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**BEFORE THE  
BOARD OF PHARMACY  
DEPARTMENT OF CONSUMER AFFAIRS  
STATE OF CALIFORNIA**

In the Matter of the Accusation Against:

**TILTON NGUYEN PHARMACEUTICAL  
INC. DBA OASIS PHARMACY, PAULINE  
MINH TILTON, PRESIDENT**  
15433 W. Sand St., Ste. 102  
Victorville, CA 92392

**Pharmacy Permit No. PHY 53859,**

**and**

**PAULINE MINH TILTON**  
15433 W. Sand St., Ste. 102  
Victorville, CA 92392

**Pharmacist License No. RPH 47537**

Respondents.

Case No. 6608

**DEFAULT DECISION AND ORDER**

[Gov. Code, §11520]

**FINDINGS OF FACT**

1  
2           1.     On or about June 7, 2019, Complainant Anne Sodergren, in her official capacity as  
3 the Interim Executive Officer of the Board of Pharmacy, Department of Consumer Affairs, filed  
4 Accusation No. 6608 against Tilton Nguyen Pharmaceutical Inc. dba Oasis Pharmacy, Pauline  
5 Minh Tilton (Respondent) before the Board of Pharmacy. (Accusation attached as Exhibit A.)

6           2.     On or about December 1, 2015, the Board of Pharmacy issued Pharmacy Permit  
7 Number PHY 53859 to Tilton Nguyen Pharmaceutical Inc. doing business as Oasis Pharmacy,  
8 with Pauline Minh Tilton as the President, 100% shareholder, Director, Secretary and  
9 Treasurer/Chief Financial Officer (“Respondent Pharmacy”). The Pharmacy Permit was in full  
10 force and effect at all times relevant to the charges brought herein but was cancelled on October  
11 10, 2019.

12           3.     On or about August 12, 1994, the Board of Pharmacy issued Pharmacist License  
13 Number RPH 47537 to Pauline Minh Tilton (“Respondent Tilton”). The Pharmacist License was  
14 in full force and effect at all times relevant to the charges brought herein and will expire on June  
15 30, 2020, unless renewed.

16           4.     On or about June 12, 2019, Respondents were served by Certified and First Class  
17 Mail copies of the Accusation No. 6608, Statement to Respondent, Notice of Defense, Request  
18 for Discovery, and Discovery Statutes (Government Code sections 11507.5, 11507.6, and  
19 11507.7) at Respondents’ address of record which, pursuant to Business and Professions Code  
20 section 4100, is required to be reported and maintained with the Board. Respondents’ address of  
21 record was and is: 15433 W. Sand St., Ste. 102, Victorville, CA 92392.

22           5.     Service of the Accusation was effective as a matter of law under the provisions of  
23 Government Code section 11505(c) and/or Business and Professions Code section 124.

24           6.     Government Code section 11506(c) states, in pertinent part:

25                   (c) The respondent shall be entitled to a hearing on the merits if the respondent  
26 files a notice of defense . . . and the notice shall be deemed a specific denial of all  
27 parts of the accusation . . . not expressly admitted. Failure to file a notice of defense  
28 . . . shall constitute a waiver of respondent's right to a hearing, but the agency in its  
discretion may nevertheless grant a hearing.



1 sections 4036.5 and 4113, Health and Safety Code section 11153, subdivision (a) and Code of  
2 Federal Regulations Title 21, section 1306.04, subdivision (a); and

3 b. Failure to Comply with Requirements for Controlled Substance Program - Code  
4 section 4301, subdivisions (d), (j) and/or (o) and CCR, title 16, section 1761, subdivisions (a) and  
5 (b), in conjunction with Code sections 4036.5 and 4113, Health and Safety Code sections 11164  
6 and 11162.1, and Code of Federal Regulations Title 21, section 1306.04, subdivision (a).

7 **ORDER**

8 IT IS SO ORDERED that Pharmacy Permit No. PHY 53859, issued to Tilton Nguyen  
9 Pharmaceutical Inc. doing business as Oasis Pharmacy, with Pauline Minh Tilton as the  
10 President, 100% shareholder, Director, Secretary and Treasurer/Chief Financial Officer, is  
11 revoked.

12 IT IS SO ORDERED that Pharmacist License No. RPH 47537 issued to Respondent  
13 Pauline Minh Tilton, is revoked.

14 Pursuant to Government Code section 11520, subdivision (c), Respondents may serve a  
15 written motion requesting that the Decision be vacated and stating the grounds relied on within  
16 seven (7) days after service of the Decision on Respondents. The agency in its discretion may  
17 vacate the Decision and grant a hearing on a showing of good cause, as defined in the statute.

18 This Decision shall become effective on January 23, 2020 at 5:00 p.m.

19 It is so ORDERED on December 24, 2019.

20  
21 BOARD OF PHARMACY  
22 DEPARTMENT OF CONSUMER AFFAIRS  
23 STATE OF CALIFORNIA

24 By 

24 53829737.DOCX  
25 DOJ Matter ID:LA2018603001

25 Attachment:  
26 Exhibit A: Accusation

25 Greg Lippe  
26 Board President

# Exhibit A

Accusation

(TILTON NGUYEN PHARMACEUTICAL INC. DBA OASIS PHARMACY and PAULINE MINH TILTON)

1 XAVIER BECERRA  
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2 LINDA K. SCHNEIDER  
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*Attorneys for Complainant*  
7

8 **BEFORE THE**  
9 **BOARD OF PHARMACY**  
10 **DEPARTMENT OF CONSUMER AFFAIRS**  
11 **STATE OF CALIFORNIA**

12 In the Matter of the Accusation Against:

Case No. 6608

13 **TILTON NGUYEN PHARMACEUTICAL**  
14 **INC. DBA OASIS PHARMACY, PAULINE**  
15 **MINH TILTON, PRESIDENT**  
15433 W. Sand St., Ste. 102  
Victorville, CA 92392

**ACCUSATION**

16 **Pharmacy Permit No. PHY 53859,**

17 **and**

18 **PAULINE MINH TILTON**  
15433 W. Sand St., Ste. 102  
Victorville, CA 92392

19 **Pharmacist License No. RPH 47537**

20 Respondents.  
21  
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23

24 Complainant alleges:

25 **PARTIES**

26 1. Anne Sodergren ("Complainant") brings this Accusation solely in her official  
27 capacity as the Interim Executive Officer of the Board of Pharmacy, Department of Consumer  
28 Affairs.



1 (b) Any device that bears the statement: 'Caution: federal law restricts this device to sale by  
2 or on the order of a \_\_\_\_\_,' 'Rx only,' or words of similar import, the blank to be filled in  
3 with the designation of the practitioner licensed to use or order use of the device.

4 (c) Any other drug or device that by federal or state law can be lawfully dispensed only on  
5 prescription or furnished pursuant to Section 4006."

6 8. Code section 4036.5 states that "'Pharmacist-in-charge' means a pharmacist proposed  
7 by a pharmacy and approved by the board as the supervisor or manager responsible for ensuring  
8 the pharmacy's compliance with all state and federal laws and regulations pertaining to the  
9 practice of pharmacy."

10 9. Code section 4059, subdivision (a), in pertinent part, prohibits furnishing of any  
11 dangerous drug or dangerous device except upon the prescription of an authorized prescriber.

12 10. Code section 4113 states, in pertinent part, that: "(c) The pharmacist-in-charge shall  
13 be responsible for a pharmacy's compliance with all state and federal laws and regulations  
14 pertaining to the practice of pharmacy."

15 11. Code section 4301 provides, in pertinent part:

16 "The board shall take action against any holder of a license who is guilty of  
17 unprofessional conduct or whose license has been issued by mistake. Unprofessional conduct  
18 shall include, but is not limited to, any of the following:

19 ...

20 (d) The clearly excessive furnishing of controlled substances in violation of subdivision (a)  
21 of Section 11153 of the Health and Safety Code.

22 ...

23 (j) The violation of any of the statutes of this state, of any other state, or of the United  
24 States regulating controlled substances and dangerous drugs.

25 ...

26 (o) Violating or attempting to violate, directly or indirectly, or assisting in or abetting the  
27 violation of or conspiring to violate any provision or term of this chapter or of the applicable  
28

1 federal and state laws and regulations governing pharmacy, including regulations established by  
2 the board or by any other state or federal regulatory agency.”

3 12. Section 4306.5 states:

4 “Unprofessional conduct for a pharmacist may include any of the following:

5 (a) Acts or omissions that involve, in whole or in part, the inappropriate exercise of his or  
6 her education, training, or experience as a pharmacist, whether or not the act or omission arises in  
7 the course of the practice of pharmacy or the ownership, management, administration, or  
8 operation of a pharmacy or other entity licensed by the board.

9 (b) Acts or omissions that involve, in whole or in part, the failure to exercise or implement  
10 his or her best professional judgment or corresponding responsibility with regard to the  
11 dispensing or furnishing of controlled substances, dangerous drugs, or dangerous devices, or with  
12 regard to the provision of services.

13 (c) Acts or omissions that involve, in whole or in part, the failure to consult appropriate  
14 patient, prescription, and other records pertaining to the performance of any pharmacy function.

15 (d) Acts or omissions that involve, in whole or in part, the failure to fully maintain and  
16 retain appropriate patient-specific information pertaining to the performance of any pharmacy  
17 function.”

18 13. Section 4307 states, in pertinent part:

19 “(a) Any person who has been denied a license or whose license has been revoked or is  
20 under suspension, or who has failed to renew his or her license while it was under suspension, or  
21 who has been a manager, administrator, owner, member, officer, director, associate, or partner of  
22 any partnership, corporation, firm, or association whose application for a license has been denied  
23 or revoked, is under suspension or has been placed on probation, and while acting as the manager,  
24 administrator, owner, member, officer, director, associate, or partner had knowledge of or  
25 knowingly participated in any conduct for which the license was denied, revoked, suspended, or  
26 placed on probation, shall be prohibited from serving as a manager, administrator, owner,  
27 member, officer, director, associate, or partner of a licensee as follows:

28 ///

1 (1) Where a probationary license is issued or where an existing license is placed on  
2 probation, this prohibition shall remain in effect for a period not to exceed five years.

3 (2) Where the license is denied or revoked, the prohibition shall continue until the license is  
4 issued or reinstated.”

5 14. Health and Safety Code section 11153 states, in pertinent part:

6 “(a) A prescription for a controlled substance shall only be issued for a legitimate medical  
7 purpose by an individual practitioner acting in the usual course of his or her professional practice.  
8 The responsibility for the proper prescribing and dispensing of controlled substances is upon the  
9 prescribing practitioner, but a corresponding responsibility rests with the pharmacist who fills the  
10 prescription. Except as authorized by this division, the following are not legal prescriptions: (1)  
11 an order purporting to be a prescription which is issued not in the usual course of professional  
12 treatment or in legitimate and authorized research; or (2) an order for an addict or habitual user of  
13 controlled substances, which is issued not in the course of professional treatment or as part of an  
14 authorized narcotic treatment program, for the purpose of providing the user with controlled  
15 substances, sufficient to keep him or her comfortable by maintaining customary use.”

16 15. Health and Safety Code section 11162.1 states:

17 “(a) The prescription forms for controlled substances shall be printed with the following  
18 features:

19 (1) A latent, repetitive “void” pattern shall be printed across the entire front of the  
20 prescription blank; if a prescription is scanned or photocopied, the word “void” shall appear in a  
21 pattern across the entire front of the prescription.

22 (2) A watermark shall be printed on the backside of the prescription blank; the watermark  
23 shall consist of the words “California Security Prescription.”

24 (3) A chemical void protection that prevents alteration by chemical washing.

25 (4) A feature printed in thermochromic ink.

26 (5) An area of opaque writing so that the writing disappears if the prescription is lightened.

27 (6) A description of the security features included on each prescription form.

28 ///

1 (7)(A) Six quantity check off boxes shall be printed on the form so that the prescriber may  
2 indicate the quantity by checking the applicable box where the following quantities shall appear:

3 1-24

4 25-49

5 50-74

6 75-100

7 101-150

8 151 and over.

9 (B) In conjunction with the quantity boxes, a space shall be provided to designate the units  
10 referenced in the quantity boxes when the drug is not in tablet or capsule form.

11 (8) Prescription blanks shall contain a statement printed on the bottom of the prescription  
12 blank that the "Prescription is void if the number of drugs prescribed is not noted."

13 (9) The preprinted name, category of licensure, license number, federal controlled  
14 substance registration number, and address of the prescribing practitioner.

15 (10) Check boxes shall be printed on the form so that the prescriber may indicate the  
16 number of refills ordered.

17 (11) The date of origin of the prescription.

18 (12) A check box indicating the prescriber's order not to substitute.

19 (13) An identifying number assigned to the approved security printer by the Department of  
20 Justice.

21 (14)(A) A check box by the name of each prescriber when a prescription form lists multiple  
22 prescribers.

23 (B) Each prescriber who signs the prescription form shall identify himself or herself as the  
24 prescriber by checking the box by his or her name.

25 (15) A uniquely serialized number, in a manner prescribed by the Department of Justice.

26 (b) Each batch of controlled substance prescription forms shall have the lot number printed  
27 on the form and each form within that batch shall be numbered sequentially beginning with the  
28 numeral one.

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...”

16. Health and Safety Code section 11164 states in pertinent part:

“Except as provided in Section 11167, no person shall prescribe a controlled substance, nor shall any person fill, compound, or dispense a prescription for a controlled substance, unless it complies with the requirements of this section.

(a) Each prescription for a controlled substance classified in Schedule II, III, IV, or V, except as authorized by subdivision (b), shall be made on a controlled substance prescription form as specified in Section 11162.1 . . . .”

**REGULATORY PROVISIONS**

17. California Code of Regulations, title 16, section 1761, states:

“(a) No pharmacist shall compound or dispense any prescription which contains any significant error, omission, irregularity, uncertainty, ambiguity or alteration. Upon receipt of any such prescription, the pharmacist shall contact the prescriber to obtain the information needed to validate the prescription.

(b) Even after conferring with the prescriber, a pharmacist shall not compound or dispense a controlled substance prescription where the pharmacist knows or has objective reason to know that said prescription was not issued for a legitimate medical purpose.”

**CODE OF FEDERAL REGULATIONS**

18. Code of Federal Regulations, Title 21, section 1306.04, subdivision (a), states:

“A prescription for a controlled substance to be effective must be issued for a legitimate medical purpose by an individual practitioner in the usual course of his professional practice. The responsibility for the proper prescribing and dispensing of controlled substances is upon the prescribing practitioner, but a corresponding responsibility rests with the pharmacist who fills the prescription. An order purporting to be a prescription issued not in the usual course of professional treatment or in legitimate and authorized research is not a prescription within the meaning and intent of section 309 of the Act (21 U.S.C. 829) and the person knowingly filling such a purported prescription as well as the person issuing it, shall be subject to the penalties provided for violations of the provisions of law relating to controlled substances.”

1 **COST RECOVERY**

2 19. Section 125.3 of the Code states, in pertinent part, that the Board may request the  
3 administrative law judge to direct a licentiate found to have committed a violation or violations of  
4 the licensing act to pay a sum not to exceed the reasonable costs of the investigation and  
5 enforcement of the case.

6 **CONTROLLED SUBSTANCES/DANGEROUS DRUGS**

7 20. Oxycodone 30 mg, under the brand name Roxicodone 30 mg, is a dangerous drug  
8 pursuant to Business and Professions Code section 4022 and a Schedule II controlled substance  
9 pursuant to Health and Safety Code section 11055, subdivision (b)(1)(M). It is used to treat pain.

10 21. Alprazolam 2 mg, under the brand name Xanax 2 mg, is a dangerous drug pursuant to  
11 Business and Professions Code section 4022 and a Schedule IV controlled substance pursuant to  
12 Health and Safety Code section 11057, subdivision (d)(1). It is used to treat anxiety.

13 22. Promethazine with codeine syrup, under the brand name Phenergan with Codeine  
14 Syrup, is a dangerous drug pursuant to Business and Professions Code section 4022 and a  
15 Schedule V controlled substance pursuant to Health and Safety Code section 11058, subdivision  
16 (c)(1). Promethazine with codeine syrup is a prescription cough syrup.

17 **FACTUAL BACKGROUND**

18 23. The Board analyzed Controlled Substance Utilization Review and Evaluation  
19 System (CURES)<sup>1</sup> data from Respondent Pharmacy and determined a need for an investigation  
20 to evaluate Respondent Pharmacy's controlled substance dispensing practices. The investigation  
21 determined Respondent Pharmacy and Respondent Tilton, while acting as pharmacist-in-charge,  
22 failed to fulfill their corresponding responsibility in dispensing prescriptions under the prescribing  
23 authority of Dr. Kevin M. Smith of Orange, California ("Dr. Smith").

24 24. The Board's investigation determined that between August 16, 2016 and June 30,  
25 2017, Respondent Pharmacy dispensed 1,026 prescriptions under the prescribing authority of Dr.

26 \_\_\_\_\_  
27 <sup>1</sup> The Controlled Substance Utilization Review and Evaluation System (CURES) is California's  
28 Prescription Drug Monitoring Program (PDMP) which requires mandatory weekly reporting of dispensed  
Schedule II-IV (CII-IV) controlled substances prescriptions across the state. It can be used by healthcare  
professionals to evaluate and determine whether patients are utilizing controlled substances correctly.

1 Smith in the presence of numerous factors indicating the prescriptions were not written for  
2 legitimate medical purposes. Factors of irregularity related to Dr. Smith's prescriptions included  
3 the following:

- 4 a. Dr. Smith's prescriptions at Respondent Pharmacy included only three commonly  
5 abused controlled substances: oxycodone 30 mg, alprazolam 2 mg, and  
6 promethazine/codeine syrup.
- 7 b. Over 99% of Dr. Smith's prescriptions at Respondent Pharmacy were purchased  
8 in cash, without the aid of prescription insurance.
- 9 c. Many of Dr. Smith's patients paid a very high price, over \$600, for 180  
10 oxycodone 30 mg tablets.
- 11 d. Dr. Smith's office was located over 77 miles away from Respondent Pharmacy.
- 12 e. Dr. Smith exclusively prescribed the highest strength of immediate release  
13 oxycodone and alprazolam.
- 14 f. All of Dr. Smith's patients received at least one prescription for oxycodone 30  
15 mg, alprazolam 2 mg, and promethazine/codeine syrup.
- 16 g. Respondent Pharmacy processed large numbers of Dr. Smith's prescriptions on  
17 the same days and frequently assigned Dr. Smith's prescriptions consecutive  
18 prescription numbers. Additionally, many of Dr. Smith's prescriptions were  
19 found filed consecutively in Respondent Pharmacy's prescription document files.
- 20 h. There were several instances when Dr. Smith prescribed multiple identical sets of  
21 prescriptions to a single patient in less than one month.
- 22 i. There were instances when Dr. Smith's patients obtained controlled substance  
23 prescriptions from Respondent Pharmacy when a previously dispensed supply of  
24 medication should not have been exhausted.
- 25 j. There were numerous instances when Respondent Pharmacy dispensed  
26 promethazine/codeine syrup prescriptions to patients of Dr. Smith more than one  
27 month after the prescriptions were written.

28 ///

- 1 k. At least 27 patients of Dr. Smith received prescriptions for promethazine/codeine  
2 syrup in four or more consecutive months, even though promethazine/codeine  
3 syrup is indicated for short term use.
- 4 l. The drug name codeine was misspelled as “codein” on Dr. Smith’s prescriptions.
- 5 m. Dr. Smith’s prescription documents did not conform to the requirements of Health  
6 and Safety Code section 11162.1 in that they lacked a lot number, a batch  
7 number, and a “California Security Prescription” watermark.

8 25. Despite these irregularities, there were no notations or documentation on the  
9 prescription documents indicating Respondent Tilton contacted Dr. Smith and attempted to  
10 validate the prescriptions or resolve any of the irregularities listed above.

11 **FIRST CAUSE FOR DISCIPLINE**

12 **(Violation of Corresponding Responsibility)**

13 26. Respondent Pharmacy and Respondent Tilton are subject to disciplinary action under  
14 Code section 4301, subdivisions (d), (j) and/or (o) and CCR, title 16, section 1761, subdivisions  
15 (a) and (b), in conjunction with Code sections 4036.5 and 4113, Health and Safety Code section  
16 11153, subdivision (a) and Code of Federal Regulations Title 21, section 1306.04, subdivision  
17 (a), in that they violated their corresponding responsibility by excessively furnishing controlled  
18 substances and repeatedly failing to resolve irregularities and red flags of illegitimacy for  
19 controlled substances prescribed by Dr. Smith. Between August 16, 2016 and June 30, 2017,  
20 Respondent Tilton, while acting as pharmacist-in-charge, and Respondent Pharmacy dispensed  
21 1,026 prescriptions under the prescribing authority of Dr. Smith in the presence of numerous  
22 objective factors indicating the prescriptions were not written for legitimate medical purposes.  
23 Complainant incorporates by reference paragraphs 23 through 25 as though fully set forth herein.

24 **SECOND CAUSE FOR DISCIPLINE**

25 **(Failure to Comply with Requirements for Controlled Substance Prescriptions)**

26 27. Respondent Pharmacy and Respondent Tilton are subject to disciplinary action under  
27 Code section 4301, subdivisions (d), (j) and/or (o) and CCR, title 16, section 1761, subdivisions  
28 (a) and (b), in conjunction with Code sections 4036.5 and 4113, Health and Safety Code sections

1 11164 and 11162.1, and Code of Federal Regulations Title 21, section 1306.04, subdivision (a),  
2 in that Respondent Pharmacy and Respondent Tilton, while acting as pharmacist-in-charge,  
3 dispensed at least 1,032 controlled substance prescriptions written on 344 prescription documents  
4 that did not conform to the requirements of Health and Safety Code section 11162.1 in that they  
5 lacked a lot number, a batch number, and a "California Security Prescription" watermark.  
6 Complainant incorporates by reference paragraphs 23 through 25 as though fully set forth herein.

7 **OTHER MATTERS**

8 34. Pursuant to section 4307, if discipline is imposed on Pharmacy Permit Number PHY  
9 53859 to Tilton Nguyen Pharmaceutical Inc. doing business as Oasis Pharmacy, with Pauline  
10 Minh Tilton as the President, 100% shareholder, Director, Secretary and Treasurer/Chief  
11 Financial Officer, Tilton Nguyen Pharmaceutical Inc. shall be prohibited from serving as a  
12 manager, administrator, owner, member, officer, director, associate, or partner of a licensee for  
13 five years if Pharmacy Permit Number PHY 53859 is placed on probation or until Pharmacy  
14 Permit Number PHY 53859 is reinstated if the license is revoked.

15 35. Pursuant to section 4307, if discipline is imposed on Pharmacy Permit Number PHY  
16 53859 issued to Tilton Nguyen Pharmaceutical Inc. doing business as Oasis Pharmacy, with  
17 Pauline Minh Tilton as the President, 100% shareholder, Director, Secretary and Treasurer/Chief  
18 Financial Officer, while Pauline Minh Tilton has been an officer or owner, and had knowledge of,  
19 or knowingly participated in, any conduct for which Tilton Nguyen Pharmaceutical Inc. was  
20 disciplined, Pauline Minh Tilton shall be prohibited from serving as a manager, administrator,  
21 owner, member, officer, director, associate, or partner of a licensee for five years if Pharmacy  
22 Permit Number PHY 53859 is placed on probation or until Pharmacy Permit Number PHY 53859  
23 is reinstated if the license is revoked.

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PRAYER

WHEREFORE, Complainant requests that a hearing be held on the matters herein alleged, and that following the hearing, the Board of Pharmacy issue a decision:

1. Revoking or suspending Pharmacy Permit Number PHY 53859, issued to Tilton Nguyen Pharmaceutical Inc. doing business as Oasis Pharmacy;
2. Prohibiting Tilton Nguyen Pharmaceutical Inc. from serving as a manager, administrator, owner, member, officer, director, associate, or partner of a licensee for five years if Pharmacy Permit Number PHY 53859 is placed on probation or until Pharmacy Permit Number PHY 53859 is reinstated if Pharmacy Permit Number PHY 53859 issued to Tilton Nguyen Pharmaceutical Inc., doing business as Oasis Pharmacy, is revoked;
3. Prohibiting Pauline Minh Tilton from serving as a manager, administrator, owner, member, officer, director, associate, or partner of a licensee for five years if Pharmacy Permit Number PHY 53859 is placed on probation or until Pharmacy Permit Number PHY 53859 is reinstated if Pharmacy Permit Number PHY 53859 issued to Tilton Nguyen Pharmaceutical Inc., doing business as Oasis Pharmacy, is revoked;
4. Revoking or suspending Pharmacist License Number RPH 47537 to Pauline Minh Tilton;
5. Ordering Tilton Nguyen Pharmaceutical Inc., doing business as Oasis Pharmacy, and Pauline Minh Tilton to pay the Board of Pharmacy the reasonable costs of the investigation and enforcement of this case, pursuant to Business and Professions Code section 125.3; and
6. Taking such other and further action as deemed necessary and proper.

DATED: June 7, 2019



ANNE SODERGREN  
Interim Executive Officer  
Board of Pharmacy  
Department of Consumer Affairs  
State of California  
*Complainant*

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