UNITED STATES
SECURITIES AND EXCHANGE COMMISSION
Washington, D.C. 20549

FORM 6-K

Report of Foreign Private Issuer
Pursuant to Rule 13a-16 or 15d-16
Under the Securities Exchange Act of 1934

For the month of August 2021

Commission File Number 001-38367

SOL-GEL TECHNOLOGIES LTD.

(Translation of registrant's name into English)

7 Golda Meir Street
Ness Ziona 7403650, Israel
(Address of principal executive offices)

Indicate by check mark whether the registrant files or will file annual reports under cover Form 20-F or Form 40-F.

Form 20-F ☒ Form 40-F ☐

Indicate by check mark if the registrant is submitting the Form 6-K in paper as permitted by Regulation S-T Rule 101(b)(1): ☐

Indicate by check mark if the registrant is submitting the Form 6-K in paper as permitted by Regulation S-T Rule 101(b)(7): ☐

INFORMATION CONTAINED IN THIS REPORT ON FORM 6-K

Sol-Gel Technologies Ltd. (the "Company") is posting on its website a corporate presentation.

Attached hereto and incorporated by reference in this Report on Form 6-K is the following exhibit:

Exhibit 99.1: Corporate Presentation.

SIGNATURES

Pursuant to the requirements of the Securities Exchange Act of 1934, the Registrant has duly caused this report to be signed on its behalf by the undersigned, thereunto duly authorized.

SOL-GEL TECHNOLOGIES LTD.

Date: August 4, 2021

By: /s/ Gilad Mamlok
Gilad Mamlok
Chief Financial Officer

Exhibit 99.1

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 forwa 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 Food and Drug Administration (FDA) approval and commercial launch of EPSOLAY®, commercial launch of TWYNEO, the strategic partnership with Galderma, progress on our innovative earlier stage programs, 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 expenditur 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 th following: risks relating to the timing of the PDUFA action date for EPSOLAY®, the timing of FDA approval, if any, of EPSOLAY; the risk that we will not receive 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 we don't progress on our innovative earlier stage programs, 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 expe 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 o 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 ke 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 forwar 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 require 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. These forward-looking statements should not be relied upon as representing our views as of any date subsequent to the dat of this presentation. This presentation contains trademarks, trade names, and service marks of other companies, which are the property of their respective owners. We do not intend our use or display of other parties' trademarks, trade names, or service marks to imply, and such use or display should not be construed to imply, a relationship with, or endorsement or sponsorship of us by, these other parties.

OVERVIEW

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 FDA Approved July 26, 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

U.S. COMMERCIAL PARTNERSHIP FOR EPSOLAY® AND TWYNEO®

STRATEGIC PARTNERSHIP WITH GLOBAL LEADER GALDERMA

- $8 million upfront payment received. Additional regulatory milestone payments of up to $7 million
- 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 at no cost to Sol-Gel

Cash-flow positive deal supporting high-value development pipeline

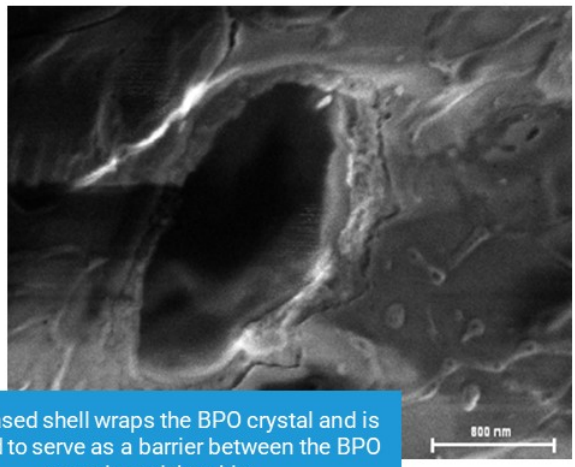
THE SCIENCE BEHIND OUR PROPRIETARY TECHNOLOGY

Aiming to provide effective and tolerable topical therapies to achieve local action

ENCAPSULATION IS DESIGNED TO ALLOW FOR CONTINUOUS FLOW

ENCAPSULATED BENZOYL PEROXIDE (E-BPO)

CRYO-SEM PICTURE



Silica-based shell wraps the BPO crystal and is intended to serve as a barrier between the BPO crystals and the skin

ENERGY-DISPERSIVE X-RAY SPECTROSCOPY MAPPING

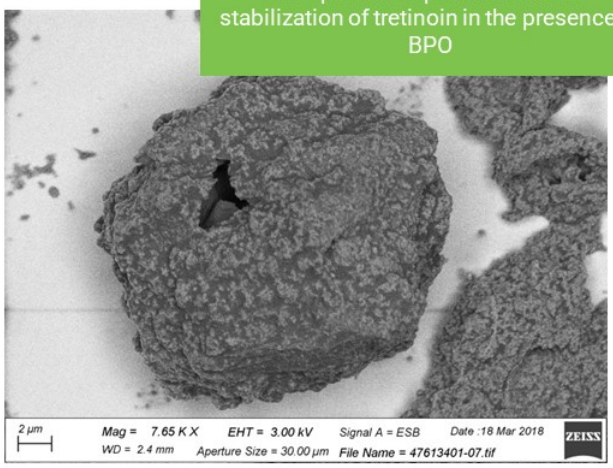
After application onto skin, BPO slowly migrates through the shell resulting in a continuous flow of BPO for up to 24 hours

ENCAPSULATION IS DESIGNED TO ENHANCE STABILITY

ENCAPSULATED TRETINOIN (E-TRETINOIN)

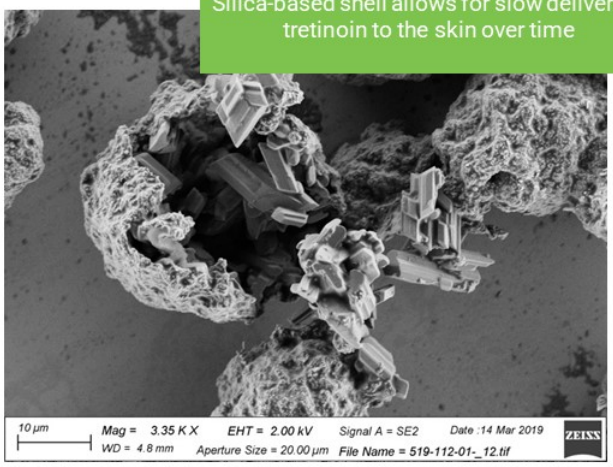
SEM PICTURE

Complete encapsulation allows stabilization of tretinoin in the presence of BPO



SEM PICTURE

Silica-based shell allows for slow delivery of tretinoin to the skin over time



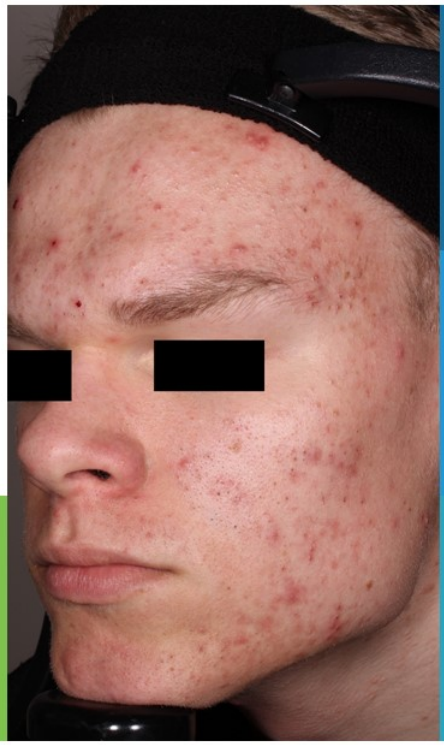
TWYNEO: FIRST PROPRIETARY DRUG APPROVAL

twyneo®
(tretinoin and benzoyl peroxide) cream, 0.1%/3%

LONG-TERM SAFETY STUDY

MULTIFACTORIAL DISEASE REQUIRING POWERFUL COMBINATION TREATMENTS

UNMET NEED IN ACNE VULGARIS



Acne Vulgaris

A multifactorial disease of the pilosebaceous unit, involving abnormalities in sebum production, follicular epithelial desquamation, bacterial proliferation, and inflammation

How is it Treated?

- Topical BPO, retinoids (such as tretinoin, adapalene), antibiotics, and their combinations
- Oral Isotretinoin and antibiotics

Current Treatment Shortfalls

- Insufficient efficacy negatively affects self-esteem
- Systemic side effects
- Contributes to antibiotic resistance

TWYNEO®: OUR FIRST BRANDED PRODUCT APPROVAL

(Tretinoin and Benzoyl Peroxide) Cream, 0.1%/3%

Indication: for the treatment of acne vulgaris in adults and pediatric patients nine years of age and older

- First acne treatment that contains a fixed-dose combination of tretinoin and benzoyl peroxide, which are separately encapsulated in silica using Sol-Gel's proprietary microencapsulation technology.
  - Tretinoin and benzoyl peroxide are widely prescribed as separate treatments for acne vulgaris; however, these products have not been available for simultaneous use in a fixed dose combination until the availability of TWYNEO.
- TWYNEO is protected until 2038 by granted patents and until 2041 by a pending patent application

SUBJECT 507-003 || 18 YEARS OLD | FEMALE | WHITE | NOT HISPANIC OR LATINO*

IMPROVEMENT IN SEVERE PATIENT

BASELINE

"Severe"; 29 inflamed lesions
31 non-inflamed lesions; 1 nodule

WEEK 12

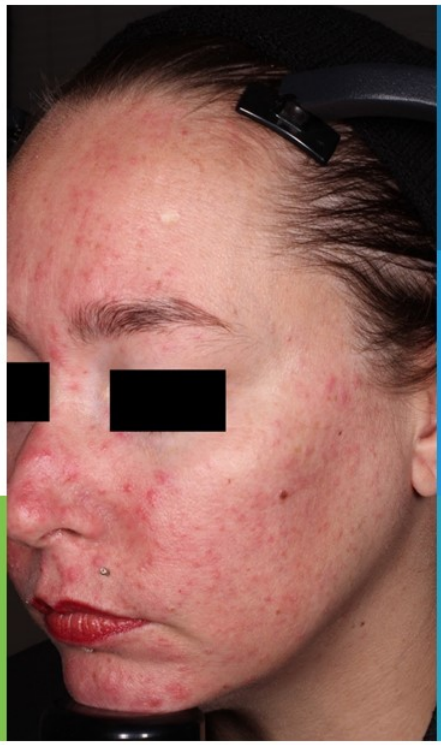
"Moderate"; 9 inflamed lesions
5 non-inflamed lesions; No nodules

* Individual results vary

THE CHALLENGE

CHRONIC CONDITION WITH POOR ADHERENCE TO CURRENT TREATMENTS

UNMET NEED IN PAPULOPUSTULAR ROSACEA



Papulopustular Rosacea

Chronic, inflammatory condition that primarily affects the face and is often characterized by flushing, redness, inflamed bumps, and pustules

How is it Treated?

- Topical antimicrobials (metronidazole, clindamycin)
- Topical anti-mite (ivermectin)
- Systemic antibiotics (minocycline, doxycycline)

Current Treatment Shortfalls

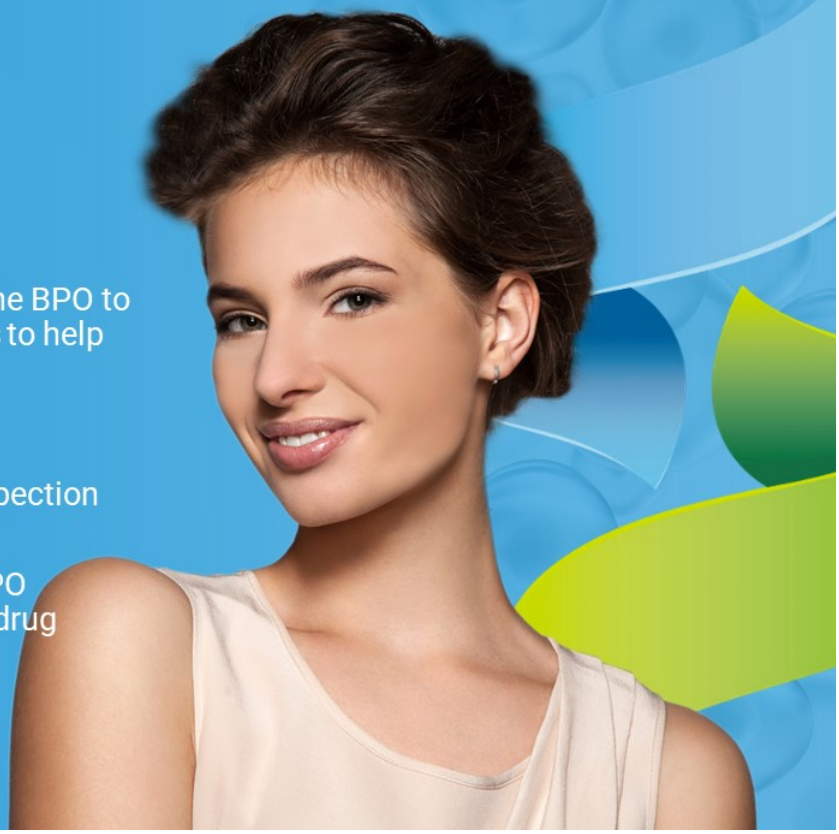
- Insufficient efficacy resulting in poor adherence
- Systemic side effects
- Contributing to antibiotic resistance

SOL-GEL SOLUTION*

EPSOLAY®

Benzoyl Peroxide Cream, 5%

- Encapsulation was designed to allow the BPO to slowly migrate from the microcapsules to help reduce irritation
- PDUFA goal date was set for April 26, 2021. Awaiting FDA's pre-approval inspection
- Potential to be the first single-active BPO approved by the FDA as a prescription drug product



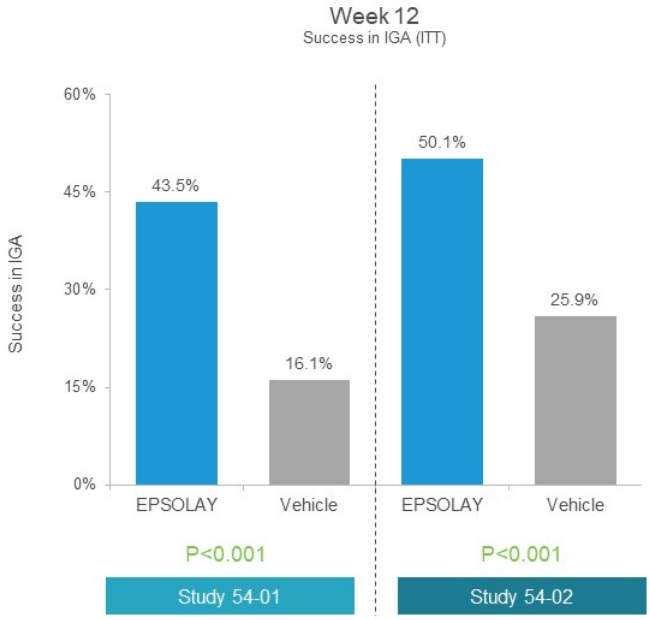
* EPSOLAY is investigational. Safety and efficacy have not been established

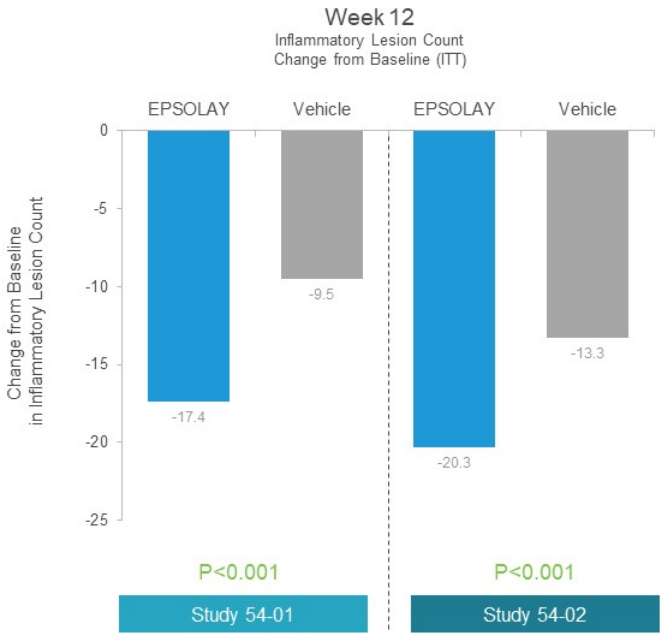
EPSOLAY®
PHASE III STUDIES

Two Parallel, Multi-Center, Double-Blinded,
Randomized, Vehicle-Controlled Studies, 2:1 Ratio, QD

PHASE III RESULTS

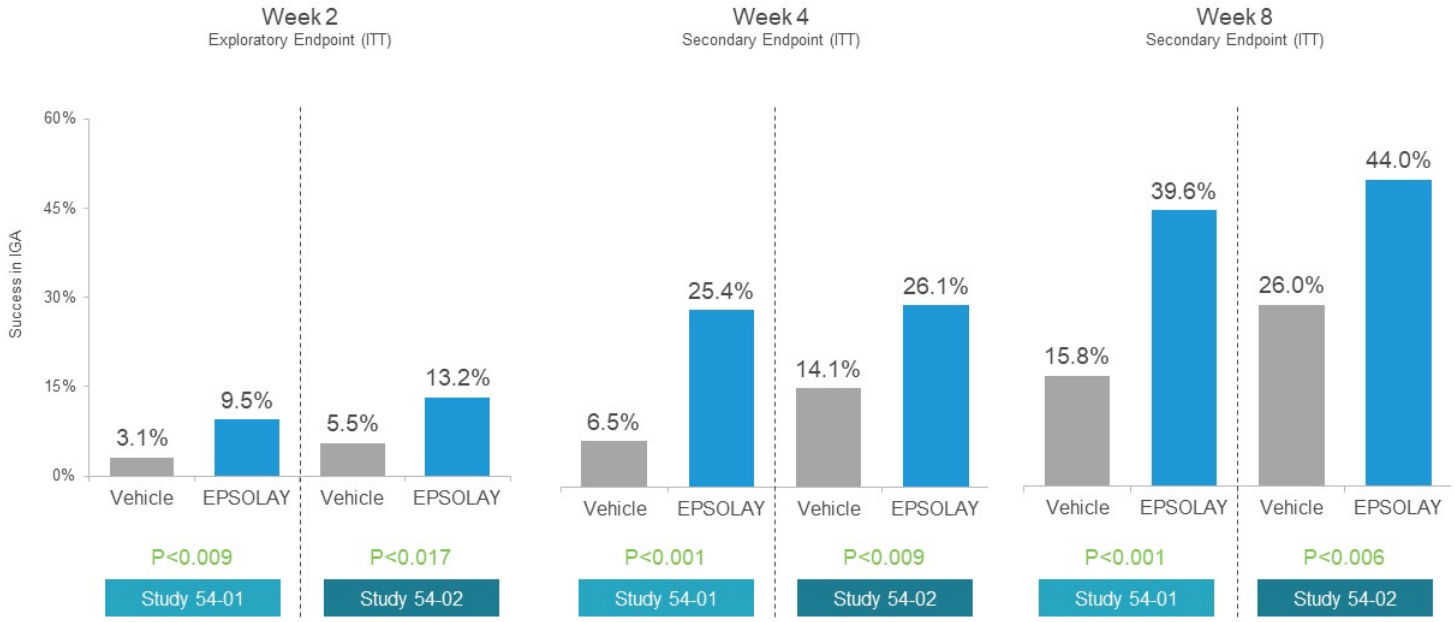
SUCCESS IN PRIMARY ENDPOINTS





SUCCESS IN IGA

IMPROVEMENT AS OF WEEK 2

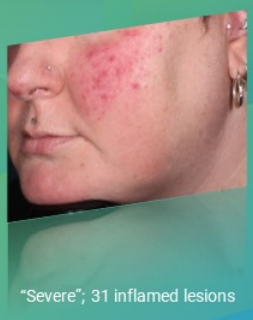
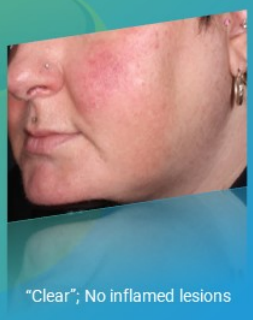
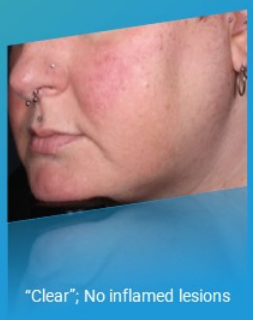
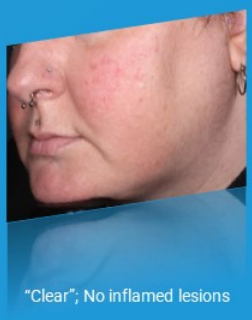
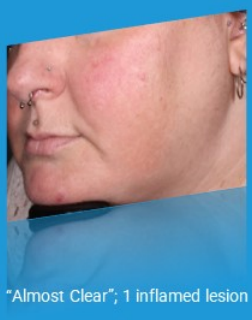


REDUCTION OF LESIONS

IMPROVEMENT AS OF WEEK 2

SUBJECT 116-009 || 41 YEARS OLD | FEMALE | WHITE | NOT HISPANIC OR LATINO*

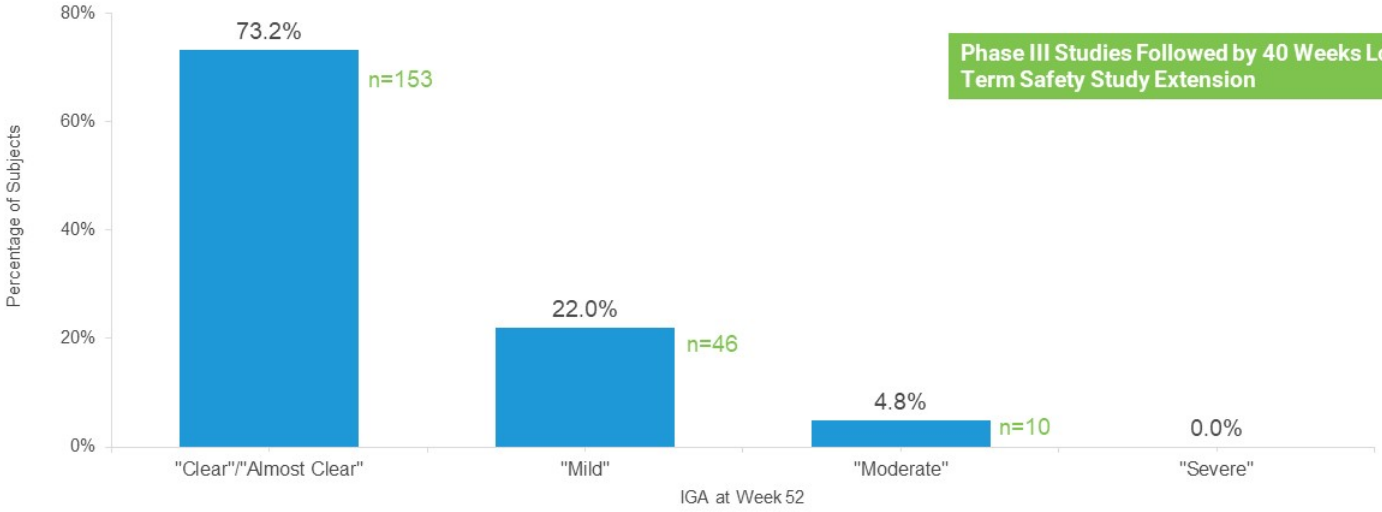
ONSET OF ACTION AS OF WEEK 2

BASELINE	WEEK 2	WEEK 4	WEEK 8	WEEK 12
				
"Severe"; 31 inflamed lesions	"Clear"; No inflamed lesions	"Clear"; No inflamed lesions	"Clear"; No inflamed lesions	"Almost Clear"; 1 inflamed lesion

* Individual results vary

LONG-TERM SAFETY STUDY

IMPROVEMENT IN IGA*



* This study was not designed for efficacy; however, efficacy was evaluated. Interpret results with caution

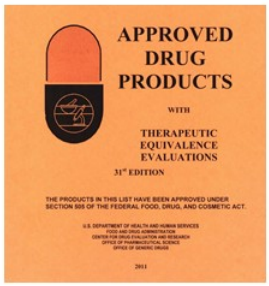
BROAD LONG-TERM INTELLECUAL PROPERTY ESTATE

25 patent applications for erlotinib, tapinarof and roflumilast in various skin conditions (as of February 26, 2021)

EPSOLAY is protected until 2040 by granted patents and until 2041 by a pending patent application

TWYNEO is protected until 2038 by granted patents and until 2041 by a pending patent application

LUCRATIVE GENERIC PRODUCTS AND PIPELINE



2015 — 12 collaborations with Perrigo with 50/50 gross profit sharing

March 2017 — In March 2017, Perrigo filed a Paragraph IV Certification for SOOLANTRA®

February 2019 — In February 2019, Perrigo launched acyclovir cream, 5%, developed in collaboration with Sol-Gel. This product generated $22.8 million in net revenues in 2019, $8.7 million in net revenues in 2020 and $0.7 million in net revenues in 1Q/21

January 2020 — In January 2020, Perrigo filed a Paragraph IV Certification for BRYHALI®

June 2020 — In June 2020, Perrigo was first-to-file a Paragraph IV Certification for DUOBRII®

June 2021 — In June 2021, Perrigo began selling a generic ivermectin cream, 1% product. In 2019, sales of the brand name product amounted to $192[1] million in the U.S.

Source: [1] IQVIA sale data

RECENT UPDATES

PIPELINE PROGRESS

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

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

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REVENUE STREAM FROM PARTNERSHIPS WILL SUPPORT DEVELOPMENT OF OUR INNOVATIVE PIPELINE

INNOVATIVE PIPELINE OF TOPICAL SKIN MEDICATIONS

Pipeline focused on large and attractive categories and two active moieties that already demonstrated positive Phase 3 results

RECENT UPDATES

FINANCIAL PROFILE

- Gross proceeds of $86.3 million raised in IPO on February 5, 2018
- Gross proceeds of $11.5 and $23 million raised in public follow-on offerings on August 12, 2019 and February 13, 2020, respectively
- Additional $5 million investment by controlling shareholder in April 2020
- 23,029,953 Ordinary Shares as of June 30, 2021
- $8.7 million net revenues from generic products in 2020 and $1.6 million net revenues from generic products in 1H/21
- $38.9 million in cash and investments as of June 30, 2021
- 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 EPSOLAY in 2021)


Sol-Gel
Advanced Topical Therapy
NASDAQ: SLGL