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9	BEFORE THE BOARD OF PHARMACY
10	DEPARTMENT OF CONSUMER AFFAIRS STATE OF CALIFORNIA
11	STATE OF CALIFORNIA
12	In the Matter of the Accusation Against: Case No. 3488
13	KULDEEP KAUR GREWAL 13313 Glencliff Way A C C U S A T I O N
14	San Diego, CA 92130
15	Pharmacist License No. RPH 40706
16	Respondent.
17	
18	Complainant alleges:
19	PARTIES
20	1. Complainant Virginia Herold brings this Accusation solely in her official capacity as
21	the Executive Officer of the Board of Pharmacy (Board), Department of Consumer Affairs.
22	2. On May 2, 1987, the Board issued Pharmacist License Number RPH 40706 to
23	Respondent Kuldeep Kaur Grewal. The License was in full force and effect at all times relevant
24	to the charges brought herein and will expire on September 30, 2010, unless renewed.
25	JURISDICTION
26	3. This Accusation is brought before the Board, Department of Consumer Affairs, under
27	the authority of the following laws. All section references are to the Business and Professions
28	Code unless otherwise indicated.
	1

- 4. Section 4300, subdivision (a) of the Business and Professions Code (Code) provides, in pertinent part, that every license issued may be suspended or revoked.
- 5. Section 118, subdivision (b), of the Code provides that the suspension, expiration, surrender, or cancellation of a license shall not deprive the Board of jurisdiction to proceed with a disciplinary action during the period within which the license may be renewed, restored, reissued or reinstated.

STATUTORY PROVISIONS

- 6. Section 480 of the Code states, in pertinent part:
- (a) A board may deny a license regulated by this code on the grounds that the applicant has one of the following:
- (3)(A) Done any act that if done by a licentiate of the business or profession in question, would be grounds for suspension or revocation of license.
- (B) The board may deny a license pursuant to this subdivision only if the crime or act is substantially related to the qualifications, functions, or duties of the business or profession for which application is made.
- 7. Section 4022 of the Code states:

"Dangerous drug" or "dangerous device" means any drug or device unsafe for self-use in humans or animals, and includes the following:

- (a) Any drug that bears the legend: "Caution: federal law prohibits dispensing without prescription," "Rx only," or words of similar import.
- (b) Any device that bears the statement: "Caution: federal law restricts this device to sale by or on the order of a _____," "Rx only," or words of similar import, the blank to be filled in with the designation of the practitioner licensed to use or order use of the device.
- (c) Any other drug or device that by federal or state law can be lawfully dispensed only on prescription or furnished pursuant to Section 4006.
- 8. Section 4081 of the Code states in pertinent part:
- (a) All records of manufacture and of sale, acquisition, or disposition of dangerous drugs or dangerous devices shall be at all times during business hours open to inspection by authorized officers of the law, and shall be preserved for at least three years from the date of making. A current inventory shall be kept by every . . . pharmacy . . . holding a currently valid and unrevoked certificate, license, permit, registration, or exemption under Division 2 (commencing with Section 1200) of the Health and Safety Code or under Part 4 (commencing with Section 16000) of Division 9 of the Welfare and Institutions Code who maintains a stock of dangerous drugs or dangerous devices.

1	11. Section 4301 of the Code states, in pertinent part:
unprofessional conduct or whose license has been procured by fraud or	The board shall take action against any holder of a license who is guilty of
	misrepresentation or issued by mistake. Unprofessional conduct shall include, but is
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6	(o) Violating or attempting to violate, directly or indirectly, or assisting in or abetting the violation of or conspiring to violate any provision or term of this chapter
7	or of the applicable federal and state laws and regulations governing pharmacy, including regulations established by the board or by any other state or federal
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10	REGULATORY PROVISIONS
11	12. California Code of Regulations, title 16 (Regulations), section 1714 states in pertinent
12	part:
13	••••
14	(b) Each pharmacy licensed by the board shall maintain its facilities, space, fixtures, and equipment so that drugs are safely and properly prepared, maintained,
15	secured and distributed. The pharmacy shall be of sufficient size and unobstructed area to accommodate the safe practice of pharmacy.
16	••••
	(d) Each pharmacist while on duty shall be responsible for the security of the
18	prescription department, including provisions for effective control against theft or diversion of dangerous drugs and devices, and records for such drugs and devices.
19	Possession of a key to the pharmacy where dangerous drugs and controlled substances are stored shall be restricted to a pharmacist.
20	13. Regulations, section 1718 states:
21	"Current Inventory" as used in Sections 4081 and 4332 of the Business and
22	Professions Code shall be considered to include complete accountability for all dangerous drugs handled by every licensee enumerated in Sections 4081 and 4332.
The controlled substances inventories required by Title 21, CFR, Section 1304	The controlled substances inventories required by Title 21, CFR, Section 1304 shall be available for inspection upon request for at least 3 years after the date of the
24	inventory.
25	COST RECOVERY
26	14. Section 125.3 of the Code states, in pertinent part, that the Board may request the
27	administrative law judge to direct a licentiate found to have committed a violation or violations of
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the licensing act to pay a sum not to exceed the reasonable costs of the investigation and enforcement of the case.

FACTS

- 15. On March 6, 2008, Drug Enforcement Administration (DEA) and Federal Bureau of Investigation (FBI) agents simultaneously served federal search warrants at White Cross Pharmacy, Park Blvd. Pharmacy, and Galloway Pharmacy in San Diego after a two-year investigation into controlled substance diversion from the three pharmacies. From June 14, 2005, through September 17, 2007, Respondent was Pharmacist-in-Charge (PIC) at White Cross Pharmacy, and from September 17, 2007, through March 6, 2008, PIC at Park Blvd. Pharmacy.
 - 16. As to White Cross Pharmacy, the warrants produced the following information:
- a. Printouts of the tabulations and summary of White Cross's purchases of Hydrocodone 5/500, Hydrocodone 10/325, and Oxycodone 80, show White Cross purchased 600,800 tablets of Hydrocodone 5/500; 139,700 tablets of Hydrocodone 10/325; and 27,100 tablets of Oxycodone 80 from February 7, 2005, through March 6, 2008.
- b. The DEA Biennial Inventory of February 7, 2005 shows 980 tablets of Oxycodone 80; 1500 tablets of Hydrocodone 5/500; and 1,640 tablets of Hydrocodone 10/325; and a closing inventory (stock on hand) of March 6, 2008, show 1,165 tablets of Oxycodone 80; 4,495 tablets of Hydrocodone 5/500; and 3,819 tablets of Hydrocodone 10/325.
- c. From February 7, 2005, through March 6, 2008, White Cross dispensed 23,068 tablets of Oxycodone 80; 309,293 tablets of Hydrocodone 5/500; and 45,247 tablets of Hydrocodone 10/325.

Together, this information revealed that White Cross was short 2,341 (8%) tablets of Oxycodone 80mg.; short 49,126 (8%) tablets of Hydrocodone 5/500; and short 51,263 (38%) of Hydrocodone 10/325.

- 17. As to Park Blvd. Pharmacy, the warrants produced the following information:
- a. Printouts of the tabulations and summary of Park's purchases of Hydrocodone 5/500, Hydrocodone 10/325 and Oxycodone 80, showing Park purchased 634,200 tablets of