

1 **BEFORE THE BOARD OF MEDICAL EXAMINERS**  
2 **OF THE STATE OF NEVADA**

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6 **In the Matter of Charges and**  
7 **Complaint Against**  
8 **Michael S. Mall, M.D.,**  
9 **Respondent.**

Case No. 15-8666-1

**FILED**  
SEP 23 2015  
NEVADA STATE BOARD OF  
MEDICAL EXAMINERS  
By: \_\_\_\_\_

10  
11 **COMPLAINT**

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13 The Investigative Committee (IC) of the Nevada State Board of Medical Examiners (Board)  
14 hereby issues this formal Complaint (Complaint) against Michael S. Mall, M.D. (Respondent), a  
15 licensed physician in Nevada. After investigating this matter, the IC has a reasonable basis to  
16 believe that Respondent has violated provisions of Nevada Revised Statutes (NRS) chapter 630 and  
17 Nevada Administrative Code (NAC) chapter 630 (collectively Medical Practice Act). The IC  
18 alleges the following facts:

19 1. Respondent is currently licensed in active status (License No. 6074), and has been  
20 licensed by the Board since July 1, 1990 pursuant to the provisions of the Medical Practice Act.

21 2. Respondent's specialty listed with the Board is family practice.

22 3. At the time of the incidents alleged herein, Patient A was being treated by  
23 Respondent for pain management. His true identity is not disclosed in this Complaint to protect  
24 his identity, but his identity is disclosed in the Patient Designation served on Respondent along  
25 with a copy of this Complaint.

26 4. Patient A first presented to Respondent on or about May 2002 with complaints of  
27 pain relating to Fibromyalgia.

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1           5.       On or about July 19, 2002, Patient A presented to Respondent with complaints of  
2 pain. Respondent increased Patient A's Oxycontin 40-milligram (mg) prescription from one twice  
3 a day to two twice a day and prescribed five mg of extended release Oxycodone. The medical  
4 records from this visit are illegible.

5           6.       On or about August 16, 2002, Patient A presented to respondent with complaints of  
6 pain. Respondent increased Patient A's Oxycontin 40-mg prescription from two twice a day to  
7 three twice a day. The medical records from this visit are illegible.

8           7.       On or about November 5, 2004, Respondent prescribed Patient A a Duragesic  
9 patch, which is the transdermal delivery method for Fentanyl, and Actiq, which is the oral delivery  
10 method for Fentanyl. The medical records from this visit are illegible.

11          8.       On July 5, 2005, Respondent increased Patient A's Actiq medication from 800  
12 micrograms (mcg) to 1,200 mcg. The medical records from this visit are illegible.

13          9.       On or about July 11, 2006, Patient A's medical records demonstrate that Patient A  
14 was taking four different benzodiazepines-Xanax, Valium, Restoril and Klonopin. On this same  
15 date, Respondent increased Patient A's Actiq medication from 1,200 mcg to the maximum  
16 allowed dose of 1,600 mcg. The medical records from this visit are illegible. There is no medical  
17 rationale for this daily combination of four benzodiazepines.

18          10.      On or about December 12, 2006, Respondent prescribed Patient A 800 mcg of  
19 Fentora, which is the highest dose available for Fentora. Fentora is another mouth-absorbing  
20 Fentanyl product. Respondent also prescribed Patient A 1,600-mcg of Actiq, which is the highest  
21 dose available for Actiq. He also prescribed Patient A two 75-mcg Duragesics and four different  
22 benzodiazepines. Thus, at this visit, Respondent prescribed Patient A the maximum allowable  
23 dose for two different Fentanyl medications, a third Fentanyl medication, and four different  
24 benzodiazepine medications. The medical records from this visit are illegible.

25          11.      Respondent's medical records for Patient A lack documentation regarding the  
26 etiology of the patient's pain, the nature and intensity of the patient's pain, the effect of pain on  
27 the patient's ability to function, the presence of recognized medical indications for the use of

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1 controlled substances, and the fact that other treatment modalities or adjuvant therapies were  
2 considered, tried and failed prior to prescribing controlled substances.

3 12. Respondent's medical records for Patient A lack documentation demonstrating that  
4 Respondent checked Patient A's prescription monitoring profile with the Nevada State Board of  
5 Pharmacy (BOP) before prescribing controlled substances and that Patient A underwent urine  
6 drug screens.

7 13. Respondent failed to refer Patient A to a pain management specialist.

8 14. Respondent's medical records for Patient A lack pain management contracts signed  
9 by the patient, drug profiles, urine toxicology screens, complete physical examinations, complete  
10 motor or neurological exams to assess for nerve root involvement, diagnostic tests, outside  
11 consultations with pain management specialists, and additional examinations/testing to determine  
12 the etiology of the alleged pain.

13 **COUNT I**  
14 **(Medical Records Violation)**

15 15. All of the allegations in the above paragraphs are hereby incorporated as if fully set  
16 forth herein.

17 16. NRS 630.3062(1) provides that the failure to maintain timely, legible, accurate and  
18 complete medical records relating to the diagnosis, treatment and care of a patient is grounds for  
19 initiating discipline against a licensee.

20 17. As demonstrated by, but not limited to, the above-outlined facts, Respondent failed  
21 to maintain accurate and/or complete medical records relating to the diagnosis, treatment and care  
22 of Patient A when he failed to document the nature and intensity of the Patient's pain, the effect of  
23 pain on the Patient's ability to function, and the presence of recognized medical indications for the  
24 use of controlled substances.

25 18. As demonstrated by, but not limited to, the above-outlined facts, Respondent failed  
26 to maintain accurate and/or complete medical records relating to the diagnosis, treatment and care  
27 of Patient A when he failed to document that other treatment modalities or adjuvant therapies had  
28 been considered.

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**COUNT III**

**(Unlawful Administration, Dispensing or Prescribing of Controlled Substances)**

27. All of the allegations contained in the above paragraphs are hereby incorporated by reference as though fully set forth herein.

28. NRS 630.306(3) provides that administering, dispensing or prescribing any controlled substance, or any dangerous drug, to others except as authorized by law is grounds for initiating discipline against a licensee.

29. As demonstrated by, but not limited to, the above-outlined facts, Respondent prescribed controlled substances in violation of the law when he failed to determine the etiology of pain for Patient A; failed to check Patient A's prescription monitoring profile with the BOP prior to prescribing Patient A controlled substances; failed to treat the nonmalignant pain of Patient A with other modalities and/or adjuvant therapies prior to prescribing controlled substances; failed to perform urine drug testing on Patient A, and failed to follow the policies set forth in the *Model Policy on the Use of Opioid Analgesics in the Treatment of Chronic Pain* (Model Policy), adopted by reference in NAC 630.18, when prescribing controlled substances to Patient A.

30. By reason of the foregoing, Respondent is subject to discipline by the Board as provided in NRS 630.352.

**COUNT IV**

**(Practicing Beyond the Scope of Training)**

31. All of the allegations contained in the above paragraphs are hereby incorporated by reference as though fully set forth herein.

32. NRS 630.306(5) provides that practicing or offering to practice beyond the scope permitted by law or performing services that the licensee knows or has reason to know that he is not competent to perform or which are beyond the scope of his or her training is grounds for initiating disciplinary action against a licensee.

33. Respondent's specialty is family medicine, not pain management. As demonstrated by, but not limited to, the above-outlined facts, Respondent was prescribing controlled substances to Patient A without first determining the etiology of Patient A's pain and considering other modalities and/or adjuvant therapies prior to prescribing Patient A controlled

1 substances. Further, as demonstrated by, but not limited to, the above-outlined facts, Respondent  
2 was practicing beyond the scope permitted by law and/or was performing services that were  
3 beyond the scope of his training.

4 34. By reason of the foregoing, Respondent is subject to discipline by the Board as  
5 provided in NRS 630.352.

6 **COUNT V**  
7 **(Continual Failure to Exercise the Skill or Diligence or Use the Methods Exercised**  
8 **by Physicians in the Same Specialty or Field)**

9 35. All of the allegations contained in the above paragraphs are hereby incorporated by  
10 reference as though fully set forth herein.

11 36. NRS 630.306(7) provides that the continual failure to exercise the skill or diligence  
12 or use the methods ordinarily exercised under the same circumstances by physicians in good  
13 standing practicing in the same specialty or field is grounds for initiating discipline against a  
14 licensee.

15 37. Respondent's specialty is family medicine, not pain management. As  
16 demonstrated by, but not limited to, the above-outlined facts, Respondent failed to exercise the  
17 skill or diligence or use the methods exercised by physicians in the same specialty or field when  
18 he prescribed controlled substances to Patient A without first checking Patient A's prescription  
19 monitoring profile with the BOP prior to prescribing Patient A controlled substances, without  
20 determining the etiology of Patient A's pain, without considering other modalities and/or adjuvant  
21 therapies prior to prescribing Patient A controlled substances and without referring Patient A to a  
22 pain management specialist.

23 38. By reason of the foregoing, Respondent is subject to discipline by the Board as  
24 provided in NRS 630.352.

25 **COUNT VI**  
26 **(Conduct That Violates the Standards of Practice)**

27 39. All of the allegations contained in the above paragraphs are hereby incorporated by  
28 reference as though fully set forth herein.

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1           40.    NRS 630.306(2)(b) provides that if the licensee engages in any conduct that is a  
2 violation of the standards of practice established by regulation of the Board it is grounds for  
3 initiating disciplinary action against the licensee.

4           41.    NAC 630.185 provides that NAC 630.185 to 630.230, inclusive, set forth the  
5 standards of practice established by the Board.

6           42.    NAC 630.230(k) provides that a licensee shall not engage in the practice of writing  
7 prescriptions for controlled substances to treat acute pain or chronic pain in a manner that deviates  
8 from the policies set forth in the Model Policy.

9           43.    As demonstrated by, but not limited to, the above-outlined facts, Respondent  
10 prescribed controlled substances in violation of the law when he failed to determine the etiology  
11 of pain for Patient A, failed to check Patient A's prescription monitoring profile with the BOP  
12 prior to prescribing Patient A controlled substances, failed to treat the nonmalignant pain of  
13 Patient A with other modalities and/or adjuvant therapies prior to prescribing controlled  
14 substances, and failed to perform urine drug testing on Patient A.

15           44.    By reason of the foregoing, Respondent is subject to discipline by the Board as  
16 provided in NRS 630.352.

17           **WHEREFORE**, the Investigative Committee prays:

18           1.    That the Board give Respondent notice of the charges herein against him and give  
19 him notice that he may file an answer to the Complaint herein as set forth in NRS 630.339(2)  
20 within twenty (20) days of service of the Complaint;

21           2.    That the Board set a time and place for a formal hearing after holding an  
22 Early Case Conference pursuant to NRS 630.339(3);

23           3.    That the Board determine the sanctions it will impose if it finds Respondent  
24 violated the Medical Practice Act;

25           4.    That the Board make, issue and serve on Respondent, in writing, its findings of  
26 fact, conclusions of law and order, which shall include the sanctions imposed; and

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OFFICE OF THE GENERAL COUNSEL


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5. That the Board take such other and further action as may be just and proper in these premises.

DATED this 23<sup>rd</sup> day of September, 2015.

INVESTIGATIVE COMMITTEE OF THE  
NEVADA STATE BOARD OF MEDICAL EXAMINERS

By:   
Erin L. Albright, Esq.  
General Counsel  
Attorney for the Investigative Committee

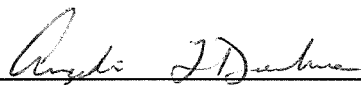


**CERTIFICATE OF MAILING**

I hereby certify that I am employed by Nevada State Board of Medical Examiners and that on 23<sup>rd</sup> day of September 2015; I served a file stamp copy of COMPLAINT, PATIENT DESIGNATION & FINGERPRINT INFORMATION via USPS e-certified return receipt mail to the following:

Michael Mall, M.D.  
c/o John Savage, Esq.  
7900 West Sahara Ste. 200  
Las Vegas, NV 89117

Dated this 23<sup>rd</sup> day of September, 2015.

  
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Angelia L. Donohoe  
Legal Assistant