill in the art that the techniques disclosed in the examples which follow represent techniques discovered by the inventor to function well in the practice of the invention, and thus can be considered to constitute preferred modes for its practice. However, those

of skill in the art should, in light of the present disclosure, appreciate that many changes can be made in the specific embodiments which are disclosed and still obtain a like or similar result without departing from the spirit and scope of the invention.

A. Methods

5 1. MICROSCOPY

A Zeiss Universal microscope configured with a polarized visible light source and polarizable analyzer was used to evaluate the optical properties of the samples. Specimens were typically mounted on a microscope slide with a cover glass. Because of the "waxy" nature of the material slide preparations were manually smeared with a spatula and appearance noted. Observations of 10 particle/crystal size and shape and birefringence were recorded.

2. HOT STAGE MICROSCOPY (HSM)

A Linkam hot stage accessory was used in tandem with the microscope. Specimens were mounted on a microscope slide with a cover glass. Samples were heated from room temperature through melting using a Linkam TMS 94 temperature control and Linksys 32 data capture software system. Observations of possible phase change, melting, recrystallization, decomposition, etc. were recorded.

3. PROTON NUCLEAR MAGNETIC RESONANCE (^1H NMR)

Samples were analyzed by ^1H NMR to determine stoichiometry (molar ratio of acid to base counter ion). Spectra of salt candidates were compared to that of the free acid and changes in 20 chemical shifts indicating salt formation observed. Samples were prepared by dissolving in a chloroform: methanol: water mixture with 0.05% (v/v) tetramethylsilane (TMS). Spectra were collected at ambient temperature on a Bruker Avance III 400 MHz FT-NMR spectrometer and Bruker Topspin software (version 2.1). Prior to each sample analysis, the magnetic field surrounding the sample was optimized by an automated shimming program.

25 4. DIFFERENTIAL SCANNING CALORIMETRY (DSC)

Differential scanning calorimetry (DSC) is a technique used to measure characteristic heat flux of a test article as it is scanned through a temperature gradient under a controlled atmosphere. Thermal phase transitions such as endothermic melting and exothermic decomposition were recorded. DSC data were collected on a TA Instruments DSC. In general, samples in the mass range of 1 to 1 0 30 mg were crimped in aluminum sample pans and scanned from 25 to approximately 300 °C at 10 °C/min using a nitrogen purge of 50 mL/min.

5. THERMOGRAVIMETRIC ANALYSIS (TGA)

Thermogravimetric analysis involves the determination of the mass of a specimen as a function of temperature. TGA data were collected on a TA Instruments Q500 TGA. In general, samples in the mass range of 2 to 10 mg were placed in an open, pre-tared platinum sample pan and attached by fine wire to a microbalance. The sample was suspended into a furnace which was heated from 25 to about 250 °C at 10 °C/min using a nitrogen purge at 100 mL/min. Sample weight change as a function of temperature was observed.

6. X-RAY POWDER DIFFRACTION (XRD)

X-ray powder diffraction patterns were obtained using a Bruker D8 Discovery diffractometer equipped with an XYZ stage, laser video microscope for positioning, and a two dimensional HiStar area Detector or a scintillation detector. A CuK α radiation 1.5406 angstrom source operating at 40 kV and 40 mA was used to irradiate samples. The X-ray optics consists of a Gobel mirror coupled with a pinhole collimator of 0.5 or 1.0 mm. Theta-theta continuous scans were employed with a sample-detector distance of approximately 30 cm, which gives an effective 20 range of 4-40 °C. Samples were mounted in low background quartz plates.

7. SOLUBILITY

Milligram size (2 mg) quantities of each sample were placed into a vial. Buffered water (pH 4, 7, and 10) was added and the vials were stirred for a few minutes, followed by visual observation for remaining solids. The solvent was incrementally added until the solids were dissolved, or a maximum volume of solvent (10 mL) was added and the experiment was terminated.

8. HYGROSCOPICITY- DYNAMIC VAPOR SORPTION (DVS)

DVS is a gravimetric screening technique that measures how quickly and how much of a solvent (water) is adsorbed by a sample. The relative humidity or vapor concentration surrounding the sample is varied while the change in mass of the sample is measured. A vapor sorption isotherm shows the equilibrium amount of vapor sorbed as a function of relativity humidity. The mass values at each relative humidity step are used to generate the isotherm. Isotherms are divided in two components: sorption for increasing humidity steps and desorption for decreasing humidity steps. A plot of kinetic data is also supplied which shows the change in mass and humidity as a function of time.

Samples were analyzed using a TA Q2000 automated dynamic vapor sorption analyzer. The samples were dried at 40 °C for 5 hours and then cooled to 25 °C with a dry nitrogen purge over them

until they no longer lost mass at 0% RH. The samples were then subjected to 0 to 95% RH, back to 0% RH at 25 °C in 5% RH steps.

9. STABILITY

The scaled up salts and free acid were challenged by heat (solids stored at 25 and 60 °C for 1 week), oxidation (solids stored in oxygen headspace at 25 °C for 1 week), light (solids exposed $\geq 1 \times$ ICH International Conference on Harmonization of Technical Requirements for Registration of Pharmaceuticals for Human Use UV confirmatory conditions), and solutions (HPLC diluent-methanol) at 25 and 40 °C for 1 week.

These samples were analyzed, along with unstressed controls, by HPLC to characterize their stability.

10. HPLC ANALYSIS

Salt candidates were analyzed by total area normalization (TAN). The samples were dissolved in methanol at a concentration of 0.8 mg/mL.

HPLC Conditions

HPLC Column: Sunfire C18, 3.5 μ m, 4.6 x 150 mm
Column Temp: 45 °C
Autosampler Flush: 25:75 Acetonitrile: reagent alcohol
Flow Rate: 1.5 mL/min
Injection Volume: 10 μ L
UV Detection: 215 nm
Mobile Phase: A – 0.1% acetic acid in acetonitrile
B – 0.1% acetic acid in reagent alcohol

Gradient Pump Program:

Step Time (minutes)	% A	% B	Curve
0.5	92.0	8.0	0.0
6.0	92.0	8.0	0.0
6.0	10.0	90.0	1.0
2.0	10.0	90.0	0.0
7.0	92.0	8.0	0.0

B. Salt Screen

1. CHARACTERIZATION OF THE FREE ACID

The α -TEA free acid, lot 54543-7-34, was analyzed by XRD, DSC, TGA, DVS and microscopy. The sample is crystalline as shown by the XRD pattern in FIG. 1. The DSC thermogram of the free acid shows a melting endotherm onset at 54.5 °C. The TGA thermogram shows the free acid to be free of volatiles, with a weight loss of 0.2% at 207.5 °C.

DVS was performed on a sample of the free acid. The free acid exhibits a total weight gain of about 0.2 weight% at 95% RH. This data indicates that the free acid does not form hydrates or exhibit deliquescence.

The free acid was examined using optical microscopy. The sample exhibited areas of both birefringent and amorphous agglomerates. Manual manipulation of the sample under magnification resulted in a smearing and a droplet like appearance. This behavior suggests that the free acid has some of the characteristics of a liquid crystal. Liquid crystals are a state of matter that have properties between those of a conventional liquid and those of a solid. Liquid crystals are substances that are not as rigidly ordered as a solid, but have some degree of alignment. Molecules which tend to be candidates for having liquid crystal phases are often long having a rigid central region and end groups that are slightly flexible.

2. PRIMARY SALT SCREEN (50 MG SCALE) PREPARATIVE PROCEDURES

The predicted pK_a value of the α -TEA free acid is 3.23. Based on these predicated pK_a values, a library of salt forming bases were chosen and is shown in Table 1.

15 *Table 1: Salt Forming Bases*

Base	pK _a	Class*
sodium hydroxide	14	I
potassium hydroxide	14	I
magnesium hydroxide	11.4	I
meglumine	8	I
diethylamine	10.9	II
ethanolamine	9.5	III
diethanolamine	9.3	III
L-Arginine	13.2	I
L-Lysine	10.8	I
Choline	>11	I
tris-hydroxymethylamine (Tris)	8	II
deanol, N,N-dimethylethanolamine	8.8	II
2-diethylaminoethanol	9.6	II
calcium hydroxide	12.6	I
Benzathine, dibenzylethylenediamine	9.99	II
procaine	8.9	II

*Safety and pharmaceutical acceptability of the counter-ions: class I-unrestricted used based on physiological ubiquitous ions or intermediate metabolites; class II-universally approved, but not naturally occurring. Low toxicity and good tolerability; class III-case by case limited approval, some reactivity, some pharmacological activity.

20 Initially, a series of experiments consisted of 7 salts prepared on an approximately 50 mg scale were carried out and are panel 1. A solution of free acid was prepared in methanol and appropriate molar equivalents of base solution and/or solids were added. Samples were mixed several hours and allowed to slowly evaporate at room temperature with about 2 psi nitrogen purge.

Characterization of the resultant salts was performed using X-ray diffraction (XRD). One sample (ethanolamine) exhibited a crystalline pattern different than the free acid starting material indicating possible salt formation. The remaining six samples produced amorphous materials. These six samples were solvent ripened in an attempt to increase the crystallinity of the resultant salts.

5 Unfortunately, this technique had limited success in increasing the crystallinity.

The crystalline salt candidate, ethanolamine, was further characterized by differential scanning calorimetry (DSC). An overview of the results is summarized in Table 2.

Table 2: Panel 1

Base	Class	Sample ID	Appearance	Crystallinity/XRD	DSC
sodium hydroxide	I	116	waxy solid	amorphous	N/A
potassium hydroxide	I	117	glassy ¹	amorphous	N/A
magnesium hydroxide	I	118	glassy ¹	amorphous	N/A
meeglumine	I	119	waxy solid	amorphous	N/A
diethylamine	II	120	glassy ¹	amorphous	N/A
ethanolamine	III	121	waxy solid	crystalline possible salt	Multiple endotherms
diethanolamine	III	122	glassy ¹	amorphous	N/A

¹ glassy materials are amorphous (not crystalline) non free flowing solids which often do not exhibit a sharp melting endotherm during DSC analysis.

The next group of experiments, panel 2, consists of 8 salts prepared on an approximately 50 mg scale. A solution of free acid was prepared in 1-butanol and appropriate molar equivalents of base solution and or solids were added. Samples were mixed several hours and allowed to slowly evaporate at room temperature with about 2 psi nitrogen purge.

15 Characterization was performed using X-ray Diffraction (XRD). One sample (ethanolamine) exhibited a crystalline pattern similar to the pattern from Panel 1, indicating possible salt formation. Two samples exhibited evidence of the base counterion, which would indicate that no salt was formed. The remaining five samples produced amorphous materials. These five samples were solvent ripened in an attempt to increase the crystallinity. Again, this technique was not successful in 20 increasing the crystallinity.

The results of Panel 2 are listed in Table 3.

Table 3: Panel 2

Base	Class	Sample ID	Appearance	Crystallinity/XRD
sodium hydroxide	I	412	glassy	amorphous
potassium hydroxide	I	413	glassy	amorphous
magnesium hydroxide	I	414	waxy solid	no salt, evidence of MgOH
meeglumine	I	415	waxy solid	amorphous
diethylamine	II	416	glassy	amorphous

Base	Class	Sample ID	Appearance	Crystallinity/XRD
ethanolamine	III	417	waxy solid	crystalline, possible salt
diethanolamine	III	418	glassy	amorphous
calcium hydroxide	I	419	waxy solid	Amorphous, no salt, evidence of Ca(OH) ₂

Finally, another set of experiments, Panel 3, consisting of 16 salts was prepared on approximately 50 mg scale. A solution of free acid was prepared in isopropyl alcohol and appropriate molar equivalents of base solution and or solids were added. Samples were mixed several hours and allowed to slowly evaporate at room temperature with about 2 psi nitrogen purge.

5 Characterization was performed using XRD. Six samples (ethanolamine, arginine, lysine, choline, tris, and deanol) exhibited crystalline XRD patterns indicating possible salt formation. The eight remaining samples produced amorphous materials. These eight samples were solvent ripened in an attempt to increase the crystallinity. This technique was not successful in increasing the crystallinity. Two samples exhibited evidence of the base counter-ion, indicating that no salt
10 formation.

The Panel 3 results are listed in Table 4.

Table 4: Panel 3

Base	Class	Sample ID	Appearance	Crystallinity/XRD	DSC
sodium hydroxide	I	510	waxy solid	amorphous	N/A
potassium hydroxide	I	511	glassy	amorphous	N/A
magnesium hydroxide	I	512	waxy solid	no salt, evidence of MgOH	N/A
meglumine	I	513	waxy solid	amorphous	N/A
diethylamine	II	514	glassy	amorphous	N/A
ethanolamine	III	515	waxy solid	crystalline, possible salt	Multiple endotherms
diethanolamine	III	516	glassy	amorphous	N/A
L-arginine	I	517	solid	crystalline, possible salt	Multiple endotherms
L-lysine	I	518	solid	crystalline, possible salt	Multiple endotherms
choline	I	519	waxy solid	low crystallinity, possible salt	Multiple endotherms
Tris-hydroxylethanolamine	II	520	waxy solid	low crystallinity, possible salt	Multiple endotherms
Deanol, N,N-dimethylethanolamine	II	521	waxy solid	crystalline, possible salt	Multiple endotherms
2-diethylaminoethanol	I	522	glassy	amorphous	N/A
calcium hydroxide	I	523	white solid	no salt, evidence of Ca(OH) ₂	N/A
Benzathine, dibenzylethylenediamine	II	524	glassy	amorphous	N/A
procaine	II	525	glassy	amorphous	N/A

3. PRIMARY SALT SCREEN (50 MG SCALE) RESULTS AND DISCUSSION

Panels 1 through 3 produced six crystalline salt compounds. The characteristics of these compounds are further described below.

a) Ethanolamine

5 The ethanolamine salt of α -TEA is a waxy crystalline solid. The unique X-ray diffraction characteristics were observed in all three panels and are shown in **FIG. 2**. The DSC thermogram exhibited multiple endothermic transitions with temperatures of about 88, 100, 115, 149 and 169 °C. The DSC plot is shown in **FIG. 3**. The total volatile content by TGA over the temperature range 25-95.5 °C was 0.4 wt%.

10 The ^1H NMR of this compound shows the stoichiometry of free acid to base to be in a 1:1 ratio. A change in chemical shifts from the free acid spectrum was observed suggesting the compound is a salt.

b) L-Arginine

15 The arginine salt of α -TEA was a partially crystalline solid that was less "waxy" than the free acid. The unique X-ray diffraction pattern of the L-arginine sample is shown in **FIG. 4**. The DSC thermogram exhibited multiple endothermic transitions at about 131 and 184 °C. The DSC plot is shown in **FIG. 5**. The total volatile content by TGA over the temperature range 25-194 °C was about 0.3 wt%.

20 The ^1H NMR of this candidate shows the stoichiometry of free acid to base to be in a 1:1 ratio. A change in chemical shifts from the free acid spectrum was observed suggesting this candidate is a salt.

25 The dynamic vapor sorption (DVS) isotherm indicates a possible hemihydrate formation (1.3% weight gain) at about 40% RH. The desorption cycle indicates that the hydration process is reversible. The particle size of this material was very small as evidenced by photomicrograph. The salt produced some liquid crystal-like behavior, but to a lesser degree than observed with the free acid. The approximate visual solubility of the arginine salt in pH 4, 7, and 10 buffers was < 0.2 mg/mL.

c) L-Lysine

30 The lysine salt of α -TEA is a crystalline solid that is less "waxy" than the free acid. The unique X-ray diffraction pattern of this sample is shown in **FIG. 6**. The DSC thermogram exhibited

multiple endothermic transitions at about 55 and 200 °C. The DSC plot is shown in **FIG. 7**. The total volatile content by TGA over the temperature range 25-168.8 °C was 0.4 wt%.

The ^1H NMR of the lysine salt shows the stoichiometry of free acid to base to be in a 1:0.7 ratio. A change in chemical shifts from the free acid spectrum was observed suggesting the compound 5 is a salt.

The DVS adsorption isotherm shows onset of weight gain at about 75% RH, with a rapid total weight gain of about 12% at 95% RH. The desorption cycle shows hysteresis, in which upon desorption (decreasing humidity) the sample does not lose the sorbed water. The residual 2 weight % corresponds to a putative hydrate of the lysine salt. The stoichiometry of the hydrated lysine salt is 10 not defined. The kinetic plot indicates the weight change on desorption does not come to equilibrium indicating that hydrate formation may be reversible given enough time. The particle size of this material appears was very small as evidenced by photomicrograph. The salt exhibits some liquid crystal-like behavior, but to a lesser degree than observed with the free acid. The approximate visual solubility of the lysine salt in pH 4, 7, and 10 buffers was < 0.2 mg/mL.

15 **d) Choline**

The choline salt of α -TEA is a partially crystalline "waxy" solid. The unique X-ray diffraction pattern of the sample was observed is shown in **FIG. 8**. The DSC thermogram exhibited multiple endothermic transitions at about 50, 80, and 150 °C. The DSC plot is shown in **FIG. 9**. The total volatile content by TGA over the temperature range 25-188.2 °C was 2.5 wt%.

20 The ^1H NMR of this candidate shows the stoichiometry of free acid to base to be in a 1:0.8 ratio. A change in chemical shifts from the free acid spectrum was observed suggesting this candidate is a salt.

e) Tris-hydroxymethyl amine (Tris)

The tris candidate was a partially crystalline "semi-waxy" solid. The unique X-ray diffraction pattern of the sample was observed is shown in **FIG. 10**. The DSC thermogram exhibited multiple endothermic transitions at about 101 and 175 °C. The DSC plot is shown in **FIG. 11**. The total volatile content by TGA over the temperature range 25-144 °C was 0.6 wt%.

25 The ^1H NMR of this candidate shows the stoichiometry of free acid to base to be in a 1:1+ ratio. A change in chemical shifts from the free acid spectrum was observed suggesting this candidate 30 is a salt.

The DVS isotherm indicates a possible hemihydrate formation (1.5% weight gain) starting at about 55% RH. The desorption cycle shows hysteresis indicating that the water is well bound. The particle size of this material was very small based upon photomicrograph images. There is some liquid crystal-like behavior, but to a lesser degree than observed with the free acid. The approximate 5 visual solubility of the tris salt in pH 4, 7, and 10 buffers was < 0.2 mg/mL.

D) *N,N*-dimethylethanolamine (deanol)

The deanol candidate is a crystalline "waxy" solid. The unique X-ray diffraction pattern of the sample was observed is shown in FIG. 12. The DSC thermogram exhibited multiple endothermic transitions at about 47, 73, and 87 °C. The DSC plot is shown in FIG. 13. The total volatile content by 10 TGA over the temperature range 25-207 °C was 7.7 wt%.

The ¹H NMR of this candidate shows the stoichiometry of free acid to base to be in a 1:0.7 ratio. A change in chemical shifts from the free acid spectrum was observed suggesting this candidate is a salt.

4. SALT SCREEN SCALE UP RESULTS AND DISCUSSION

15 Based on results of the primary salt screening panels, the lysine, arginine and tris monosalts were selected for further investigation. This analysis served two purposes; firstly to provide a sufficient quantity of material for further testing, and secondly, to gather information on the processibility and physical handling characteristics of the salt candidates. Samples were analyzed by XRD, DSC, Hot Stage Microscopy (HSM), HPLC for purity, static humidity, stoichiometry, and 20 stability.

a) Lysine (Sample 901)

The scaled-up lysine salt had more desirable handling properties compared to the free acid, including being less "tacky" than the free acid. Furthermore, the salt was a crystalline powder and easier to formulate into a capsule than the free acid.

25 The XRD pattern of the scaled-up lysine salt candidate was consistent with the pattern of the small scale sample. The DSC profile of this sample is very similar to the small scale sample, both exhibiting endothermic transitions at about 55 and 200°C. Hot stage microscopy (HSM) was carried out on this salt candidate in order to aid in explaining the multiple endotherms observed by DSC. Visual onset of melting was observed at about 200 °C with complete melting by about 215 °C. No 30 obvious transition was observed at about 55 °C.

The HPLC total area normalization (TAN) purity of the salt was determined to be 98 area% given a relative purity of the compound.

Formation of hydrates can be a very slow kinetic process. A second approach to identify hydrate formation is performing static humidity experiments and monitor gravimetrically for weight change. Samples of lysine salt were placed in 75 and 84% RH environments for 33 and 9 days respectively. After 33 days at 75% RH the lysine sample gained 0.6 weight %. At 84% RH, after 9 days, the lysine salt exhibited a weight gain of 1.8%. The X-ray diffraction pattern of this 84% RH sample showed no change from the prehumidified sample. Given the DVS behavior of the small scale sample, these studies suggest an isostructural hemihydrate exists.

The ¹H NMR analysis showed this candidate to have a monomolar ratio of free acid to base. The chemical shift of the methylene group in the polar part of the molecule suggests that the compound appears to be a salt.

b) Arginine (Sample 903)

The scaled up arginine salt exhibited better handling relative to the free acid.

The XRD pattern of the scaled up arginine salt matches the pattern of the small scale sample pattern. The DSC profile of this sample matches small scale exhibiting 2 endothermic transitions at about 136, and 187 °C respectively. Hot stage microscopy (HSM) on this salt candidate showed visual melting at about 180 °C and above. No changes in the sample were visually observed in the 130 to 140 °C temperature range. The endothermic transition observed in the DSC profile in this temperature region may be due to a thermally induced liquid crystal phase change. Some variable temperature XRD studies may be useful in studying this transition further.

The HPLC-Total Area Normalization (TAN purity) of the salt was found to be 98 area% given a relative purity of the compound.

Samples of arginine salt were placed in 75 and 84% RH environments for 33 and 9 days respectively. After 33 days at 75% RH the arginine sample gained 0.7 weight%. At 84% RH, after 9 days, the arginine salt exhibited a weight gain of 2.3%. The X-ray diffraction pattern of this 84% RH sample showed no change from the pre-humidified sample, suggesting that if a hydrate is forming, it has the same XRD pattern.

The ¹H NMR analysis showed this salt to have a mole ratio of free acid to base of one. The chemical shift of the methylene group in the polar part of the molecule suggests that a salt has been formed.

c) Tris (Sample 902)

The scaled up tris salt appears to have more normal handling than the free acid but is slightly waxier than both the lysine and arginine salts.

The XRD pattern of the scaled up tris salt matches small scale sample pattern. The DSC profile is very similar to the small scale sample. The tris scaled up salt exhibits an endotherm with an onset on 106 °C. Visual melting was observed using HSM in the 95 to 105 °C temperature range, which corresponds to the endotherm observed in the DSC.

The HPLC-Total Area Normalization (TAN purity) of the salt was found to be 97.5 area% given a relative purity of the compound.

A sample of tris salt was placed in 75% RH environment for 33 days. The sample exhibited a weight gain of 4.0%. This fact is consistent with the DVS data collected earlier. These data suggest that this salt forms a hydrate upon exposure to sufficient moisture.

The ¹H NMR analysis showed this salt to have a mole ratio of free acid to base of about one. The chemical shift of the methylene group in the polar part of the molecule suggests that the compound formed is likely a salt.

A stacked plot of the NMR spectra of the free acid, lysine, arginine, and tris salts shows the movement of the methylene group supporting the conclusion of the formation of salts with the compound.

d) STABILITY OF SALTS

The three salts (lysine, arginine, and tris) were challenged using heat (solids stored at 25 and 60 °C for 1 week), oxidation (solids stored in oxygen headspace at 25 °C for 1 week), light (UV source under ICH confirmatory conditions > 200 Whr/m²), and solutions (in HPLC diluent) at 25 and 40 °C for 1 week. Stressed samples were analyzed using HPLC to determine their impurity profiles.

A summary of the stability data generated using HPLC analysis is shown below in Table 5.

Table 5: Summary of HPLC Stability Data on Stressed Salts

Conditions	HPLC Total Area Normalization – Area % Purity			
	Free Acid Sample 734	Lysine Sample 901	Arginine Sample 903	Tris Sample 902
Solid State Ambient	97.8	98.8	98.6	98.6
Solid State 60 °C	98.0	98.7	98.6	98.4
Solution Ambient Methanol	96.0	98.8	98.4	98.5
Solution 40 °C Methanol	94.6	98.2	98.2	98.2
Oxidation Ambient	97.8	98.8	98.4	98.5

Conditions	HPLC Total Area Normalization – Area % Purity			
	Free Acid Sample 734	Lysine Sample 901	Arginine Sample 903	Tris Sample 902
Photo Stability Dark Control	97.7	98.7	97.9	98.2
Photo Stability Exposed	98.2	98.8	98.5	98.7

The stability results shown in Table 5 represent the averages of two injections of duplicate sample preparations. While the data can only suggest the short term stability, they do provide some insight into the stability of the salt forms which could be extrapolated to the salts long term stability. FIGS. 14-17 show the chromatogram overlay plots for samples analyzed during the stability portion 5 of the study. The HPLC stability data show that the salts exhibited little to no degradation under the stress conditions used in the study. All three salt candidates show improved solution stability in methanol compared to the free acid.

5. SUMMARY OF SALT SCREENING STUDIES

A salt screening study was performed to identify suitable salt candidates of α -TEA. During 10 the course of the study, 16 different potential bases were examined. Of these bases, 6 bases formed crystalline salts. The most promising of the group, salts of lysine, arginine, and tris were isolated and studied in detail. These three salts were scaled-up to 2 grams and evaluated using multiple analytical techniques.

Based on the data obtained during this study, all three of these salts appear to be viable 15 candidates. These salts exhibit improved physical properties over the free acid. All of the salts are crystalline and have less "waxy" physical properties relative to the free acid. The lysine and arginine salts exhibit the most improvement in this characteristic. The visual melting temperature of all three salts is higher than the free acid. All three salts exhibit equal stability under all conditions tested with improved stability over the free acid in solution. The visual solubility of each salt was comparable to 20 the free acid in buffered water (< 0.2 mg/mL).

The salt samples all exhibited the potential to form putative hydrates all which appear to be reversible. The arginine and tris salts exhibited DVS water uptake near critical humidities, 40% and 55% RH respectively. The lysine salt shows DVS water uptake above about 75% RH.

Table 6: Summary of Final Salt Characteristics

Salt	Physical handling as compared to free acid	XRD	Visual melting compared to free acid	Stability compared to free acid	Hydration properties		Counter Ion Class
					DVS	Static 75% RH 30 Days	
lysine	better	crystalline	higher, ~200 °C	more stable in solution	~75% RH hydrate	0.6 wt% gain	II
arginine	better	crystalline	higher, ~180 °C	more stable in solution	~40% RH hemihydrate	0.7 wt% gain	II
tris	better	crystalline	higher, ~100 °C	more stable in solution	~55% RH hemihydrate	4.0 wt% gain	II

C. Anti-Tumor Efficacy of α -TEA Salts in Animal Models

Studies of the anti-tumor efficacy of the α -TEA salts were carried out using a well characterized but difficult to treat 4T1 mouse model of metastatic mammary cancer with the α -TEA lysine salt (α -TEA-Lys). The lysine salt was incorporated into standard mouse chow at an amount of 5 3 g/kg chow (0.3% w/w) by Harlan Teklad, Madison WI. Half of the mice were placed on an α -TEA-Lys diet after the tumors were established (9 days post tumor injection.) and the rest placed on an analogous nutrient diet without α -TEA-Lys which served as the control group. The average tumor size of mice that ate a diet containing α -TEA-Lys showed a significant decrease in average tumor size compared to mice feed a control diet without the α -TEA salt (FIG. 18). Furthermore, mice 10 consuming a diet which contains the α -TEA salt showed an increased survival compared to mice who consumed a control diet of standard chow (FIG. 19). These data suggest that converting α -TEA into a salt appear to have little change on the efficacy of the compound with significant tumor growth suppression, prolongation of overall survival, and complete tumor regression in about 25% of the mice treated.

15

* * * * *

All of the compounds, polymorphs, formulations, and methods disclosed and claimed herein can be made and executed without undue experimentation in light of the present disclosure. While the compounds, polymorphs, formulations, and methods of this invention have been described in terms of preferred embodiments, it will be apparent to those of skill in the art that variations may be applied to 20 the compounds, polymorphs, formulations, and methods, as well as in the steps or in the sequence of steps of the method described herein without departing from the concept, spirit, and scope of the invention. More specifically, it will be apparent that certain agents which are both chemically and physiologically related may be substituted for the agents described herein while the same or similar results would be achieved. All such similar substitutes and modifications apparent to those skilled in 25 the art are deemed to be within the spirit, scope and concept of the invention as defined by the appended claims.

REFERENCES

The following references to the extent that they provide exemplary procedural or other details supplementary to those set forth herein, are specifically incorporated herein by reference.

US 6,417,223

US 6,703,384

US 6,770,672

US 7,312,232

Anderson, *et al.*, *Cancer Res*, 64:4263-4269, 2004.

Hahn, *et al.*, *Cancer Res*, 66:9374-9378, 2006.

Hahn, *et al.*, *Mol Cancer Ther*, 8:1570-1578, 2009.

Hahn, *et al.*, *BMC Cancer*, 11:471, 2011.

Kline, *et al.*, *Vitam Horm*, 76:435-461, 2007.

Lawson, *et al.*, *Cancer Chemother Pharmacol*, 54:421-431, 2004.

March's Advanced Organic Chemistry: Reactions, Mechanisms, and Structure, 2007.

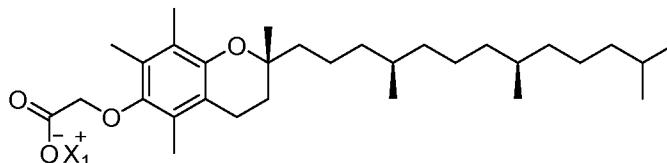
Neuzil, *et al.*, *Mol Aspects Med*, 28:607-645, 2007.

Yu, *et al.*, *Mol Carcinog*, 49:964-973, 2010.

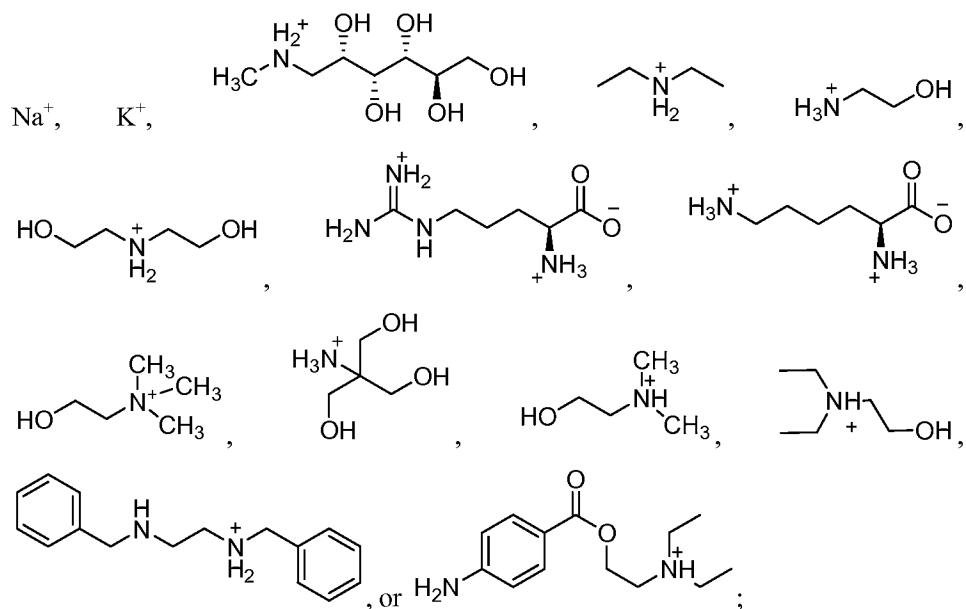
CLAIMS

What Is Claimed Is:

1. A compound of the formula:

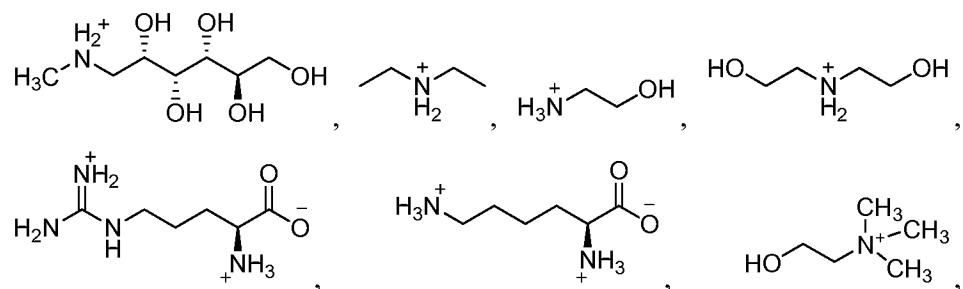


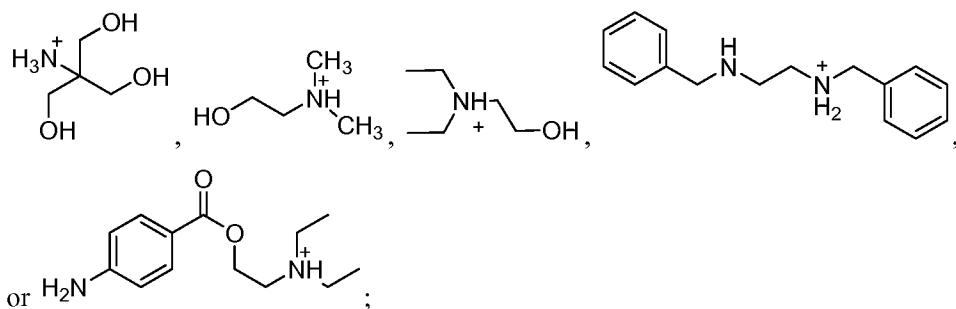
wherein X_1^+ is:



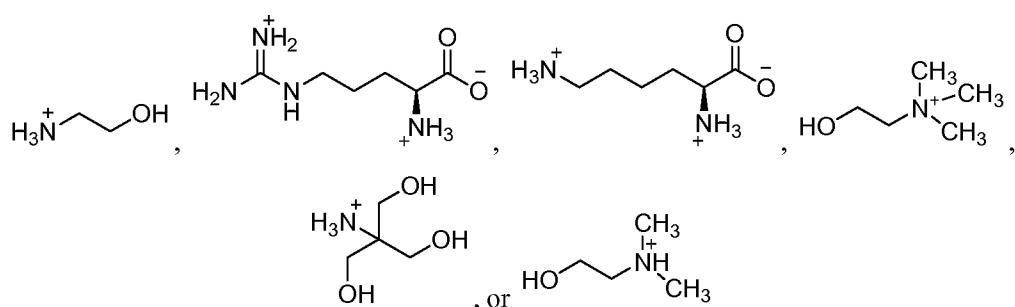
or a hydrate thereof.

2. The compound of claim 1, wherein X_1^+ is:

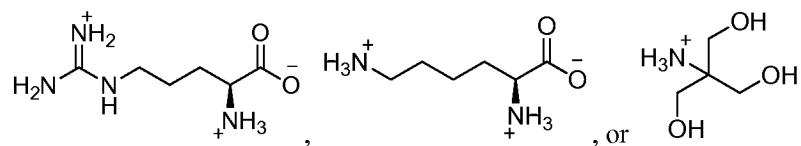




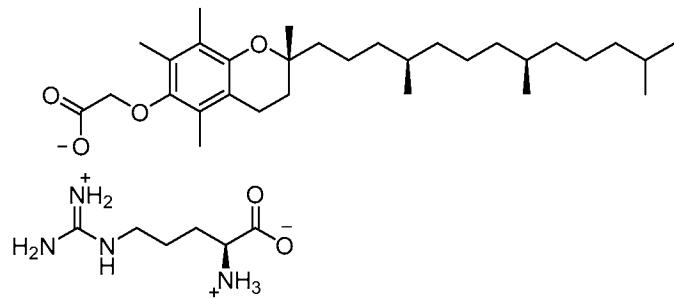
3. The compound of either claim 1 or claim 2, wherein the X_1^+ is:



4. The compound according to any one of claims 1-3, wherein the X_1^+ is

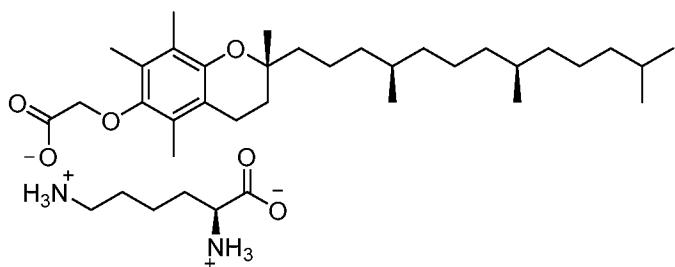


5. The compound according to any one of claims 1-4, wherein the formula is further defined as:



6. The compound of claim 5, wherein the compound is present as a polymorphic form having an X-ray powder diffraction pattern (CuK α) comprising two peaks at about 11.84 $^{\circ}$ 2 θ and 19.74 $^{\circ}$ 2 θ or a hydrate, thereof.

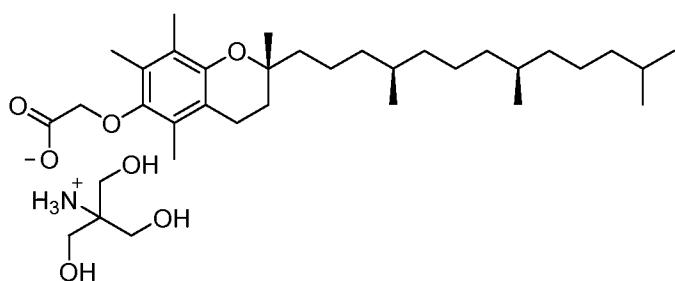
7. The compound of claim 6, wherein the polymorphic form has an X-ray powder diffraction pattern (CuK α) further comprising a peak at about 20.73 $^{\circ}$ 20.
8. The compound of either claim 6 or 7, wherein the polymorphic form has an X-ray powder diffraction pattern (CuK α) further comprising 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, or all of the peaks at about 6.24, 9.35, 14.92, 15.51, 17.03, 18.49, 21.70, 22.81, 23.51, 24.85, 27.69, 28.31, 30.93, 32.30, 33.21, 35.02, and 39.30 $^{\circ}$ 20.
9. The compound according to any one of claims 6-8, wherein the polymorphic form has an X-ray powder diffraction pattern (CuK α) substantially as shown in FIG. 4.
10. The compound according to any one of claims 6-9, wherein the polymorphic form has a differential scanning calorimetry (DSC) isotherm comprising two endotherms from about 125 $^{\circ}$ C to about 140 $^{\circ}$ C and from about 170 $^{\circ}$ C to about 195 $^{\circ}$ C.
11. The compound of claim 10, wherein the polymorphic form has a differential scanning calorimetry (DSC) isotherm comprising two endotherms centered at about 131 $^{\circ}$ C and at around 184 $^{\circ}$ C, respectively.
12. The compound of either claim 10 or claim 11, wherein the polymorphic form has a differential scanning calorimetry isotherm substantially as shown in FIG. 5.
13. The compound according to any one of claims 6-12, wherein the polymorphic form is anhydrous.
14. The compound according to any one of claims 6-12, wherein the polymorphic form is a hydrate.
15. The compound of claim 14, wherein the polymorphic form is a hemihydrate.
16. The compound according to any one of claims 1-4, wherein the formula is further defined as:



or a hydrate, thereof.

17. The compound of claim 16, wherein the compound is present as a polymorphic form having an X-ray powder diffraction pattern (CuK α) comprising a peak at about 9.69 °2 θ or a hydrate, thereof.
18. The compound of claim 17, wherein the X-ray powder diffraction pattern (CuK α) further comprises a peak at about 20.10 °2 θ .

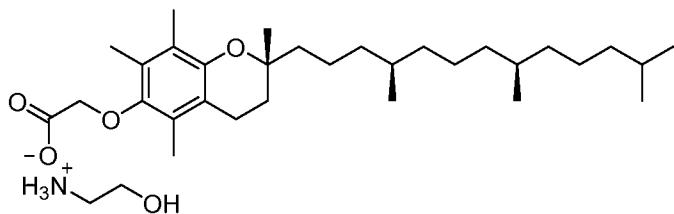
19. The compound of either claim 17 or 18, wherein the X-ray powder diffraction pattern (CuK α) further comprises a peak at about 19.78 $^{\circ}$ 20.
20. The compound according to any one of claims 17-19, wherein the X-ray powder diffraction pattern (CuK α) further comprises 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, or all of the peaks at about 5.90, 7.77, 14.08, 17.58, 17.91, 18.31, 21.26, 22.17, 23.16, 24.77, 28.57, 29.71, 31.10, 31.69, 34.45, 34.99, and 36.24 $^{\circ}$ 20.
21. The compound according to any one of claims 17-20, wherein the polymorphic form has an X-ray powder diffraction pattern (CuK α) that is substantially as shown in FIG. 6.
22. The compound according to any one of claims 17-21, wherein the polymorphic form has a differential scanning calorimetry (DSC) isotherm comprising two endotherms from about 53 $^{\circ}$ C to about 60 $^{\circ}$ C and from about 180 $^{\circ}$ C to about 235 $^{\circ}$ C.
23. The compound of claim 22, wherein the polymorphic form has a differential scanning calorimetry (DSC) isotherm comprising two endotherms centered at about 55.8 $^{\circ}$ C and at about 205 $^{\circ}$ C, respectively.
24. The compound of either claim 22 or claim 23, wherein the polymorphic form has a differential scanning calorimetry isotherm that is substantially as shown in FIG. 7.
25. The compound according to any one of claims 17-24, wherein the polymorphic form is anhydrous.
26. The compound according to any one of claims 17-24, wherein the polymorphic form is a hydrate.
27. The compound according to any one of claims 1-4, wherein the formula is further defined as:



or a hydrate, thereof.

28. The compound of claim 27, wherein the compound is present as a polymorphic form having an X-ray powder diffraction pattern (CuK α) comprising a peak at about 20.14 $^{\circ}$ 20 or a hydrate, thereof.
29. The compound of claim 28, wherein the X-ray powder diffraction pattern (CuK α) further comprises peaks at about 13.63, 14.40, 19.31, and 22.34 $^{\circ}$ 20.

30. The compound of either claim 28 or 29, wherein the X-ray powder diffraction pattern (CuK α) further comprises two peaks at about 18.25 and 18.49 $^{\circ}$ 20.
31. The compound according to any one of claims 28-30, wherein the X-ray powder diffraction pattern (CuK α) further comprises 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, or all of peaks at about 9.67, 15.60, 21.47, 23.57, 27.07, 28.39, 30.49, 32.21, 33.30, 34.29, 34.96, 35.89, and 36.39 $^{\circ}$ 20.
32. The compound according to any one of claims 28-31, wherein the polymorphic form has an X-ray powder diffraction pattern (CuK α) substantially as shown in FIG. 10.
33. The compound according to any one of claims 28-32, wherein the polymorphic form has a differential scanning calorimetry (DSC) isotherm comprising two endotherms from about 97.5 $^{\circ}$ C to about 120 $^{\circ}$ C and from about 185 $^{\circ}$ C to about 225 $^{\circ}$ C.
34. The compound of claim 33, wherein the polymorphic form has a differential scanning calorimetry (DSC) isotherm comprising two endotherms at about 106 $^{\circ}$ C and at about 192 $^{\circ}$ C, respectively.
35. The compound of either claim 33 or claim 34, wherein the polymorphic form has a differential scanning calorimetry isotherm substantially as shown in FIG. 11.
36. The compound according to any one of claims 28-35, wherein the polymorphic form is anhydrous.
37. The compound according to any one of claims 28-35, wherein the polymorphic form is a hydrate.
38. The compound of claim 37, wherein the polymorphic form is a hemihydrate.
39. The compound according to any one of claims 1-3, wherein the formula is further defined as:



or a hydrate, thereof.

40. The compound of claim 39, wherein the compound is present as a polymorphic form having an X-ray powder diffraction pattern (CuK α) comprising a peak at about 19.4 $^{\circ}$ 20 or a hydrate, thereof.
41. The compound of claim 40, wherein the X-ray powder diffraction pattern (CuK α) further comprises peaks at about 10.4, 14.1, and 17.8 $^{\circ}$ 20.

42. The compound of either claim 40 or claim 41, wherein the X-ray powder diffraction pattern (CuK α) further comprises two peaks at about 20.8 $^{\circ}$ 20.

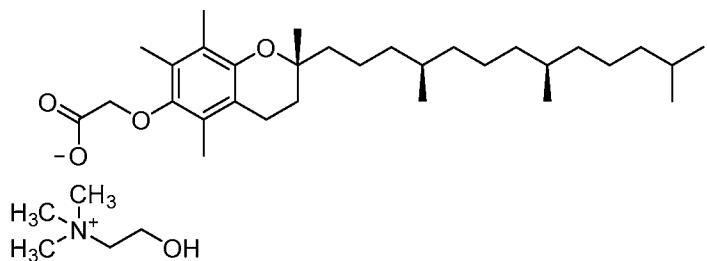
43. The compound according to any one of claims 40-42, wherein the polymorphic form has an X-ray powder diffraction pattern (CuK α) substantially as shown in FIG. 2.

44. The compound according to any one of claim 40-43, wherein the polymorphic form has a differential scanning calorimetry (DSC) isotherm comprising five endotherms from about 85 $^{\circ}$ C to about 97 $^{\circ}$ C, from about 97 $^{\circ}$ C to about 105 $^{\circ}$ C, from about 110 $^{\circ}$ C to about 125 $^{\circ}$ C, from about 145 $^{\circ}$ C to about 155 $^{\circ}$ C and from about 160 $^{\circ}$ C to about 210 $^{\circ}$ C.

45. The compound of claim 44, wherein the polymorphic form has a differential scanning calorimetry (DSC) isotherm comprising five endotherms at about 90.1 $^{\circ}$ C, at about 100.7 $^{\circ}$ C, at about 117.9 $^{\circ}$ C, at about 149.7 $^{\circ}$ C, and at about 181.6 $^{\circ}$ C, respectively.

46. The compound of either claim 44 or claim 45, wherein the polymorphic form has a differential scanning calorimetry isotherm substantially as shown in FIG. 3.

47. The compound according to any one of claims 1-3, wherein the formula is further defined as:



or a hydrate, thereof.

48. The compound of claim 47, wherein the compound is present as a polymorphic form having an X-ray powder diffraction pattern (CuK α) comprising two peaks at about 18.40 and 19.77 $^{\circ}$ 20 or hydrate, thereof.

49. The compound of claim 48, wherein the polymorphic form has an X-ray powder diffraction pattern (CuK α) further comprising peaks at about 4.28 and 10.75 $^{\circ}$ 20.

50. The compound of either claim 48 or claim 49, wherein the polymorphic form has an X-ray powder diffraction pattern (CuK α) further comprising two peaks at about 19.00 $^{\circ}$ 20.

51. The compound according to any one of claims 48-50, wherein the polymorphic form has an X-ray powder diffraction pattern (CuK α) further comprising 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, or all of the peaks at about 6.45, 12.90, 16.17, 16.73, 17.31, 17.72, 20.96, 21.72, 28.08, 28.71, 29.70, 31.05, 31.63, 32.25, 34.60, 35.18, 36.60, and 39.02 $^{\circ}$ 20.

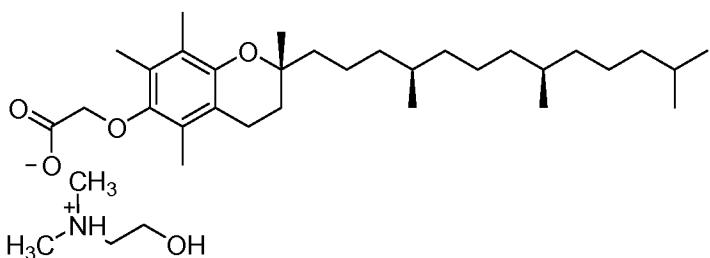
52. The compound according to any one of claims 48-51, wherein the polymorphic form has an X-ray powder diffraction pattern (CuK α) substantially as shown in FIG. 8.

53. The compound according to any one of claims 48-52, wherein the polymorphic form has a differential scanning calorimetry (DSC) isotherm comprising three endotherms from about 50 °C to about 60 °C, from about 75 °C to about 85 °C, and from about 150 °C to about 250 °C, respectively.

54. The compound of claim 53, wherein the polymorphic form has a differential scanning calorimetry (DSC) isotherm comprising three endotherms at about 50 °C, at about 80 °C, and at about 150 °C, respectively.

55. The compound of either claim 53 or claim 54, wherein the polymorphic form has a differential scanning calorimetry isotherm substantially as shown in FIG. 9.

56. The compound according to any one of claims 1-3, wherein the formula is further defined as:



or a hydrate, thereof.

57. The compound of claim 56, wherein the compound is present as a polymorphic form having an X-ray powder diffraction pattern (CuK α) comprising a peak at about 21.73 °2 θ or a hydrate, thereof.

58. The compound of claim 57, wherein the polymorphic form has an X-ray powder diffraction pattern (CuK α) further comprising peaks at about 9.34, 14.90, 16.06, 17.33, 18.51, and 22.94 °2 θ .

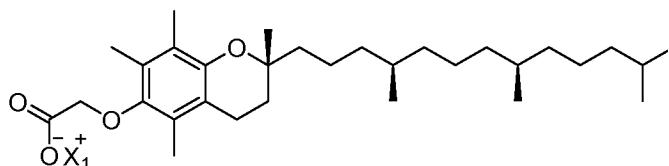
59. The compound of either claim 57 or claim 58, wherein the polymorphic form has an X-ray powder diffraction pattern (CuK α) further comprising two peaks centered at about 20.22 °2 θ .

60. The compound according to any one of claims 57-59, wherein the polymorphic form has an X-ray powder diffraction pattern (CuK α) further comprising 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, or all of peaks at about 10.05, 11.19, 14.13, 24.04, 27.65, 28.14, 29.73, 30.30, 31.06, 32.02, 33.45, and 34.49 °2 θ .

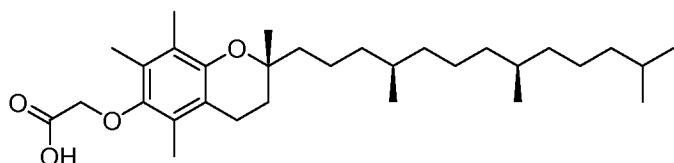
61. The compound according to any one of claims 57-60, wherein the polymorphic form has an X-ray powder diffraction pattern (CuK α) substantially as shown in FIG. 12.

62. The compound according to any one of claims 57-61, wherein the polymorphic form has a differential scanning calorimetry (DSC) isotherm comprising three endotherms from about 45 °C to about 55 °C, from about 65 °C to about 80 °C, and from about 80 °C to about 95 °C.

63. The compound of claim 62, wherein the polymorphic form has a differential scanning calorimetry (DSC) isotherm comprising three endotherms at about 47 °C, at about 73 °C, and at about 87 °C, respectively.
64. The compound of either claim 62 or claim 63, wherein the polymorphic form has a differential scanning calorimetry isotherm that is substantially as shown in FIG. 13.
65. A method of making a salt of the formula:



wherein: X_1^+ is a protonated base; comprising reacting a compound of the formula:

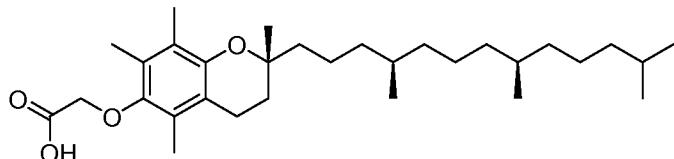


with the base sufficient to form a salt; or a hydrate thereof.

66. The method of claim 65, wherein the base is: sodium hydroxide, potassium hydroxide, magnesium hydroxide, meglumine, diethylamine, ethanolamine, diethanolamine, L-arginine, L-lysine, choline, tris-hydroxymethylamine, N,N-dimethylethanolamine, 2-diethylethanolamine, calcium hydroxide, dibenzylethylenediamine, or procaine.
67. The method of either claim 65 or claim 66, wherein the base is ethanolamine, L-arginine, L-lysine, choline, tris-hydroxymethylamine, or N,N-dimethylethanolamine.
68. The method according to any one of claims 65-67, wherein the base is L-arginine, L-lysine, or tris-hydroxymethylamine.
69. A pharmaceutical composition comprising a compound, a salt, or a polymorphic form according to any one of claims 1-68 and an excipient.
70. The pharmaceutical composition of claim 69, wherein the pharmaceutical composition is formulated for administration: orally, intraadiposally, intraarterially, intraarticularly, intracranially, intradermally, intralesionally, intramuscularly, intranasally, intraocularly, intrapericardially, intraperitoneally, intrapleurally, intraprostatically, intrarectally, intrathecally, intratracheally, intratumorally, intraumbilically, intravaginally, intravenously, intravesicularly, intravitreally, liposomally, locally, mucosally, parenterally, rectally, subconjunctival, subcutaneously, sublingually, topically, transbuccally, transdermally, vaginally, in crèmes, in lipid compositions, via a catheter, via a lavage, via continuous

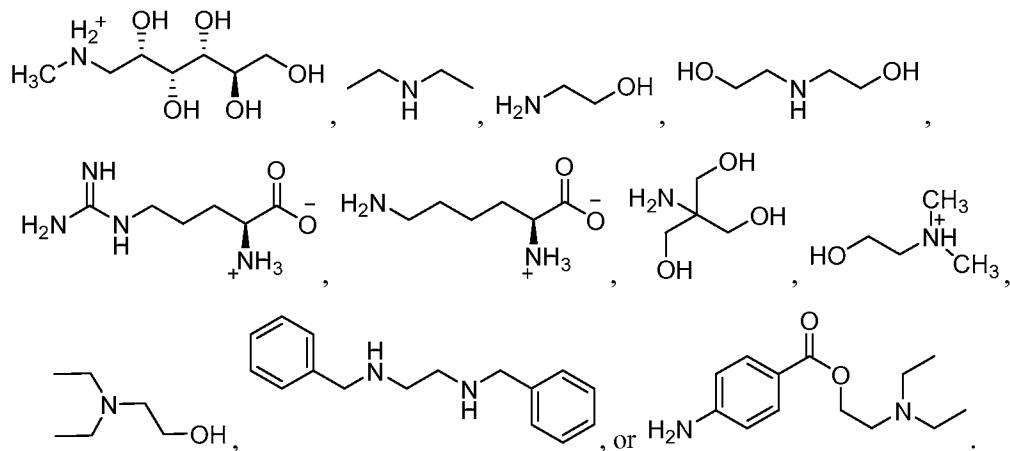
infusion, via infusion, via inhalation, via injection, via local delivery, or via localized perfusion.

71. The pharmaceutical composition of claim 70, wherein the pharmaceutical composition is formulated for oral, intraarterial, intravenous, intraperitoneal, topical, or inhalational administration.
72. A method of reacting a compound of the formula:

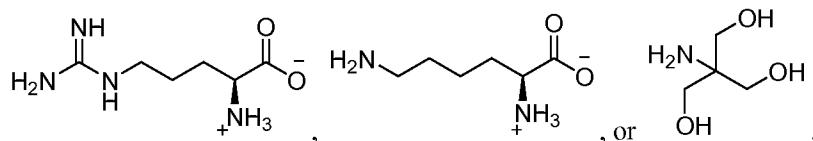


to obtain a polymorphic form comprising combining an about equal molar amount of a base and the compound in an alcoholic solvent and mixing for a time period from about 1 to about 24 hours.

73. The method of claim 72, wherein the base is NaOH, KOH,



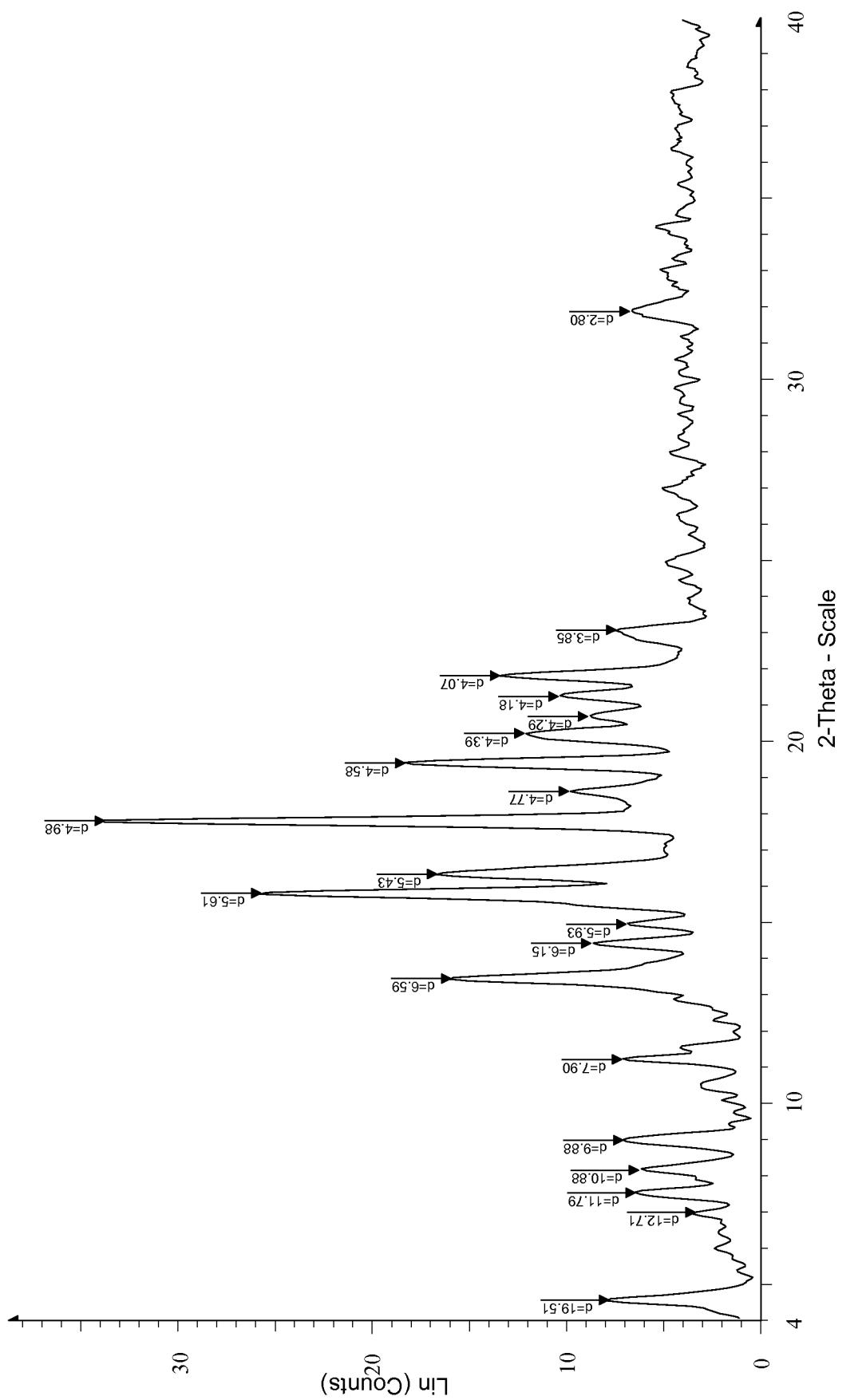
74. The method of either claim 72 or claim 73, wherein the base is



75. The method according to any one of claims 72-74, wherein the alcoholic solvent is of the formula: HO-R₁, wherein R₁ is alkyl_(C≤6).
76. The method according to any one of claims 72-75, wherein the alcoholic solvent is methanol, 1-butanol, or isopropyl alcohol.

77. The method according to any one of claims 72-76, wherein the alcoholic solvent is isopropyl alcohol.
78. The method according to any one of claims 72-77, further comprising allowing the solution to slowly evaporate at room temperature.
79. The method according to any one of claims 72-78, further comprising allowing the evaporation to occur under a constant flow of nitrogen gas.
80. The method of claim 79, wherein the constant flow of nitrogen has a pressure of about 2 psi.
81. A method of treating or preventing a hyperproliferative disease, comprising administering to the patient a therapeutically effective amount of the compound, salt, polymorphic form, or pharmaceutical composition according to any one of claims 1-71.
82. The method of claim 81, wherein the hyperproliferative disease is a cancer.
83. The method of claim 81 or claim 82, wherein the cancer is breast cancer, lung cancer, colon cancer, an endometrial tumor, cervical cancer, ovarian cancer, mammary cancer, prostate cancer, liver cancer, melanoma, lymphoma, or myeloma.
84. The method of claim 83, wherein the cancer is breast cancer.
85. The method of any one of claims 81- 83, further comprising administering to the patient a second therapy.
86. The method of claim 85, wherein the second therapy is an immunotherapy or surgery.
87. The method of claim 86, wherein the second therapy is an immunotherapy.
88. The method according to any one of claims 85 and 86, wherein the immunotherapy comprises administering a pharmaceutically effective amount of trastuzumab, adoptively transferred T lymphocytes, or a therapeutic antibody.
89. The method of claim 88, wherein the therapeutic antibody selectively binds HER2/neu.
90. The method of claim 88, wherein the immunotherapy is trastuzumab.
91. The method of claim 88, wherein the immunotherapy comprises administering adoptively transferred T lymphocytes to the patient.
92. The method of claim 91, wherein the adoptively transferred T lymphocytes are engineered to express a chimeric antigen receptor (CAR) to the patient.
93. The method of claim 88, wherein the immunotherapy comprises administering an immune modulating antibody to the patient.

94. The method of claim 93, wherein the immune modulating antibody is an anti-PD-1, anti-4-1-BB, anti-GITR, anti-TIM3, anti-LAG3, anti-TIGIT, anti-CTLA-4 or an anti-LIGHT antibody.
95. The method of claim 85, wherein the second therapy is radiotherapy or a chemotherapeutic agent.
96. The method of claim 86, wherein the second therapy is surgery.
97. The method of claim 95, wherein the second therapy is a chemotherapeutic agent.
98. The method of claim 97, wherein the chemotherapeutic agent is anthracyclines, taxanes, methotrexate, mitoxantrone, estramustine, doxorubicin, etoposide, vinblastine, carboplatin, vinorelbine, 5-fluorouracil, cisplatin, topotecan, ifosfamide, cyclophosphamide, epirubicin, gemcitabine, vinorelbine, irinotecan, etoposide, vinblastine, pemetrexed, melphalan, capecitabine, oxaliplatin, BRAF inhibitors, and TGF-beta inhibitors.

**FIG. 1**

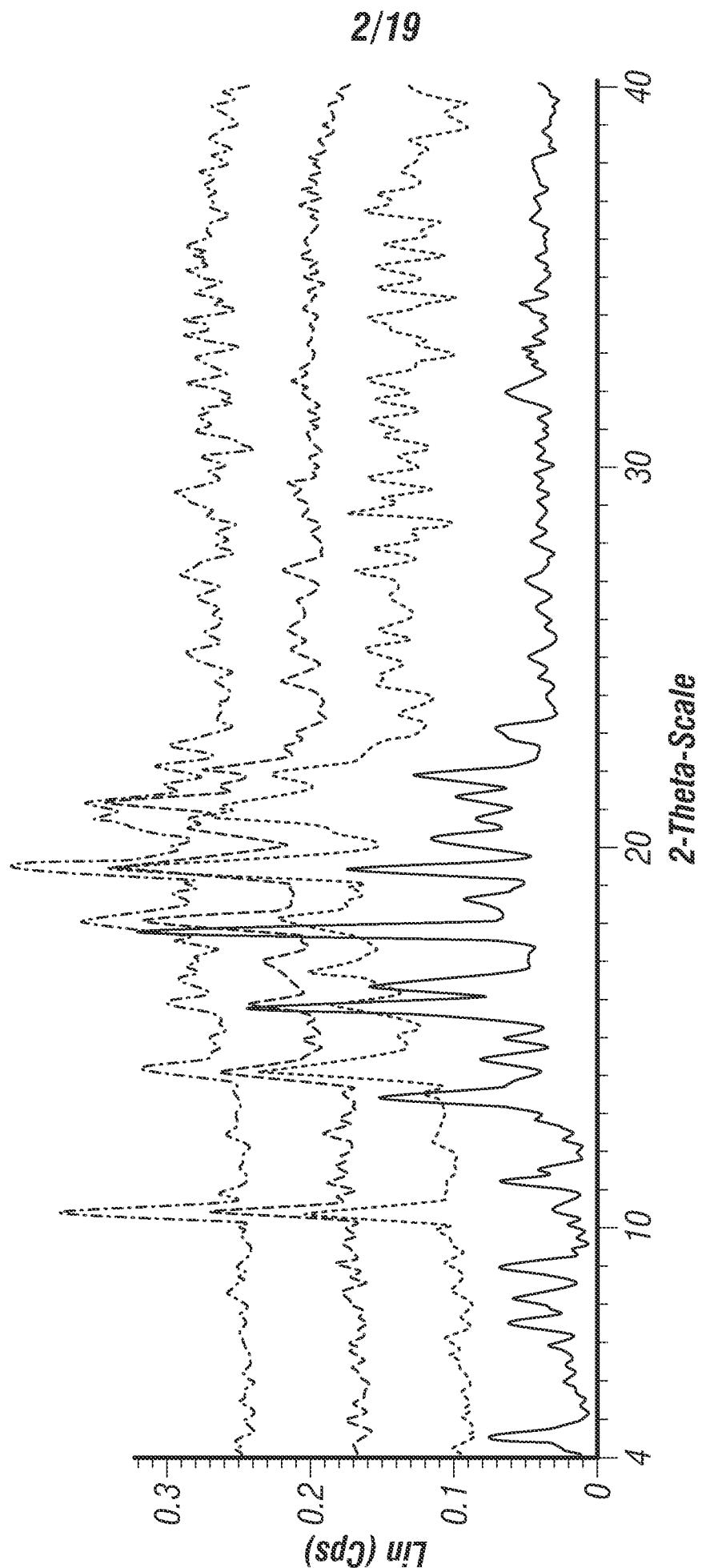


FIG. 2

DSC

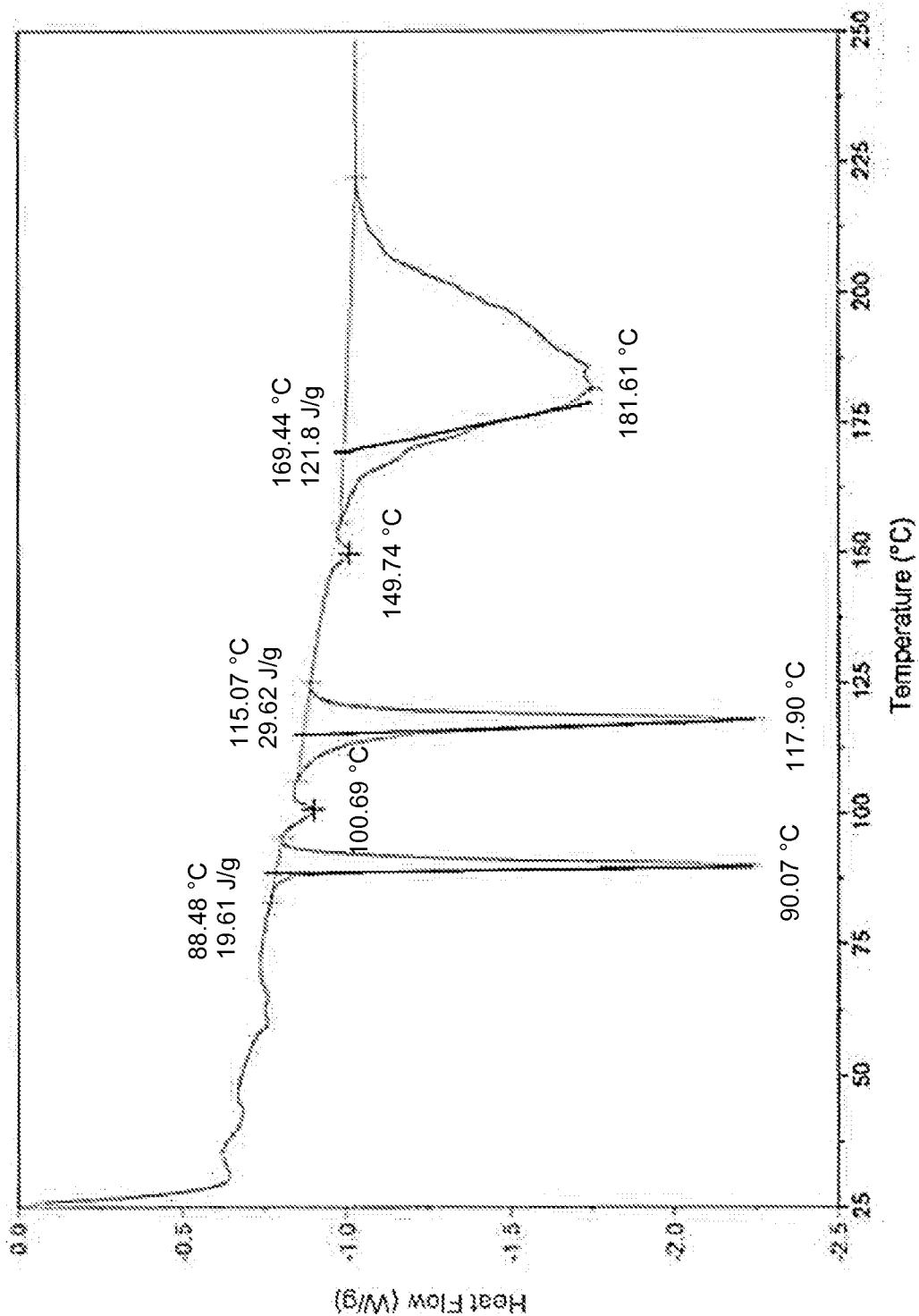


FIG. 3

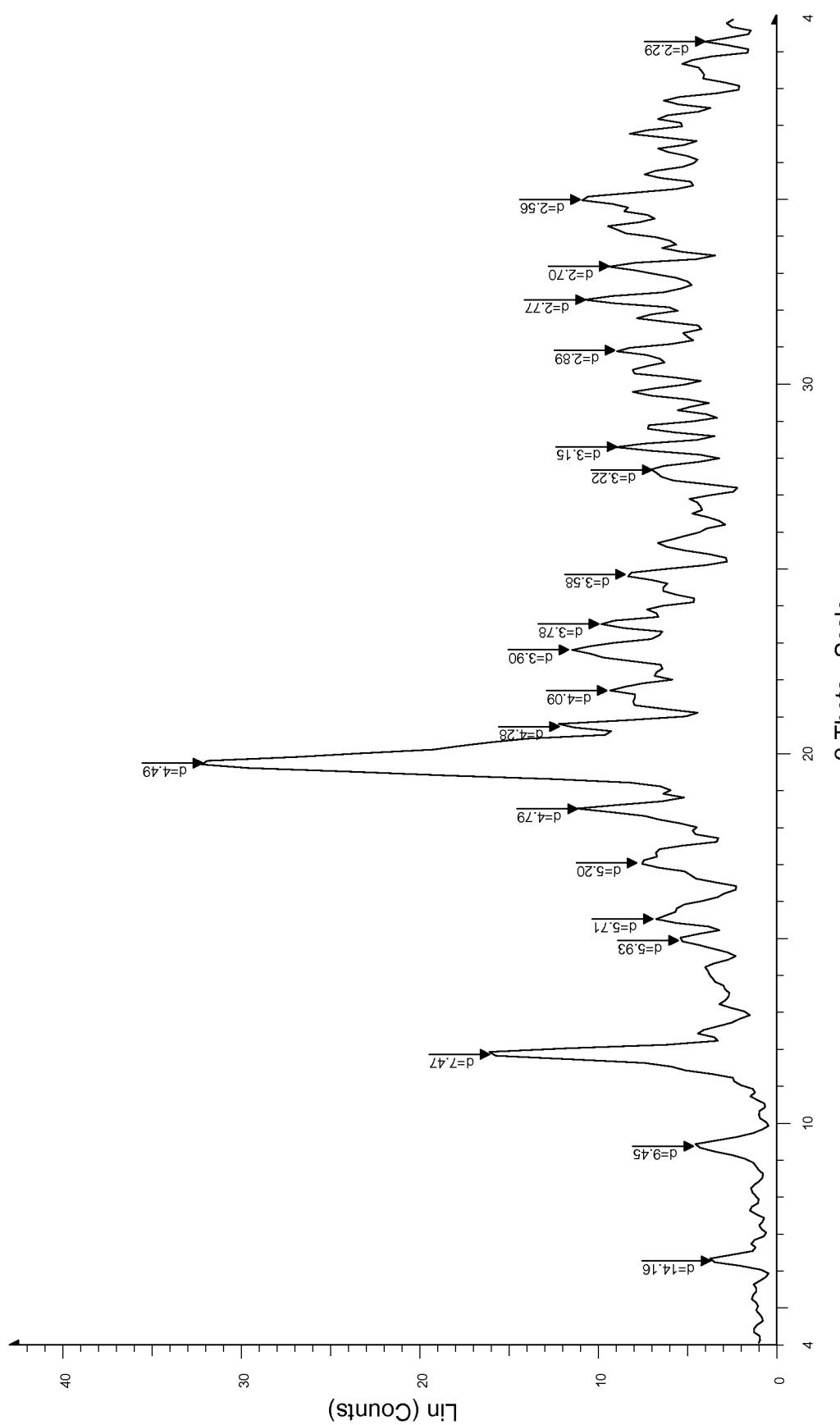


FIG. 4

DSC

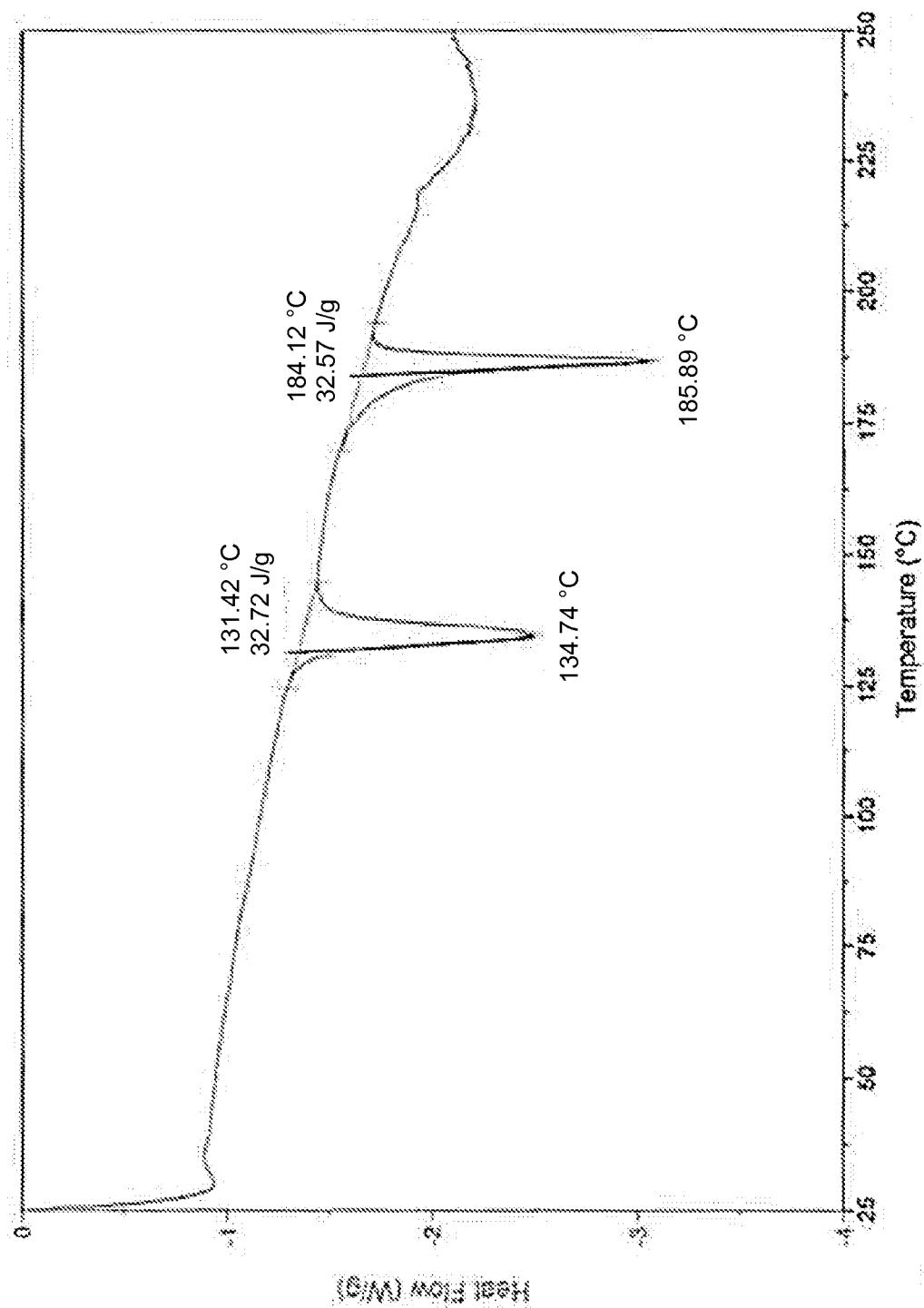
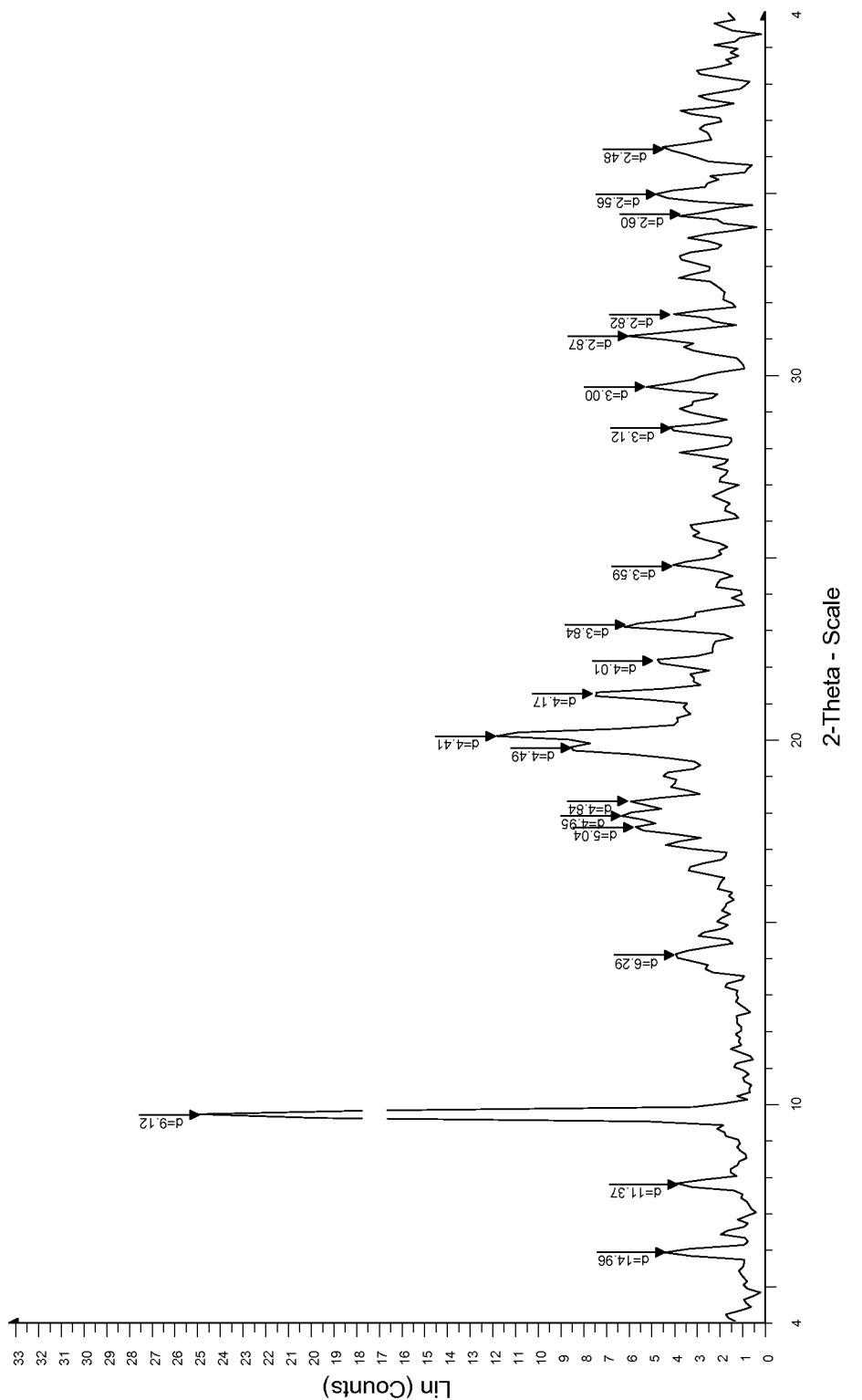


FIG. 5



DSC

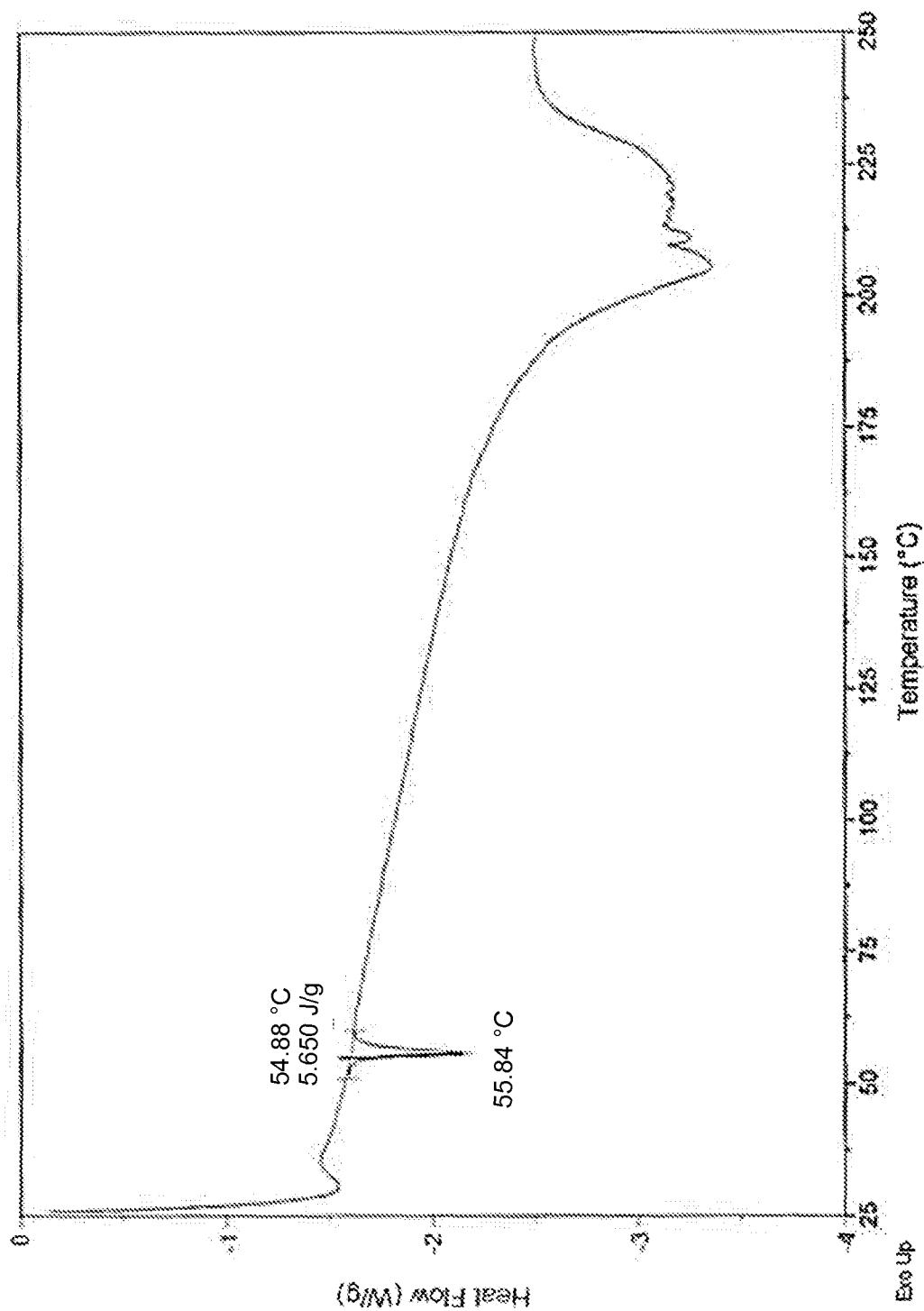
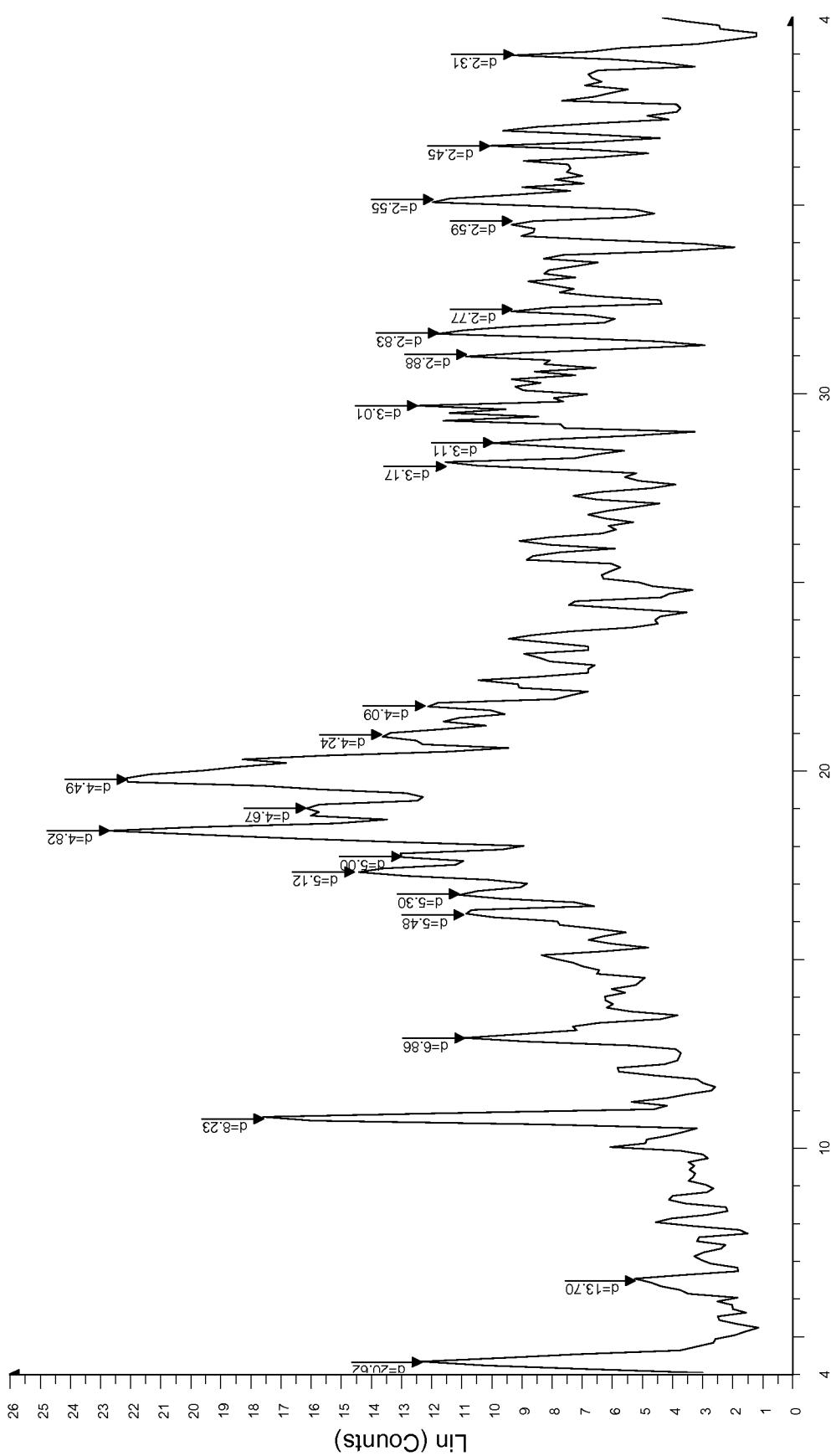


FIG. 7



2-Theta - Scale

FIG. 8

DSC

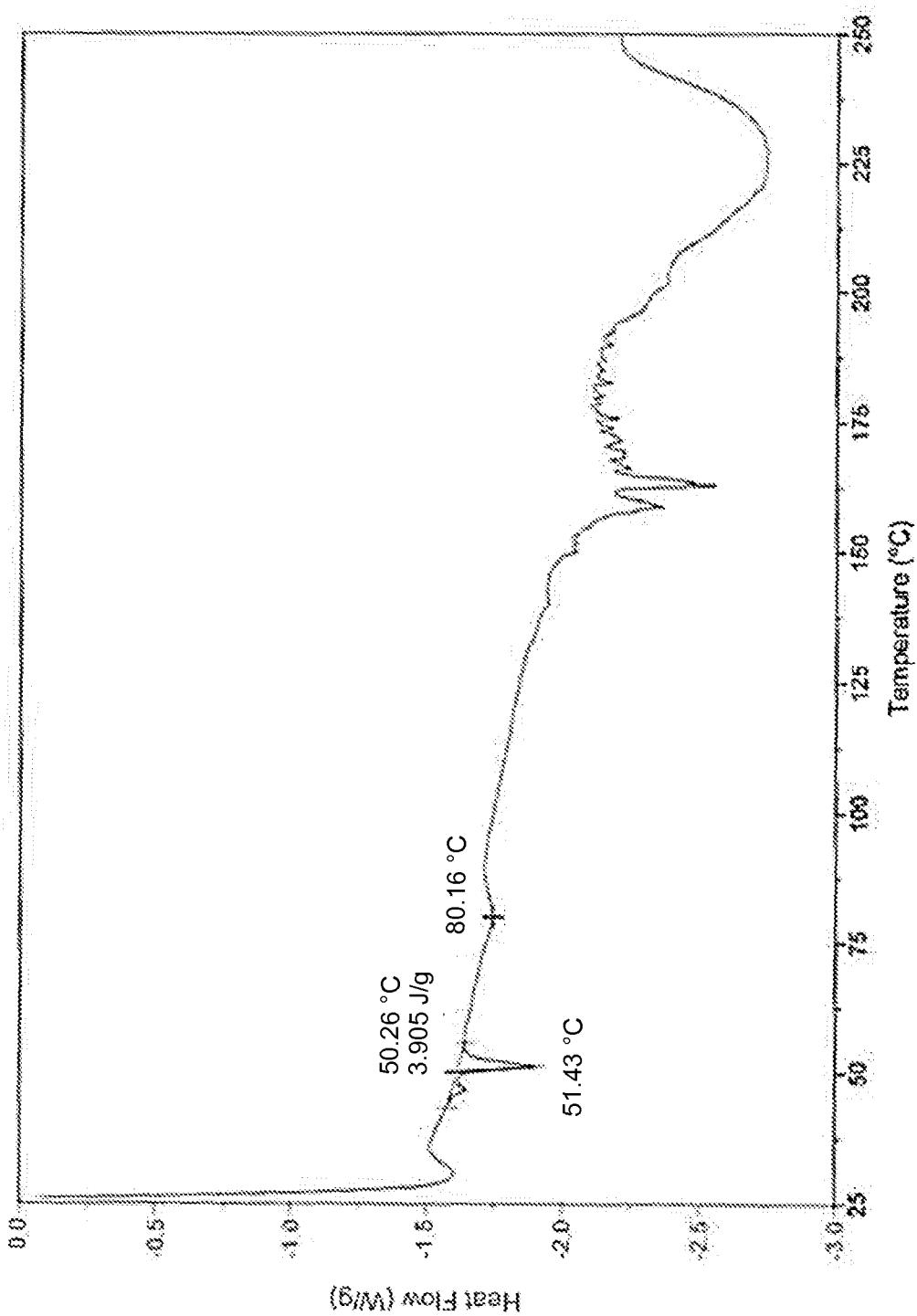
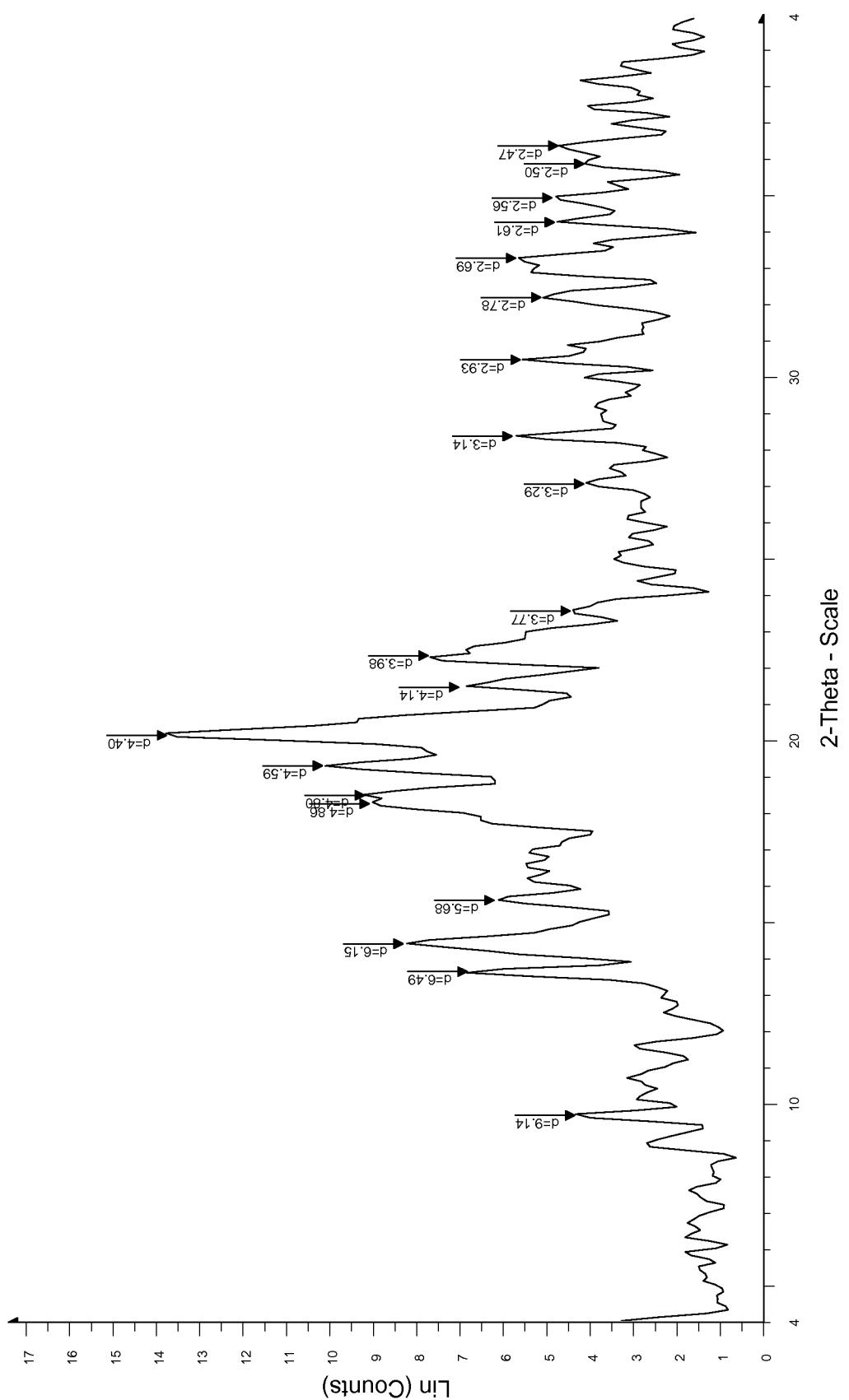


FIG. 9

**FIG. 10**

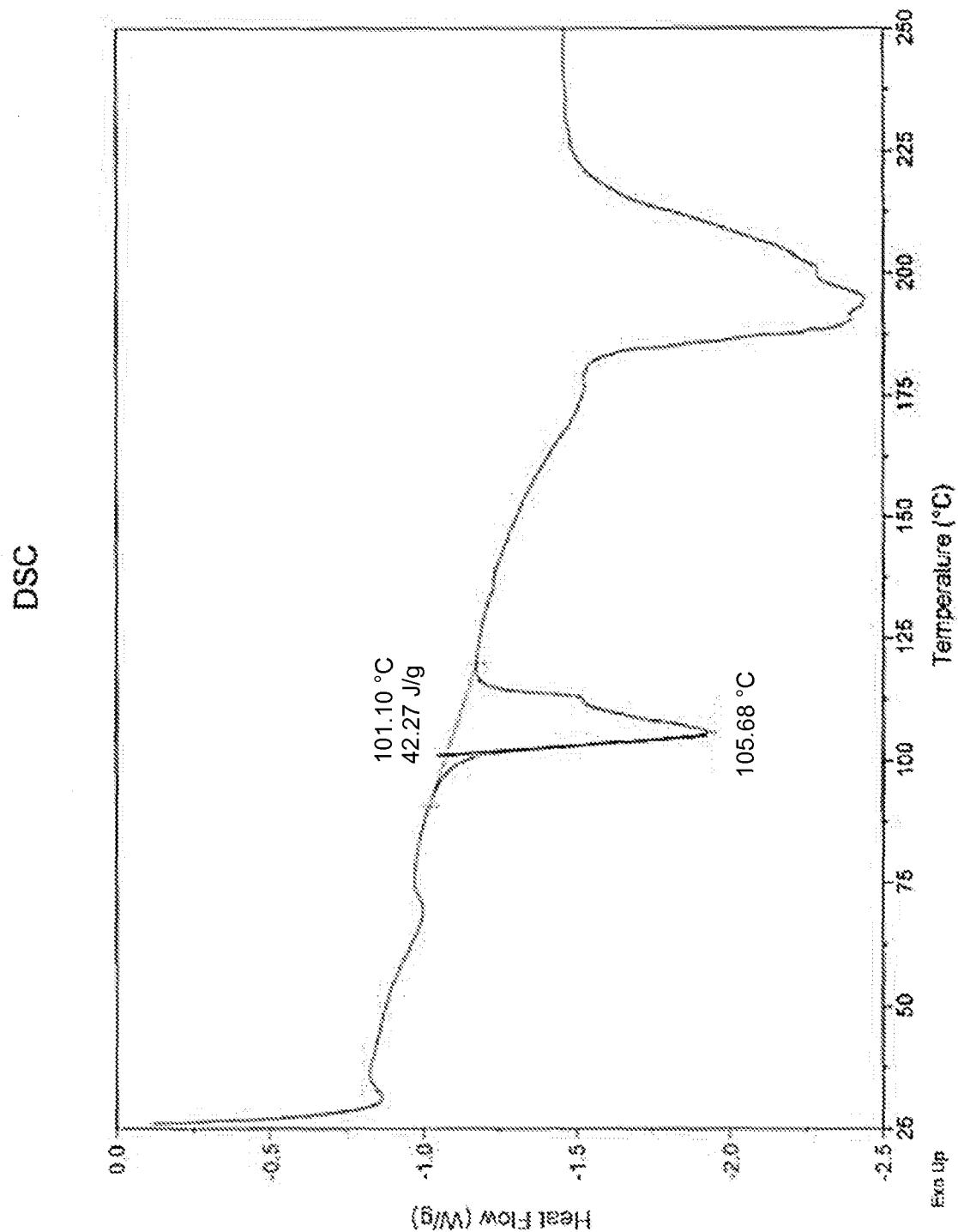


FIG. 11

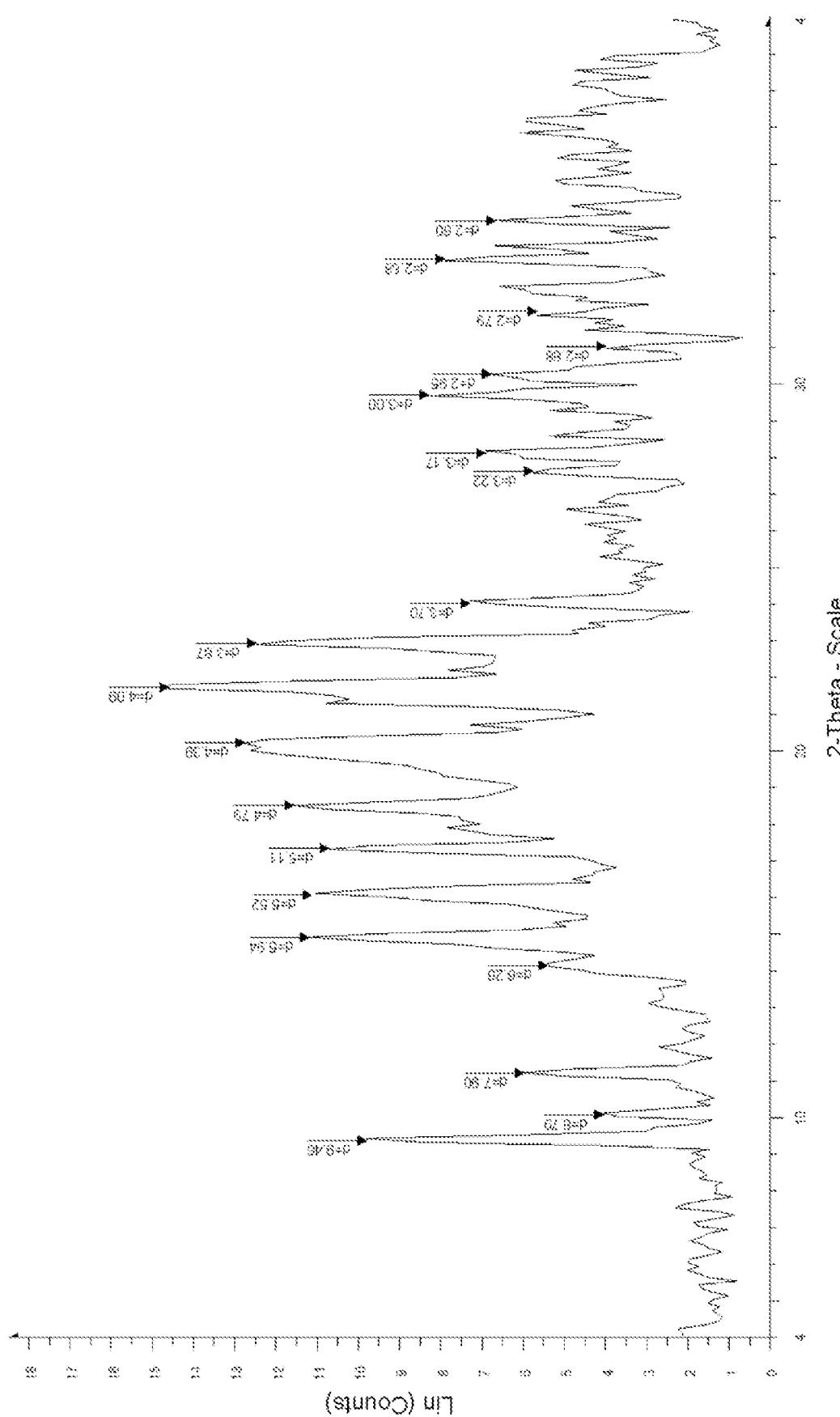


FIG. 12

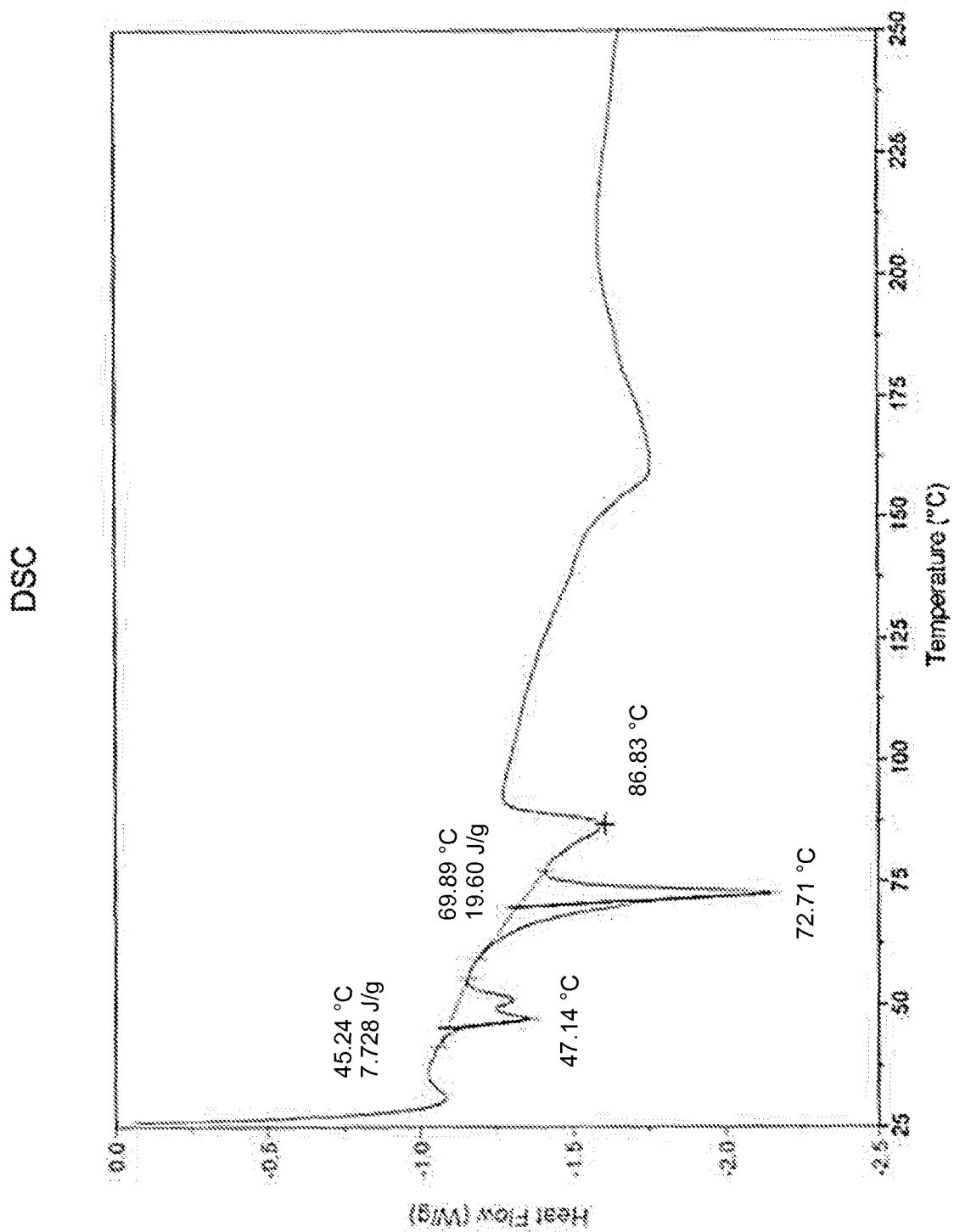
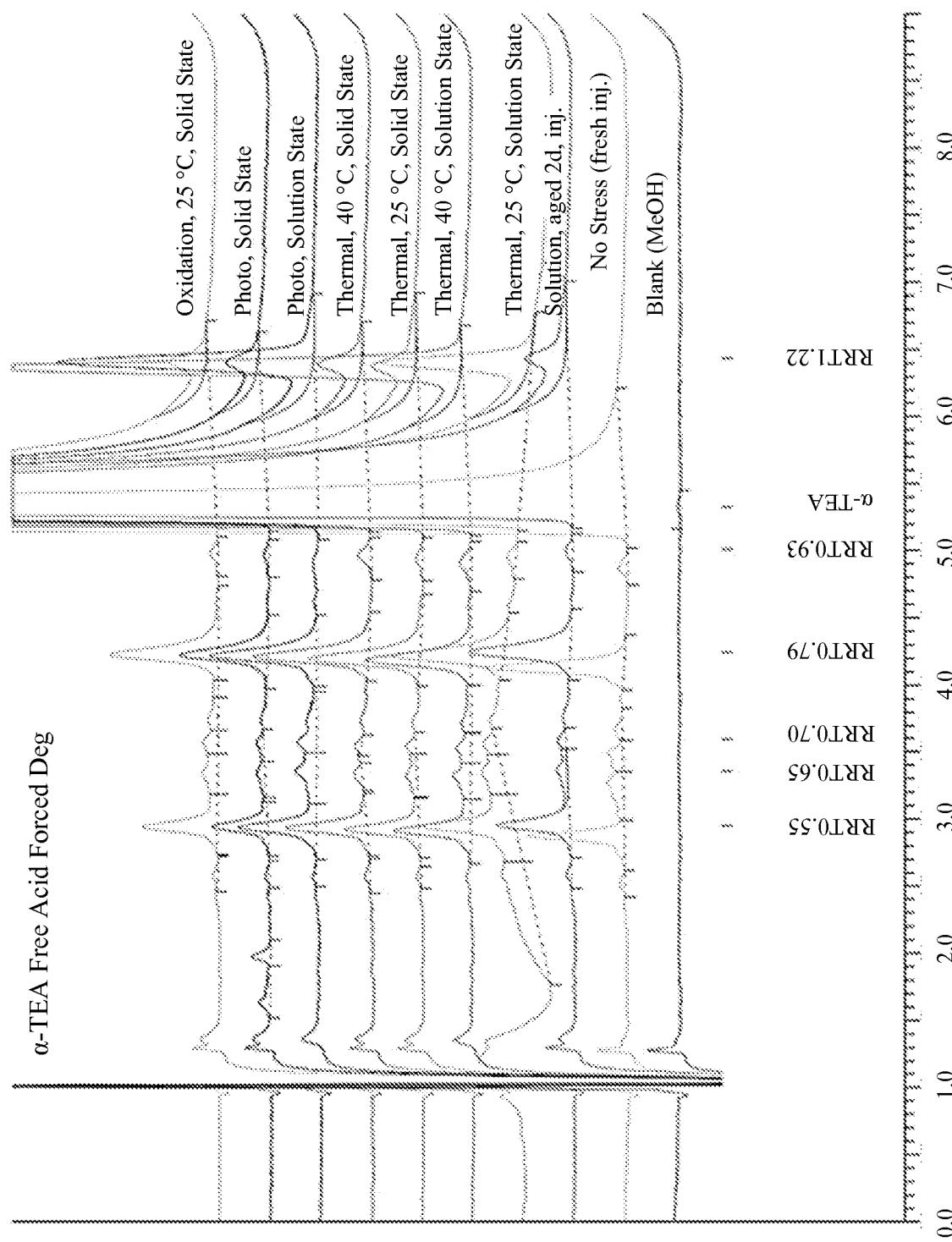
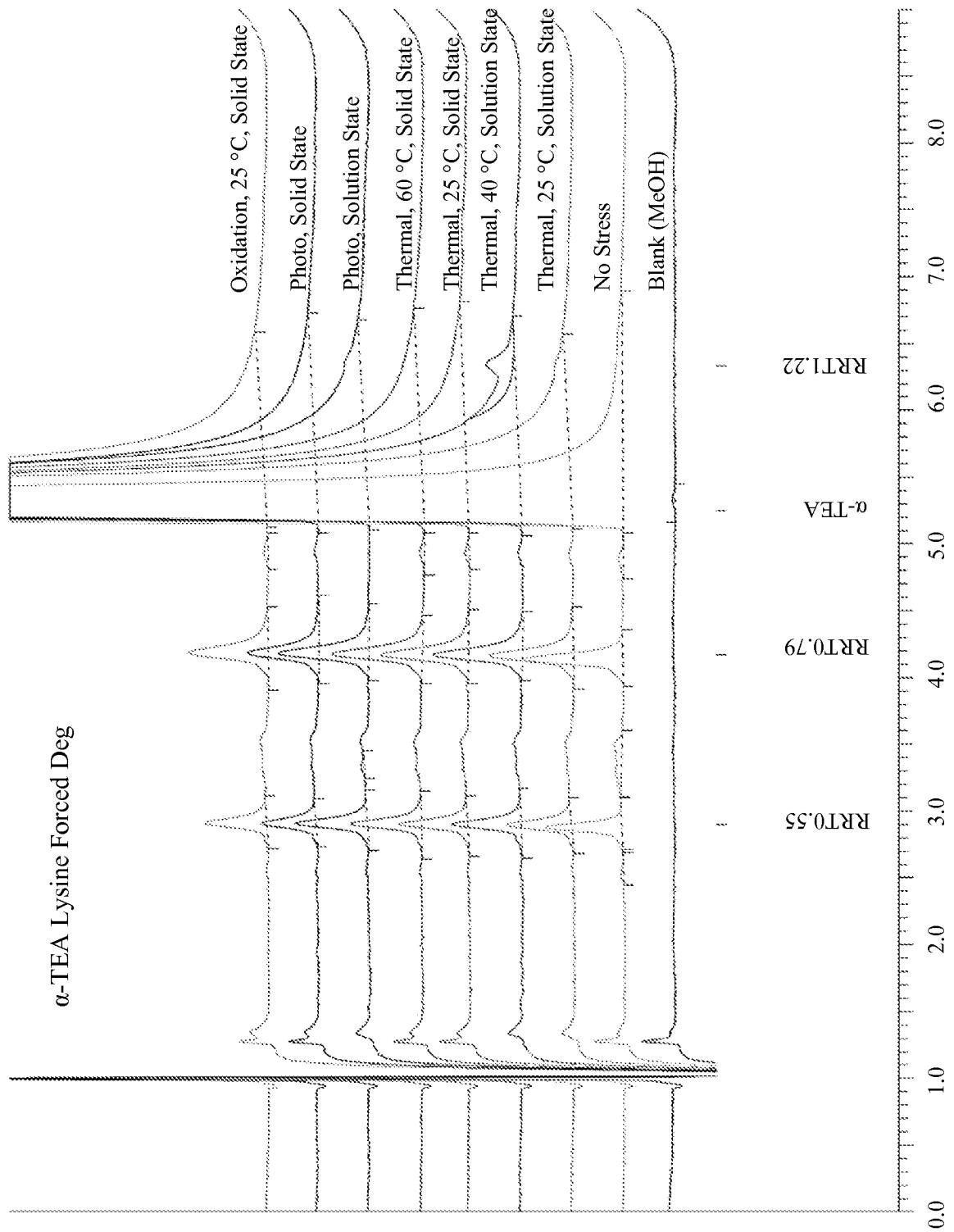
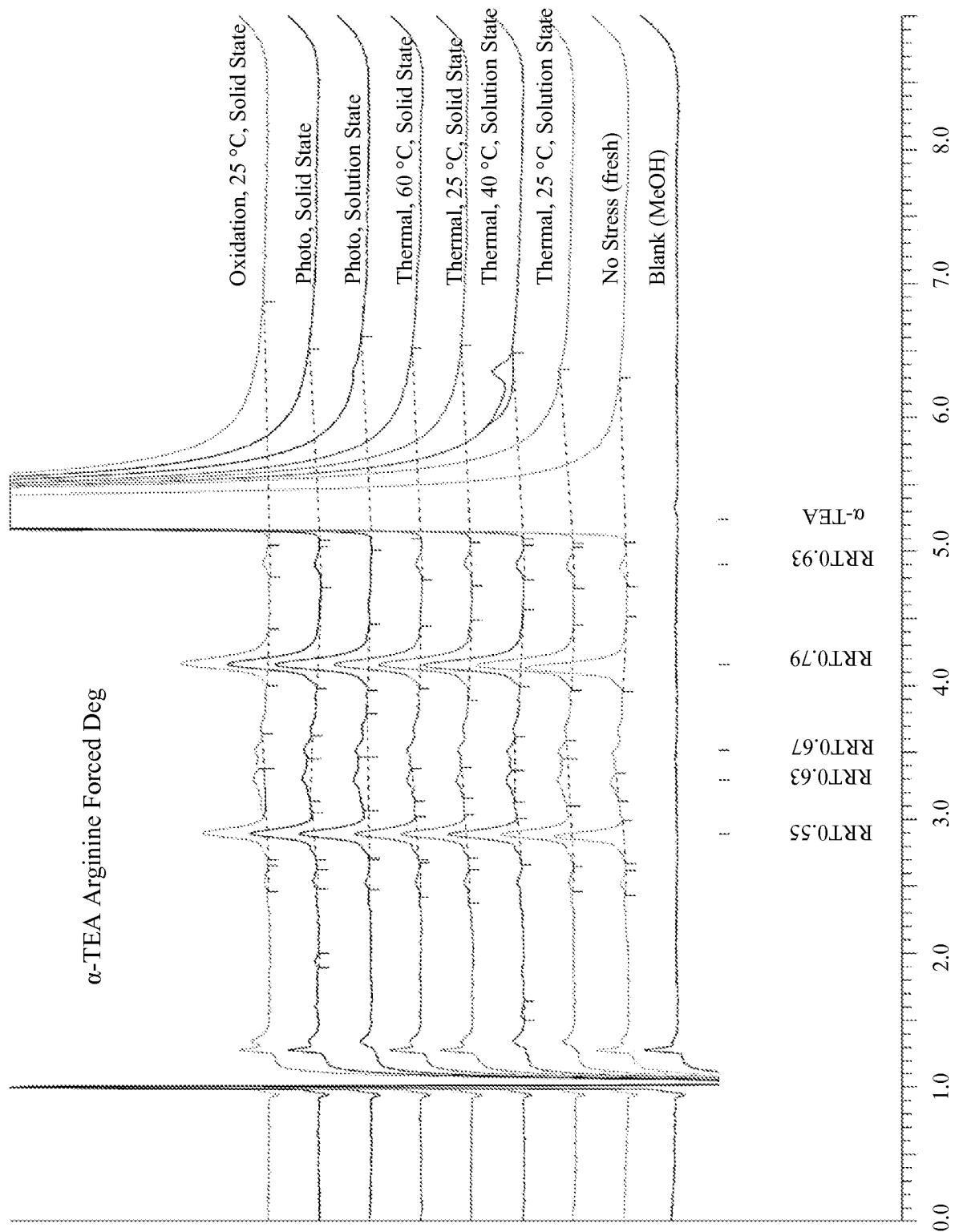
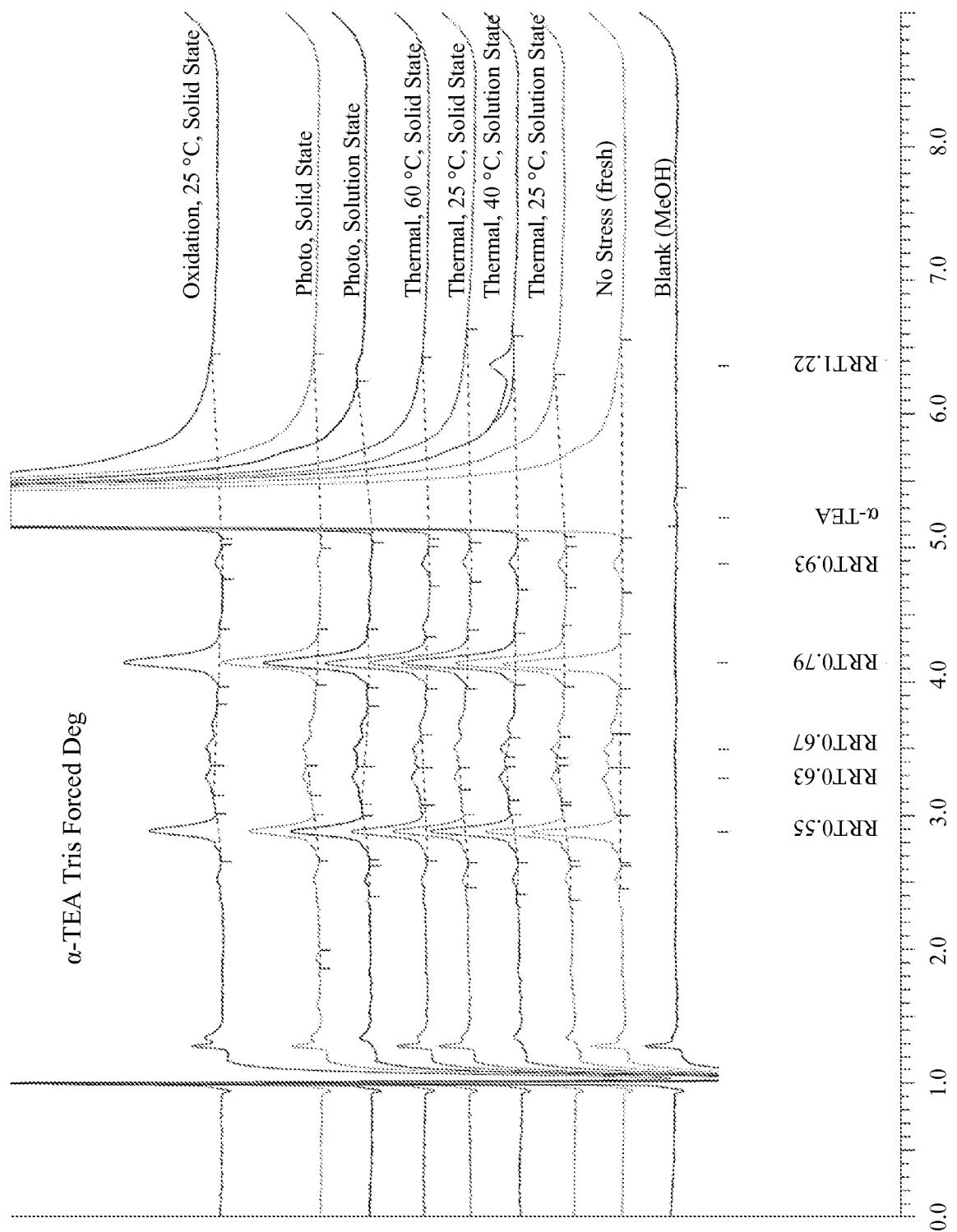


FIG. 13

**FIG. 14**

**FIG. 15**

**FIG. 16**

**FIG. 17**

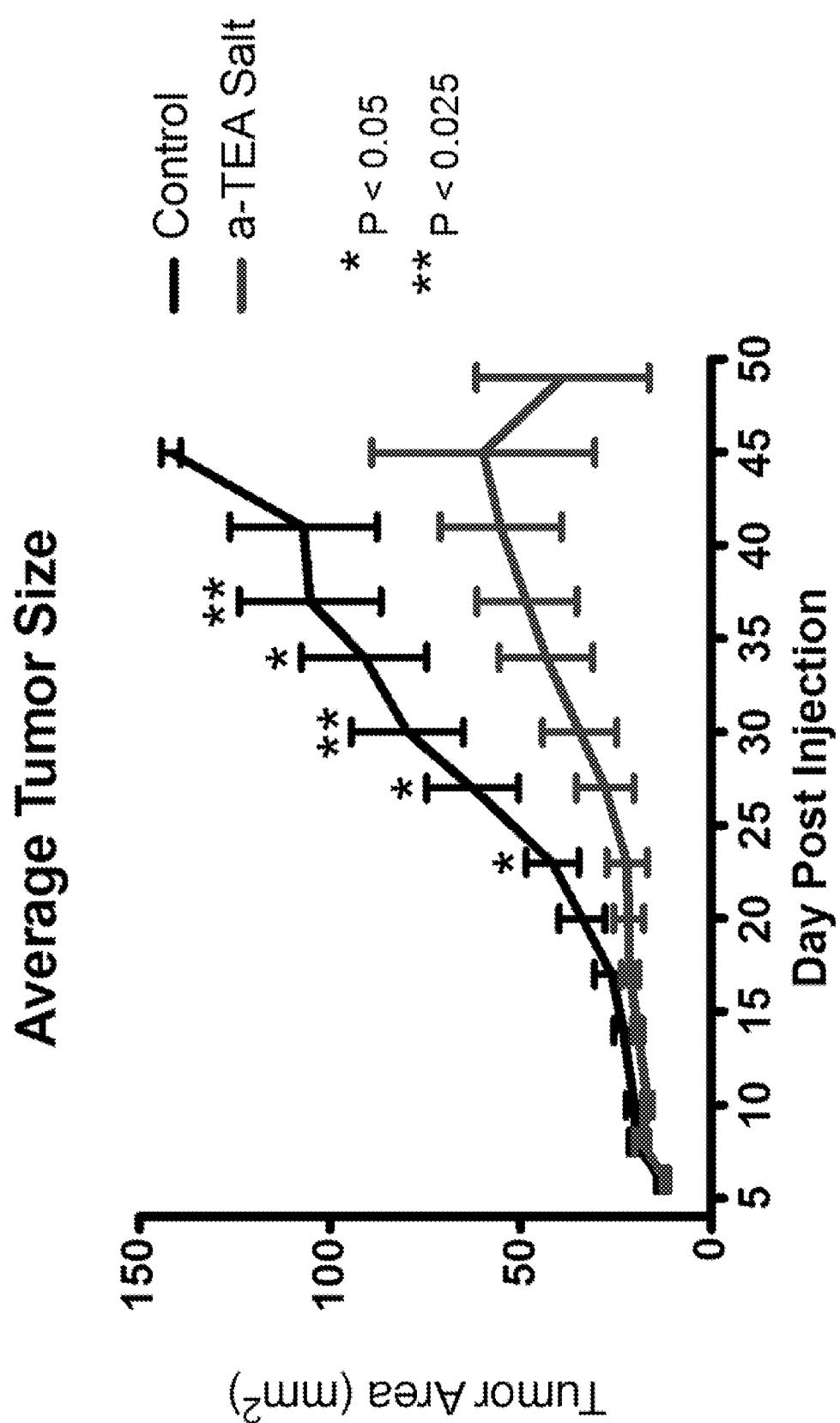


FIG. 18

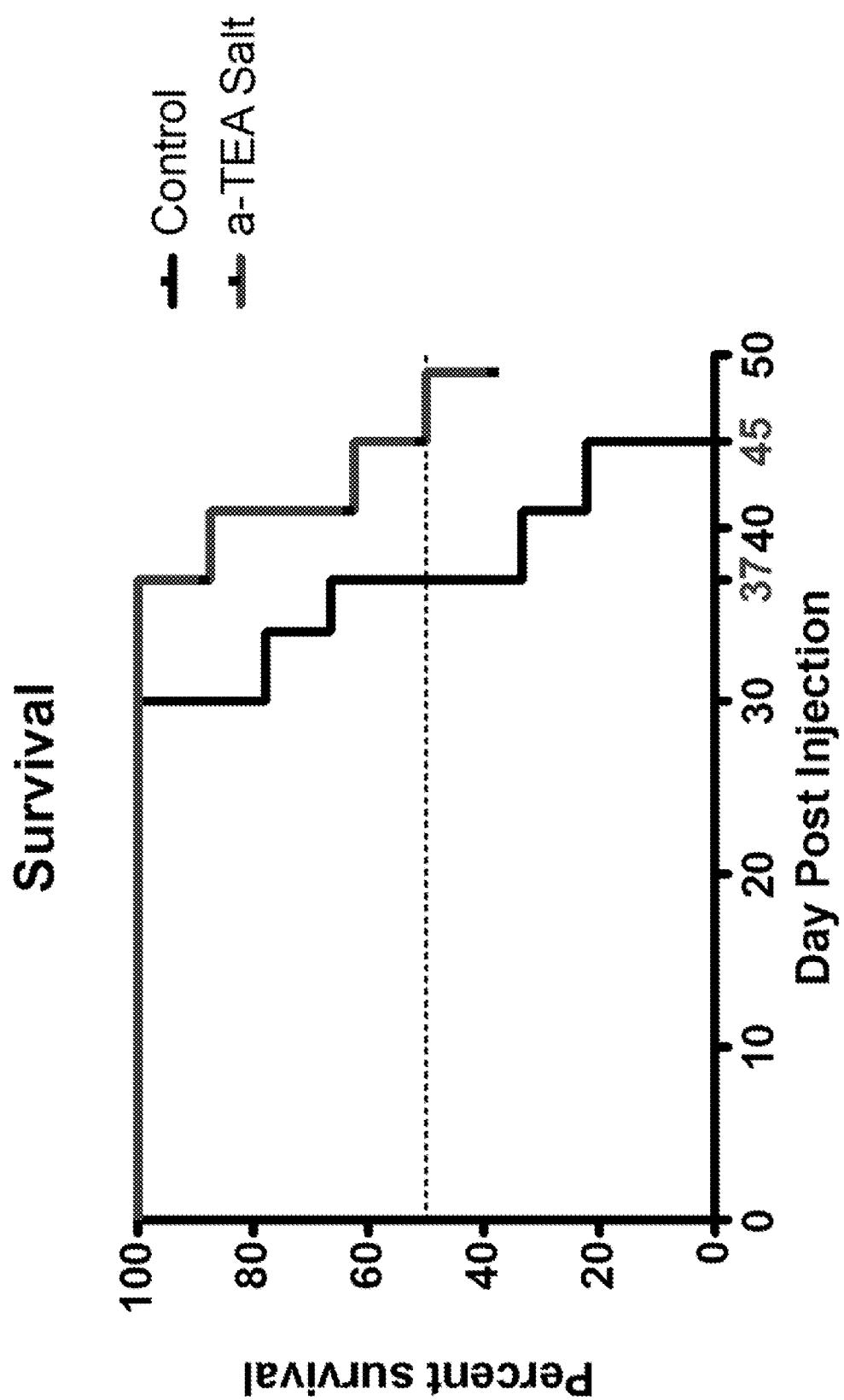


FIG. 19

INTERNATIONAL SEARCH REPORT

International application No
PCT/US2014/054354

A. CLASSIFICATION OF SUBJECT MATTER
INV. C07D311/72 A61K31/355 A61P31/00
ADD.

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
C07D A61K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

EPO-Internal, CHEM ABS Data, WPI Data, PAJ, BIOSIS

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO 00/16772 A1 (RES DEV FOUNDATION [US]) 30 March 2000 (2000-03-30) claims; example 2	1,66,72
Y	-----	1-98
X	WO 03/039461 A2 (RES DEV FOUNDATION [US]) 15 May 2003 (2003-05-15) claims; example 2	1,66,72
Y	-----	1-98
X	WO 01/58889 A1 (RES DEV FOUNDATION [US]) 16 August 2001 (2001-08-16) claims; example 2	1,66,72
Y	-----	1-98
		-/-

Further documents are listed in the continuation of Box C.

See patent family annex.

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- "A" document defining the general state of the art which is not considered to be of particular relevance
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Date of the actual completion of the international search	Date of mailing of the international search report
4 December 2014	17/12/2014
Name and mailing address of the ISA/ European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Fax: (+31-70) 340-3016	Authorized officer Frelon, Didier

INTERNATIONAL SEARCH REPORT

International application No
PCT/US2014/054354

C(Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	<p>FARISS M ET AL: "THE SELECTIVE ANTIPROLIFERATIVE EFFECTS OF ALPHA-TOCOPHERYL HEMISUCCINATE AND CHOLESTERYL HEMISUCCINATE ON MURINE LEUKEMIA CELLS RESULT FROM THE ACTION OF THE INTACT COMPOUNDS", CANCER RESEARCH, vol. 54, no. 13, 1 July 1994 (1994-07-01), pages 3346-3351, XP000965041, ISSN: 0008-5472 page 3347 - page 3348</p> <p>-----</p> <p>WANG, XIU-FANG ET AL: "Vitamin E analogues as anticancer agents: lessons from studies with .alpha.-tocopheryl succinate", MOLECULAR NUTRITION & FOOD RESEARCH, vol. 50, no. 8, 2006, pages 675-685, XP002733367, DOI: 10.1002/MNFR.200500267 page 676</p> <p>-----</p> <p>SHUN M-C ET AL: "PRO-APOPTOTIC MECHANISMS OF ACTION OF A NOVEL VITAMIN E ANALOG (ALPHA-TEA) AND A NATURALLY OCCURRING FORM OF VITAMIN E (DELTA-TOCOTRIENOL) IN MDA-MB-435 HUMAN BREAST CANCER CELLS", NUTRITION AND CANCER, vol. 48, no. 1, 1 January 2004 (2004-01-01), pages 95-105, XP009070526, ISSN: 0163-5581, DOI: 10.1207/S15327914NC4801_13 page 103</p> <p>-----</p> <p>NEUZIL J ET AL: "Vitamin E Analogues: A New Class of Inducers of Apoptosis with Selective Anti-Cancer Effects", CURRENT CANCER DRUG TARGETS, [Online] vol. 4, no. 4, 1 January 2004 (2004-01-01), pages 355-372, XP008105093, ISSN: 1568-0096, DOI: 10.2174/1568009043332943 Retrieved from the Internet: URL:http://www.bentham.org/ccdt/contabs/ccdt4-4.htm#link5</p> <p>page 360</p> <p>-----</p>	1-98
Y		1-98
Y		1-98
Y		1-98

INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No

PCT/US2014/054354

Patent document cited in search report	Publication date	Patent family member(s)		Publication date
WO 0016772	A1 30-03-2000	AT 466850 T		15-05-2010
		AU 757013 B2		30-01-2003
		AU 6155399 A		10-04-2000
		CA 2345079 A1		30-03-2000
		CN 1325303 A		05-12-2001
		CN 1706838 A		14-12-2005
		DK 1115398 T3		16-08-2010
		EP 1115398 A1		18-07-2001
		IL 142082 A		18-12-2005
		JP 2002526446 A		20-08-2002
		NZ 510732 A		30-01-2004
		RU 2232758 C2		20-07-2004
		TW 592695 B		21-06-2004
		US 6417223 B1		09-07-2002
		US 2002156024 A1		24-10-2002
		WO 0016772 A1		30-03-2000
		ZA 200102057 A		19-03-2002
<hr/>				
WO 03039461	A2 15-05-2003	AU 2002353971 A1		19-05-2003
		US 2002107207 A1		08-08-2002
		US 2004097431 A1		20-05-2004
		US 2008161349 A1		03-07-2008
		US 2010303899 A1		02-12-2010
		US 2012251616 A1		04-10-2012
		WO 03039461 A2		15-05-2003
<hr/>				
WO 0158889	A1 16-08-2001	AT 382615 T		15-01-2008
		AU 3680501 A		20-08-2001
		AU 2001236805 B2		14-07-2005
		CA 2399802 A1		16-08-2001
		CN 1529701 A		15-09-2004
		EP 1254130 A1		06-11-2002
		IL 151108 A		01-08-2006
		JP 2004504268 A		12-02-2004
		KR 20030005186 A		17-01-2003
		NZ 520798 A		28-05-2004
		US 2004235938 A1		25-11-2004
		US 2008119514 A1		22-05-2008
		WO 0158889 A1		16-08-2001
		ZA 200206243 A		28-10-2009
<hr/>				