



Scilex Holding Company Appoints Kasowitz LLP as Litigation and Intellectual Property Counsel

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PALO ALTO, Calif., Jan. 02, 2026 (GLOBE NEWSWIRE) -- Scilex Holding Company ("Scilex" or the "Company") (Nasdaq: SCLX), an innovative revenue-generating company focused on acquiring, developing and commercializing non-opioid pain management products for the treatment of acute and chronic pain and neurodegenerative and cardiometabolic disease, announced that it has appointed Kasowitz LLP and its founding partner Marc Kasowitz as the Company's litigation and intellectual property counsel.

Kasowitz LLP will advise Scilex on complex commercial litigation, patent litigation, intellectual property enforcement and defense, licensing matters, and strategic intellectual property governance as the Company continues to advance and protect its proprietary pharmaceutical assets and development pipeline. Mr. Kasowitz is a highly experienced U.S. trial lawyer with decades of experience leading high-stakes intellectual property and complex commercial litigation in federal and state courts. His background includes representation in intellectual property disputes involving innovative technologies and life sciences.

Marc E. Kasowitz is a veteran American trial lawyer and the founding managing partner of Kasowitz LLP, a New York-based litigation firm he co-founded in 1993. He is widely recognized for his expertise in complex commercial litigation, having been named among the country's top trial lawyers by Benchmark Litigation and honored as a Litigation Trailblazer by the National Law Journal. Mr. Kasowitz regularly serves as lead counsel in high-stakes disputes involving antitrust, securities, banking, and product liability matters, representing major corporate clients across a range of industries, and has earned recognition from Chambers USA, The Legal 500, and other legal ranking organizations for his trial experience and advocacy. For more information, click [Kasowitz LLP](#)

"Protecting and enforcing our intellectual property portfolio is central to Scilex's long-term strategy and shareholder value," said a spokesperson for Scilex. "The appointment of Marc Kasowitz and Kasowitz LLP strengthens our legal capabilities as we continue to commercialize our products and advance our late-stage pipeline."

For more information on Scilex Holding Company, refer to www.scilexholding.com

For more information on Semnur Pharmaceuticals, Inc., refer to www.semnurpharma.com

For more information on ZTlido[®] including Full Prescribing Information, refer to www.ztlido.com.

For more information on ELYXYB[®], including Full Prescribing Information, refer to www.elyxyb.com.

For more information on Gloperba[®], including Full Prescribing Information, refer to www.gloperba.com.

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About Scilex Holding Company

Scilex is an innovative revenue-generating company focused on acquiring, developing and commercializing non-opioid pain management products for the treatment of acute and chronic pain and neurodegenerative and cardiometabolic disease. Scilex targets indications with high unmet needs and large market opportunities with non-opioid therapies for the treatment of patients with acute and chronic pain and is dedicated to advancing and improving patient outcomes. Scilex's commercial products include: (i) ZTlido[®] (lidocaine topical system) 1.8%, a prescription lidocaine topical product approved by the U.S. Food and Drug Administration (the "FDA") for the relief of neuropathic pain associated with postherpetic neuralgia, which is a form of post-shingles nerve pain; (ii) ELYXYB[®], a potential first-line treatment and the only FDA-approved, ready-to-use oral solution for the acute treatment of migraine, with or without aura, in adults; and (iii) Gloperba[®], the first and only liquid oral version of the anti-gout medicine colchicine indicated for the prophylaxis of painful gout flares in adults.

In addition, Scilex has three product candidates: (i) SP-102 (10 mg, dexamethasone sodium phosphate viscous gel) ("SEMDEXA[™]" or "SP-102"), which is owned by Semnur (a majority owned subsidiary of Scilex) and is a novel, viscous gel formulation of a widely used corticosteroid for epidural injections to treat lumbosacral radicular pain, or sciatica, for which Scilex has completed a Phase 3 study and was granted Fast Track status from the FDA in 2017; (ii) SP-103 (lidocaine topical system) 5.4%, ("SP-103"), a next-generation, triple-strength formulation of ZTlido, for the treatment of acute pain and for which Scilex has recently completed a Phase 2 trial in acute low back pain. SP-103 has been granted Fast Track status from the FDA in low back pain; and (iii) SP-104 (4.5 mg, low-dose naltrexone hydrochloride delayed-release capsules) ("SP-104"), a novel low-dose delayed-release naltrexone hydrochloride being developed for the treatment of fibromyalgia.

Scilex is headquartered in Palo Alto, California.

Forward-Looking Statements

This press release includes forward-looking statements that involve risks and uncertainties. Forward-looking statements are statements that are not historical facts and may be accompanied by words that convey projected future events or outcomes, such as "believe," "may," "will," "estimate," "continue," "anticipate," "intend," "expect," "should," "would," "plan," "predict," "potential," "seem," "seek," "future," "outlook" or variations of such words or by expressions of similar meaning. These forward-looking statements include, but are not limited to, statements regarding future events, Scilex's intellectual property goals and processes, future opportunities for Scilex and its subsidiaries, the future business strategies, long-term objectives and commercialization plans of Scilex and its subsidiaries, the current and prospective product candidates, planned clinical trials and preclinical activities

and potential product approvals, as well as the potential for market acceptance of any approved products and the related market opportunity of Scilex and its subsidiaries, statements regarding SP-102, if approved by the FDA, Scilex's potential to attract new capital and avoid the effects of negative debt leverage and other statements that are not historical facts. These statements are based on management's current expectations of and are not predictions of actual performance. These forward-looking statements are provided for illustrative purposes only and are not intended to serve as, and must not be relied on, by any investor as a guarantee, an assurance, a prediction or a definitive statement of fact or probability. Actual events and circumstances are difficult or impossible to predict and will differ from assumptions. Many actual events and circumstances are beyond the control of Scilex. These statements are subject to a number of risks and uncertainties regarding Scilex's business, and actual results may differ materially. These risks and uncertainties include, but are not limited to, general economic, political and business conditions; the ability of Scilex and its subsidiaries to develop and successfully market products; the ability of Scilex and its subsidiaries to grow and manage growth profitably and retain its key employees; the risk that the potential product candidates that Scilex develops may not progress through clinical development or receive required regulatory approvals within expected timelines or at all; risks relating to uncertainty regarding the regulatory pathway for Scilex's product candidates; the risk that Scilex's product candidates may not be beneficial to patients or successfully commercialized; the risk that Scilex has overestimated the size of the target patient population, their willingness to try new therapies and the willingness of physicians to prescribe these therapies; risks that the prior results of the clinical trials may not be replicated; regulatory and intellectual property risks; the risk of failure to realize the anticipated benefits of the transactions contemplated with Datavault and other risks and uncertainties indicated from time to time and other risks set forth in Scilex's filings with the SEC. There may be additional risks that Scilex presently does not know or that Scilex currently believes are immaterial that could also cause actual results to differ from those contained in the forward-looking statements. In addition, forward-looking statements provide Scilex's expectations, plans or forecasts of future events and views as of the date of the communication. Scilex anticipates that subsequent events and developments will cause such assessments to change. However, while Scilex may elect to update these forward-looking statements at some point in the future, Scilex specifically disclaims any obligation to do so. These forward-looking statements should not be relied upon as representing Scilex's assessments as of any date subsequent to the date of this communication. Accordingly, investors are cautioned not to place undue reliance on these forward-looking statements.

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SEMDEXA™ (SP-102) is a trademark owned by Semnur Pharmaceuticals, Inc., a majority-owned subsidiary of Scilex Holding Company. A proprietary name review by the FDA is planned.

ZTlido® is a registered trademark owned by Scilex Pharmaceuticals Inc., a wholly-owned subsidiary of Scilex Holding Company.

Gloperba® is the subject of an exclusive, transferable license to use the registered trademark by Scilex Holding Company.

ELYXYB® is a registered trademark owned by Scilex Holding Company.

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