Sol-Gel Technologies to Host Conference Call to Discuss Top-Line Results from Pivotal Phase 3 Clinical Trial of SGT-610 in Gorlin Syndrome

November 27, 2026

Conference call to take place Monday, November 30, 2026, at 8:00am Eastern Time

NESS ZIONA, Israel, Nov. 27, 2026 (GLOBE NEWSWIRE) -- **Sol-Gel Technologies, Ltd.** (NASDAQ: SLGL), a dermatology company, pioneering treatments for patients with rare and severe skin conditions, today announced that it will host a conference call on Monday, November 30, 2026, at 8:00am Eastern Time to discuss top-line results from its pivotal Phase 3 clinical trial (SGT-610-01) evaluating SGT-610 (patidegib gel, 2%) for the prevention of basal cell carcinoma lesions in adult patients with Gorlin syndrome. The Company plans to issue a press release announcing the top-line results at 7:00am Eastern Time on Monday, November 30, 2026, prior to the start of the conference call.

To access the live webcast of the call, please click here or visit the "Events" section of Sol-Gel's website. To access the call by phone, please use this registration link, and you will be provided with dial in details. A replay of the call will be available approximately 2 hours after the conclusion of the call and archived for 90 days under the "Events " section of the Company's website at http://www.sol-gel.com.

About Sol-Gel Technologies

Sol-Gel is a specialized dermatology company advancing innovative therapies for rare and serious skin diseases. Its lead investigational candidate, SGT-610 (patidegib gel, 2%), is a Phase 3, orphan- and breakthrough-designated topical hedgehog inhibitor being developed for the prevention of new basal cell carcinoma (BCC) lesions in patients with Gorlin syndrome, with the potential to offer an improved safety profile relative to oral hedgehog inhibitors. Subject to regulatory approval in Gorlin syndrome, SGT-610 may also represent a future opportunity in high-frequency BCC. Sol-Gel is also advancing SGT-210, an investigational topical EGFR inhibitor for indications with significant unmet need, and has developed two FDA-approved dermatology products, TWYNEO® and EPSOLAY®.

Forward Looking Statements

This press release contains "forward-looking statements" within the meaning of the Private Securities Litigation Reform Act of 1995. All statements contained in this press release that do not relate to matters of historical fact should be considered forward-looking statements. including, but not limited to statements regarding the timing of the release of top-line results for the Phase 3 program of SGT-610 in Gorlin syndrome. In some cases, you can identify forward-looking statements by terminology such as "believe," "may," "estimate," "continue," "anticipate," "intend," "should," "plan," "expect," "predict," "potential," or the negative of these terms or other similar expressions. Forward-looking statements are based on information we have when those statements are made or our management's current expectations and are subject to risks and uncertainties that could cause actual performance or results to differ materially from those expressed in or suggested by the forward-looking statements. Important factors that could cause such differences include, but are not limited to, the risk of a delay in reporting the top-line results of Sol-Gel's pivotal Phase 3 clinical trial SGT-610-01 evaluating SGT-610 (patidegib gel, 2%) for Gorlin syndrome, that we will not successfully complete such Phase 3 clinical trial or that such clinical trial will fail to meet its primary endpoint and/or one or more of its secondary endpoints as well as the following factors: (i) the adequacy of our financial and other resources, particularly in light of our history of recurring losses; (ii) our ability to complete the development of our product candidates; (iii) our ability to find suitable co-development partners; (iv) our ability to obtain and maintain regulatory approvals for our product candidates in our target markets, the potential delay in receiving such regulatory approvals and the possibility of adverse regulatory or legal actions relating to our product candidates even if regulatory approval is obtained; (v) our collaborators' ability to commercialize our pharmaceutical product candidates; (vi) our ability to obtain and maintain adequate protection of our intellectual property; (vii) our collaborators' ability to manufacture our product candidates in commercial quantities, at an adequate quality or at an acceptable cost; (viii) our collaborators' ability to establish adequate sales, marketing and distribution channels; (ix) acceptance of our product candidates by healthcare professionals and patients; (x) the possibility that we may face third-party claims of intellectual property infringement; (xi) the timing and results of clinical trials that we may conduct or that our competitors and others may conduct relating to our or their products; (xii) intense competition in our industry, with competitors having substantially greater financial, technological, research and development, regulatory and clinical, manufacturing, marketing and sales, distribution and personnel resources than we do; (xiii) potential product liability claims; (xiv) potential adverse federal, state and local government regulation in the United States, China, Europe or Israel; and (xv) loss or retirement of key executives and research scientists; (xvi) general market, political and economic conditions in the countries in which the Company operates; and, (xvii) the security situation in Israel. These factors and other important factors discussed in the Company's Annual Report on Form 20-F filed with the Securities and Exchange Commission ("SEC") on March 19, 2026, and our other reports filed with the SEC, could cause actual results to differ materially from those indicated by the forward-looking statements made in this press release. Except as required by law, we undertake no obligation to update any forward-looking statements in this press release.

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