

New Data from the RECOVER™ Study Reveal Two-Year Outcomes in People with Opioid Use Disorder Following Transition from Pivotal Phase 3 Clinical Trials to a Real-World Setting



The ongoing RECOVER™ study is assessing real-world, patient-centered outcomes including self-reported sustained illicit opioid-free weeks and life changes, such as improved health, employment status and connection to community following treatment with SUBLOCADE®

Slough, UK and Richmond, VA, June 24, 2020 – Indivior PLC (LON: INDV) announces new data from the two-year analysis of the 24-month real-world observational study, RECOVER™ (Remission from Chronic Opioid Use—Studying Environmental and SocioEconomic Factors on Recovery).

The RECOVER study examines long-term recovery in individuals with moderate to severe opioid use disorder (OUD) following their transition from two Phase 3 clinical trials of SUBLOCADE® (buprenorphine extended-release) injection, for subcutaneous use (CIII) into a real-world setting.¹ This is the first analysis to examine patient outcomes two years after receiving up to 18 months of treatment with SUBLOCADE®.

The main findings from this analysis were as follows:

- Of the 533 participants enrolled in RECOVER, 396 (74%) completed the 24-month assessment.^{2,3}
- 218 participants (44%) self-reported sustained illicit opioid-free weeks for the entire 24-month period.³
- Nearly half of the participants were employed over the assessment period; percentages increased from 35% at pre-trial screening to 45-48% at 24-months.³

- As it relates to healthcare resource utilization over the entire 24-month period, 15% of participants reported staying overnight in a hospital and 12% reported having an emergency department visit.³
- Among the 495 participants who contributed data over the 24-month period, 69% of the cohort received substance use disorder treatment, of which buprenorphine medication assisted treatment was the most prevalent.³

These results were reported at the 82nd Annual Scientific Meeting of the College on Problems of Drug Dependence (CPDD) held virtually from June 22-24, 2020.

“Opioid use disorder is a chronic disease and its treatment is complex and multifaceted,” said the study’s lead author, Walter Ling, MD, Research Professor, Department of Family Medicine, David Geffen School of Medicine, UCLA. “These findings show that long-term treatment with counseling may positively assist patients in focusing on their recovery, including discontinuation of illicit opioid use. Over the 24-month RECOVER observation period, participants also achieved and maintained positive effects including improved employment rates and lower healthcare system utilization.”³

“Studies such as RECOVER™ can help bridge the knowledge gap between the efficacy of medications as seen in the controlled clinical trial environment, and the use and effect of medications outside of a research setting and their long-term impact on patients’ health,” said Warren K. Bickel, a professor at Fralin Biomedical Research Institute at Virginia Tech Carilion.

Dr. Bickel is leading the next phase of the RECOVER study, which researchers hope will provide further information to health care systems and policymakers on how continuity of care can help address the nation’s opioid crisis.

RECOVER 24 Month Study Limitations

It is important to note that self-reported data were used to define sustained and past-week illicit opioid-free use. Although urine drug screen data were also collected at each 3-month follow-up, drug use is only captured over a limited time span of a few days. Participant self-report of illicit opioid use may be subject to recall bias. However, asking about illicit opioid-free use in the past week, in addition to over longer periods of time, minimizes this potential bias. While this is acknowledged, self-reported illicit opioid use is a frequently used methodology for reporting opioid-free use. Illicit opioid-free use observed during RECOVER study may be attributable to additional OUD pharmacotherapy beyond BUP-XR received during clinical trial period, which warrants further investigation.

ABOUT SUBLOCADE

INDICATION AND USAGE

SUBLOCADE® (buprenorphine extended-release) injection, for subcutaneous use (CIII) is indicated for the treatment of moderate to severe opioid use disorder in patients who have initiated treatment with a transmucosal buprenorphine-containing product followed by a dose adjustment period for a minimum of seven days.

SUBLOCADE should be used as part of a complete treatment program that includes counseling and psychosocial support.

IMPORTANT SAFETY INFORMATION

**WARNING: RISK OF SERIOUS HARM OR DEATH WITH INTRAVENOUS ADMINISTRATION;
SUBLOCADE RISK EVALUATION AND MITIGATION STRATEGY**

- **Serious harm or death could result if administered intravenously. SUBLOCADE forms a solid mass upon contact with body fluids and may cause occlusion, local tissue damage, and thrombo-embolic events, including life threatening pulmonary emboli, if administered intravenously.**
- **Because of the risk of serious harm or death that could result from intravenous self-administration, SUBLOCADE is only available through a restricted program called the SUBLOCADE REMS Program. Healthcare settings and pharmacies that order and dispense SUBLOCADE must be certified in this program and comply with the REMS requirements.**
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Prescription use of this product is limited under the Drug Addiction Treatment Act.

SUBLOCADE should only be prepared and administered by a healthcare provider.

CONTRAINDICATIONS

SUBLOCADE should not be administered to patients who have been shown to be hypersensitive to buprenorphine or any component of the ATRIGEL® delivery system.

WARNINGS AND PRECAUTIONS

Addiction, Abuse, and Misuse: SUBLOCADE contains buprenorphine, a Schedule III controlled substance that can be abused in a manner similar to other opioids. Buprenorphine is sought by people with opioid use disorder and is subject to criminal diversion. Monitor patients for conditions indicative of diversion or progression of opioid dependence and addictive behaviors.

Risk of Life-Threatening Respiratory Depression and Concomitant Use of Benzodiazepines or Other CNS Depressants with Buprenorphine: Buprenorphine has been associated with life-threatening respiratory depression, overdose, and death, particularly when misused by self-injection or with concomitant use of benzodiazepines or other CNS depressants, including alcohol. Warn patients of the potential danger of self-administration of benzodiazepines, other CNS depressants, opioid analgesics, and alcohol while under treatment with SUBLOCADE. Counsel patients that such medications should not be used concomitantly unless supervised by a healthcare provider.

Use with caution in patients with compromised respiratory function (e.g., chronic obstructive pulmonary disease, cor pulmonale, decreased respiratory reserve, hypoxia, hypercapnia, or pre-existing respiratory depression).

Opioids can cause sleep-related breathing disorders; e.g., central sleep apnea (CSA), sleep-related hypoxemia. Opioid use increases the risk of CSA in a dose-dependent fashion. Consider decreasing the opioid using best practices for opioid taper if CSA occurs.

Neonatal Opioid Withdrawal Syndrome: Neonatal opioid withdrawal syndrome (NOWS) is an expected and treatable outcome of prolonged use of opioids during pregnancy. NOWS may be life-threatening if not recognized and treated in the neonate. Newborns should be observed for signs of NOWS and managed accordingly. Advise pregnant women receiving opioid addiction treatment with SUBLOCADE of the risk of neonatal opioid withdrawal syndrome.

Adrenal Insufficiency: Adrenal insufficiency has been reported with opioid use. If adrenal insufficiency is diagnosed, treat with physiologic replacement doses of corticosteroids. Wean the patient off the opioid.

Discontinuation of SUBLOCADE Treatment: Due to the long-acting nature of SUBLOCADE, if treatment is discontinued, monitor patients for several months for withdrawal and treat appropriately.

Inform patients that they may have detectable levels of buprenorphine for a prolonged period of time after treatment with SUBLOCADE. Considerations of drug-drug interactions, buprenorphine effects, and analgesia may continue to be relevant for several months after the last injection.

Risk of Hepatitis, Hepatic Events: Because cases of cytolytic hepatitis and hepatitis with jaundice have been observed in individuals receiving buprenorphine, monitor liver function tests prior to treatment and monthly during treatment.

Hypersensitivity Reactions: Hypersensitivity to buprenorphine-containing products have been reported most commonly as rashes, hives, and pruritus. Some cases of bronchospasm, angioneurotic edema, and anaphylactic shock have also been reported.

Precipitation of Opioid Withdrawal in Patients Dependent on Full Agonist Opioids: Buprenorphine may precipitate opioid withdrawal signs and symptoms in persons who are currently physically dependent on full opioid agonists such as heroin, morphine, or methadone before the effects of the full opioid agonist have subsided. Verify that patients have tolerated and are dose adjusted on transmucosal buprenorphine before subcutaneously injecting SUBLOCADE.

Risks Associated With Treatment of Emergent Acute Pain: When patients need acute pain management, or may require anesthesia, treat patients receiving SUBLOCADE currently or within the last 6 months with a non-opioid analgesic whenever possible. If opioid therapy is required, patients may be treated with a

high-affinity full opioid analgesic under the supervision of a physician, with particular attention to respiratory function, as higher doses may be required for analgesic effect and therefore, a higher potential for toxicity exists with opioid administration.

Advise patients of the importance of instructing their family members, in the event of emergency, to inform the treating healthcare provider or emergency room staff that the patient is physically dependent on an opioid and that the patient is being treated with SUBLOCADE.

Use in Opioid Naïve Patients: Because death has been reported for opioid naïve individuals who received buprenorphine sublingual tablet, SUBLOCADE is not appropriate for use in opioid naïve patients.

Use in Patients With Impaired Hepatic Function: Because buprenorphine levels cannot be rapidly decreased, SUBLOCADE is not recommended for patients with preexisting moderate to severe hepatic impairment. Patients who develop moderate to severe hepatic impairment while being treated with SUBLOCADE should be monitored for several months for signs and symptoms of toxicity or overdose caused by increased levels of buprenorphine.

Use in Patients at Risk for Arrhythmia: Buprenorphine has been observed to prolong the QTc interval in some patients participating in clinical trials. Avoid use of buprenorphine in patients with a history of Long QT Syndrome or an immediate family member with this condition or those taking Class IA antiarrhythmic medications (e.g., quinidine, procainamide, disopyramide) or Class III antiarrhythmic medications (e.g., sotalol, amiodarone, dofetilide), or other medications that prolong the QT interval.

Impairment of Ability to Drive or Operate Machinery: SUBLOCADE may impair the mental or physical abilities required for the performance of potentially dangerous tasks such as driving a car or operating machinery. Caution patients about driving or operating hazardous machinery until they are reasonably certain that SUBLOCADE does not adversely affect their ability to engage in such activities.

Orthostatic Hypotension: Buprenorphine may produce orthostatic hypotension.

Elevation of Cerebrospinal Fluid Pressure: Buprenorphine may elevate cerebrospinal fluid pressure and should be used with caution in patients with head injury, intracranial lesions, and other circumstances when cerebrospinal pressure may be increased. Buprenorphine can produce miosis and changes in the level of consciousness that may interfere with patient evaluation.

Elevation of Intracholedochal Pressure: Buprenorphine has been shown to increase intracholedochal pressure, as do other opioids, and thus should be administered with caution to patients with dysfunction of the biliary tract.

Effects in Acute Abdominal Conditions: Buprenorphine may obscure the diagnosis or clinical course of patients with acute abdominal conditions.

Unintentional Pediatric Exposure: Buprenorphine can cause severe, possibly fatal, respiratory depression in children who are accidentally exposed to it.

ADVERSE REACTIONS: Adverse reactions commonly associated with SUBLOCADE (≥5% of subjects) during clinical trials were constipation, headache, nausea, vomiting, increased hepatic enzymes, fatigue, and injection site pain and pruritus. This is not a complete list of potential adverse events. Please see the full Prescribing Information for a complete list.

DRUG INTERACTIONS

CYP3A4 Inhibitors and Inducers: Monitor patients starting or ending CYP3A4 inhibitors or inducers for potential over- or under-dosing.

Serotonergic Drugs: If concomitant use with serotonergic drugs is warranted, monitor for serotonin syndrome, particularly during treatment initiation, and during dose adjustment of the serotonergic drug.

Consult the full Prescribing Information for SUBLOCADE for more information on potentially significant drug interactions.

USE IN SPECIFIC POPULATIONS

Pregnancy: Opioid-dependent women on buprenorphine maintenance therapy may require additional analgesia during labor.

Lactation: Buprenorphine passes into the mother's milk. Advise breastfeeding women to monitor the infant for increased drowsiness and breathing difficulties.

Fertility: Chronic use of opioids may cause reduced fertility. It is not known whether these effects on fertility are reversible.

Geriatric Patients: Monitor geriatric patients receiving SUBLOCADE for sedation or respiratory depression.

To report pregnancy or side effects associated with taking SUBLOCADE, please call 1-877-782-6966.

For more information about SUBLOCADE, see the full [Prescribing Information](#) including **BOXED WARNING**, and [Medication Guide](#). For REMS information visit www.sublocadeREMS.com.

About Opioid Use Disorder (OUD)

Opioid addiction isn't a moral weakness. Opioid addiction is a chronic disease called Opioid Use Disorder (OUD)⁵ in which people develop a pattern of using opioids that can lead to negative consequences.⁶ Opioid addiction may affect the parts of the brain that control impulses, judgment, and decision-making.^{6,7} Patients become trapped in a cycle of opioid use, which produces changes in brain function that can reduce their ability to control their use.^{6,8}

In 2018, an estimated 10.3 million people aged 12 or older misused opioids in the past year, including 9.9 million prescription pain reliever misusers and 808,000 heroin users.

Approximately 506,000 people misused prescription pain relievers and used heroin in the past year.⁹ SUBLOCADE is not indicated for use in children younger than 18 years of age.

Buprenorphine, the active ingredient of SUBLOCADE can cause severe, possibly fatal, respiratory depression in children who are accidentally exposed to it.⁴

About the RECOVER™ Study¹

The RECOVER (Remission from Chronic Opioid Use-Studying Environmental and Socio-Economic Factors on Recovery; NCT03604861) study is a multisite, non-interventional cohort study examining long-term recovery in individuals with moderate to severe OUD who received at least one dose of study treatment during the Phase III clinical trials (NCT02357901 and NCT02510014) for SUBLOCADE. Participants (n=533) were eligible to join the RECOVER study 28 days after completing or terminating participation in the SUBLOCADE Phase III trials.

Results are being analyzed to understand the clinical, socio-economic and environmental factors associated with continuous effects of medications for OUD after a clinical trial.

About Indivior

Indivior is a global pharmaceutical company working to help change patients' lives by developing medicines to treat addiction and serious mental illnesses. Our vision is that all patients around the world will have access to evidence-based treatment for the chronic conditions and co-occurring disorders of addiction. Indivior is dedicated to transforming addiction from a global human crisis to a recognized and treated chronic disease.

Building on its global portfolio of opioid dependence treatments, Indivior has a pipeline of product candidates designed to both expand on its heritage in this category and potentially address other chronic conditions and co-occurring disorders of addiction, including alcohol use disorder. Headquartered in the United States in Richmond, VA, Indivior employs more than 800 individuals globally and its portfolio of products is available in over 40 countries worldwide. Visit www.indivior.com to learn more. Connect with Indivior on LinkedIn by visiting www.linkedin.com/company/indivior.

Forward-Looking Statements

This press release contains certain statements that are forward-looking, and which should be considered, amongst other statutory provisions, in light of the safe harbor provisions of the United States Private Securities Litigation Reform Act of 1995. By their nature, forward-looking statements involve risk and uncertainty as they relate to events or circumstances that may or may not occur in the future. Actual results may differ materially from those

expressed or implied in such statements because they relate to future events. Forward-looking statements include, among other things, statements regarding the Indivior Group's financial guidance for 2020 and its medium- and long-term growth outlook, its operational goals, its product development pipeline and statements regarding ongoing litigation. Various factors may cause differences between Indivior's expectations and actual results, including: factors affecting sales of Indivior Group's products; the outcome of research and development activities; decisions by regulatory authorities regarding the Indivior Group's drug applications; the speed with which regulatory authorizations, pricing approvals and product launches may be achieved; the outcome of post-approval clinical trials; competitive developments; difficulties or delays in manufacturing; the impact of existing and future legislation and regulatory provisions on product exclusivity; trends toward managed care and healthcare cost containment; legislation or regulatory action affecting pharmaceutical product pricing, reimbursement or access; claims and concerns that may arise regarding the safety or efficacy of the Indivior Group's products and product candidates; risks related to legal proceedings, including the ongoing investigative and antitrust litigation matters; the Indivior Group's ability to protect its patents and other intellectual property; the outcome of patent infringement litigation relating to Indivior Group's products, including the ongoing ANDA lawsuits; changes in governmental laws and regulations; issues related to the outsourcing of certain operational and staff functions to third parties; uncertainties related to general economic, political, business, industry, regulatory and market conditions; and the impact of acquisitions, divestitures, restructurings, internal reorganizations, product recalls and withdrawals and other unusual items. This press release does not constitute an offer to sell or the solicitation of an offer to subscribe for or otherwise acquire or dispose of shares in the Company to any person in any jurisdiction to whom it is unlawful to make such offer or solicitation.

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