



Sol-Gel

Advanced Topical Therapy

NASDAQ: SLGL

June 2021



Forward-Looking Statements

This presentation contains forward-looking statements within the meaning of the Private Securities Litigation Reform Act of 1995. All statements other than statements of historical facts are forward-looking statements. In some cases, you can identify forward-looking statements by terms such as “may,” “will,” “should,” “expect,” “plan,” “anticipate,” “could,” “future,” “outlook,” “intend,” “target,” “project,” “contemplate,” “believe,” “estimate,” “predict,” “potential,” “continue,” or the negative of these terms or other similar expressions, although not all forward-looking statements contain these words. The forward-looking statements in this presentation relate to, among other things, statements regarding the PDUFA goal date for TWYNEO, approval and commercial launch of EPSOLAY and TWYNEO, the strategic partnership with Galderma, anticipated timing of the initiation of clinical trials for SGT-510, the intellectual property protection that would be provided by patents for SGT-210 and our tapinarof drug product, the timing of the launch of our tapinarof drug product, the future markets for various skin diseases, the timing of a test and a second POC study of erlotinib, projected profit margins, our expectations regarding our liquidity and ability to fund operational and capital expenditure requirements. These statements are neither promises nor guarantees, but involve known and unknown risks, uncertainties, and other important factors that may cause our actual results, performance, or achievements to be materially different from any future results, performance, or achievements expressed or implied by the forward-looking statement, including but not limited to the following: risks relating to the timing of the PDUFA action date for EPSOLAY, the timing of FDA approval, if any, of EPSOLAY and TWYNEO; the risk that we not receive any or all of the anticipated benefits of the strategic partnership with Galderma, the risk of a delay in the clinical trials for SGT-510; the risk that patents for SGT-210 and our tapinarof drug product will not provide the anticipated intellectual property protection; the risk of a delay in the launch of our tapinarof drug product; the risk that our estimate of the markets for psoriasis, atopic dermatitis and for hyperkeratotic skin diseases are inaccurate; the risk that our tapinarof drug product will not be the only other player besides the brand for a number of years; the risk of a delay in the timing of a test of erlotinib with a much higher concentrations in an animal model and the risk of a delay of a second POC study on PPK patients; the fact that we have and expect to continue to incur significant losses; our need for additional funding, which may not be available; our ability to complete the development of, and obtain marketing approval for, our product candidates; our ability to obtain and maintain regulatory approvals for our product candidates in our target markets and the possibility of adverse regulatory or legal actions relating to our product candidates even if regulatory approval is obtained; our ability to commercialize and launch our product candidates at all or on a timely basis; our ability to obtain and maintain adequate protection of our intellectual property; our ability to manufacture our product candidates in commercial quantities, at an adequate quality or at an acceptable cost; acceptance of our product candidates by healthcare professionals and patients; the possibility that we may face third-party claims of intellectual property infringement; 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; delays in the launch of product candidates and generic drugs; intense competition in our industry; potential product liability claims; potential adverse federal, state, and local government regulation in the United States, Europe, or Israel; the impact of pandemics, such as COVID-19 (coronavirus); and loss or retirement of key executives and research scientists. These 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 4, 2021, and in 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 presentation. Any such forward-looking statements represent management’s estimates as of the date of this presentation. While we may elect to update such forward-looking statements at some point in the future, unless required by applicable law, we disclaim any obligation to do so, even if subsequent events cause our views to change. Thus, one should not assume that our silence over time means that actual events are bearing out as expressed or implied in such forward-looking statements.

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Strategic Partnership With Galderma

- Exclusive commercialization agreements in the US for TWYNEO and EPSOLAY
- Up to \$15 million in upfront and product approval payments (assuming 2021 approvals of both products)
- Tiered double-digit royalties (mid- to high-teen percentage) of net sales
- Up to an additional \$9 million in sales milestone payments

- Option to regain commercialization rights 5 years following first sale
- We expect that by then markets will be well-established
- EPSOLAY and TWYNEO are patent protected until 2040 and 2038, respectively

- Deal is immediately cash flow positive, with capital to be redeployed towards our pipeline



Our Dermatology Company

IPO

\$86.3M raised in February 2018

GALDERMA PARTNERSHIP

5-year license, with option to regain brands. EPSOLAY PDUFA goal date was set for April 26, 2021 (awaiting FDA's pre-approval inspection). TWYNEO PDUFA goal date set for August 1, 2021

PERRIGO PARTNERSHIP

Twelve 50/50 gross profit-sharing collaborations

Our Pipeline

ROFLUMILAST (SGT-510)

Our innovative investigational topical formulation of roflumilast (SGT-510) was found to be more effective than roflumilast cream, 0.3%, that was formulated by Sol-Gel according to conventional methods of cream formulation, in a human xenograft psoriasis animal model

ERLOTINIB (SGT-210)

Our proof-of-concept study for erlotinib gel (SGT-210) in palmoplantar keratoderma patients was completed and indicated a possible modest improvement. We plan to investigate higher concentrations of erlotinib

TAPINAROF (SGT-310)

We are currently developing an innovative investigational formulation of tapinarof (SGT-310) aiming to offer product formulation innovations and increased affordability for patients compared to the brand expected to be launched



Recent Updates

Financial Profile

01

Gross proceeds of \$86.3 million raised in IPO on February 5, 2018

02

Gross proceeds of \$11.5 and \$23 million raised in public follow-on offerings on August 12, 2019 and February 13, 2020, respectively

03

Additional \$5 million investment by controlling shareholder in April 2020

04

23,028,264 Ordinary Shares as of March 31, 2021

05

\$8.7 million net revenues from generic products in 2020 and \$0.7 million net revenues from generic products in 1Q/21

06

\$42.1 million in cash and investments as of May 31, 2021

07

Based on Galderma's upfront and milestone payments, we expect that our cash resources will enable funding of operational and capital expenditure requirements into the first quarter of 2023 (assuming timely approval of both products in 2021)



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