

**AWARD NUMBER:** W81XWH-21-1-0057

**TITLE:** Precision Antibody-Based Synergistic Drug Combinations for Refractory Triple-Negative Breast Cancer

**PRINCIPAL INVESTIGATOR:** Dr. Leif Ellisen, MD, PhD

**CONTRACTING ORGANIZATION:** MASSACHUSETTS GENERAL HOSPITAL, THE  
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<b>14. ABSTRACT</b> Metastatic triple-negative breast cancer (mTNBC) represents an unmet clinical need. Our group recently led the clinical development of a new therapy for TNBC called sacituzumab govitecan (SG). Herein, we propose to capitalize on this treatment innovation. First is a laboratory-based component that tests the mechanisms and effectiveness of specific drug combinations with SG, asking in part whether the combinations will work in multiple different TNBC models. Second is an animal-based component, in which we will test how effective the most promising combinations with SG are, how well-tolerated these combinations are, and whether staggered dosing schedules might preserve efficacy while minimizing side effects. Third is analysis of specimens from SG-treated patients, testing predicted biomarkers of response and resistance to SG, and using molecular analysis to discover new biomarkers. This analysis will highlight resistance mechanisms to SG and will help guide selection of TNBC patients who will benefit most from either single-agent SG or the new combinations.					
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**1. INTRODUCTION:** *Narrative that briefly (one paragraph) describes the subject, purpose and scope of the research.*

Our group recently led the clinical development of a highly promising new therapy for TNBC called sacituzumab govitecan (SG). We seek to capitalize on this treatment innovation by identifying select targeted drugs for combination with SG that i) will uniquely complement/enhance its efficacy and ii) can be rapidly moved into clinical trials for TNBC patients. First is a laboratory-based component that tests the mechanisms and effectiveness of specific drug combinations with SG, asking in part whether the combinations will work in multiple different TNBC models. This is a critical test since TNBCs from different patients vary substantially in their behavior. Second is an animal-based component, in which we will test how effective the most promising combinations with SG are against TNBC tumors, how well-tolerated these combinations are, and whether staggered dosing schedules might preserve efficacy while minimizing side effects. Third is analysis of specimens from SG-treated patients, testing predicted biomarkers of response and resistance to SG, and using molecular analysis to discover new biomarkers.

**2. KEYWORDS:** *Provide a brief list of keywords (limit to 20 words).*

Triple-negative breast cancer; topoisomerase 1 inhibitor; antibody-drug conjugate; TROP2; Combination therapy; cancer therapy resistance;

**3. ACCOMPLISHMENTS:**

**What were the major goals of the project?**

***Aim 1: Establish the Optimal Drug Sequencing and Therapeutic Window for SG/PARPi Combination Therapy***

Major Task 1: Systematic analysis of staggered dosing of SG and PARPi in TNBC cells versus normal cells to gauge optimal sequencing in vitro

Major Task 2: Perform direct comparison of simultaneous and staggered dosing in vivo, aiming to maximize tumor kill while mitigating systemic toxicity

Major Task 3: Test hypothesized biomarkers of SG versus SG/PARPi sensitivity to identify patients requiring combination therapy

***Aim 2: Discover combination therapy strategies that address TNBC inter-tumor heterogeneity and SG resistance***

Major Task 4: Carry out focused secondary CRISPR screens to reveal inter-tumor heterogeneity of response/resistance pathways to SG in TNBC and the specific pathways related to the ADC mechanism versus SN-38

Major Task 5: Identify novel biomarkers and targetable resistance mechanism to SG in patients with mTNBC

Major Task 6: Test selected synergistic pathways including checkpoint kinases and CDKs in gain and loss of function studies in vitro & in vivo

***Aim 3: Employ in vivo and patient-derived models for pre-clinical trials of novel synergistic combinations with SG***

Major Task 7: Establish in vivo efficacy and tolerability of select SG/clinical drug combinations selected based on Aim 2 using in vivo models

Major Task 8: Test promising combinations in organoid and PDX models from TNBC patients exhibiting resistance to SG and SG/PARPi

## What was accomplished under these goals?

We are pleased to share the accomplishments under this award to date. In order to facilitate review, accomplishments are organized and enumerated based on tasks corresponding to the approved SOW for this award. Accomplishments are supported by figures and graphs as appropriate. Note: the Principal Investigator responsible for each Major Task is listed for each instance.

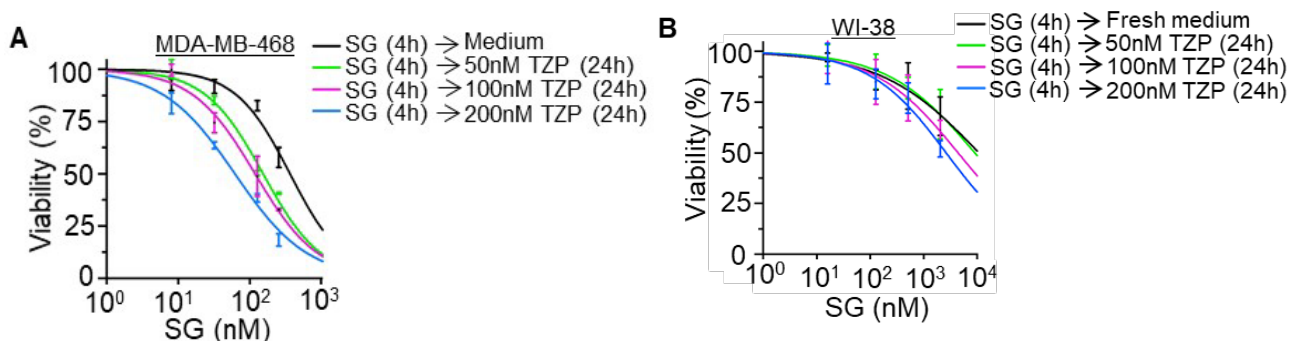
### *Aim 1: Establish the Optimal Drug Sequencing and Therapeutic Window for SG/PARPi Combination Therapy*

Major Task 1 (Ellisen): Systematic analysis of staggered dosing of SG and PARPi in TNBC cells versus normal cells to gauge optimal sequencing in vitro

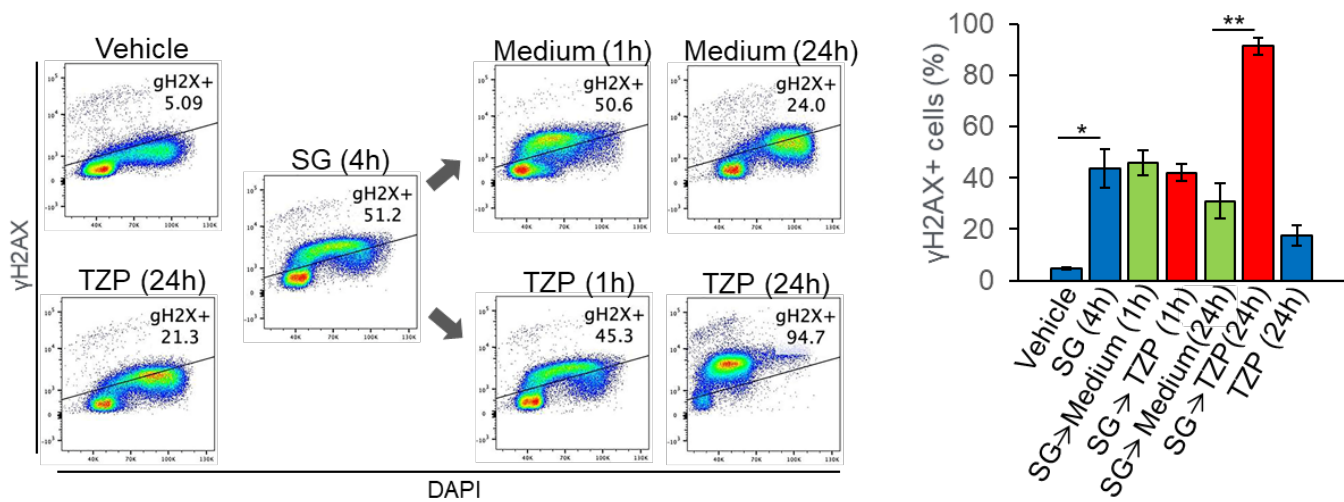
Activities: Here we carried out in vitro studies testing different schedules, dosing and timing of sequential SG and PARPi (PARP inhibitor, Talazoparib) therapy. We determined optimal in vitro sequencing.

Objectives: We sought to understand the kinetics required for optimal sequential dosing of SG and PARPi in vitro, as well as biomarkers for SG/PARPi induced DNA damage.

Methodology, Results, and Conclusions: We carried out a series of in vitro experiments employing different doses and schedules of sequential SG/PARPi treatment. We further compared these doses and schedules in TNBC cells and normal cells. We assessed cell viability quantitatively employing cell titre glow. Select results from these experiments are shown in Figure 1. These data demonstrate that rapid addition of PARPi post SG washout provides optimal synergy for sequential therapy. Furthermore, these data demonstrate that synergy is selective for tumor cells and does not occur in normal cells. This finding illustrates the potential therapeutic window for sequential SG/PARPi therapy, and suggests that normal cells will be spared synergistic toxicity. A second focus of this Task was to identify biomarkers of SG-induced damage. Figure 2 demonstrates assessment of the double-strand DNA break indicator gamma-H2.AX by flow cytometry with SG, PARPi or the combination delivered in a sequential manner. The findings demonstrate dramatic accumulation of damage with the sequential SG/PARPi dosing, compared to repair/resolution of damage with single-agent SG or TZP alone.



**Figure 1. Cell viability with staggered SG/PARPi dosing vs. single-agent SG in normal and tumor cells.** We tested multiple schedules and doses of SG in (A) TNBC cells and (B) normal WI-38 cells. (A) Four-hour (4h) SG dosing followed by washout, followed by 24 hour (24h) Talazoparib (TZP), demonstrating potent synergy for sequential SG/TZP dosing even at non-lethal TZP doses. (B) Comparable sequential dosing of SG/TZP in normal cells is not associated with cytotoxic synergy.



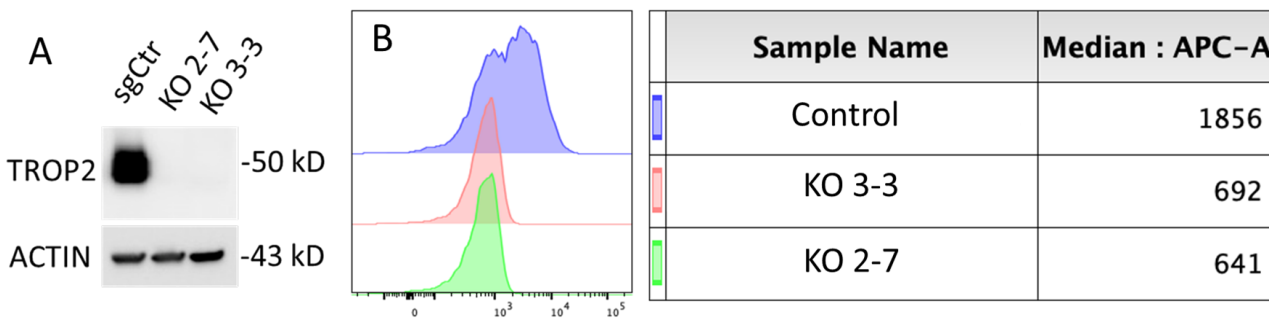
**Figure 2. DNA damage assessed by gamma-H2A.X (gH2X) with staggered SG/PARPi dosing vs. single-agent SG.** Representative flow cytometry plots of MDA-MB-468 cells treated with vehicle, single-agent Talazoparib (TZIP) or SG, or sequential SG (4 hours) followed by washout then addition of either medium or TZIP (1 hour or 24 hours). Y-axis shows gamma-H2A.X staining, and X-axis shows DNA staining. Bar graph at right summarizes results from triplicate samples of two biological replicate experiments. These data show that damage is resolving at 24 hours post SG alone, whereas damage accumulates at 24 hours when is SG is followed by sequential TZIP \*  $p < 0.05$ ; \*\*  $p < 0.01$ .

Major Task 2 (Ellisen): Perform direct comparison of simultaneous and staggered dosing *in vivo*, aiming to maximize tumor kill while mitigating systemic toxicity

Activities: We are in process of carrying out *in vivo* studies of SG with single-agent vs. staggered PARPi dosing *in vivo*, employing xenograft TNBC models.

Objectives: We seek to determine whether staggered SG/PARPi dosing will mitigate systemic toxicity while preserving anti-tumor efficacy *in vivo*.

Methodology, Results, and Conclusions: To date, we have successfully obtained DOD ACURO approval for our IACUC-approved protocol. Experiments described in this Task are currently being carried out/are in process. Furthermore, we have expanded the scope of these experiments to include effects of SG in a syngeneic TNBC model. As SG is non-reactive with murine TROP2, the syngeneic experiments requires generation of a humanized TROP2 in the mouse TNBC cells. Figure 3 shows progress toward this goal, demonstrating successful ablation of endogenous TROP2 expression in murine (4T1) TNBC cells. The next step involves reconstitution of these lines with human TROP2.



**Figure 3. CRISPR knockout of TROP2 in murine TNBC cells.** (A) Western blot and (B) Flow cytometry with APC-linked anti-TROP2 antibody staining of control guide RNA (sgCtrl) or TROP2 knockout (KO) cells in two independent clones, KO 3-3 and KO 2-7. Both KO clones show absence of TROP2 protein.

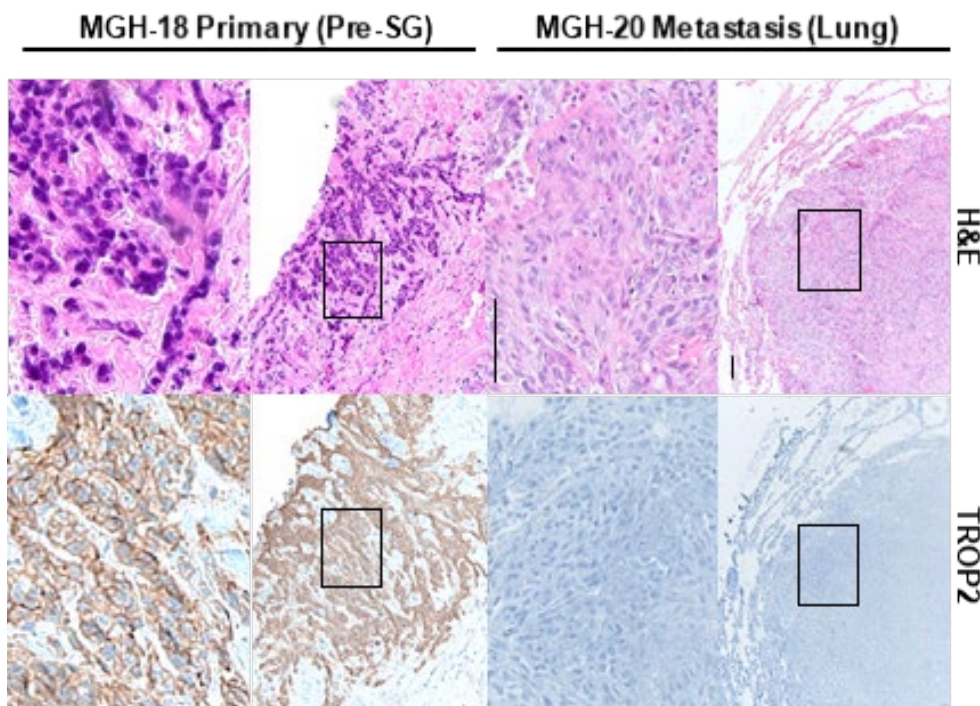
Major Task 3 (Bardia): Test hypothesized biomarkers of SG versus SG/PARPi sensitivity to identify patients requiring combination therapy

Activities: We are testing pre-treatment specimens from patients undergoing SG and SG/PARPi treatment for specified biomarkers to establish response predictors.

Objectives: We seek to identify biomarkers that may indicate patients requiring combination therapy versus single-agent SG therapy.

Methodology, Results, and Conclusions: We have successfully obtained DOD HRPO approval for our IRB-approved protocol for testing pre-treatment patient specimens. While approval was in process, we carried out pilot studies funded by separate non-DOD funding sources to test biomarkers. Some of the findings were recently published: Coates, Bardia, Ellisen et al, *Parallel Genomic Alterations of Antigen and Payload Targets Mediate Polyclonal Acquired Clinical Resistance to Sacituzumab Govitecan in Triple-Negative Breast Cancer*. *Cancer Discovery* 2021 11(10):2436-2445 (PMID:34404686). Among the findings were that pre-treatment TROP2 levels were correlated with extent and duration of response to SG in patients, as shown in Figure 4 below,

Participant ID	Molecular Subtype	Age at Diagnosis	Days on IMMU-132	Days from Last Dose SG to Death	Treatments Before SG	Best Response (per RECIST)	Extent of Best Response (%)
MGH-18	TNBC	41	253	138	2	PR	-45.0
MGH-19	TNBC	59	150	305	5	SD	-21.9
MGH-20	TNBC	62	34	56	4	PD	+78.0



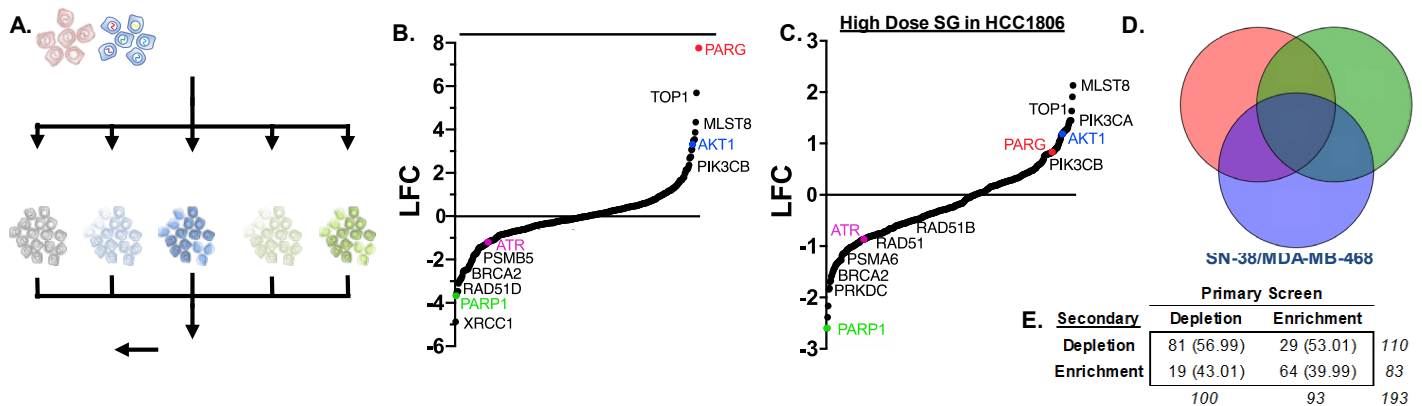
**Figure 4. Pre-treatment assessment of TROP2 levels correlates with extent and duration of response to SG in patients.** Top table shows relevant clinical data for 3 patients undergoing single-agent SG treatment. Best response is shown: PR, partial response; SD, stable disease; PD, progressive disease. Bottom photomicrographs of hematoxylin and eosin (H&E) and TROP2 immunohistochemical staining for two of these patients. Patient MGH-18 tumor shows high-level membranous TROP2 staining and had a PR to SG lasting >250 days, while patient MGH-20 tumor shows absence of TROP2 staining, and had de novo PD.

Major Task 4 (Ellisen): Carry out focused secondary CRISPR screens to reveal inter-tumor heterogeneity of response/resistance pathways to SG in TNBC and the specific pathways related to the ADC mechanism versus SN-38

Activities: We carried out focused secondary CRISPR screens, employing key hits from the primary CRISPR screen described in the application. reveal inter-tumor heterogeneity of response/resistance pathways to SG in TNBC and the specific pathways related to the ADC mechanism versus SN-38

Objectives: We sought to reveal inter-tumor heterogeneity of response/resistance pathways to SG in TNBC and the specific pathways related to the ADC mechanism versus SN-38

Methodology, Results, and Conclusions: We generated a custom sgRNA library by selecting hits and related pathway gene from the primary CRISPR screen described in the application. We then employed this library to screen multiple TNBC models as shown in Figure 5 below. These data demonstrated a high level of reproducibility between primary and secondary screens, and high level of overlap between hits in multiple TNBC models. Collectively, these findings support the robustness of our approach, and they speak to the likely validity of genes and pathways we identified in these screens as relevant to SG sensitivity and resistance.



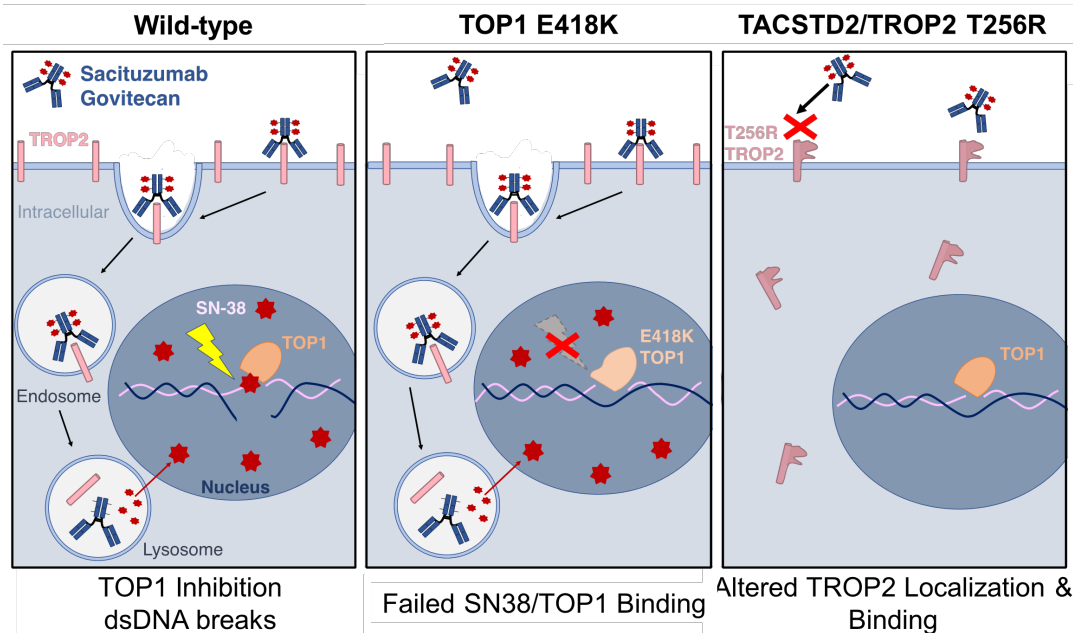
**Figure 5. Multi-Cell Line Secondary Pooled CRISPR Screens.** (A) Schematic depicting the pooled CRISPR/Cas9-based secondary screens in multiple TNBC models. (B, C) S-plots of high dose SG (3 nM) CRISPR screen results for the indicated models. Each dot represents one gene (average of 6 guide RNAs from duplicate experiments). Key depletion (sensitizing) hits for either cell line (Log-Fold Change (LFC)<0) are highlighted as well as key enrichment (resistance; LFC>0) hits, which confirm key hypotheses from our primary CRISPR screen. (D) Venn diagram depicting sensitizing hits in secondary screen for the indicated models and treatment. Note the large proportion of sensitizing hits with SG are shared between the two cell lines (N=145) and, as anticipated, a majority of sensitizing hits with SN-38 (144 of 156) are shared with SG. A high level of correlation between hits in the two cell lines is shown by non-parametric analysis ( $\rho=0.453$ ;  $P<0.0001$ ) suggesting common pathways for resistance and sensitivity. (E) A high level of reproducibility between primary and secondary screens shown by contingency table analysis of the top 33% enriching or depleting genes ( $\chi^2=47.91$ ,  $P<0.00001$ , Yates correction).

Major Task 5 (Bardia): Identify novel biomarkers and targetable resistance mechanism to SG in patients with mTNBC

Activities: We wished to perform post-treatment molecular analysis of specimens from patients demonstrating acquired resistance to SG.

Objectives: We sought to identify novel biomarkers and targetable resistance mechanism to SG in patients with mTNBC.

Methodology, Results, and Conclusions: While IRB approval was in process, we carried out pilot studies funded by separate non-DOD funding sources, which led to the identification of the first in vivo molecular mechanisms of acquired SG drug resistance. These findings were described on our recent publication cited above: Coates, Bardia, Ellisen et al, *Parallel Genomic Alterations of Antigen and Payload Targets Mediate Polyclonal Acquired Clinical Resistance to Sacituzumab Govitecan in Triple-Negative Breast Cancer*. *Cancer Discovery* 2021 11(10):2436-2445. As demonstrated in Figure 6, we found parallel mechanisms of acquired resistance involving mutations of genes encoding topoisomerase 1 (TOP1, the target of the SG cytotoxic payload SN-38), and TROP2 (the target of SG, encoded by the TACSTD2 gene). These findings have important implications for future sequential and combination therapies incorporating SG.



**Figure 6. Model for parallel acquired resistance mechanisms to SG.** Left, SG induces tumor cell killing through TROP2-dependent delivery of SN-38, which binds at the TOP1/DNA interface and induces double-strand DNA (dsDNA) breaks. Middle, TOP1 E418K mutation has altered DNA binding and sequence specificity, resulting in failure of SN-38 binding and SN-38-induced DNA damage. Right, the novel TROP2 T256R mutant shows attenuated cell membrane localization and decreased binding to hRS7, impeding SG binding and SN-38 delivery. (Fig. 3G in Coates, Bardia, Ellisen et al, *Cancer Discovery* 2021 11(10):2436.)

Major Task 6 (Ellisen): Test selected synergistic pathways including checkpoint kinases and CDKs in gain and loss of function studies in vitro & in vivo

Major Task 7 (Ellisen): Establish in vivo efficacy and tolerability of select SG/clinical drug combinations selected based on Aim 2 using in vivo models

Major Task 8 (Bardia): Test promising combinations in organoid and PDX models from TNBC patients exhibiting resistance to SG and SG/PARP<sub>i</sub>

Activities: These aims seek to test novel pathways and therapeutic agents in combination with SG.

Objectives: We seek to find targets and therapeutic approaches that may synergize to overcome de novo and acquired resistance to SG.

Methodology, Results, and Conclusions: We have successfully obtained DOD HRPO approval for collection and analysis of cadaveric specimens. Now, all these Tasks are currently in planning and/or in process, as described in the original application.

## **What opportunities for training and professional development has the project provided?**

While this is not a training grant, multiple opportunities for professional development were provided by this award. Dr. James Coates presented findings at the Ludwig Center at Harvard annual seminar. He also presented findings at the annual MGH Cancer Center Retreat and poster session. He has graduated to an independent scientific consulting position in the UK as of 2/1/2022. His assigned duties going forward will be carried out by Dr. Bogang Wu, a highly talented postdoctoral fellow who recently received his PhD degree from George Washington University.

## **How were the results disseminated to communities of interest?**

Findings were presented:

- a) At the Ludwig Center at Harvard annual seminar.
- b) At the annual MGH Cancer Center Retreat and poster session.
- c) At the annual “Science Monday” seminar for the MGH/Harvard Community
- d) At the annual MGH Breast Program “Latest updates in breast cancer science at MGH”.
- e) At the annual Faculty Showcase for the Cancer Center at MGH.

*Describe briefly what you plan to do during the next reporting period to accomplish the goals and objectives.*

We plan to continue studies of mechanisms and specimens related to sequential SG/PARP $\alpha$ 1 therapy and its application in the clinic. We plan to study CRSIPR screen hits that revealed potential SG resistance mechanisms and collaborating therapeutic targets. We plan to continue systematic studies of pre- and post-treatment patient specimens to validate and uncover additional relevant mechanisms. We plan to test identified mechanisms of synergy and their potential therapeutic relevance in vitro and in vivo.

## **4. IMPACT:**

### **What was the impact on the development of the principal discipline(s) of the project?**

Currently there are nine antibody drug conjugates (ADCs) approved by the FDA in the field of oncology, and more than 100 being investigated in clinical trials. Our findings highlight that molecular alterations in both antibody binding target and payload target may mediate resistance to ADCs, a finding that is likely to have implications for therapeutic sequencing of ADCs. For example, resistance mechanisms to SG due to drug payload target alterations would likely confer cross-resistance to other ADCs with similar (TOP1 inhibitor) payloads currently approved by the FDA. In such a scenario, therapeutic sequencing with the same antibody but a different payload could provide therapeutic benefit. On the other hand, resistance mechanisms due to TROP2 alteration would likely confer cross-resistance to other TROP2-directed ADCs, but not to non-TROP2 directed ADCs regardless of payload.

### **What was the impact on other disciplines?**

As discussed under section (4) above, the findings to date are likely to have a potentially broad impact on the field of cancer therapy, as ADCs emerge as groundbreaking therapeutics in a variety of settings.

**What was the impact on technology transfer?**

Nothing to report

**What was the impact on society beyond science and technology?**

Nothing to Report

**5. CHANGES/PROBLEMS:**

No major changes or alterations. Minor modification of experiments and scope are mentioned above. No major problems to report, although DOD approval for all proposed specimen analysis took 7-8 from the initiation of the award.

**Actual or anticipated problems or delays and actions or plans to resolve them**

*Describe problems or delays encountered during the reporting period and actions or plans to resolve them.*

We do not anticipate further delays or problems going forward.

**Changes that had a significant impact on expenditures**

*Describe changes during the reporting period that may have had a significant impact on expenditures, for example, delays in hiring staff or favorable developments that enable meeting objectives at less cost than anticipated.*

The fact that DOD approval for all proposed specimen analysis took 7-8 from the initiation of the award may have delayed some aspects of anticipated spending in Year 1. For this reason, the overall budget may be somewhat underspent for this reporting period. However, with all approvals now in place, we anticipate full and complete expenditure over the entire award period as spelled out in the detailed budget.

**Significant changes in use or care of human subjects, vertebrate animals, biohazards, and/or select agents**

*Describe significant deviations, unexpected outcomes, or changes in approved protocols for the use or care of human subjects, vertebrate animals, biohazards, and/or select agents during the reporting period. If required, were these changes approved by the applicable institution committee (or equivalent) and reported to the agency? Also specify the applicable Institutional Review Board/Institutional Animal Care and Use Committee approval dates.*

**Significant changes in use or care of human subjects**

No significant deviations, unexpected outcomes, or changes in approved protocols for the use of human tissues/human subjects. Most recent IRB approval date is 12/28/2021.

### Significant changes in use or care of vertebrate animals

No significant deviations, unexpected outcomes, or changes in approved protocols for the use or care of vertebrate animals. Annual animal protocol was re-approved per institutional policy **for 2 years on 2/9/2022.**

### Significant changes in use of biohazards and/or select agents

No significant deviations, unexpected outcomes, or changes in approved protocols for biohazard and/or select agent use. Annual approval was certified 6/26/2021.

**6. PRODUCTS:** *List any products resulting from the project during the reporting period. If there is nothing to report under a particular item, state "Nothing to Report."*

- **Publications, conference papers, and presentations**

**Journal publications.** *List peer-reviewed articles or papers appearing in scientific, technical, or professional journals. Identify for each publication: Author(s); title; journal; volume: year; page numbers; status of publication (published; accepted, awaiting publication; submitted, under review; other); acknowledgement of federal support (yes/no).*

Published manuscript: Coates, Bardia, Ellisen et al, Parallel Genomic Alterations of Antigen and Payload Targets Mediate Polyclonal Acquired Clinical Resistance to Sacituzumab Govitecan in Triple-Negative Breast Cancer. Cancer Discovery 2021 11(10):2436-2445 (PMID:34404686; PMCID: PMC8495771  
Acknowledgement of federal support: yes.

**Books or other non-periodical, one-time publications.**

None

**Other publications, conference papers and presentations.** *Identify any other publications, conference papers and/or presentations not reported above. Specify the status of the publication as noted above. List presentations made during the last year (international, national, local societies, military meetings, etc.). Use an asterisk (\*) if presentation produced a manuscript.*

None

- **Website(s) or other Internet site(s)**

None

- **Technologies or techniques**

*Identify technologies or techniques that resulted from the research activities. Describe the technologies or techniques were shared.*

None

- **Inventions, patent applications, and/or licenses**

*Identify inventions, patent applications with date, and/or licenses that have resulted from the research. Submission of this information as part of an interim research performance progress report is not a substitute for any other invention reporting required under the terms and conditions of an award.*

None

- **Other Products**

None to date

## 7. PARTICIPANTS & OTHER COLLABORATING ORGANIZATIONS

### What individuals have worked on the project?

Name: Leif Ellisen  
Project Role: PI  
ORCID ID: 0000-0003-0444-7910  
Nearest person month worked: 1  
Contribution to project: Oversight of all research activities reviewing data and writing manuscripts.

Name: Aditya Bardia  
Project Role: Partnering PI  
ORCID ID: 0000-0003-4885-1157  
Nearest person month worked: 1  
Contribution to project: Oversight of all research activities reviewing data and writing manuscripts.

Name: Veerle Bossuyt  
Project Role: Pathologist  
ORCID ID: 0000-0003-0403-6791  
Nearest person month worked: 1  
Contribution to project: Aid in tissue collection and analyze tissue biomarkers.

Name: Ryan, Phoebe K  
Project Role: Clinical Research Coordinator  
ORCID ID: n/a  
Nearest person month worked: 12  
Contribution to project: Collect clinical data and participate in informed consent process.

Name: Nayana Thimmiah,  
Project Role: Research Technician  
ORCID ID: n/a  
Nearest person month worked: 5  
Contribution to project: Assist in all laboratory-based analyses.

Name: James Coates  
Project Role: Postdoctoral Research Fellow  
ORCID ID: n/a  
Nearest person month worked: 9  
Contribution to project: Oversee laboratory-based analyses and help writing manuscripts.

Name: Bogang Wu  
Project Role: Postdoctoral Research Fellow  
ORCID ID: n/a  
Nearest person month worked: 3  
Contribution to project: Oversee laboratory-based analyses and help writing manuscripts.

Name: Sheng Sun  
Project Role: Postdoctoral Research Fellow  
ORCID ID: n/a  
Nearest person month worked: 1  
Contribution to project: Assist in specialized laboratory-based analyses.

**Has there been a change in the active other support of the PD/PI(s) or senior/key personnel since the last reporting period?**

*If there is nothing significant to report during this reporting period, state “Nothing to Report.”*

*If the active support has changed for the PD/PI(s) or senior/key personnel, then describe what the change has been. Changes may occur, for example, if a previously active grant has closed and/or if a previously pending grant is now active. Annotate this information so it is clear what has changed from the previous submission. Submission of other support information is not necessary for pending changes or for changes in the level of effort for active support reported previously. The awarding agency may require prior written approval if a change in active other support significantly impacts the effort on the project that is the subject of the project report.*

**PI: Leif Ellisen**

R21EB025270-01A1 has ended as of 5/31/21.

R01CA260890-01A1 began – project dates 01/01/2022 - 12/31/2026, 1.2 CM effort

**PI: Aditya Bardia**

5P50CA168504-08 began – project dates 06/01/2021 – 05/31/2022, 1.2 CM effort

1R01CA260890-01A1 began – project dates 01/01/2022 - 12/31/2026, 1.2 CM effort

5R01CA255602-02 began – project dates 01/08/2021 - 12/31/2025, 0.6 CM effort

**What other organizations were involved as partners?**

Nothing to Report.

**COLLABORATIVE AWARDS:** *For collaborative awards, independent reports are required from BOTH the Initiating Principal Investigator (PI) and the Collaborating/Partnering PI. A duplicative report is acceptable; however, tasks shall be clearly marked with the responsible PI and research site. A report shall be submitted to <https://ebrap.org/eBRAP/public/index.htm> for each unique award.*

**QUAD CHARTS:** *If applicable, the Quad Chart (available on <https://www.usamraa.army.mil/Pages/Resources.aspx>) should be updated and submitted with attachments.*

- 9. APPENDICES:** *Attach all appendices that contain information that supplements, clarifies or supports the text. Examples include original copies of journal articles, reprints of manuscripts and abstracts, a curriculum vitae, patent applications, study questionnaires, and surveys, etc.*

## RESEARCH BRIEF

# Parallel Genomic Alterations of Antigen and Payload Targets Mediate Polyclonal Acquired Clinical Resistance to Sacituzumab Govitecan in Triple-Negative Breast Cancer



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## ABSTRACT

Sacituzumab govitecan (SG), the first antibody–drug conjugate (ADC) approved for triple-negative breast cancer, incorporates the anti-TROP2 antibody hRS7 conjugated to a topoisomerase-1 (TOP1) inhibitor payload. We sought to identify mechanisms of SG resistance through RNA and whole-exome sequencing of pretreatment and postprogression specimens. One patient exhibiting *de novo* progression lacked TROP2 expression, in contrast to robust TROP2 expression and focal genomic amplification of *TACSTD2/TROP2* observed in a patient with a deep, prolonged response to SG. Analysis of acquired genomic resistance in this case revealed one phylogenetic branch harboring a canonical *TOP1*<sup>E418K</sup> resistance mutation and subsequent frameshift *TOP1* mutation, whereas a distinct branch exhibited a novel *TACSTD2/TROP2*<sup>T256R</sup> missense mutation. Reconstitution experiments demonstrated that *TROP2*<sup>T256R</sup> confers SG resistance via defective plasma membrane localization and reduced cell-surface binding by hRS7. These findings highlight parallel genomic alterations in both antibody and payload targets associated with resistance to SG.

**SIGNIFICANCE:** These findings underscore TROP2 as a response determinant and reveal acquired SG resistance mechanisms involving the direct antibody and drug payload targets in distinct metastatic subclones of an individual patient. This study highlights the specificity of SG and illustrates how such mechanisms will inform therapeutic strategies to overcome ADC resistance.

## INTRODUCTION

Metastatic triple-negative breast cancer (mTNBC), defined by absence of estrogen and progesterone receptors (ER and PR) and human epidermal growth factor receptor 2 (*HER2*) gene amplification, is associated with very poor prognosis (1, 2). Standard chemotherapy, the mainstay of

management, is associated with low response rates (5%–10%) and poor median progression-free survival (2–3 months) in the relapsed/refractory setting (3, 4). Similarly, for patients with endocrine-resistant hormone receptor-positive (HR<sup>+</sup>) metastatic breast cancer (MBC), chemotherapy is the mainstay of management but has limited efficacy and has not

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been demonstrated to improve overall survival (OS). Antibody–drug conjugates (ADC), generated by linking cancer-directed antibodies to potent payloads, can selectively deliver toxic payloads to cancer cells, thereby maximizing the therapeutic window (5).

Although TNBC lacks the traditional actionable receptors (ER, PR, and HER2), the majority (85%–90%) express TROP2, making it an attractive target for ADCs (6). Sacituzumab govitecan (SG) is a first-in-class TROP2 ADC in which SN-38 (the active metabolite of irinotecan), a topoisomerase I (TOP1) inhibitor, is coupled to the humanized anti-TROP2 monoclonal antibody hRS7 IgG1 $\kappa$  via the hydrolysable CL2A linker. In a phase I/II clinical trial, patients with metastatic TNBC (median of 3 prior regimens) treated with SG had remarkable therapeutic responses (33%), including complete responses (7). The pivotal randomized phase III ASCENT trial demonstrated comparable clinical activity and a near-doubling of OS versus physicians' choice chemotherapy (8). Based on its clinical activity in advanced refractory TNBC, SG received accelerated FDA approval for mTNBC in April 2020 and full approval in April 2021. Substantial clinical efficacy has also been observed in HR<sup>+</sup> MBC and other solid tumors that express TROP2 (9).

Despite multiple clinical trials of SG, little is known about predictors of therapeutic response and mechanisms of *de novo* and acquired resistance. Thus, we carried out genomic and transcriptomic analysis of tumor tissue from patients with mTNBC treated with SG for whom we had access to pretreatment and multisite postprogression (rapid autopsy) specimens.

## RESULTS

### Association of TROP2 Levels and Gene Copy Number with SG Response/Resistance

As TROP2 is the protein bound by the antibody moiety of SG, we first analyzed TROP2 gene expression and genomic copy number in pre- and posttreatment specimens from the three-case autopsy series (Fig. 1A; Supplementary Table S1). Each of these patients received SG following progression on at least two prior therapies for mTNBC (Fig. 1A; Supplementary Table S2). Case MGH-20 showed evidence of clinical *de novo* progression by the time of first interval radiologic assessment, resulting in drug discontinuation. MGH-19 experienced stable disease (SD) by Response Evaluation Criteria in Solid Tumors (RECIST) version 1.1 criteria for a period of 5 months, whereas MGH-18 had a substantial and prolonged partial response according to RECIST 1.1 (45% tumor regression, >8 months), prior to multisite progressive disease (PD; Fig. 1A; Supplementary Fig. S1). RNA-sequencing (RNA-seq) analysis of all analyzed tumor specimens from MGH-20 demonstrated essentially undetectable *TROP2* RNA expression, and this was corroborated by the absence of TROP2 protein expression on IHC (Fig. 1B and C). MGH-18 and MGH-19 both exhibited detectable *TROP2* RNA expression in all specimens analyzed, including in metastatic foci that had progressed on SG treatment (Fig. 1B). MGH-18 was distinguished by the presence of genomic amplification of the *TACSTD2* locus (encoding TROP2) in multiple specimens tested, and as anticipated the primary tumor showed robust levels of

membrane-localized TROP2 protein (Fig. 1C and D). These findings are in keeping with recent data demonstrating that absence of TROP2 expression is commonly associated with *de novo* clinical resistance to SG (8).

### Parallel Genomic Alterations of *TACSTD1/TROP2* and *TOP1* Denote Acquired Resistance to SG

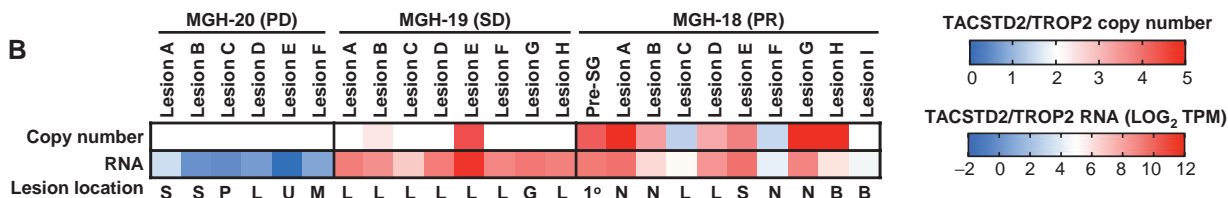
We focused our analysis of acquired resistance on MGH-18 because this case was associated with the most profound and prolonged response, and a relatively short interval (<4 months) between progression on SG and tissue harvest (autopsy). This case involved a 42-year-old female who underwent standard preoperative chemotherapy for primary ER/PR/HER2-negative invasive ductal carcinoma (i.e., TNBC), but within weeks of completion of neoadjuvant therapy was found to have metastatic disease to liver and other organs. The patient underwent palliative mastectomy and then received two sequential investigational therapeutic combinations that were short-lived, and there was no objective response (Supplementary Table S2). The patient then initiated treatment with SG, and restaging scans after two months demonstrated a radiologic partial response with 45% reduction, as per RECIST (Fig. 2A). However, after eight months, restaging scans revealed disease progression at multiple sites, and following brief treatment with standard chemotherapy, the patient expired (Fig. 2A; Supplementary Table S2).

In order to understand the evolution of acquired resistance to SG in this case, we carried out whole-exome sequencing (WES) of the pre-SG treatment and nine postprogression rapid autopsy tumor lesions. From these data, we then reconstructed the phylogenetic relationships and clonal architecture of the resistant lesions using PhylogNDT, an integrated suite of tools designed for jointly modeling evolution of many samples from the same patient (Fig. 2B; ref. 10). This analysis revealed shared clonal (truncal) mutations common to all samples and private mutations within a given branch (Supplementary Table S3). A truncal *TP53* mutation (K132R) was present in the pre-SG primary tumor, as is commonly observed in TNBC, and was present in all postprogression lesions (Fig. 2B and C). In most cases, multiple branches were detected in each metastatic lesion (Fig. 2D; Supplementary Fig. S2A and S2B). Most notable in the phylogenetic tree of resistance were two major branches that demonstrated distinct, branch-restricted mutations of *TOP1* (encoding the SN-38 drug target topoisomerase 1) and *TACSTD2* (encoding TROP2; Fig. 2B; Supplementary Fig. S2C). These two phylogenetic branches were present in multiple, mutually exclusive metastatic lesions at high frequency. One branch, which was present at high frequency within abdominal lesions in the liver and periaortic lymph nodes, harbored the *TOP1* missense mutation E418K (Fig. 2C and D; Supplementary Fig. S2C). This specific *TOP1* mutation has been previously described and is established to confer resistance to clinical TOP1 inhibitors, most likely by altering sequence specificity for DNA cleavage by TOP1 (11, 12). Further evolution of this *TOP1*<sup>E418K</sup>-containing branch involves an additional subclonal 8-bp frameshift insertion in *TOP1* (*TOP1p*-122fs), which is present in these same metastases (Fig. 2C and D; Supplementary Fig. S2C). Of note, because the *TOP1* frameshift mutation appears to arise from the

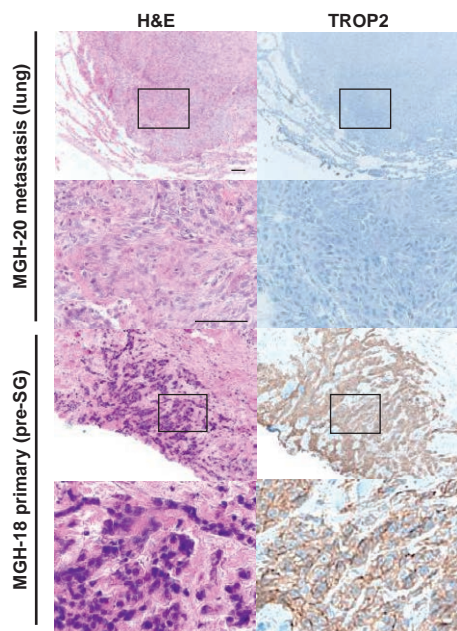
A

Participant ID	Molecular subtype	Age at diagnosis	Days on IMMU-132	Days from last dose SG to death	Treatments before SG	Treatments after SG	Lesions sequenced at autopsy	Best response (per RECIST)	Extent of best response (%)
MGH-18	TNBC	41	253	138	2	2	9	PR	-45.0
MGH-19	TNBC	59	150	305	5	4	8	SD	-21.9
MGH-20	TNBC	62	34	56	4	1	6	PD	+78.0

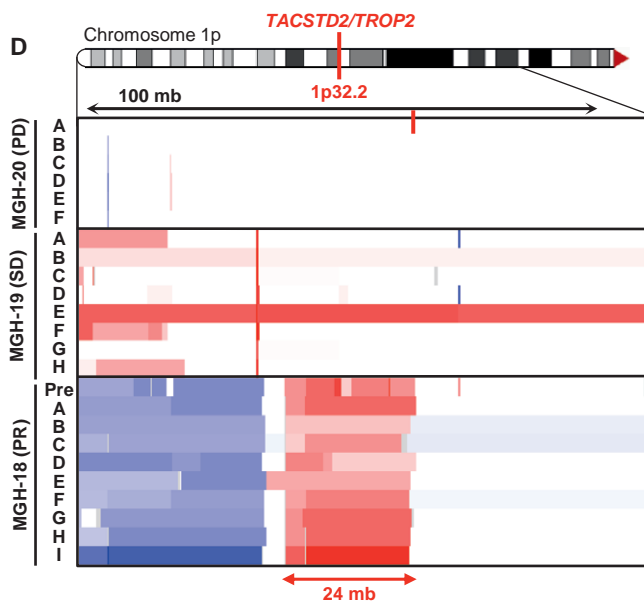
B



C



D



**Figure 1.** TROP2 expression and gene copy number and response to SG. **A**, Clinical characteristics, treatment history, and SG response data for the three autopsy series patients. PR, partial response; SD, stable disease; PD, progressive disease. **B**, *TACSTD2/TROP2* copy number (top) and RNA expression (bottom) for all analyzed tumor specimens from each case. Note, white indicates diploid copy number. Key for lesion location: S, subcutaneous; P, pericardium; L, liver; U, lung; M, mediastinum; G, gallbladder; N, lymph node; B, brain; 1°, primary breast tumor. **C**, Hematoxylin and eosin (H&E) and TROP2 IHC show tumor TROP2 protein expression is absent in MGH-20 but shows intense membrane staining in MGH-18. Scale bars, 50  $\mu$ m. **D**, Focal amplification of *TACSTD2/TROP2* in pretreatment primary tumor and multiple metastatic lesions from MGH-18. Copy-number scale per **B** is shown for the indicated region of chromosome 1p of each lesion, indicating up to  $N=5$  *TACSTD2/TROP2* copies. See also Supplementary Fig. S1 and Supplementary Table S1.

clone harboring the *TOP1*<sup>E418K</sup> mutation, this suggests that the frameshift could represent a “second hit” promoting enhanced resistance to SG. Furthermore, analysis of RNA-seq data demonstrated that both *TOP1* mutations were expressed in the metastatic lesions (Supplementary Fig. S2D).

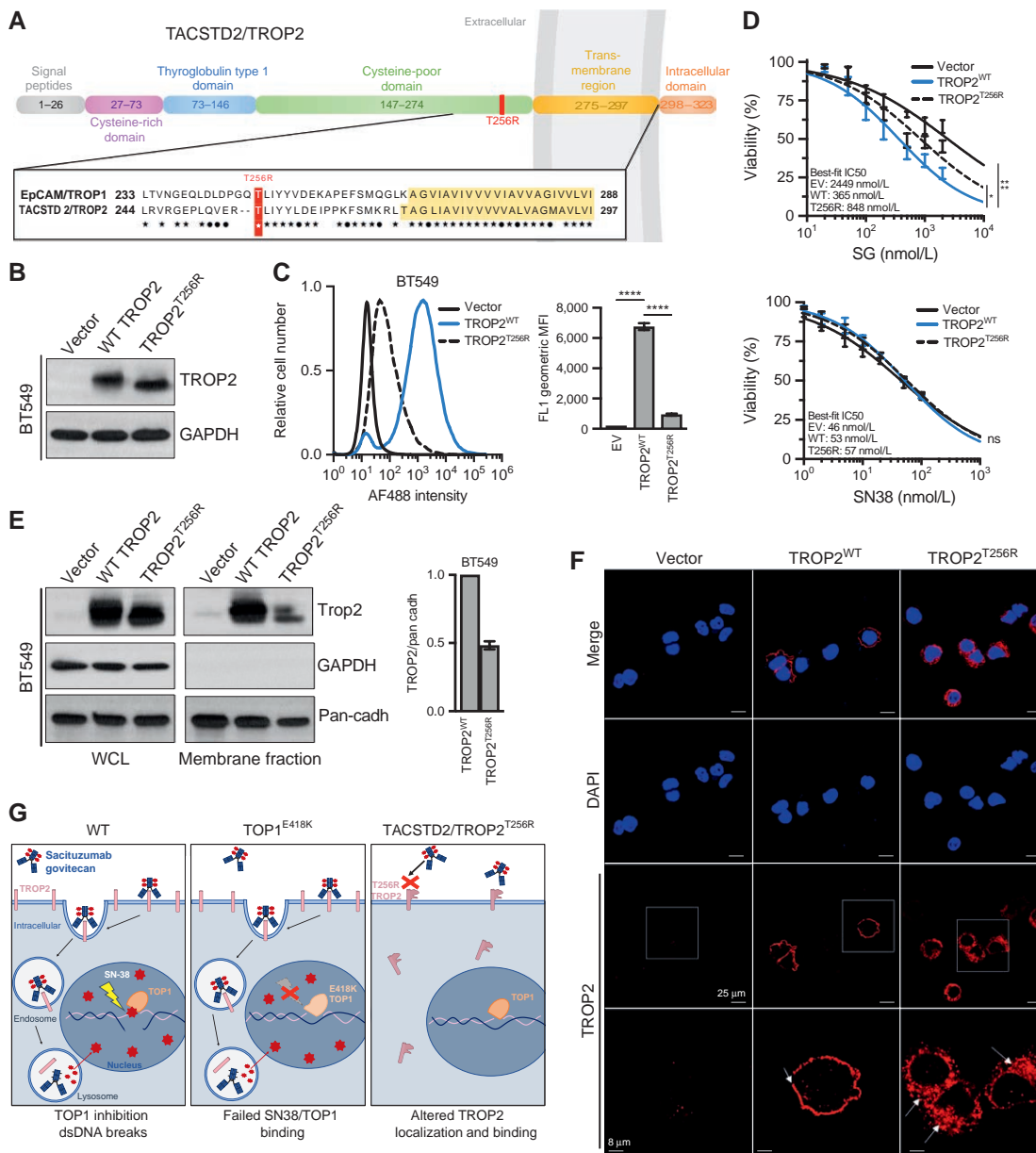
A second major phylogenetic branch of resistance was hallmarked by a missense mutation of *TACSTD2/TROP2*, T256R. As noted, this mutation occurred in a distinct phylogenetic branch from the *TOP1* mutations, and this *TACSTD2/TROP2*-mutant branch was present at high frequency in multiple thoracic metastatic lesions, including hilar lymph node and chest wall (Fig. 2C and D; Supplementary Fig. S2C). RNA bearing this mutation was also expressed in the respective lesions (Supplementary Fig. S2D). To our knowledge, this mutation of the gene encoding TROP2 has not been

previously described in the literature in any context. However, its acquisition in the setting of tumor progression on SG therapy suggested the possibility of parallel mechanisms of resistance to SG involving both the antibody target, TROP2, and its SN-38 payload target, TOP1.

### The Membrane-Proximal *TACSTD2/TROP2* Mutation T256R Alters Plasma Membrane Localization and Cell-Surface Binding of hRS7

To determine the potential functional significance of the *TACSTD2/TROP2*<sup>T256R</sup> mutation for SG treatment, we synthesized cDNAs encoding either the wild-type or mutant TROP2 and reconstituted them into multiple TROP2-negative cell-based models. TROP2 is a cell-surface glycoprotein that exhibits high homology to epithelial cell adhesion





**Figure 3.** *TACSTD2/TROP2*<sup>T256R</sup> confers resistance to SG via altered localization and decreased binding to hRS7. **A**, Domain structure and homology between TROP1/EPCAM and TROP2/TACSTD2. EPCAM and TROP2 share 49% sequence identity and 67% sequence similarity. Homology is highest in the membrane-proximal region and transmembrane region (yellow highlight). ★, identical amino acids; \*, conservative differences. T256R mutation site indicated in red. **B**, Western blot of lentivirally expressed control vector or TROP2 cDNAs in TROP2-negative BT549 cells, showing equivalent expression levels of wild-type (WT) and mutant proteins. **C**, Live-cell flow cytometry of BT549 cells shown in **B** stained with hRS7, the anti-TROP2 antibody backbone of SG, reveals a marked decrease in signal intensity for cells harboring TROP2<sup>T256R</sup> compared with WT. Bar graphs to the right represent geometric mean fluorescent intensity (MFI). Data are representative of three independent experiments, and error bars show SD. \*\*\*\*,  $P < 0.0001$ . **D**, TROP2<sup>T256R</sup>-mutant confers resistance to SG but not SN38. BT549 cells expressing empty vector (EV), WT TROP2, or T256R-mutant TROP2 were treated with SG or SN-38 at the indicated doses for 4 hours and then cultured in fresh medium for 96 hours before assessing cell viability by using CellTiter-Glo. Values plotted represent the mean of quadruplicate wells and an average of three independent experiments. Error bars indicate SD.  $P$  value calculated by two-way ANOVA (cell line, concentration). \*,  $P < 0.05$ ; ns, nonsignificant. **E**, TROP2<sup>T256R</sup> has significantly decreased plasma membrane association, shown by comparing whole-cell lysate (WCL) and plasma membrane fraction of BT549 cells expressing the vector, WT TROP2, or T256R-mutant TROP2. GAPDH and pan-cadherin serve as controls for cytosol and membrane fractions, respectively. Bar graphs at right represent the mean of at least two independent experiments. Error bars indicate range. **F**, Confocal immunofluorescence using rabbit monoclonal clone EPR20043 reveals plasma membrane staining of WT TROP2 and intracellular staining of TROP2<sup>T256R</sup>. Vector, WT, or T256R TROP2-expressing BT549 cells were stained for TROP2 and then examined at 40× using a Nikon A1R confocal microscope. Bottom, magnified images corresponding to white boxes. Scale bars, 25 μm and 8 μm (magnified). **G**, Model for parallel polyclonal acquired resistance to SG. Left, SG induces tumor cell killing through TROP2-dependent delivery of SN-38, which binds at the TOP1/DNA interface and induces double-strand DNA (dsDNA) breaks. Middle, TOP1<sup>E418K</sup> has altered DNA binding and sequence specificity, resulting in failure of SN-38 binding and SN-38-induced DNA damage. Right, the novel TROP2<sup>T256R</sup> mutant shows attenuated cell membrane localization and decreased binding to hRS7, impeding SG binding and SN-38 delivery. See also Supplementary Figs. S3 and S4.

assayed cell-surface binding to TROP2 in these cells by hRS7, the (humanized) antibody incorporated into SG. Although control transfected cells showed no binding as anticipated, reconstitution of wild-type TROP2 conferred robust cell-surface binding by hRS7. In contrast, in cells reconstituted with the TROP2 mutant, the binding was reduced by >80% compared with wild-type TROP2 in TNBC cells, and similar results were observed in 3T3 cells (Fig. 3C; Supplementary Fig. S3B). Binding of an unrelated anti-TROP2 antibody was also highly significantly reduced in these same cells (Supplementary Fig. S3C and S3D). Thus, *TACSTD2/TROP2*<sup>T256R</sup> encodes a stable protein associated with diminished cell-surface binding by multiple antibodies including the SG backbone hRS7.

We next tested directly whether attenuated binding of the mutant TROP2 protein by hRS7 was associated with decreased sensitivity to SG. As expected, reconstitution of wild-type TROP2 into TROP2-negative cells conferred highly significant increases in sensitivity to SG, in both TNBC and 3T3 cells (Fig. 3D; Supplementary Fig. S3E). In contrast, reconstitution with mutant TROP2 conferred only minimal sensitivity that was only marginally distinguishable from that of nonreconstituted cells. As a control, we tested sensitivity to the SG payload SN-38 itself in these cells. As anticipated, reconstitution of either wild-type or mutant TROP2 had no impact on SN-38 sensitivity in either TNBC or 3T3 cells (Fig. 3D; Supplementary Fig. S3E). Thus, expression of the mutant TROP2 compared with the wild-type protein is associated with decreased cellular sensitivity to SG.

Given attenuated cell-surface binding of TROP2<sup>T256R</sup> to hRS7 despite expression levels comparable to wild-type protein, we hypothesized that the mutant protein might exhibit altered subcellular localization. We therefore isolated plasma membrane fractions from TROP2-expressing TNBC and 3T3 cells and compared localization of mutant and wild-type TROP2 proteins. In both cell types, TROP2<sup>T256R</sup> consistently displayed decreased plasma membrane localization compared with the wild-type protein (Fig. 3E; Supplementary Fig. S3F). We then used confocal immunofluorescence (IF) to corroborate these findings. Staining TNBC cells with a rabbit monoclonal antibody, we observed nearly exclusive plasma membrane staining of wild-type TROP2. In contrast, TROP2<sup>T256R</sup> showed substantial localization to the cytosol (Fig. 3F). Furthermore, confocal IF in these same cells using the hRS7 antibody revealed both mislocalized and decreased staining, possibly suggesting a selective decrease in binding of the mutant protein to hRS7 (Supplementary Fig. S3G). Finally, we carried out IHC for TROP2 in parallel on the patient MGH-18 hilar lymph node metastasis containing the mutant TROP2<sup>T256R</sup> and a brain metastasis harboring only wild-type TROP2. Consistent with IF analysis, the hilar lesion showed a largely diffuse cytoplasmic staining pattern for TROP2, whereas the brain metastasis showed a membrane pattern similar to the primary tumor (Supplementary Fig. S4A and S4B). Altogether, this study suggests the emergence of parallel genetic mechanisms of acquired resistance under selective pressure from SG treatment in this patient. A subset of metastatic lesions harbor an established resistance mutation to the SG TOP1 inhibitor (SN-38) payload, *TOP1*<sup>E418K</sup>, whereas a nonoverlapping set of lesions harbor

the *TACSTD2/TROP2*<sup>T256R</sup> mutation, which encodes a protein with altered subcellular localization, attenuated cell-surface binding to the therapeutic antibody backbone, and ability to confer SG resistance (Fig. 3G).

## DISCUSSION

To our knowledge, this is the first report to describe the emergence of acquired resistance to SG hallmarked by mutually exclusive somatic mutations in genes encoding both TROP2 (target of antibody) and TOP1 (target of payload) in distinct metastatic lesions of the same patient. We use the term “parallel” to describe this pattern of polyclonal acquired genetic resistance, in contrast to multiple recently described “convergent” mechanisms that mediate resistance to a single pathway inhibitor (13, 14). Parallel resistance seems naturally applicable to ADCs, given they have dual molecular targets for the antibody and drug payload. The parallel resistance mechanisms we describe highlight the specificity of this ADC, and they may have direct implications for the choice of subsequent therapies, particularly as additional ADCs with potentially overlapping antibody and payload targets become available for this and other patient populations (5, 6).

Comparing three autopsy cases with multiple tumor lesions available for analysis, we find an association between absence of TROP2 expression and a lack of clinical response to SG. In general, reduced expression of an ADC target antigen could affect both binding of the antibody as well as intracellular uptake and payload release (15, 16). In the pivotal EMILIA trial, which led to approval of the HER2-directed ADC trastuzumab emtansine (T-DM1) for metastatic HER2<sup>+</sup> breast cancer, the survival benefit with T-DM1 compared with control arm was greater in patients whose tumors had high HER2 mRNA levels (hazard ratio = 0.53; median OS, 34.1 vs. 26.5 months), compared with lower HER2 mRNA levels (hazard ratio, 0.80; median OS, 24.8 vs. 23.7 months; ref. 17). Similarly, in the confirmatory phase III clinical trial with SG (ASCENT), patients with mTNBC who had high or medium TROP2 expression had higher objective response rates (44% and 39%, respectively), as compared with low TROP2 expression (22%; refs. 8, 18). Furthermore, out of seven patients who had tumors with complete absence of TROP2 expression in ASCENT, only one had a response (8). These observations are consistent with our findings and highlight how lower expression of TROP2 could modulate therapeutic efficacy and ultimately might influence patient selection for SG.

By focusing on a single patient with a deep and prolonged response to SG who subsequently progressed, we were able to identify multiple potentially important mechanisms of acquired genetic resistance to this ADC. The point mutation in *TOP1* we describe, E418K, is one of several reported missense mutations that are known to induce resistance to clinical TOP1 inhibitors but do not alter intrinsic enzyme catalytic activity (11, 19). Instead, this mutation is believed to alter the DNA sequence specificity and/or DNA binding affinity of the enzyme and thereby hinder the stereotactic binding of drug to the enzyme/DNA interface (12). Although such mutations are established to induce resistance in the heterozygous state, the potential relevance of the mutation in this case is further underscored by the development of a coexisting, subclonal

frameshift *TOP1* mutation in the same metastatic lesions. We are not certain that the frameshift mutation exists in *trans* with *TOP1*<sup>E418K</sup>, though the resulting loss of the wild-type protein would seem potentially advantageous under the selective pressure of SG. In theory, these genetic events would be expected to induce cross-resistance to multiple emerging ADCs that use TOP1 inhibitors as payloads. However, future treatment strategies using different payloads such as maytansine (microtubule inhibitor), or possibly a more potent TOP1 inhibitor payload such as deruxtecan (>10 times potent than SN-38), could potentially overcome therapeutic resistance induced by such *TOP1* mutations (20, 21).

Although germline loss-of-function mutations of *TACSTD2/TROP2* are associated with a hereditary human corneal dystrophy, to our knowledge no prior somatic mutations of this gene have been described in the context of acquired resistance to SG or in any other context (22). We find that *TACSTD2/TROP2*<sup>T256R</sup> encodes a protein that can readily be expressed at equivalent total cellular levels to the wild-type protein but is dramatically attenuated in its cell-surface binding to the SG backbone antibody hRS7. Furthermore, *TROP2*<sup>T256R</sup> is mislocalized from the plasma membrane and instead appears to be primarily intracellular. Of note, we consistently observe altered migration of the mutant *TROP2* protein on Western blot analysis, potentially suggesting a posttranslational modification that may contribute to altered localization and/or attenuated hRS7 binding. Although it remains to be seen how common such mutations will prove to be as a mechanism of acquired resistance to SG, our results provide strong proof-of-principle that genetic alteration of the ADC target *TROP2* is a relevant resistance mechanism. In contrast, resistance to T-DM1 has been shown to occur through downregulation of *HER2* expression in cell culture models, but the acquisition of missense mutations that confer resistance has not been well documented (23). Whether *TROP2* alteration is a preferred mechanism of cellular resistance to SG could depend on its expression level and its functional contribution in tumor cells that may vary, both between TNBCs of different patients and between different tumor types for which SG is in clinical development (6, 24).

This study utilized genomic analyses of pretreatment and postprogression lesions to identify acquired alterations, a strategy that is increasingly being used to identify novel alterations mediating therapeutic resistance. Comparison of molecular alterations from pre- and posttreatment biopsies led to discovery of acquired *ESR1* mutations emerging under selective pressure from aromatase inhibitors (25–27), acquired *PTEN* loss in response to PI3K inhibition (13), acquired *RB1* mutations in response to CDK4/6 inhibition (28, 29), and acquired T790M *EGFR* mutations that circumvent inhibition from first- and second-generation *EGFR* inhibitors (30). Whether the emergence of such alterations represents newly acquired mutations versus expansion of a preexisting, low-frequency subclone is a long-standing question. Future improvement in sequencing strategies, including ultra-deep sequencing, will further illuminate this issue.

Currently, there are nine ADCs approved by the FDA in the field of oncology, and more than 100 being investigated in clinical trials. Our study highlights that molecular alterations in both antibody binding target and payload target may

mediate resistance to ADCs, a finding that is likely to have implications for therapeutic sequencing of ADCs. For example, resistance mechanisms to SG due to drug payload target alterations would likely confer cross-resistance to other ADCs with similar (TOP1 inhibitor) payloads, such as trastuzumab deruxtecan and datopotomab deruxtecan (DS1062). In such a scenario, therapeutic sequencing with the same antibody but a different payload could provide therapeutic benefit. On the other hand, resistance mechanisms due to *TROP2* alteration would likely confer cross-resistance to other *TROP2* ADCs such as DS1062, but not to non-*TROP2*-directed ADCs regardless of payload. Another important consideration related to antibody target-associated therapeutic resistance is the bystander effect, which describes the ability of the ADC to kill adjacent non-target-expressing cells. Intratumoral heterogeneity in expression of the target would be predicted to affect ADCs with limited bystander effect more than those with bystander effect, as seen with T-DM1 versus trastuzumab deruxtecan (21). Finally, combination therapies that could increase ADC internalization via effects on the antibody target could potentially overcome resistance and provide synergistic effects, such as seen with T-DM1 and neratinib in the setting of *HER2*-mutant tumors (31).

The study has a few limitations. First, the molecular alterations were seen in a single patient, and the findings and implications for clinical management need to be evaluated in larger studies. Second, patient MGH-18 received two additional therapies after SG (vinorelbine and capecitabine). However, these are not TOP1 inhibitors, and it is therefore unlikely that the TOP1 mutations emerged due to selective pressure from these agents. Third, MGH-18 had a tumor with focal *TACSTD2/TROP2* amplification, which may have increased the chance of developing resistance via acquisition of a mutation in *TACSTD2/TROP2* itself. Whether such mutations would also be seen in tumors without *TACSTD2/TROP2* amplification is unclear and requires further investigation.

In conclusion, our findings highlight the role of parallel genetic alterations in both antibody and payload targets in mediating resistance to ADCs such as SG. Future studies are needed to determine the prevalence of the key resistance mutations we have identified across large numbers of patients and to establish their detailed mechanisms *in vivo*. Additional work will also be required to identify how these mutations temporally emerge under selective pressure from ADCs, in order to enable development of rational, mechanism-based sequential strategies to improve outcomes of patients treated with these agents.

## METHODS

### Patients and Specimen Collection

All biopsies and tumor specimens were collected in accordance with institutional review board (IRB)-approved protocols. Patients provided written informed consent, and all studies were conducted in accordance with the Declaration of Helsinki. Rapid autopsies were performed within the first 3 hours postmortem with informed consent of patient and/or designee under IRB-approved protocol 13-416. Imaging studies, including CT scans, were obtained as part of routine clinical care and/or clinical trial participation. Imaging response was determined by independent radiology review using RECIST version

1.1. Sex, gender, age, and weight were not used as selection or exclusion factors for this work.

### RNA-seq and WES Analysis

For WES, the AllPrep DNA/RNA Mini Kit (Qiagen) was used for dual extraction of both genomic DNA and RNA. DNA was quantified in triplicate using a standardized PicoGreen dsDNA Quantitation Reagent (Invitrogen) assay. The quality control identification check was performed using fingerprint genotyping of 95 common single-nucleotide polymorphisms by Fluidigm Genotyping (Fluidigm). Library construction from double-stranded DNA was performed using the KAPA Library Prep Kit with palindromic forked adapters from Integrated DNA Technologies. Libraries were pooled before hybridization. Hybridization and capture were performed using the relevant components of Illumina's Rapid Capture Enrichment Kit, with a 37-Mb target. All library construction, hybridization, and capture steps were automated on the Agilent Bravo liquid handling system. After postcapture enrichment, library pools were denatured using 0.1 N NaOH on the Hamilton STARlet. Cluster amplification of DNA libraries was performed according to the manufacturer's protocol (Illumina) using HiSeq 4000 exclusion amplification chemistry and HiSeq 4000 flow cells. Flow cells were sequenced utilizing sequencing-by-synthesis chemistry for HiSeq 4000 flow cells. The flow cells were then analyzed using RTA version 2.7.3 or later. Each pool of whole-exome libraries was sequenced on paired 76-cycle runs with two eight-cycle index reads across the number of lanes needed to meet coverage for all libraries in the pool.

### Phylogenetic Analysis of Multiple Samples from the Same Patient

WES data were analyzed on the FireCloud cloud-based analysis platform (<https://portal.firecloud.org/>). Somatic mutations for tumor/normal pairs were detected using the Cancer Genome Analysis WES Characterization Pipeline available on FireCloud. The WES Characterization Pipeline includes multiple steps, including MuTect (for detection of somatic single-nucleotide variants; ref. 32), Strelka (for detecting small insertions and deletions; ref. 33), deTiN (estimates potential tumor-in-normal contamination; ref. 34), ContEst (for detecting cross-patient contamination; ref. 35), AllelicCapSeg (for assessing allele specific copy-number alterations; ref. 36), and ABSOLUTE (for estimating tumor purity, ploidy, absolute allelic copy number, and cancer cell fractions; ref. 36). The data for this study consisted of multiple samples collected from each patient (pre- and posttreatment), as well as autopsy samples. Phylogenetic analysis, subclonal reconstruction, and tree building were done with the PhylogN package (10). Purity of samples was inferred using the established ABSOLUTE method (36).

### Cell Lines, Cell Culture, and Drug Sensitivity Studies

All cell lines were obtained from the MGH Center for Molecular Therapeutics cell bank and underwent high-density SNP typing to confirm their identity. All experiments shown were performed within less than 6 months' passage of all lines since acquisition. Cells were maintained at 37°C in 5% CO<sub>2</sub>. BT549 (ATCC) cells were grown in RPMI (Lonza) supplemented with 10% FBS (SAFC), 1% penicillin (Gibco), and streptomycin (Gibco). NIH3T3 (ATCC) cells were grown in DMEM (Lonza) supplemented with 10% FBS (SAFC), 1% penicillin (Gibco), and streptomycin (Gibco).

pReceiver-based wild-type TROP2 lentiviral vector and empty control vector were purchased from GeneCopoeia. T256R-mutant TROP2 was constructed by introducing the mutation using QuikChange Lightning Site-Directed Mutagenesis Kit (Agilent). The primers for site-directed mutagenesis is: 5'-CTGCAGGTGGAGCGCAGGCTCA TCTATTACCTG-3'.

The lentiviral constructs were transfected into HEK293T cells with LV-MAX Lentiviral Packaging Mix (Gibco) by using the CalPhos Mammalian Transfection Kit (Clontech Laboratories) according to the manufacturer's instructions. The conditioned media containing lentiviral particles were collected 36 hours after transfection and filtered using 0.45- $\mu$ m pore filter (Millipore). The filtered media were then used to infect target cells. Polybrene (Sigma) was added into filtered media at a final concentration of 10  $\mu$ g/mL to increase the infection efficiency. The infected cells were selected with 1  $\mu$ g/mL puromycin (Sigma) 72 hours after infection. Expression of wild-type and T256R-mutant TROP2 was confirmed via Western blotting.

For SG and SN-38 sensitivity studies, cells were seeded in 96-well plates (5,000 cells per well) and exposed to varying doses of SG or SN-38 for 4 hours before changing to fresh medium. Cell viability was assayed after 96 hours. To determine cell viability upon drug treatment, CellTiter-Glo Luminescent Cell Viability Assay (Promega) was used according to the manufacturer's recommendations.

### Protein Extraction, Membrane Fractionation, and Western Blot Analysis

For total protein extraction, cells were lysed in RIPA buffer [10 mmol/L Tris-HCl pH 7.5, 150 mmol/L NaCl, 1 mmol/L EDTA, 1% (w/v) sodium deoxycholate, 0.1% (w/v) SDS, 1% (v/v) NP40, proteinase inhibitor cocktail, phosphatase inhibitor cocktail] for 30 minutes at 4°C.

Cells cultured in 10-cm dish were collected with 500  $\mu$ L fractionation buffer [250 mmol/L sucrose, 20 mmol/L HEPES (pH 7.4), 10 mmol/L KCl, 2 mmol/L MgCl<sub>2</sub>, 1 mmol/L EDTA, 1 mM EGTA, 1 mmol/L DTT, proteinase inhibitor cocktail] and passed through a 25-gauge needle 10 times. Cell suspension was centrifuged at 720  $\times$  g for 5 minutes, and the supernatant was then centrifuged at 10,000  $\times$  g for 5 minutes. The supernatant was then centrifuged at 100,000  $\times$  g for 1 hour. Pellets were washed by adding 500  $\mu$ L of fractionation buffer and resuspended by pipetting and passing through a 25-G needle. The sample was centrifuged for 1 hour. The membrane pellets were resuspended with SDS sample buffer.

Protein samples were mixed with SDS sample buffer and boiled for 10 minutes before being subjected to SDS-PAGE. The protein samples on the SDS-PAGE gel were then transferred onto PVDF membrane (Millipore), which was blocked by 5% nonfat milk in PBST (PBS plus 0.02% Tween 20) at room temperature for 1 hour. Then, the PVDF membrane was incubated with primary antibodies diluted in 3% BSA in PBST at 4°C overnight and horseradish peroxidase-conjugated secondary antibodies (Sigma) diluted in 3% BSA in PBST at room temperature for 2 hours. The signal was detected by enhanced chemiluminescence solution (PerkinElmer).

### Live-Cell Flow Cytometry

BT549 or NIH3T3 cells were plated, allowed to attach overnight, and then harvested by trypsinization the following day. Cell pellets were washed with 1 mL of ice-cold 1 $\times$  PBS and maintained at 4°C for further processing. Cells were blocked for 30 minutes with 1% BSA + 5% normal goat serum (Sigma) in 1 $\times$  PBS, incubated with 5  $\mu$ g/mL hRs7 (Immunomedics/Gilead, Immunomedics, Inc.), and then incubated with a goat anti-human F(ab)<sub>2</sub> immunoglobulin secondary (Jackson ImmunoResearch). Samples were examined using a FACS Aria flow cytometer. Analysis was conducted with FlowJo using Cell Quest software (Becton Dickinson). Data were representative of three biological repeats with at least triple technical replicates.

### Immunofluorescence and Confocal Microscopy

BT549 or NIH3T3 cells were lifted and plated at subconfluent densities into 96-well plates having #1.5 glass coverslips to facilitate confocal microscopy (Cellvis, P96-1.5H-N). Cells were washed with 1 $\times$  PBS and then fixed at room temperature using 4% PFA (Chem-Cruz) for 15 minutes at a gentle shake. Thereafter, cells were washed

thrice with 1× PBS and then incubated with 0.5% Triton X-100 (Sigma) for 5 minutes to permeabilize. Blocking then used 1% BSA with 5% normal goat serum and 0.1% Triton X-100, followed by one of two anti-TROP2 antibodies (as indicated) for 1 hour at room temperature. Cells were then washed with 1% BSA in 1× PBS thrice for 20 minutes and incubated with a fluorescent FITC secondary antibody (Jackson ImmunoResearch, 1:500) for 1 hour. After incubation with secondary, cells were again washed thrice for 20 minutes with 1% BSA, 5% normal goat serum, and 0.1% Triton X-100 in 1× PBS. Nuclei were counterstained with 0.5 μg/mL 4',6-diamidino-2-phenylindole (DAPI; Sigma) and maintained in 200 μL PBS for imaging. Images were acquired using a Nikon A1R confocal microscope with a 60× oil immersion objective, a physical magnification of 1.5×, and a lens magnification of 1.3×, the latter of which was to eliminate distal warping. Images were converted and compiled using ImageJ software.

Antibodies used are listed in Supplementary Table S4.

### Statistical Analysis

Tests to determine significance are detailed in the relevant figure legends. Briefly, *t* tests or one-way ANOVAs were used to determine statistical significance of treatment conditions and two-way ANOVA to discriminate between the contributions of both dose and transformation.

Upload of data from WES and RNA-seq is available in dbGaP, accession number phs2555.v1.

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### Authors' Contributions

**J.T. Coates:** Data curation, supervision, validation, investigation, visualization, methodology, writing–review and editing. **S. Sun:** Data curation, supervision, validation, investigation, methodology,

writing–review and editing. **I. Leshchiner:** Formal analysis, supervision, investigation, writing–review and editing. **N. Thimmiah:** Investigation, writing–review and editing. **E.E. Martin:** Formal analysis, writing–review and editing. **D. McLoughlin:** Supervision, project administration, writing–review and editing. **B.P. Danysh:** Software, project administration, writing–review and editing. **K. Slowik:** Software and project administration. **R.A. Jacobs:** Formal analysis, project administration, writing–review and editing. **K. Rhrissorrakrai:** Funding acquisition. **F. Utro:** Funding acquisition and writing–review and editing. **C. Levovitz:** Funding acquisition. **E. Denault:** Project administration. **C.S. Walmsley:** Conceptualization, formal analysis, supervision, funding acquisition, writing–original draft, project administration, writing–review and editing, and specimen acquisition. **A. Kambadakone:** Investigation. **J.R. Stone:** Conceptualization, software, formal analysis, supervision, funding acquisition, writing–original draft, project administration, writing–review and editing, and contributed patients. **S.J. Isakoff:** Conceptualization, software, formal analysis, supervision, funding acquisition, writing–original draft, project administration, writing–review and editing, contributed patients. **L. Parida:** Conceptualization, formal analysis, supervision, funding acquisition, writing–original draft, project administration, writing–review and editing. **D. Juric:** Conceptualization, formal analysis, supervision, funding acquisition, writing–original draft, project administration, writing–review and editing. **G. Getz:** Conceptualization, software, formal analysis, supervision, funding acquisition, writing–original draft, project administration, writing–review and editing. **A. Bardia:** Conceptualization, software, funding acquisition, writing–original draft, project administration, writing–review and editing. **L.W. Ellisen:** Conceptualization, software, funding acquisition, writing–original draft, project administration, writing–review and editing.

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### REFERENCES

- Bianchini G, Balko JM, Mayer IA, Sanders ME, Gianni L. Triple-negative breast cancer: challenges and opportunities of a heterogeneous disease. *Nat Rev Clin Oncol* 2016;13:674–90.
- Carey L, Winer E, Viale G, Cameron D, Gianni L. Triple-negative breast cancer: disease entity or title of convenience? *Nat Rev Clin Oncol* 2010;7:683–92.
- Anders CK, Zagar TM, Carey LA. The management of early-stage and metastatic triple-negative breast cancer: a review. *Hematol Oncol Clin North Am* 2013;27:737–49.
- Gradishar WJ, Anderson BO, Abraham J, Aft R, Agnese D, Allison KH, et al. Breast cancer, version 3.2020, NCCN clinical practice guidelines in oncology. *J Natl Compr Canc Netw* 2020;18:452–78.

5. Nicolo E, Zagami P, Curigliano G. Antibody-drug conjugates in breast cancer: the chemotherapy of the future? *Curr Opin Oncol* 2020;32:494–502.
6. Nagayama A, Vidula N, Ellisen L, Bardia A. Novel antibody-drug conjugates for triple negative breast cancer. *Ther Adv Med Oncol* 2020;12:1758835920915980.
7. Bardia A, Mayer IA, Vahdat LT, Tolaney SM, Isakoff SJ, Diamond JR, et al. Sacituzumab govitecan-hziy in refractory metastatic triple-negative breast cancer. *N Engl J Med* 2019;380:741–51.
8. Bardia A, Hurvitz SA, Tolaney SM, Loirat D, Punie K, Oliveira M, et al. Sacituzumab govitecan in metastatic triple-negative breast cancer. *N Engl J Med* 2021;384:1529–41.
9. Kalinsky K, Diamond JR, Vahdat LT, Tolaney SM, Juric D, O'Shaughnessy J, et al. Sacituzumab govitecan in previously treated hormone receptor-positive/HER2-negative metastatic breast cancer: final results from a phase I/II, single-arm, basket trial. *Ann Oncol* 2020;31:1709–18.
10. Leshchiner I, Livitz D, Gainor JF, Rosebrock D, Spiro O, Martinez A, et al. Comprehensive analysis of tumour initiation, spatial and temporal progression under multiple lines of treatment. *bioRxiv* 2018.
11. Chang JY, Liu JF, Juang SH, Liu TW, Chen LT. Novel mutation of topoisomerase I in rendering cells resistant to camptothecin. *Cancer Res* 2002;62:3716–21.
12. Fiorani P, Chillemi G, Losasso C, Castelli S, Desideri A. The different cleavage DNA sequence specificity explains the camptothecin resistance of the human topoisomerase I Glu418Lys mutant. *Nucleic Acids Res* 2006;34:5093–100.
13. Juric D, Castel P, Griffith M, Griffith OL, Won HH, Ellis H, et al. Convergent loss of PTEN leads to clinical resistance to a PI(3)Kalpha inhibitor. *Nature* 2015;518:240–4.
14. Tanaka N, Lin JJ, Li C, Ryan MB, Zhang J, Kiedrowski LA, et al. Clinical acquired resistance to KRASG12C inhibition through a novel KRAS switch-II pocket mutation and polyclonal alterations converging on RAS-MAPK reactivation. *Cancer Discov* 2021;11:1913–22.
15. Bon G, Pizzuti L, Laquintana V, Loria R, Porru M, Marchio C, et al. Loss of HER2 and decreased T-DM1 efficacy in HER2 positive advanced breast cancer treated with dual HER2 blockade: the SeP-HER study. *J Exp Clin Cancer Res* 2020;39:279.
16. Garcia-Alonso S, Ocana A, Pandiella A. Resistance to antibody-drug conjugates. *Cancer Res* 2018;78:2159–65.
17. Baselga J, Lewis Phillips GD, Verma S, Ro J, Huober J, Guardino AE, et al. Relationship between tumor biomarkers and efficacy in EMILIA, a phase III study of trastuzumab emtansine in HER2-positive metastatic breast cancer. *Clin Cancer Res* 2016;22:3755–63.
18. Bardia A, Tolaney SM, Punie K, Loirat D, Oliveira M, Kalinsky K, et al. Biomarker analyses in the phase III ASCENT study of sacituzumab govitecan versus chemotherapy in patients with metastatic triple-negative breast cancer. *Ann Oncol* 2021;32:1148–56.
19. Urasaki Y, Laco GS, Pourquier P, Takebayashi Y, Kohlhagen G, Gioffre C, et al. Characterization of a novel topoisomerase I mutation from a camptothecin-resistant human prostate cancer cell line. *Cancer Res* 2001;61:1964–9.
20. Ogitani Y, Aida T, Hagihara K, Yamaguchi J, Ishii C, Harada N, et al. DS-8201a, a novel HER2-targeting ADC with a novel DNA topoisomerase I inhibitor, demonstrates a promising antitumor efficacy with differentiation from T-DM1. *Clin Cancer Res* 2016;22:5097–108.
21. Yver A, Agatsuma T, Soria JC. The art of innovation: clinical development of trastuzumab deruxtecan and redefining how antibody-drug conjugates target HER2-positive cancers. *Ann Oncol* 2020;31:430–4.
22. McDougall AR, Tolcos M, Hooper SB, Cole TJ, Wallace MJ. Trop2: from development to disease. *Dev Dyn* 2015;244:99–109.
23. Hunter FW, Barker HR, Lipert B, Rothe F, Gebhart G, Piccart-Gebhart MJ, et al. Mechanisms of resistance to trastuzumab emtansine (T-DM1) in HER2-positive breast cancer. *Br J Cancer* 2020;122:603–12.
24. Weiss J, Glode A, Messersmith WA, Diamond J. Sacituzumab govitecan: breakthrough targeted therapy for triple-negative breast cancer. *Expert Rev Anticancer Ther* 2019;19:673–9.
25. Robinson DR, Wu YM, Vats P, Su F, Lonigro RJ, Cao X, et al. Activating ESR1 mutations in hormone-resistant metastatic breast cancer. *Nat Genet* 2013;45:1446–51.
26. Toy W, Shen Y, Won H, Green B, Sakr RA, Will M, et al. ESR1 ligand-binding domain mutations in hormone-resistant breast cancer. *Nat Genet* 2013;45:1439–45.
27. Yu M, Bardia A, Aceto N, Bersani F, Madden MW, Donaldson MC, et al. Cancer therapy: ex vivo culture of circulating breast tumor cells for individualized testing of drug susceptibility. *Science* 2014;345:216–20.
28. Condorelli R, Spring L, O'Shaughnessy J, Lacroix L, Bailleux C, Scott V, et al. Polyclonal RB1 mutations and acquired resistance to CDK 4/6 inhibitors in patients with metastatic breast cancer. *Ann Oncol* 2018;29:640–5.
29. O'Leary B, Cutts RJ, Liu Y, Hrebien S, Huang X, Fenwick K, et al. The genetic landscape and clonal evolution of breast cancer resistance to palbociclib plus fulvestrant in the PALOMA-3 trial. *Cancer Discov* 2018;8:1390–403.
30. Kobayashi S, Boggon TJ, Dayaram T, Janne PA, Kocher O, Meyerson M, et al. EGFR mutation and resistance of non-small-cell lung cancer to gefitinib. *N Engl J Med* 2005;352:786–92.
31. Li BT, Michelini F, Misale S, Cocco E, Baldino L, Cai Y, et al. HER2-mediated internalization of cytotoxic agents in ERBB2 amplified or mutant lung cancers. *Cancer Discov* 2020;10:674–87.
32. Cibulskis K, Lawrence MS, Carter SL, Sivachenko A, Jaffe D, Sougnez C, et al. Sensitive detection of somatic point mutations in impure and heterogeneous cancer samples. *Nat Biotechnol* 2013;31:213–9.
33. Kim S, Scheffler K, Halpern AL, Bekritsky MA, Noh E, Källberg M, et al. Strelka2: fast and accurate calling of germline and somatic variants. *Nat Methods* 2018;15:591–4.
34. Taylor-Weiner A, Stewart C, Giordano T, Miller M, Rosenberg M, Macbeth A, et al. DeTiN: overcoming tumor-in-normal contamination. *Nat Methods* 2018;15:531–4.
35. Cibulskis K, McKenna A, Fennell T, Banks E, DePristo M, Getz G. ContEst: estimating cross-contamination of human samples in next-generation sequencing data. *Bioinformatics* 2011;27:2601–2.
36. Carter SL, Cibulskis K, Helman E, McKenna A, Shen H, Zack T, et al. Absolute quantification of somatic DNA alterations in human cancer. *Nat Biotechnol* 2012;30:413–21.

# CANCER DISCOVERY

## Parallel Genomic Alterations of Antigen and Payload Targets Mediate Polyclonal Acquired Clinical Resistance to Sacituzumab Govitecan in Triple-Negative Breast Cancer

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