novœure°

Novocure

July 2024





forward-looking statements

In addition to historical facts or statements of current condition, this presentation may contain forward-looking statements. Forward-looking statements provide Novocure's current expectations or forecasts of future events. These may include statements regarding anticipated scientific progress on its research programs, clinical trial progress, development of potential products, interpretation of clinical results, prospects for regulatory approval, manufacturing development and capabilities, market prospects for its products, coverage, collections from third-party payers and other statements regarding matters that are not historical facts. You may identify some of these forward-looking statements by the use of words in the statements such as "anticipate," "estimate," "expect," "project," "intend," "plan," "believe" or other words and terms of similar meaning. Novocure's performance and financial results could differ materially from those reflected in these forward-looking statements due to general financial, economic, environmental, regulatory and political conditions and other more specific risks and uncertainties facing Novocure such as those set forth in its Annual Report on Form 10-K filed on February 22, 2024, and subsequent filings with the U.S. Securities and Exchange Commission. Given these risks and uncertainties, any or all of these forward-looking statements may prove to be incorrect. Therefore, you should not rely on any such factors or forward-looking statements. Furthermore, Novocure does not intend to update publicly any forward-looking statement, except as required by law. Any forward-looking statements herein speak only as of the date hereof. The Private Securities Litigation Reform Act of 1995 permits this discussion.

The statements contained in this presentation are made as at the date of this presentation, unless some other time is specified in relation to them, and service of this presentation shall not give rise to any implication that there has been no change in the facts set out in this presentation since such date. Nothing contained in this presentation shall be deemed to be a forecast, projection or estimate of the future financial performance of Novocure, except where expressly stated.

As of the date of this presentation, Optune Gio and Optune Lua are FDA-approved for the treatment of adults with supratentorial glioblastoma, or GBM, and for the treatment of adults with malignant pleural mesothelioma or pleural mesothelioma (MPM), respectively, and the approval for use in other indications is not certain. Novocure can provide no assurances regarding market acceptance of Optune Gio or Optune Lua or their successful commercialization and can provide no assurances regarding the company's results of operations or financial condition in the future. This presentation is for informational purposes only and may not be relied upon in connection with the purchase or sale of any security.

2023 achievements and 2024 milestones ahead

DRIVING COMMERCIAL **ADOPTION**

ADVANCING CLINICAL TRIALS

DELIVERING PRODUCT INNOVATION

2023 **ACHIEVEMENTS**

France reimbursement achieved and launch

LUNAR data presented **LUNAR** U.S. FDA PMA. CE Mark and Japanese PMDA submitted **METIS** enrollment completed PANOVA-3 enrollment completed **TRIDENT** last patient enrolled*

New array launched in Europe **New array** FDA PMA supplement submitted

2024 **EXPECTED MILESTONES** **LUNAR** PMA approval **LUNAR** CE mark approval **NSCLC** launch

METIS top-line data released PANOVA-3 top-line data **LUNAR-2** open and enrolling **KEYNOTE D58** IND approved **New array** U.S. approval and launch



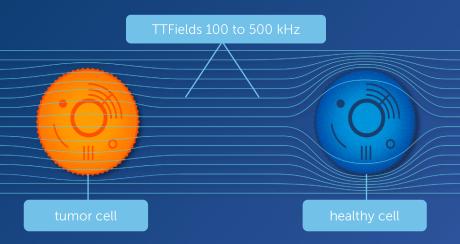
together with our patients, we strive to extend survival in some of the most aggressive forms of cancer





Tumor Treating Fields (TTFields) are selectively tuned electric fields that exert physical forces to kill cancer cells

TUNED ELECTRIC FIELDS DISRUPT PROTEINS DURING CELL DIVISION **CAUSING CANCER CELL DEATH**



Optune Gio® wearable cancer therapy system

DELIVERS CONTINUOUS TUMOR TREATING FIELDS THERAPY TO SOLID TUMORS



electric field generator and transducer arrays

patientforward®

Optune Gio® is intended as a treatment for adult patients (22 years of age or older) with histologically-confirmed glioblastoma multiforme (GBM).





strategy for long-term growth



drive commercial adoption

in approved indications



advance clinical trials

to reach new patient populations



deliver product innovation

to increase dose and duration of therapy



three focused objectives in 2024

GROW GBM

LAUNCH LUNG

DELIVER PIPELINE

patientforward® GBM, glioblastoma.



Optune Gio: established in glioblastoma



\$500M+

ANNUAL NET REVENUE (2023)

3,963

ACTIVE PATIENTS ON THERAPY*

NCCN Category 1

GUIDELINE RECOMMENDATION

30-40%

PENETRATION IN KEY COUNTRIES

reimbursement **ACROSS MAJOR GLOBAL MARKETS** robust intellectual property

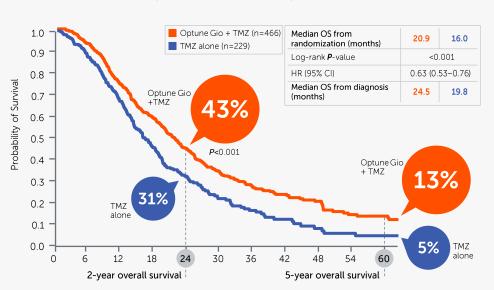
PORTFOLIO WITH MATERIAL PRODUCT DEVELOPMENTS



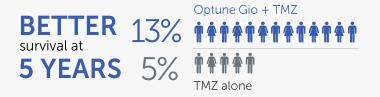
Optune Gio is proven to extend patient survival

EF-14 PHASE 3 PIVOTAL STUDY IN NEWLY DIAGNOSED GBM

Overall survival (5-year survival analysis)





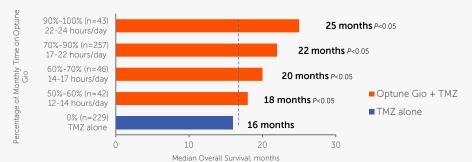


Overall Survival (months)

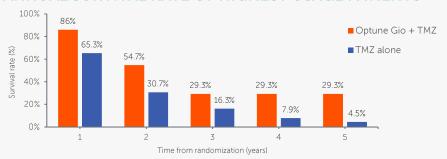


Optune Gio: greater exposure increased survival

MEDIAN OVERALL SURVIVAL BY PERCENT OF TIME ON OPTUNE GIO



ANNUAL SURVIVAL RATE OF HIGHEST USAGE PATIENTS



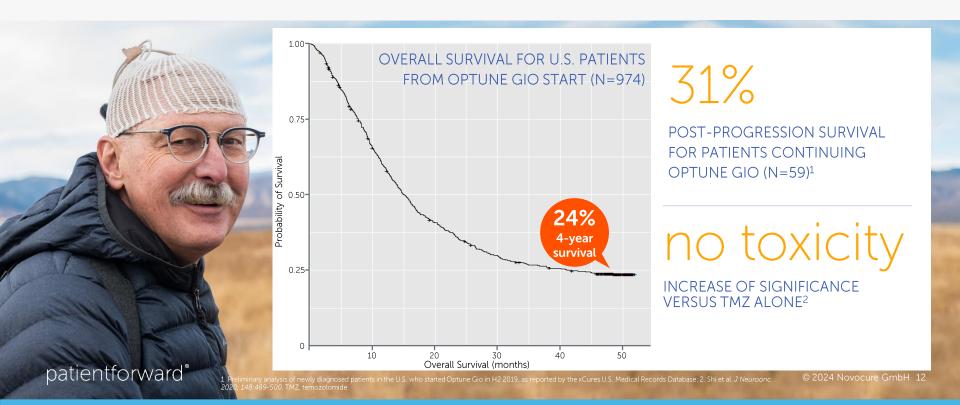
OF PATIENTS RECEIVED A SURVIVAL BENEFIT FROM OPTUNE GIO BECAUSE THEY USED IT >50% OF THE TIME

29.3% vs 4.5%

5-YEAR PROBABILITY OF SURVIVAL WITH 90% USAGE (N=43) VS SURVIVAL WITH TMZ ALONE

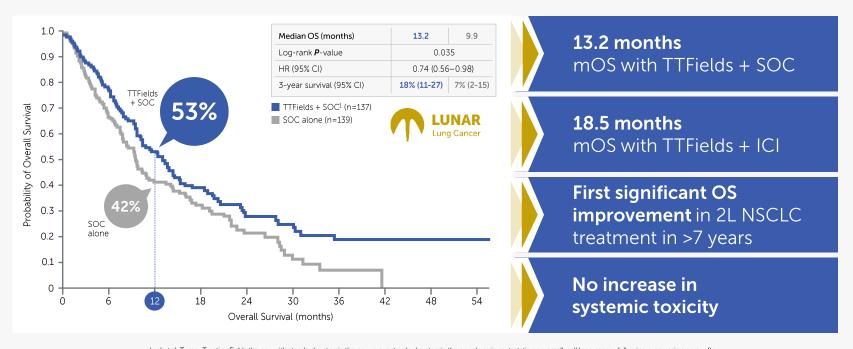


real-world evidence demonstrates long and durable survival benefit with Optune Gio, consistent with EF-14



phase 3 LUNAR trial in NSCLC met primary endpoint

STATISTICALLY SIGNIFICANT AND CLINICALLY MEANINGFUL IMPROVEMENT IN OS



significant opportunity to treat NSCLC patients

114K

stage IV NSCLC 1L patients in the U.S.

60%

receive 1L platinum-based chemotherapy

50%

progress and seek 2L treatment

~30,000

seek treatment for metastatic NSCLC post platinum

>7 YEARS

since any therapy has shown a significant improvement in overall survival in 2L NSCLC

preparing for 2024 NSCLC launch

2023 2024 2025 COMMERCIAL LAUNCH ✓ HCP and patient campaigns **REGULATORY PATHWAY** ✓ DTC campaign ✓ Technical file submitted in EU CLINICAL DATA ✓ Global advisory boards ✓ PMA accepted for review at FDA ✓ Announced top-line results ✓ KOL engagements ✓ Application submitted to Japan PMDA ✓ Data at ASCO Launch in U.S. and Germany ✓ Day 100 Meeting with FDA ✓ Published in Lancet Oncology Establish reimbursement CE Mark (expected 2H 2024) ✓ Data at ESMO, WCLC FDA PMA (expected 2H 2024)

platform technology driving robust clinical pipeline

		PHASE 3	PHASE 2
	GLIOBLASTOMA	TRIDENT TTFields therapy + TMZ + radiation treating ndGBM	
		KEYNOTE D58 TTFields therapy + pembrolizumab + TMZ treating ndGBM	
1	NON-SMALL CELL LUNG CANCER	METIS TTFields monotherapy treating brain metastases from NSCLC	KEYNOTE-B36 TTFields therapy + pembrolizumab treating 1L advanced or metastatic NSCLC
		LUNAR-2 TTFields + pembrolizumab + chemotherapy treating 1L metastatic NSCLC	LUNAR-4 TTFields + ICI treating 2L metastatic NSCLC following prior ICI treatment
e	PANCREATIC CANCER	PANOVA-3 TTFields therapy + nab-paclitaxel + gemcitabine treating 1L locally advanced pancreatic cancer	PANOVA-4 TTFields therapy + atezolizumab + nab-paclitaxel + gemcitabine treating 1L metastatic pancreatic cancer



significant pipeline catalysts on foundation of positive cashflow business

METIS DATA

LUNG APPROVALS & LAUNCH

PANOVA-3 DATA

PROFITABLE GBM BUSINESS

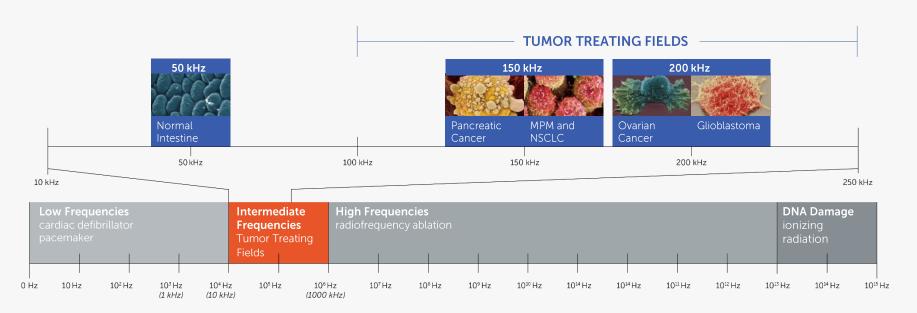


appendix





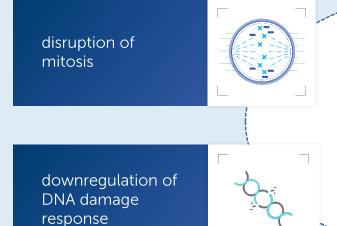
therapy is frequency-tuned to target dividing cancer cells



MPM: malignant pleural mesothelioma NSCLC: non-small cell lung cancer



TTFields have multiple, distinct mechanisms of action



mechanisms work together to selectively target and disrupt the progression of cancer cells, which can lead to their death

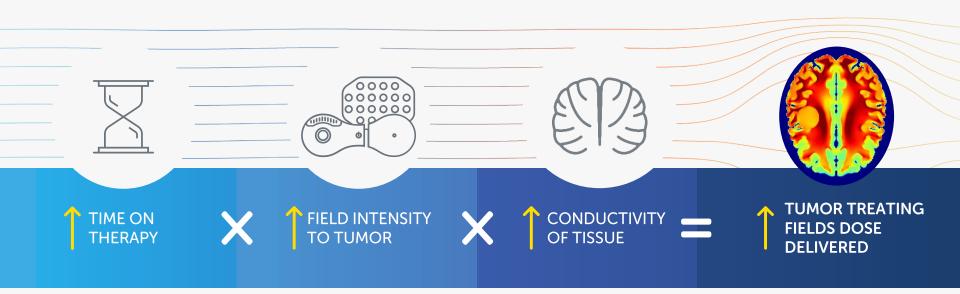


interference of cell movement and migration



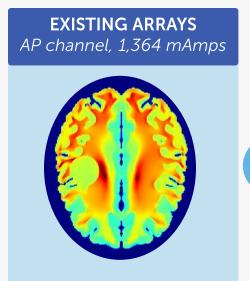
downstream enhancement of antitumor immunity

optimized dose delivered can lead to increased efficacy

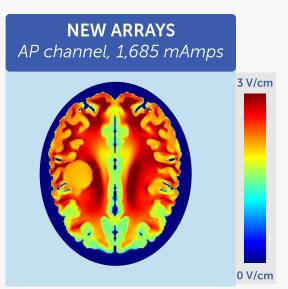




new lighter, thinner arrays deliver greater intensity



VS.





PMA SUPPLEMENT SUBMITTED IN Q4 2023

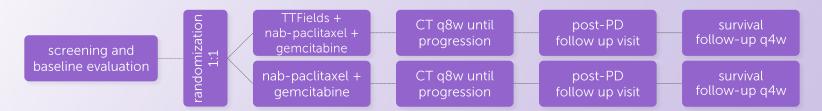
ongoing trial designs





PANOVA-3: phase 3 trial in locally advanced pancreatic cancer

OPEN-LABEL, RANDOMIZED TRIAL DESIGN¹



- 556 patients with 18-month minimum follow-up
- Primary endpoint: overall survival
- Enrollment complete (February 2023)
- Top-line data anticipated in Q4 2024



PANOVA-4: phase 2 trial in metastatic pancreatic cancer

PILOT, SINGLE-ARM TRIAL DESIGN¹



- 76 patients with 12-month minimum follow-up
- Primary endpoint: disease control rate
- Screening and enrollment ongoing

TRIDENT: phase 3 trial in newly diagnosed glioblastoma

OPEN-LABEL, RANDOMIZED TRIAL DESIGN¹



- 950 patients with 24-month minimum follow-up

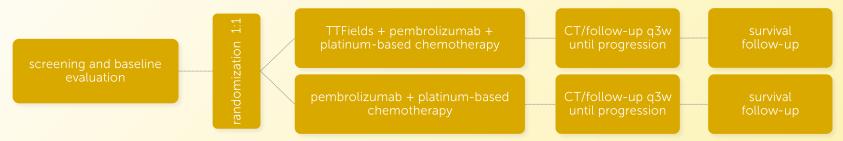
- Data anticipated in 2026





LUNAR-2: phase 3 trial in metastatic non-small cell lung cancer

OPEN-LABEL RANDOMIZED TRIAL DESIGN

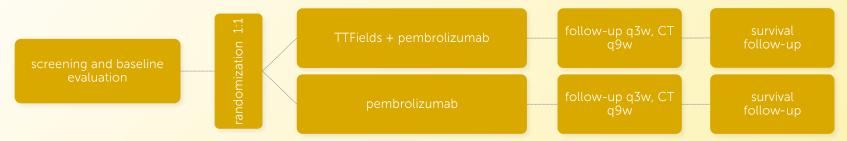






KEYNOTE B36: phase 2 trial in locally advanced or metastatic non-small cell lung cancer

OPEN-LABEL RANDOMIZED TRIAL DESIGN¹



- Screening and enrollment ongoing





Optune Lua® and Optune Gio® indications for use and important safety information

INDICATIONS

- Optune Gio is intended as a treatment for adult patients (22 years of age or older) with histologically-confirmed glioblastoma multiforme (GBM).
 - Optune Gio with temozolomide is indicated for the treatment of adult patients with newly diagnosed, supratentorial glioblastoma following maximal debulking surgery, and completion of radiation therapy together with concomitant standard of care chemotherapy.
 - For the treatment of recurrent GBM, Optune Gio is indicated following histologically-or radiologically-confirmed recurrence in the supratentorial region of the brain after receiving chemotherapy. The device is intended to be used as a monotherapy and is intended as an alternative to standard medical therapy for GBM after surgical and radiation options have been exhausted.
- Optune Lua is indicated for the treatment of adult patients with unresectable, locally advanced or metastatic, malignant pleural mesothelioma (MPM) to be used concurrently with pemetrexed and platinum-based chemotherapy.

CONTRAINDICATIONS

- Do not use Optune Gio in patients with GBM with an implanted medical device, a skull defect (such as, missing bone with no replacement), or bullet fragments. Use of Optune Gio together with skull defects or bullet fragments has not been tested and may possibly lead to tissue damage or render Optune Gio ineffective. Do not use Optune Lua in patients with MPM with implantable electronic medical devices, such as pacemakers or implantable automatic defibrillators, etc.
- Use of Optune Gio for GBM or Optune Lua for MPM together with implanted electronic devices has not been tested and may lead to malfunctioning of the implanted device.
- Do not use Optune Gio for GBM or the Optune Lua for MPM in patients known to be sensitive to conductive hydrogels. Skin contact with the gel used with Optune Gio or Optune Lua may commonly cause increased redness and itching and may rarely lead to severe allergic reactions such as shock and respiratory failure.





Optune Lua® and Optune Gio® indications for use and important safety information

WARNINGS AND PRECAUTIONS

- Optune Gio and Optune Lua can only be prescribed by a healthcare provider that has completed the required certification training provided by Novocure[®].
- The most common (>10%) adverse events involving Optune Gio in combination with chemotherapy in patients with GBM were thrombocytopenia, nausea, constipation, vomiting, fatigue, convulsions, and depression.
- The most common (>10%) adverse events related to Optune Gio treatment alone in patients with GBM were medical device site reaction and headache. Other less common adverse reactions were malaise, muscle twitching, and falls related to carrying the device.
- The most common (>10%) adverse events involving Optune Lua in combination with chemotherapy in patients with MPM were anemia, constipation. nausea, asthenia, chest pain, fatigue, device skin reaction, pruritus, and cough.
- Other potential adverse effects associated with the use of Optune Lua include: treatment related skin toxicity, allergic reaction to the plaster or to the gel, electrode overheating leading to pain and/or local skin burns, infections at sites of electrode contact with the skin, local warmth and tingling sensation beneath the electrodes, muscle twitching, medical site reaction and skin breakdown/skin ulcer.
- If the patient has an underlying serious skin condition on the treated area, evaluate whether this may prevent or temporarily interfere with Optune Gio or Optune Lua treatment.
- Do not prescribe Optune Gio or Optune Lua for patients that are pregnant, you think might be pregnant or are trying to get pregnant, as the safety and effectiveness of Optune Gio and Optune Lua in these populations have not been established.
- Please go to OptuneGio.com to see the Optune Gio Instructions For Use (IFU) for complete information regarding the device's indications, contraindications, warnings, and precautions.
- Please go to OptuneLua.com to see the Optune Lua IFU for complete information regarding the device's indications, contraindications, warnings, and precautions.