

1 **BEFORE THE ARIZONA MEDICAL BOARD**

2 In the Matter of

3 **MONEIL M. PATEL, M.D.**

4 Holder of License No. **44593**
5 For the Practice of Allopathic Medicine
6 In the State of Arizona.

Case No. MD-20-1037A

**INTERIM CONSENT AGREEMENT
FOR PRACTICE RESTRICTION**

7 **INTERIM CONSENT AGREEMENT**

8 Moneil M. Patel, M.D. (“Respondent”) elects to permanently waive any right to a
9 hearing and appeal with respect to this Interim Consent Agreement for Practice Restriction
10 and consents to the entry of this Order by the Arizona Medical Board (“Board”).

11 **INTERIM FINDINGS OF FACT**

12 1. The Board is the duly constituted authority for the regulation and control of
13 the practice of allopathic medicine in the State of Arizona.

14 2. Respondent is the holder of License No. 44593 for the practice of allopathic
15 medicine in the State of Arizona.

16 3. The Board initiated case number MD-20-1037A after receiving notification
17 from a pharmacy chain that locations would no longer fill prescriptions written by
18 Respondent for Schedule II-IV controlled substances.

19 4. Based on the complaint, Board staff requested Medical Consultant (“MC”)
20 review of Respondent’s care and treatment of three patients (LP, PMC and DA).

21 5. LP was a 62 year-old female who initiated care with Respondent in May,
22 2017. LP had a past medical history (“PMH”) of chronic pain syndrome, anxiety,
23 hypertension, diabetes, thyroid disease, endometriosis, and psoriasis for which when she
24 had been prescribed Effexor XR, zonisamide, bupropion, hydromorphone and Morphine
25 ER. At her initial visit, Respondent prescribed LP a 7 day supply of buprenorphine. LP had

1 a severe reaction to the buprenorphine that required emergency treatment. Subsequently,
2 Respondent prescribed LP medications including morphine ER 100mg, every eight hours,
3 oxycodone 15mg every five hours, and Voltaren. Through February 21, 2018, LP was
4 being prescribed clonazepam by her primary care provider.

5 6. PMC was a 40 year-old male who initiated care with Respondent in March,
6 2017. PMC's PMH included chronic cluster/migraine headaches and insomnia.
7 Respondent prescribed PMC medications including oxycodone 15mg every six hours,
8 Ambien 12.5mg at bedtime, clonazepam 1mg three times daily, and sumatriptan.

9 7. DA was a 32 year-old female who initiated care with Respondent in June,
10 2018. DA had a reported PMH of degenerative disc disease of the cervical and lumbar
11 spine with chronic pain and radiculopathy. Respondent prescribed DA medications
12 including buprenorphine 8mg twice daily, oxycodone 10mg every 5 hours, Movantik 25mg
13 daily, tizanidine 4mg daily, and Voltaren gel.

14 8. A Medical Consultant ("MC") who reviewed Respondent's care of LP, PMC
15 and DA noted deviations from the standard of care regarding Respondent's care and
16 treatment of all three patients, including inappropriately initiating Suboxone treatment on
17 an outpatient basis for Patient LP, who was on high dose opioids, by prescribing to failing
18 to trial non-pharmacologic interventions for insomnia prior to prescribing Ambien to PMC,
19 and by prescribing buprenorphine to DA without adequate informed consent and
20 medication administration instructions. For all patients, the MC opined that Respondent
21 deviated from the standard of care by prescribing controlled substances without adequate
22 clinical justification, by failing to monitor the patients by querying the CSPMP and
23 obtaining urinary drug screens.

24 9. Actual harm was identified in that LP had a severe reaction to the
25 buprenorphine that required emergency care, PMC experienced inadequately treated

1 cluster headaches and DA experienced inadequately managed pain. All patients were at
2 risk of opioid use disorder, overdose and death.

3 10. The aforementioned information was presented to the investigative staff, the
4 medical consultant and the lead Board member. All reviewed the information and concur
5 that the interim consent agreement to restrict Respondent's controlled substance
6 prescribing pending the outcome of a formal interview or formal hearing is appropriate.

7 11. The investigation into this matter is pending Board review.

8 **INTERIM CONCLUSIONS OF LAW**

9 1. The Board possesses jurisdiction over the subject matter hereof and over
10 Respondent.

11 2. Pursuant to A.R.S. § 32-1405(C)(25) the Executive Director has authority to
12 enter into a consent agreement when there is evidence of danger to the public health and
13 safety.

14 3. Pursuant to A.A.C. R4-16-504, the Executive Director may enter into an
15 interim consent agreement when there is evidence that a restriction is needed to mitigate
16 imminent danger to the public's health and safety. Investigative staff, the Board's medical
17 consultant and the lead Board member have reviewed the case and concur that an interim
18 consent agreement is appropriate.

19 **INTERIM ORDER**

20 IT IS HEREBY ORDERED THAT:

21 1. Respondent is prohibited from prescribing controlled substances in the State
22 of Arizona pending the outcome of a formal interview or formal hearing in this matter.

23 2. Respondent may request, in writing, release and/or modification of this
24 Interim Consent Agreement. The Executive Director, in consultation with and agreement of
25 the lead Board member and the Chief Medical Consultant, has the discretion to determine

1 whether it is appropriate to release Respondent from this Interim Consent Agreement.

2 3. The Board retains jurisdiction and may initiate new action based upon any
3 violation of this Interim Consent Agreement, including, but not limited to, summarily
4 suspending Respondent's license.

5 4. Because this is an Interim Consent Agreement and not a final decision by
6 the Board regarding the investigation, it is subject to further consideration by the Board.

7 5. This Interim Consent Agreement shall be effective on the date signed by the
8 Board's Executive Director.

9 **RECITALS**

10 Respondent understands and agrees that:

11 1. The Board, through its Executive Director, may adopt this Interim Consent
12 Agreement, or any part thereof, pursuant to A.R.S. § 32-1405(C)(25) and A.A.C. R4-16-
13 504.

14 2. Respondent has read and understands this Interim Consent Agreement as
15 set forth herein, and has had the opportunity to discuss this Interim Consent Agreement
16 with an attorney or has waived the opportunity to discuss this Interim Consent Agreement
17 with an attorney. Respondent voluntarily enters into this Interim Consent Agreement and
18 by doing so agrees to abide by all of its terms and conditions.

19 3. By entering into this Interim Consent Agreement, Respondent freely and
20 voluntarily relinquishes all rights to an administrative hearing on the matters set forth
21 herein, as well as all rights of rehearing, review, reconsideration, appeal, judicial review or
22 any other administrative and/or judicial action, concerning the matters related to the
23 Interim Consent Agreement.
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1 4. Respondent understands that this Interim Consent Agreement does not
2 constitute a dismissal or resolution of this matter or any matters that may be currently
3 pending before the Board and does not constitute any waiver, express or implied, of the
4 Board's statutory authority or jurisdiction regarding this or any other pending or future
5 investigations, actions, or proceedings. Respondent also understands that acceptance of
6 this Interim Consent Agreement does not preclude any other agency, subdivision, or
7 officer of this State from instituting civil or criminal proceedings with respect to the conduct
8 that is the subject of this Interim Consent Agreement. Respondent further does not
9 relinquish his/her rights to an administrative hearing, rehearing, review, reconsideration,
10 judicial review or any other administrative and/or judicial action, concerning the matters
11 related to a final disposition of this matter, unless Respondent affirmatively does so as part
12 of the final resolution of this matter.
13

14 5. Respondent acknowledges and agrees that upon signing this Interim
15 Consent Agreement and returning it to the Board's Executive Director, Respondent may
16 not revoke acceptance of this Interim Consent Agreement or make any modifications to it.
17 Any modification of this original document is ineffective and void unless mutually approved
18 by the parties in writing.

19 6. Respondent understands that this Interim Consent Agreement shall not
20 become effective unless and until it is signed by the Board's Executive Director.

21 7. Respondent understands and agrees that if the Board's Executive Director
22 does not adopt this Interim Consent Agreement, he will not assert in any future
23 proceedings that the Board's consideration of this Interim Consent Agreement constitutes
24 bias, prejudice, prejudgment, or other similar defense.
25

1 8. Respondent understands that this Interim Consent Agreement is a public
2 record that may be publicly disseminated as a formal action of the Board, and that it shall
3 be reported as required by law to the National Practitioner Data Bank.

4 9. Respondent understands that this Interim Consent Agreement does not
5 alleviate Respondent's responsibility to comply with the applicable license-renewal
6 statutes and rules. If this Interim Consent Agreement remains in effect at the time
7 Respondent's allopathic medical license comes up for renewal, Respondent must renew
8 the license if Respondent wishes to retain the license. If Respondent elects not to renew
9 the license as prescribed by statute and rule, Respondent's license will not expire but
10 rather, by operation of law (A.R.S. § 32-3202), become suspended until the Board takes
11 final action in this matter. Once the Board takes final action, in order for Respondent to be
12 licensed in the future, Respondent must submit a new application for licensure and meet
13 all of the requirements set forth in the statutes and rules at that time.

14 10. Respondent understands that any violation of this Interim Consent
15 Agreement constitutes unprofessional conduct under A.R.S. § 32-1401(27)(s) ("[v]iolating
16 a formal order, probation, consent agreement or stipulation issued or entered into by the
17 board or its executive director under this chapter.").

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20 
MONEIL M. PATEL, M.D.

DATED: 3/12/22

21 DATED this 15th day of March, 2022.

22 ARIZONA MEDICAL BOARD

23 By 
24 Patricia E. McSorley
25 Executive Director

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EXECUTED COPY of the foregoing e-
mailed this 15th day of March, 2022 to:

Sara Stark, Esq.
Chelle Law
11811 North Tatum, Suite 4000
Phoenix, Arizona 85028
Attorney for Respondent

ORIGINAL of the foregoing filed
this 15th day of March, 2022 with:

Arizona Medical Board
1740 West Adams, Suite 4000
Phoenix, Arizona 85007



Board staff