

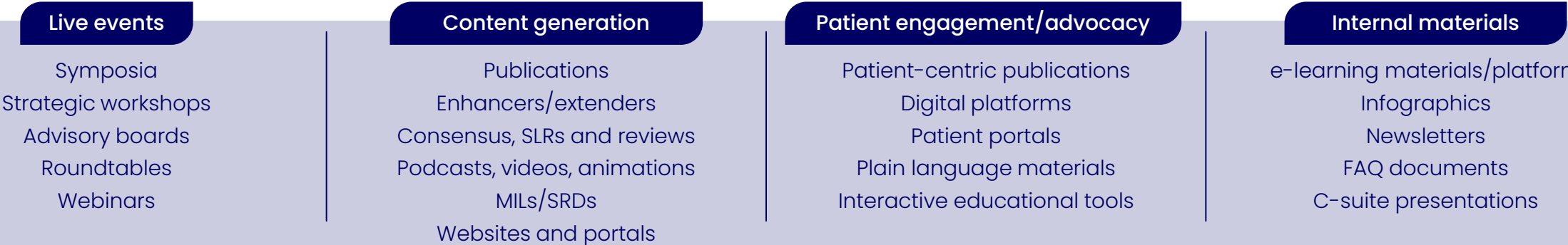
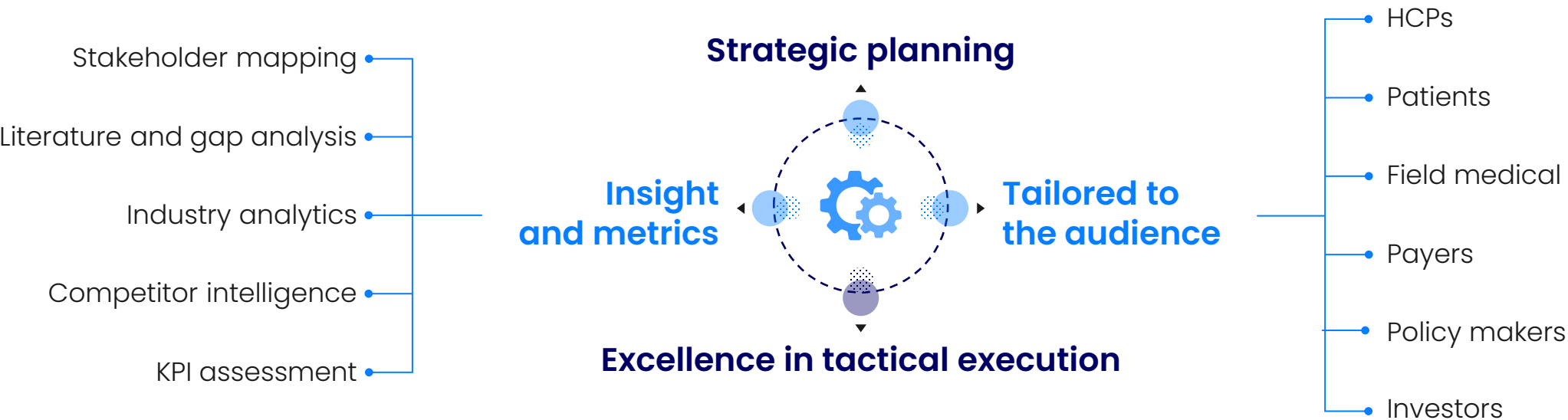
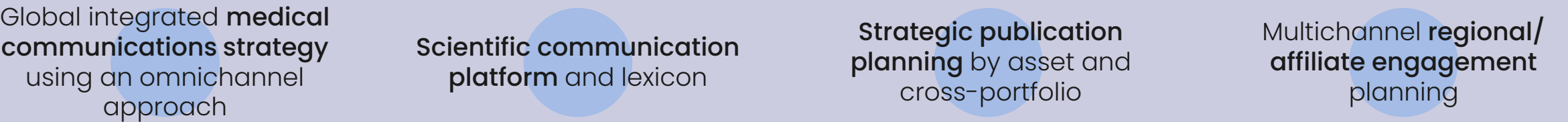
A woman's profile is shown in silhouette, looking towards the right. Her face is partially obscured by a series of glowing blue and purple light trails that curve around her head. Overlaid on her face and the light trails are numerous small, semi-transparent icons representing various aspects of technology, healthcare, and business, such as a heart with a pulse line, a car, a shopping cart, a smartphone, a magnifying glass, a bar chart, a globe, and a brain.

Accelerating
life-changing
solutions



<https://primeglobalpeople.com/>

We can support you across the spectrum of Scientific Communications





An elevated Publication Powerhouse

Heritage in strategy and execution, powered by evidence, data and technology



Interactive digital posters

Adding tislelizumab to chemotherapy significantly improved the survival of patients with advanced or metastatic esophageal squamous cell carcinoma, without compromising patient safety

Abstract #123

Lead Presenter: Jane Doe¹; Authors: Gregory House¹, Allison Cameron¹, Lisa Cuddy¹, James Wilson¹, Eric Foreman¹, Robert Chase¹, Chris Taub¹, Lawrence Kutner¹, Chi Park¹, Amber Volakis¹, Stacy Warner¹, and Edward Vogler¹; Affiliation: 1. Princeton Plainsboro Hospital.

The options for first-line treatment of advanced esophageal squamous cell carcinoma are scarce, and the outcomes remain poor

More than two-thirds of patients with esophageal cancer are diagnosed with advanced or metastatic disease.

Esophageal cancer is the 10th most common cancer and the 6th most prevalent cause of death due to cancer.

5-year survival rate is poor. Median overall survival (OS) rarely surpassing 10 months.

Tislelizumab is an anti-PD-1 antibody with antitumour activity in patients with esophageal squamous cell carcinoma

PD-1 is a T-cell receptor that recognizes healthy cells and shut down T-cell-mediated attacks, thereby preventing inappropriate immune responses damaging healthy tissues.

Tislelizumab is an anti-PD-1 antibody that blocks the PD-1 receptor, preventing it from sending the "shut down" signal to T-cells, allowing them to kill the cancer cells.

The RATIONALE-306 study assessed tislelizumab plus chemotherapy versus placebo plus chemotherapy as first-line treatment for advanced or metastatic esophageal squamous cell carcinoma

RATIONALE-306 is the first international study investigating the effects of an anti-PD-1 immunotherapy in combination with different chemotherapy regimens.

324 patients received tislelizumab + chemotherapy group.

649 patients received placebo + chemotherapy group.

162 patients received tislelizumab + chemotherapy group.

Efficacy was assessed using multiple measures

Primary Endpoints: Overall survival (OS) from start of study to death due to any cause.

Secondary Endpoints: Progression-free survival (PFS) from start of study to disease progression or death. Objective response rate (ORR) was defined as the proportion of patients with a complete or partial response to treatment.

Safety

Treatment-related treatment-emergent adverse events

Percentage of patients experiencing adverse events

12% and 5% of patients in the tislelizumab group and the placebo group remained on study treatment as of data cutoff

Treatment-emergent adverse events leading to death occurred in the same percentage (5%) of patients in both the experimental and placebo groups

Tislelizumab group

Placebo group

Efficacy

6.6 month improvement in median OS observed for groups with tislelizumab (0.66 hazard ratio)

2.7 month improvement in median PFS was observed with tislelizumab (0.60 hazard ratio)

ORR was higher in patients in the tislelizumab group when combined with either chemotherapy regimen

INTERACTIVE POSTER

VIEW WORLDWIDE VERSION OF POSTER

CONTACT AUTHOR HERE

READ FULL LANGUAGE SUMMARY

AUTHORS

Safety

Treatment-related treatment-emergent adverse events

Efficacy

6.6 month improvement in median OS

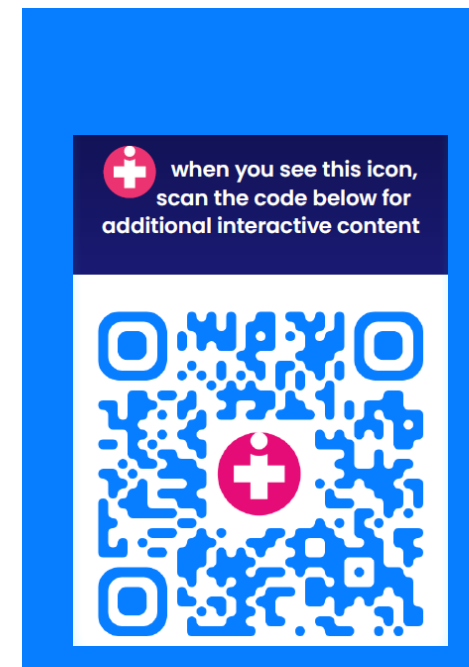
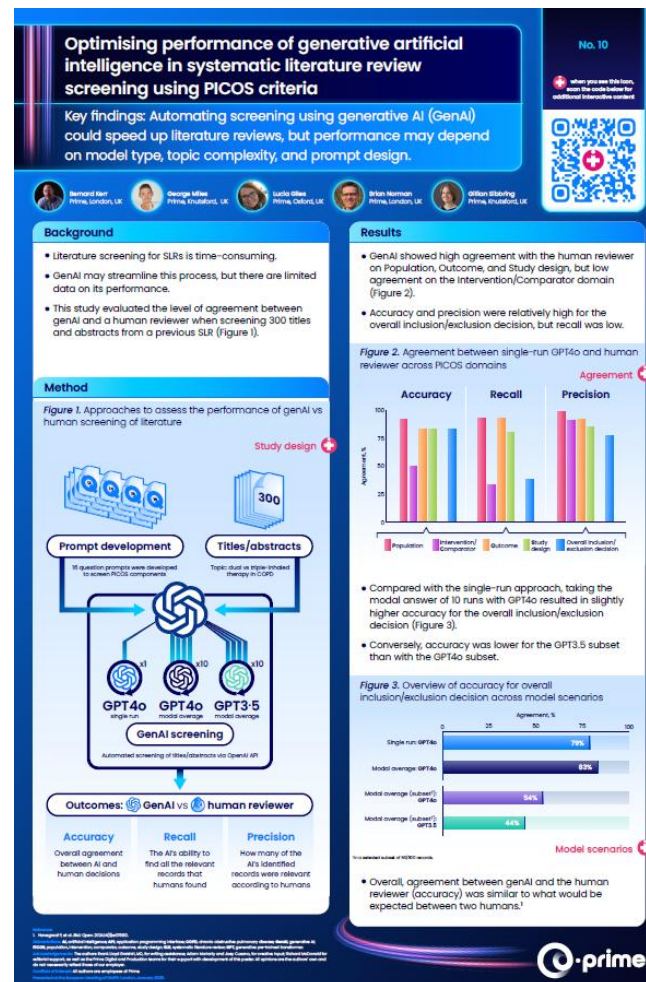
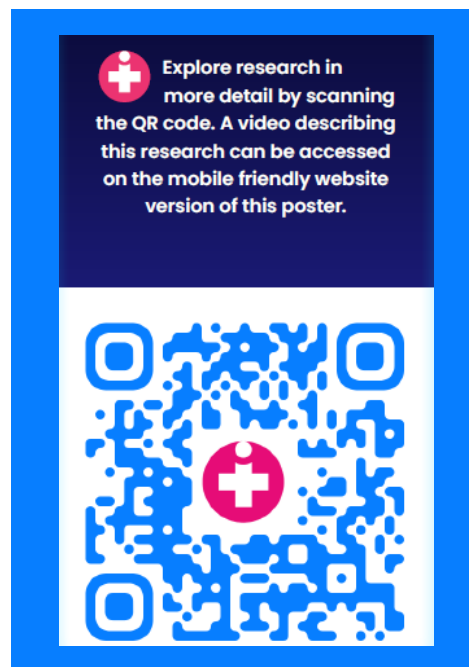
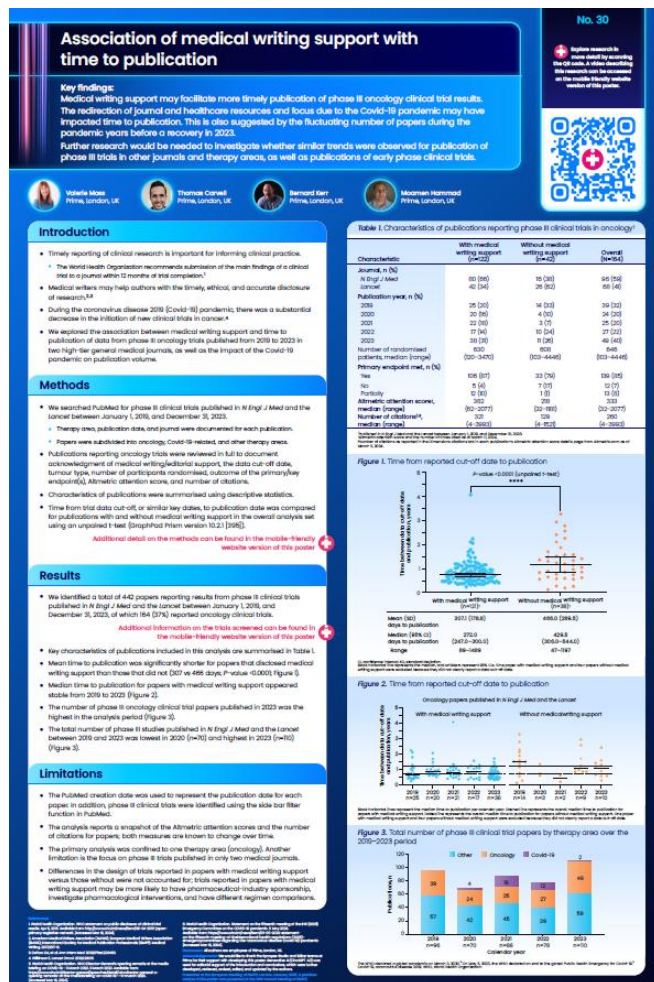
2.7 month improvement in median PFS

ORR was higher in patients in the tislelizumab group when combined with either chemotherapy regimen

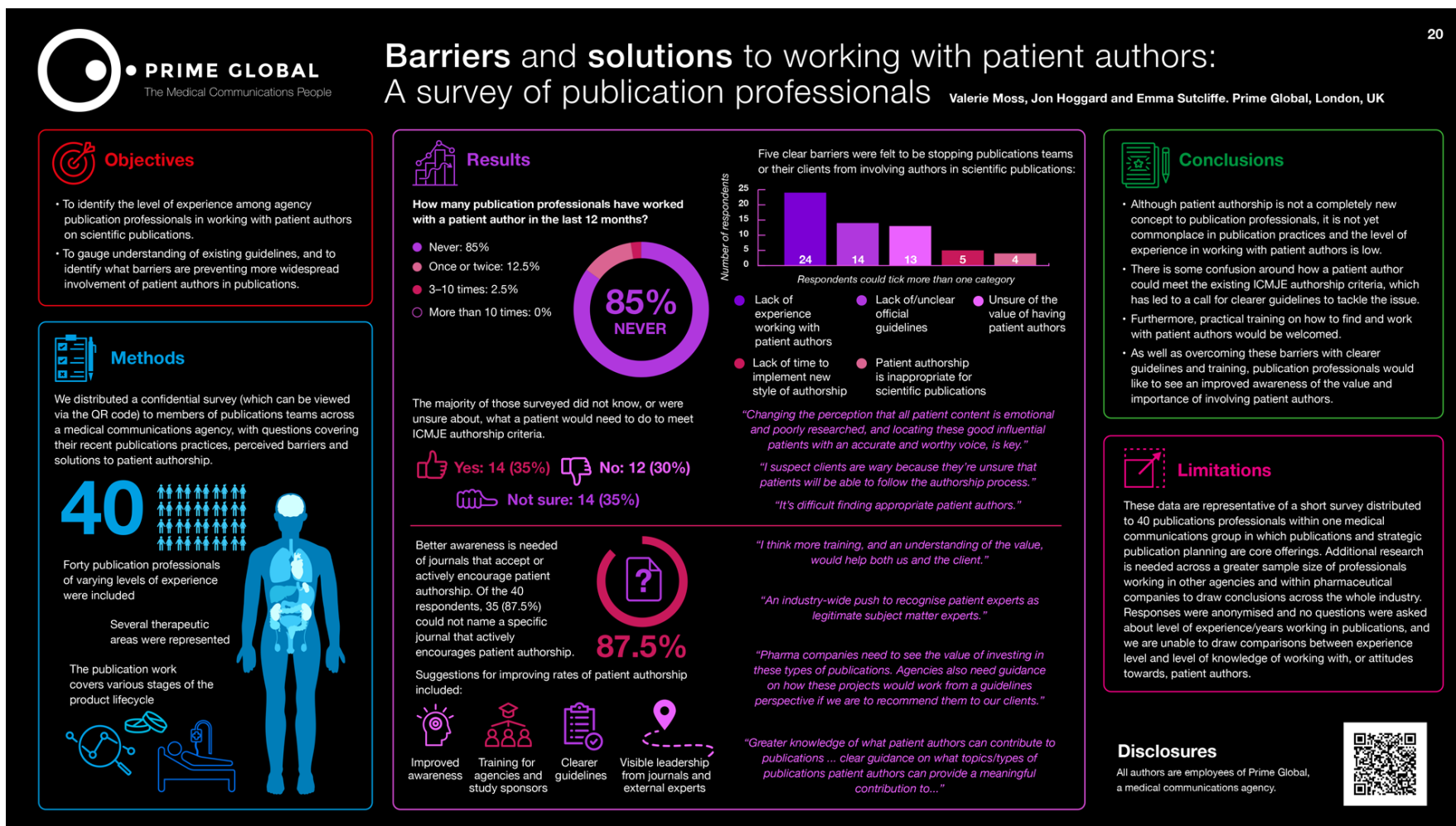
Please note that the version available here is for demonstration purposes only today and does not support full functionality

Scan Me

Our industry-leading digital posters



Bring poster data and content to life with Augmented Reality



AR overlay of presenter is available 24/7 to give a recorded introduction to the poster to in-person and remote attendees

Scan Me



Interactive publication content

frontiers | Frontiers in Oncology

TYPE Review
PUBLISHED 04 January 2023
DOI 10.3389/fonc.2022.975473

Check for updates

OPEN ACCESS

EDITED BY
Jessica Desiree Menis,
Integrated University Hospital Verona,
Italy

REVIEWED BY
Martin Proescholdt,
University Medical Center Regensburg,
Germany
Socrates Dokos,
University of New South Wales,
Australia

*CORRESPONDENCE
Milan J. Anadkat
manadkat@wustl.edu

[†]These authors share first authorship

SPECIALTY SECTION
This article was submitted to
Thoracic Oncology,
a section of the journal
Frontiers in Oncology

RECEIVED 22 June 2022
ACCEPTED 23 September 2022
PUBLISHED 04 January 2023

CITATION
Anadkat MJ, Lacouture M,
Friedman A, Horne ZD, Jung J,
Kaffenberger B, Kalmadi S,
Ovington L, Kotecha R, Abdullah H
and Grosso F (2023) Expert guidance
on prophylaxis and treatment of
dermatologic adverse events with
Tumor Treating Fields (TTFields) therapy
in the thoracic region.
Front. Oncol. 12:975473.
doi: 10.3389/fonc.2022.975473

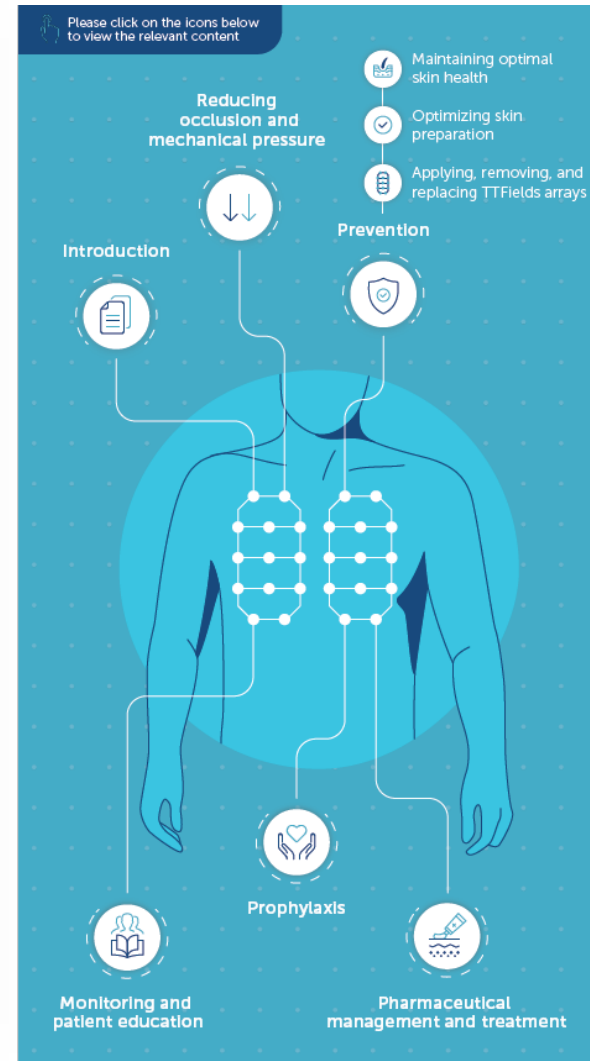
COPYRIGHT
© 2023 Anadkat, Lacouture, Friedman,
Horne, Jung, Kaffenberger, Kalmadi,
Ovington, Kotecha, Abdullah and
Grosso. This is an open-access article
distributed under the terms of the
Creative Commons Attribution License
(CC BY). The use, distribution or
reproduction in other forums is
permitted, provided the original
author(s) and the copyright owner(s)
are credited and that the source is
acknowledged.

Expert guidance on prophylaxis and treatment of dermatologic adverse events with Tumor Treating Fields (TTFields) therapy in the thoracic region

Milan J. Anadkat^{1*}, Mario Lacouture^{2†}, Adam Friedman³, Zachary D. Horne⁴, Jae Jung⁵, Benjamin Kaffenberger⁶, Sujith Kalmadi⁷, Liza Ovington⁸, Rupesh Kotecha⁹, Huda Ismail Abdullah¹⁰ and Federica Grosso¹¹

¹Division of Dermatology, Department of Medicine, Washington University, St. Louis, MO, United States, ²Dermatology Service, Department of Medicine, Memorial Sloan Kettering Cancer Center, New York, NY, United States, ³Division of Dermatology, Department of Medicine, George Washington University School of Medicine and Health Sciences, Washington, DC, United States, ⁴Department of Radiation Oncology, Allegheny Health Network Cancer Institute, Pittsburgh, PA, United States, ⁵Department of Dermatology, Norton Healthcare, Louisville, KY, United States, ⁶Wexner Medical Center, Ohio State University, Columbus, OH, United States, ⁷Oncology and Hematology Department, Ironwood Cancer & Research Center, Chandler, AZ, United States, ⁸Ovington & Associates, Walnutport, PA, United States, ⁹Miami Cancer Institute, Baptist Health South Florida, Miami, FL, United States, ¹⁰Novocure Inc., New York, NY, United States, ¹¹Mesothelioma Unit, SS Antonio e Biagio General Hospital, Alessandria, Italy

Tumor Treating Fields (TTFields) are electric fields, delivered via wearable arrays placed on or near the tumor site, that exert physical forces to disrupt cellular processes critical for cancer cell viability and tumor progression. As a first-in-class treatment, TTFields therapy is approved for use in newly diagnosed glioblastoma, recurrent glioblastoma, and pleural mesothelioma. Additionally, TTFields therapy is being investigated in non-small cell lung cancer (NSCLC), brain metastases from NSCLC, pancreatic cancer, ovarian cancer, hepatocellular carcinoma, and gastric adenocarcinoma. Because TTFields therapy is well tolerated and delivery is locoregional, there is low risk of additive systemic adverse events (AEs) when used with other cancer treatment modalities. The most common AE associated with TTFields therapy is mild-to-moderate skin events, which can be treated with topical agents and may be managed without significant treatment interruptions. Currently, there are no guidelines for



Bring opinion piece and review publications to life with enhanced supplemental content

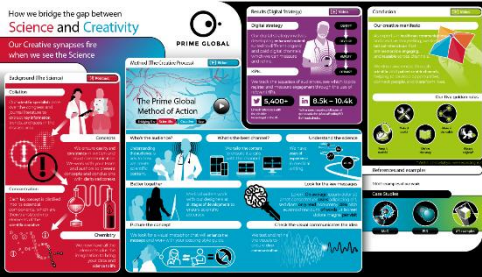
An interactive infographic overlaid with videos of the first author discussing the topics covered in the opinion piece, provided bite-sized content for readers to engage with

Scan Me

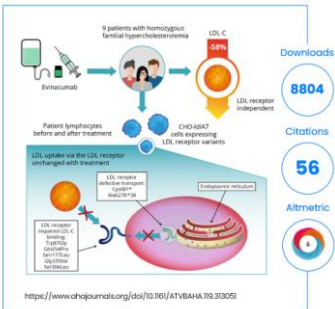


We utilize new technologies to produce publication extenders in formats preferred by your audience

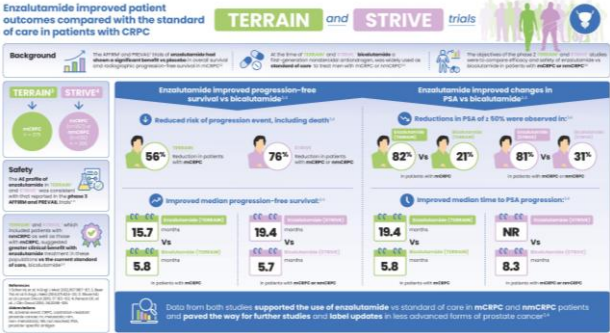
INTERACTIVE POSTERS



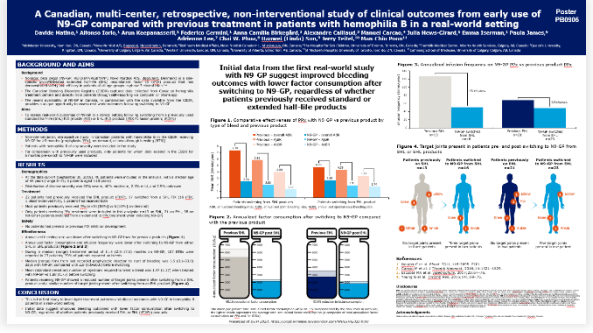
GRAPHICAL SUMMARIES



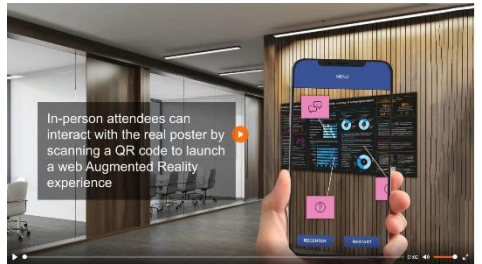
INFOGRAPHIC PLAIN LANGUAGE SUMMARIES



INFOGRAPHIC SCIENTIFIC POSTERS



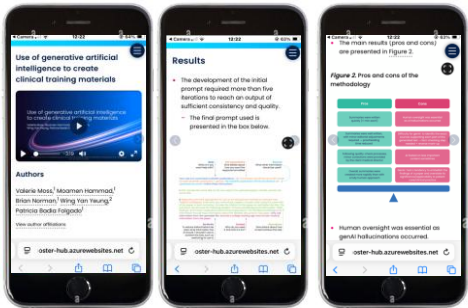
POSTER COMMENTING TOOL



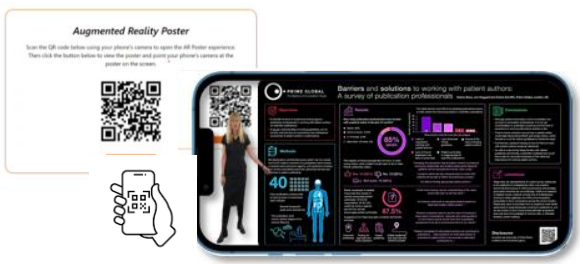
AUGMENTED REALITY VIDEOS



MOBILE-FRIENDLY POSTER FORMAT

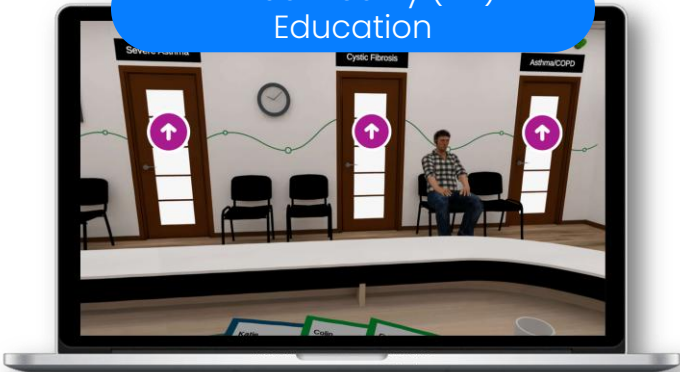


AUGMENTED REALITY AVATAR POSTER



Award winning digital creativity and innovation

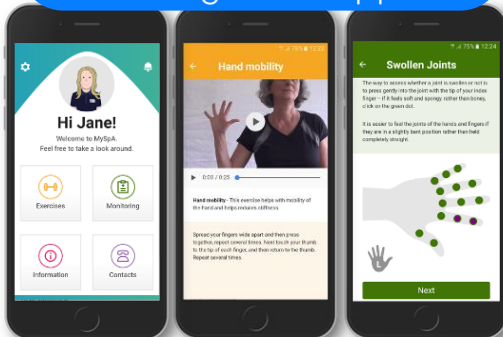
Virtual Reality (VR)
Education



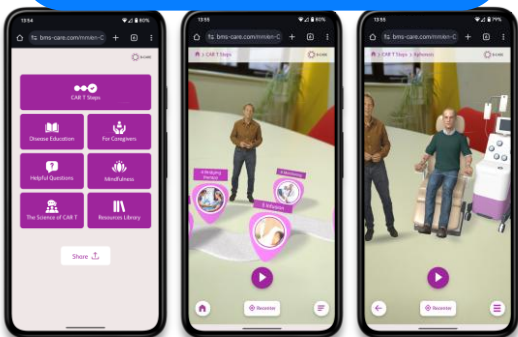
Augmented Reality (AR)
Content



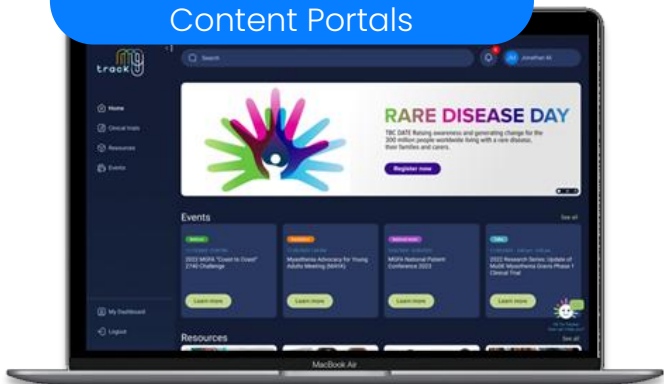
Patient Disease
Management Apps



Patient Education



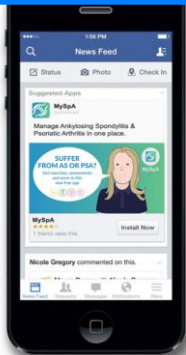
Medical Affairs
Content Portals



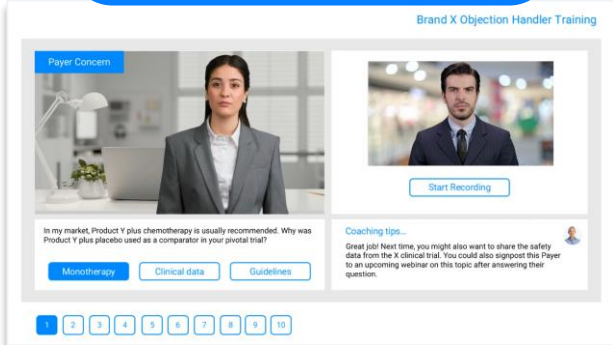
HCP & Patient Disease
Management Tools



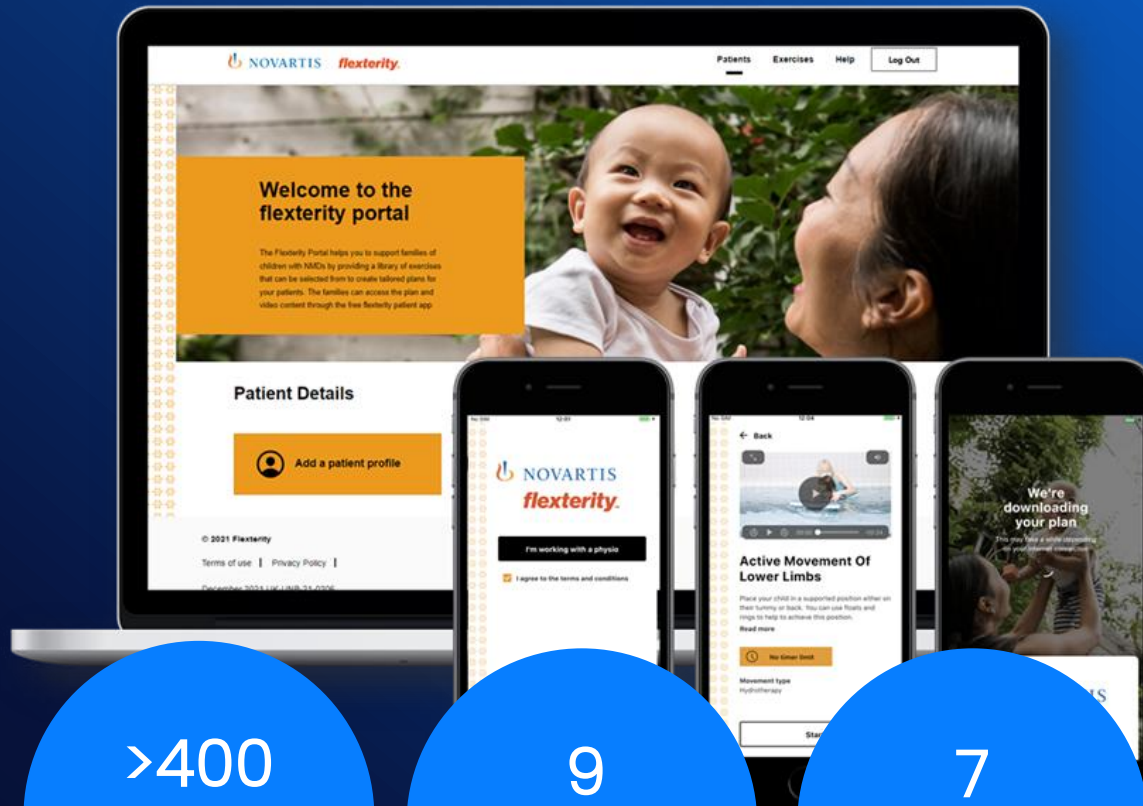
Search and Social Media
Campaigns



AI Powered Field Training
Tools



"Flexterity" app and physiotherapist portal – Spinal Muscular Atrophy



"We are delighted that the Flexterity app has been so well received, and we would like to thank everyone who contributed to this ground-breaking project which helps to enable access to physiotherapy for children affected by NMD. The future is now looking brighter for those families, whose need for this app was a priority in the NMD community."

Client, VP and General Manager

>400

UK Physios
registered in pilot

9

Countries

7

Languages

Winner

PMEA awards
Excellence in Patient
Education and
Support

Silver

PMEA awards
Excellence in
Innovation

Silver

PM Society Digital
awards
Patient
Programs



DermaPro – Revolutionizing PASI assessments in psoriasis



176%

HCP users per year
vs target

316%

User sessions per
year vs target

Gold

PM Society Digital
awards
HCP Education

Finalist

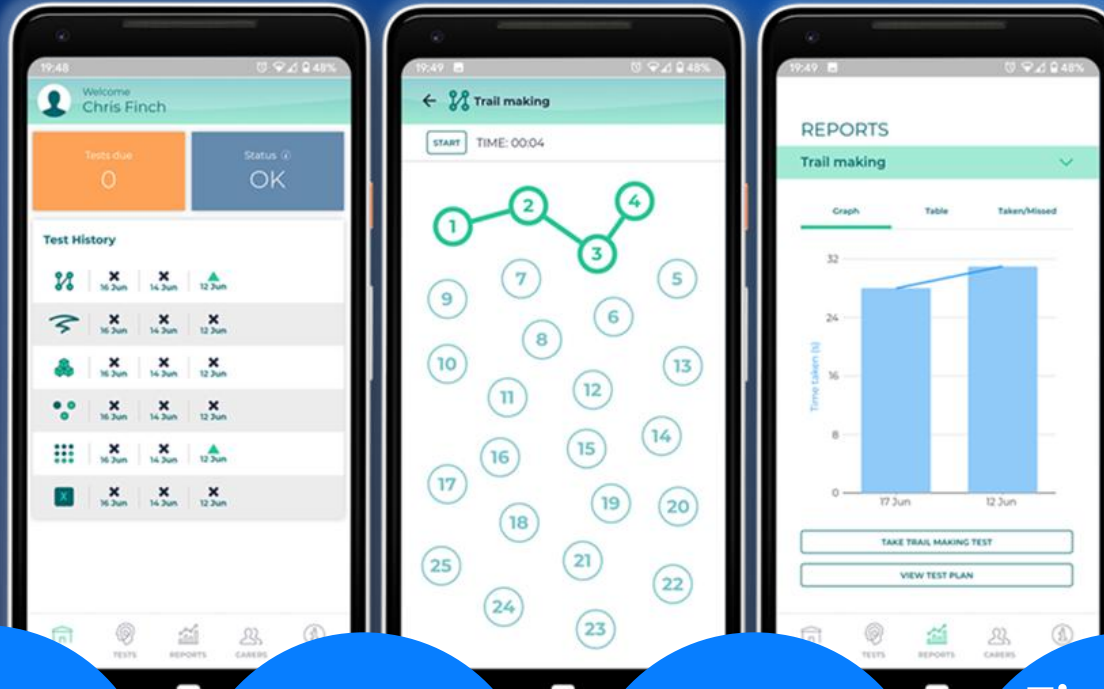
Communique
awards
Progress in
Healthcare and
Sci Comms

“DermaPro’s user-friendly interface and design have proved to be a great resource for colleagues who do not use PASI scoring very often. Its ability to precisely color in affected areas gives more consistent and reliable results, which provides confidence when discussing treatment options with patients.”

Dr Thomas King – Consultant Dermatologist, Sheffield Teaching Hospitals NHS Foundation Trust



Brain function monitoring medical device for early detection of Hepatic Encephalopathy (HE)



7

Countries

5

Languages

Gold

PM Society Digital
awards
Patient Programs

Finalist

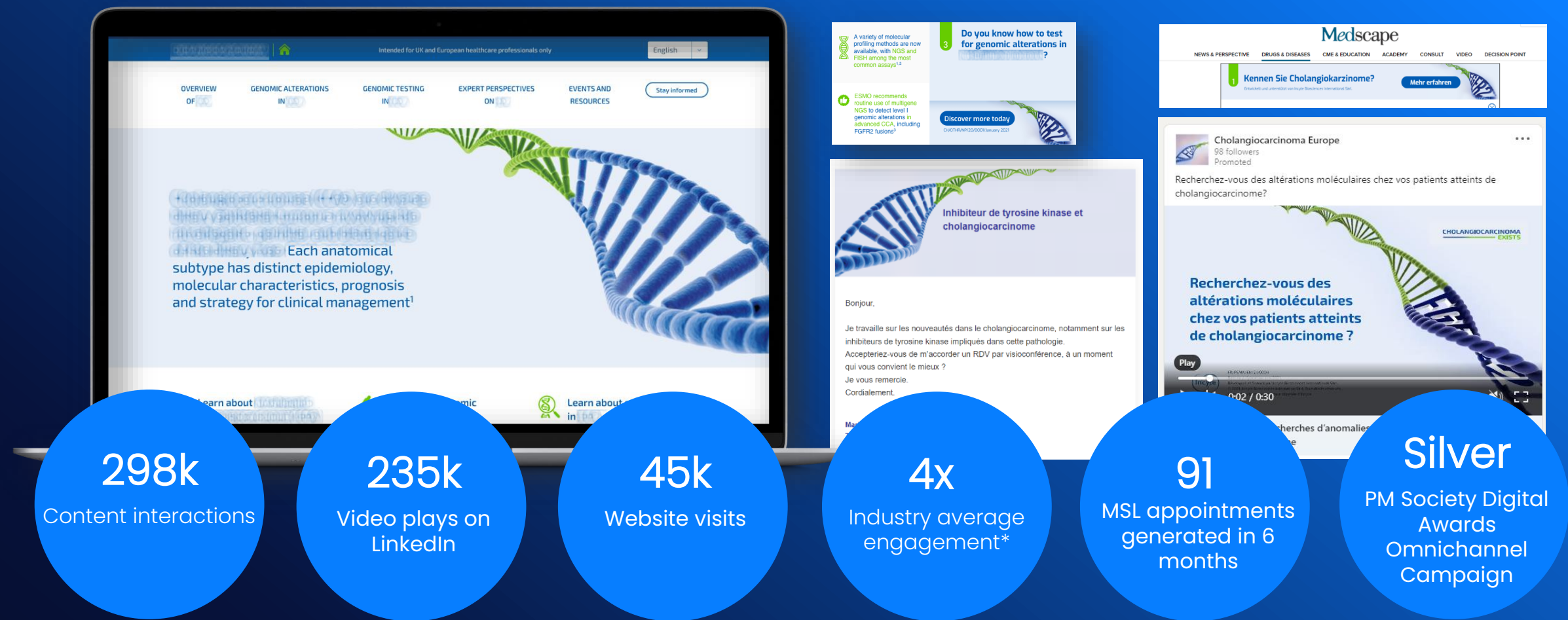
Communique
awards
Progress in
Healthcare and
Sci Comms

"This was our first software as a medical device. We chose Prime to be our partners because I know I can trust them to be honest and helpful throughout. We have a great relationship with the team that allows us to get things done efficiently as a true partnership."

**Client – Head of Global Digital
Centre of Excellence**



Rare Oncology launch – European omnichannel campaign across multiple affiliates



*Benchmarks provided by Four Health Media



For more information contact:

Valerie Moss, Chief Science Officer

valerie.moss@primeglobalpeople.com