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Ensysce Biosciences Posts Video Updates from the IASP World Congress on Pain 2024

~ PF614 and PF614-MPAR Highlighted at Symposium on Severe Pain ~

SAN DIEGO, CA / ACCESSWIRE / September 13, 2024 /Ensysce Biosciences, Inc. (NASDAQ:ENSC) ("Ensysce" or the "Company"), a clinical-stage pharmaceutical company developing innovative solutions for severe pain relief while reducing the potential for opioid abuse and overdose, has made available two video segments from its August 8th symposium at the International Association for the Study of Pain (IASP) 2024 World Congress on Pain in Amsterdam, Netherlands. The Company's symposium emphasized opioid use worldwide, the current landscape for treating severe pain, including post-surgical and cancer pain treatment, and discussed Ensysce's innovative new class of opioid drug products.

The discussion allowed for feedback and questions from experts in attendance facilitating proper introduction and explanation of the differentiated Ensysce approach and the possible needs from countries outside the United States for these unique opioids.

Two short videos from the session have been posted to Media / Science Centerpage on the Company's website. Dr. William Schmidt, Chief Medical Officer of Ensysce explains the government support the Company has received from both the National Institutes of Health and the FDA, as well as described how the profile of Ensysce's lead agent, PF614, checks all the categories of what could be perceived as an ideal pain product.

Dr. Lynn Kirkpatrick, Chief Executive Officer of Ensysce Biosciences, who moderated the symposium, commented, "Developing a novel opioid is not easy and we have been interested in sharing and discussing our technology with experts in the field who treat patients with pain on a daily basis. Our aim is to provide a new approach to treat severe pain with our novel class of opioid products that use our TAAP and MPAR platforms. These we believe will result in less abuse and overdose that have plagued traditional opioid products. We announced our plans to begin a Phase 3 pivotal trial shortly, as well as our intent to explore the use of PF614 in other indications such as cancer. Both trials are moving us one step closer to offering prescribers a new option to treat their patients more effectively."

The IASP has a mission to deliver pain relief throughout the world. Please find more information at the IASP website: https://www.iasp-pain.org.

About Ensysce Biosciences
Ensysce Biosciences is a clinical-stage company using its proprietary technology platforms to develop safer prescription drugs. Leveraging its Trypsin-Activated Abuse Protection (TAAP™) and Multi-Pill Abuse Resistance (MPAR®) platforms, the Company is developing

unique, tamper-proof treatment options for pain that minimize the risk of both drug abuse and overdose. Ensysce's products are anticipated to provide safer options to treat patients suffering from severe pain and assist in preventing deaths caused by medication abuse. The platforms are covered by an extensive worldwide intellectual property portfolio for a wide array of prescription drug compositions. For more information, please visit www.ensysce.com.

Forward-Looking Statements
Statements contained in this press release that are not purely historical may be deemed to be forward-looking statements for the purposes of the safe harbor provisions under The Private Securities Litigation Reform Act of 1995 and other federal securities laws. Without limiting the foregoing, the use of words such as "may," "intends," "can," "might," "will," "expect," "plan," "possible," "believe" and other similar expressions are intended to identify forward-looking statements. The product candidates discussed are in clinic and not approved and there can be no assurance that the clinical programs will be successful in demonstrating safety and/or efficacy, that Ensysce will not encounter problems or delays in clinical development, or that any product candidate will ever receive regulatory approval or be successfully commercialized. All forward-looking statements are based on estimates and assumptions by Ensysce's management that, although Ensysce believes to be reasonable, are inherently uncertain. All forward-looking statements are subject to risks and uncertainties that may cause actual results to differ materially from those that Ensysce expected. In addition, Ensysce's business is subject to additional risks and uncertainties, including among others, the initiation and conduct of preclinical studies and clinical trials; the timing and availability of data from preclinical studies and clinical trials; expectations for regulatory submissions and approvals; potential safety concerns related to, or efficacy of, Ensysce's product candidates; the availability or commercial potential of product candidates; the ability of Ensysce to fund its continued operations, including its planned clinical trials; the dilutive effect of stock issuances from our fundraising; and Ensysce's and its partners' ability to perform under their license, collaboration and manufacturing arrangements. These statements are also subject to a number of material risks and uncertainties that are described in Ensysce's most recent quarterly report on Form 10-Q and current reports on Form 8-K, which are available, free of charge, at the SEC's website at www.sec.gov. Any forward-looking statement speaks only as of the date on which it was made. Ensysce undertakes no obligation to publicly update or revise any forward-looking statement, whether as a result of new information, future events or otherwise, except as required under applicable law.

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