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Jongsma, M.L.M.; Waard, A.A. de; Raaben, M.; Zhang, T.; Cabukusta, B.; Platzer, R.; ... ; Spaapen, R.M.

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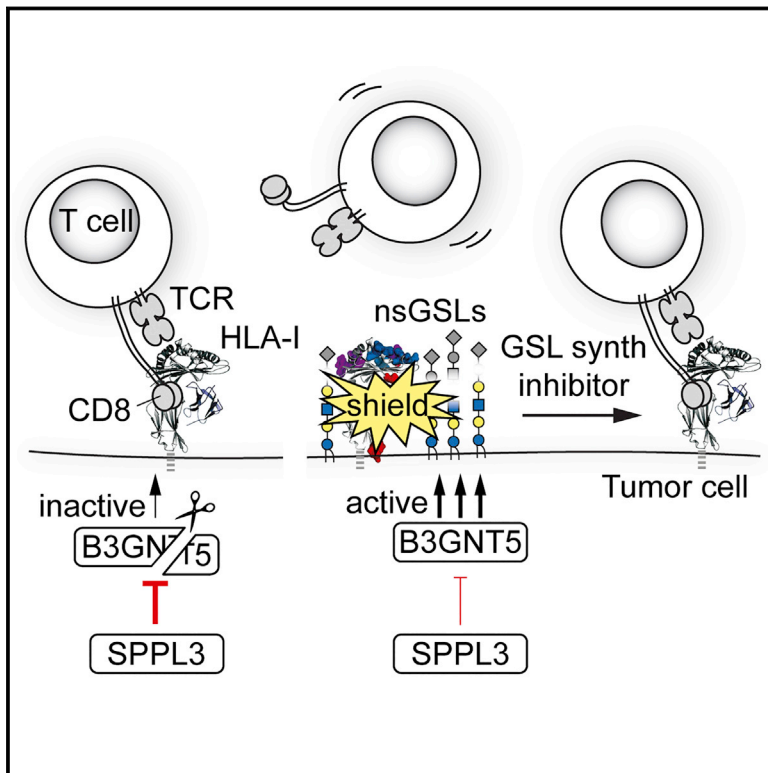
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# Immunity

## The SPPL3-Defined Glycosphingolipid Repertoire Orchestrates HLA Class I-Mediated Immune Responses

### Graphical Abstract



### Authors

Marlieke L.M. Jongsma,  
 Antonius A. de Waard,  
 Matthijs Raaben, ...,  
 Thijn R. Brummelkamp,  
 Jacques Neefjes, Robbert M. Spaapen

### Correspondence

r.spaapen@sanquin.nl

### In Brief

Numerous glycosphingolipids (GSLs) are expressed at the cell surface; however, their functions remain mostly unknown. Jongsma et al. identify that the protease SPPL3 destroys the GSL synthesis enzyme B3GNT5, whose GSL products negatively affect HLA-I-mediated immune responses. This pathway represents a potential therapeutic target in cancer, infection, and autoimmunity.

### Highlights

- Iterative KO screens reveal a new pathway controlling HLA-I antigen presentation
- SPPL3 suppresses B3GNT5 activity affecting the cell surface GSL repertoire
- B3GNT5-generated GSLs limit the capacity of HLA-I to interact with natural ligands
- Inhibition of GSL synthesis in glioma enhances anti-tumor immune activation



## Article

# The SPPL3-Defined Glycosphingolipid Repertoire Orchestrates HLA Class I-Mediated Immune Responses

Marlieke L.M. Jongsma,<sup>1,2,3,4</sup> Antonius A. de Waard,<sup>1,2,3,18</sup> Matthijs Raaben,<sup>5,18</sup> Tao Zhang,<sup>6,18</sup> Birol Cabukusta,<sup>4</sup> René Platzer,<sup>7</sup> Vincent A. Blomen,<sup>5</sup> Anastasia Xagara,<sup>1,2,3</sup> Tamara Verkerk,<sup>1,2,3</sup> Sophie Bliss,<sup>1,2,3</sup> Xiangrui Kong,<sup>1,2,3</sup> Carolin Gerke,<sup>8,9,10,11</sup> Lennert Janssen,<sup>4</sup> Elmer Stickel,<sup>5</sup> Stephanie Holst,<sup>6</sup> Rosina Plomp,<sup>6</sup> Arend Mulder,<sup>12</sup> Soldano Ferrone,<sup>13</sup> Frans H.J. Claas,<sup>12</sup> Mirjam H.M. Heemskerk,<sup>14</sup> Marieke Griffioen,<sup>14</sup> Anne Halenius,<sup>8,9</sup> Hermen Overkleef,<sup>15</sup> Johannes B. Huppa,<sup>7</sup> Manfred Wuhrer,<sup>6</sup> Thijn R. Brummelkamp,<sup>5,16,17</sup> Jacques Neefjes,<sup>4</sup> and Robbert M. Spaapen<sup>1,2,3,19,\*</sup>

<sup>1</sup>Department of Immunopathology, Sanquin Research, Amsterdam, the Netherlands

<sup>2</sup>Landsteiner Laboratory, Amsterdam UMC, University of Amsterdam, Amsterdam, the Netherlands

<sup>3</sup>Cancer Center Amsterdam, Amsterdam, the Netherlands

<sup>4</sup>Oncode Institute and Department of Cell and Chemical Biology, LUMC, Leiden, the Netherlands

<sup>5</sup>Oncode Institute, Division of Biochemistry, the Netherlands Cancer Institute, Amsterdam, the Netherlands

<sup>6</sup>Center for Proteomics and Metabolics, LUMC, Leiden, the Netherlands

<sup>7</sup>Institut für Hygiene und Angewandte Immunologie, Vienna, Austria

<sup>8</sup>Institute of Virology, Medical Center University of Freiburg, Freiburg, Germany

<sup>9</sup>Faculty of Medicine, University of Freiburg, Freiburg, Germany

<sup>10</sup>Spemann Graduate School of Biology and Medicine, University of Freiburg, Freiburg, Germany

<sup>11</sup>Faculty of Biology, University of Freiburg, Freiburg, Germany

<sup>12</sup>Department of Immunology, LUMC, Leiden, the Netherlands

<sup>13</sup>Department of Surgery, Massachusetts General Hospital, Harvard Medical School, Boston, MA, USA

<sup>14</sup>Department of Hematology, Leiden University Medical Center, Leiden, the Netherlands

<sup>15</sup>Leiden Institute of Chemistry, Leiden University, Leiden, the Netherlands

<sup>16</sup>CeMM Research Center for Molecular Medicine of the Austrian Academy of Sciences, Vienna, Austria

<sup>17</sup>Cancer Genomics Center, Amsterdam, the Netherlands

<sup>18</sup>These authors contributed equally

<sup>19</sup>Lead Contact

\*Correspondence: [r.spaapen@sanquin.nl](mailto:r.spaapen@sanquin.nl)

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## SUMMARY

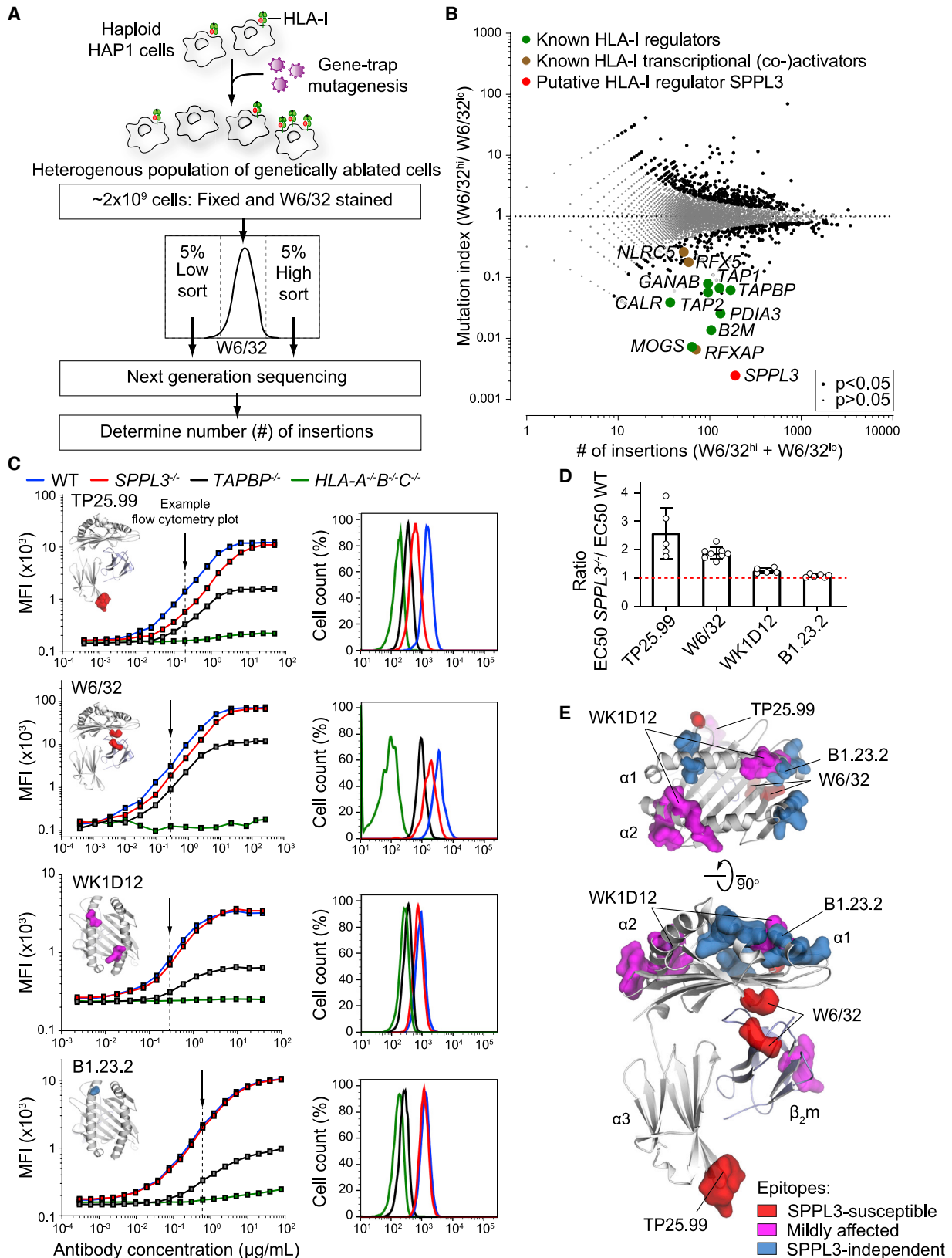
HLA class I (HLA-I) glycoproteins drive immune responses by presenting antigens to cognate CD8<sup>+</sup> T cells. This process is often hijacked by tumors and pathogens for immune evasion. Because options for restoring HLA-I antigen presentation are limited, we aimed to identify druggable HLA-I pathway targets. Using iterative genome-wide screens, we uncovered that the cell surface glycosphingolipid (GSL) repertoire determines effective HLA-I antigen presentation. We show that absence of the protease SPPL3 augmented B3GNT5 enzyme activity, resulting in upregulation of surface neolacto-series GSLs. These GSLs sterically impeded antibody and receptor interactions with HLA-I and diminished CD8<sup>+</sup> T cell activation. Furthermore, a disturbed SPPL3-B3GNT5 pathway in glioma correlated with decreased patient survival. We show that the immunomodulatory effect could be reversed through GSL synthesis inhibition using clinically approved drugs. Overall, our study identifies a GSL signature that inhibits immune recognition and represents a potential therapeutic target in cancer, infection, and autoimmunity.

## INTRODUCTION

Human leukocyte antigen class I (HLA-I) glycoproteins are the primary modules recognized by CD8<sup>+</sup> T cells determining both the induction and the effector phases of adaptive immune responses. Their primary function is to present peptide fragments from degraded proteins to the T cell receptor (TCR) of cytotoxic CD8<sup>+</sup>

T cells, leading to T cell-mediated elimination of target cells (Neefjes et al., 2011; Unanue and Cerottini, 1989). Because HLA-I molecules on tumor cells also present tumor antigen-derived peptides to cognate T cells, they play a major role in the anti-tumor activity of T cells unleashed by current immunotherapeutic strategies (Schumacher and Schreiber, 2015). As a consequence, tumors often escape from immune surveillance by downregulating





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one or multiple molecules critical in HLA-I antigen presentation (Chowell et al., 2018; Gettinger et al., 2017; Restifo et al., 1996; Sade-Feldman et al., 2017; Zaretsky et al., 2016). This reduction is often reversible, for example, by interferon (IFN) stimulation, ionizing radiation, or inhibition of histone deacetylases, which has led to various experimental therapies aimed at increasing tumor HLA-I surface expression (Haworth et al., 2015; Reits et al., 2006; Thor Straten and Garrido, 2016). Moreover, sensitizing tumor cells as immune targets can act synergistically with T cell activating and re-activating strategies, thereby increasing the therapeutic potential of enhancing HLA-I availability (Hähnel et al., 2008). Active suppression of HLA-I surface expression to escape T cell surveillance is also employed by an array of pathogens, such as Epstein-Barr virus, cytomegalovirus, and SARS-CoV-2 (Hansen and Bouvier, 2009; Yewdell and Hill, 2002; Zhang et al., 2020). These examples underscore the broad relevance of HLA-I-based interventions, necessitating a thorough understanding of the molecular mechanisms underpinning the HLA-I pathway.

Over the last 35 years, several key elements in HLA-I expression and antigen presentation have been identified and extensively studied. A protein complex directed by the transcriptional regulator *NLRC5* drives HLA-I expression in selected tissues, whereas HLA-I translation and glycosylation are thought to be executed by general enzymes and mechanisms (Jongsma et al., 2019; Ryan and Cobb, 2012). In the endoplasmic reticulum (ER), the HLA-I heavy chain and its light chain  $\beta_2m$  microglobulin ( $\beta_2m$ ) assemble and are stabilized by a unique combination of the ER chaperone proteins: tapasin (TAPBP), Erp57 (PDIA3), and calreticulin (CALR) (Rock et al., 2016). These HLA-I chaperone complexes bind the peptide transporter TAP to form the peptide-loading complex (PLC), which drives efficient ER import and loading of peptides into the HLA-I peptide binding groove (Blees et al., 2017). Subsequently, mature trimeric complexes of HLA-I heavy chain,  $\beta_2m$ , and peptide are released from the PLC and transported to the cell surface for peptide presentation to T cells (Garstka et al., 2015; Wearsch and Cresswell, 2008). Given the multifactorial complexity of the HLA-I antigen presentation pathway, we hypothesized that additional regulatory mechanisms of this central process in adaptive immunity must exist.

To uncover additional components involved in successful HLA-I antigen presentation, we performed a series of genome-wide haploid genetic screens. We identified the enzyme signal peptide peptidase-like 3 (SPPL3) as a positive regulator of HLA-I antigen presentation. We found that SPPL3 controlled the composition of the cell surface glycosphingolipid (GSL) repertoire by inhibiting the glycosyltransferase B3GNT5. In the absence of SPPL3, an in-

crease in B3GNT5 activity led to a high amount of complex negatively charged neolacto-series GSLs (nsGSLs), preventing HLA-I from being accessed by several immune cell receptors and interfering with activation of CD8<sup>+</sup> T cells. GSL synthesis in several tumors, including glioma, was skewed toward the nsGSL synthesis pathway. High activation of this pathway negatively correlated with survival of glioma patients. We show that intervention of GSL synthesis in glioma cells by U.S. Food and Drug Administration (FDA)-approved drugs led to improved anti-tumor immunity *in vitro*. In conclusion, nsGSL synthesis constitutes a targetable layer of immune regulation through HLA-I shielding at the cell surface.

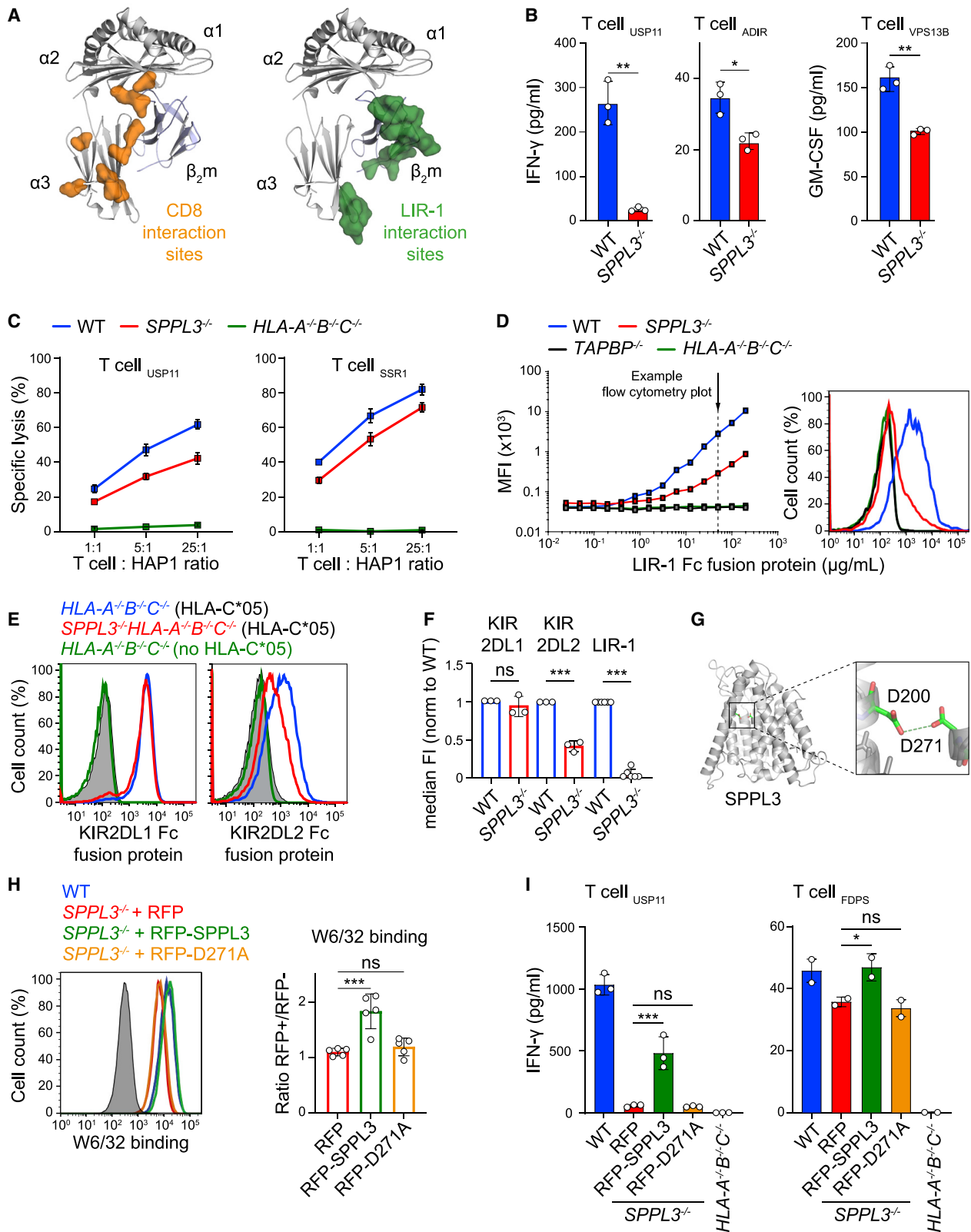
## RESULTS

### A Haploid Genetic Screen Provides a High-Resolution Map of the HLA-I Antigen Presentation Pathway

To identify unknown factors regulating HLA-I antigen presentation, we performed a genome-wide insertional mutagenesis screen in haploid human fibroblast-like HAP1 cells endogenously expressing HLA-I (Brockmann et al., 2017; Carette et al., 2009). A heterogeneous pool of millions of cells, each genetically ablated for a random gene or set of genes, was generated by retroviral gene trap insertion and expanded. Mutagenized cells that were poorest or best recognized by the HLA-I-specific W6/32 antibody were sorted by flow cytometry (Figure 1A). Subsequently, we determined the relative enrichment of unique disruptive integrations per gene between the sorted populations using deep sequencing. This provided an unbiased overview of genes involved in HLA-I antigen presentation (Figure 1B). Among the highly significant positive modifiers were the known key genes directing HLA-I expression, such as the transcriptional activators and coactivators *NLRC5*, *RFXAP*, and *RFX5* (Jongsma et al., 2019) and essential components for post-transcriptional assembly, glycosylation, and peptide loading: *B2M*, *MOGS* ( $\alpha$ -glucosidase I), *GANAB* ( $\alpha$ -glucosidase II), *TAP1*, *TAP2*, *TAPBP* (tapasin), *PDIA3* (Erp57), and *CALR* (Figure 1B) (Wearsch and Cresswell, 2008). Thus, this single genetic map identified components of the HLA-I pathway previously discovered through decades of research. In addition, HLA-I regulators identified in human B cell lymphoma CRISPR-Cas9 screens by Dersh et al., such as *SUSD6*, *SND1*, *ANKRD33B*, and *EZH2*, were confirmed as highly significant hits in our haploid genetic screen (Dersh et al., 2020). The most prominent hit from our screen in the W6/32<sup>lo</sup>-sorted population was the gene encoding SPPL3, a protease that had not been previously described in the context of antigen presentation (Figure 1B). SPPL3 is an ER- and Golgi-localized transmembrane

### Figure 1. A Haploid Genetic Screen Reveals SPPL3 as a Regulator of Antibody Accessibility to Membrane-Proximal HLA-I Regions

(A) Schematic overview of a genome-wide haploid genetic screen using HLA-I-specific W6/32 antibody.  
 (B) Fish-tail plot showing per gene the mutation index (ratio of integrations mapped in the specified populations) against the amount of mapped integrations. A two-sided false discovery rate (FDR) (Benjamini-Hochberg) corrected Fisher's exact test was applied. The symbol legend is indicated.  
 (C) (Left) Titration curves of four HLA-I-specific antibodies on mixed barcoded HAP1 cells (see Figure S1E). The individual antibody binding epitopes are shown on the HLA-I structure. (Right) Flow cytometry histograms of nonsaturating antibody stain as indicated by the arrow (close to EC<sub>50</sub> values). MFI, mean fluorescence intensity.  
 (D) Quantification of (C) using the ratio of WT and SPPL3<sup>-/-</sup> EC<sub>50</sub> values. Mean  $\pm$  SD is plotted (n = 5–8 independent experiments).  
 (E) SPPL3-susceptibility of different epitopes is plotted on the HLA-I/ $\beta_2m$  crystal structure (individual epitopes in Figures 1C and S2A). See also Figures S1 and S2.



**Figure 2. SPPL3 Expression Promotes LIR-1 Binding to HLA-I and Enhances CD8<sup>+</sup> T Cell Activation**

(A) CD8 and LIR-1 interaction sites mapped on the HLA-I/ $\beta_2m$  crystal structure.

(B) IFN- $\gamma$  production by T cells recognizing the specified endogenously derived antigens after coculture with the indicated HAP1 cells (n = 3).

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protein of the family of intramembrane-cleaving aspartyl proteases (Fluhrer and Haass, 2007).

### SPPL3 Enables Antibody Binding to Membrane-Proximal Regions of HLA-I

To validate that HLA-I cell surface expression was altered by SPPL3, we created *SPPL3*<sup>-/-</sup> HAP1 cells using CRISPR-Cas9 (Table S1) and performed flow cytometry using the W6/32 antibody. HLA-I surface expression in the absence of SPPL3 turned out identical to those of wild-type (WT) cells (Figure S1A). Likewise, the total HLA-I content of *SPPL3*<sup>-/-</sup> and WT cells was similar, as determined by immunoblot analysis (Figure S1B). Furthermore, SPPL3 deficiency did not alter HLA-I turnover or stability of the peptide-HLA-I interaction (Figures S1C and S1D). Because the anti-HLA antibody W6/32 was used under nonsaturating staining conditions in the genome-wide screen, these seemingly contradictory outcomes may have resulted from reduced accessibility of the W6/32 epitope in the absence of SPPL3.

To test this hypothesis, we titrated the W6/32 antibody for binding to WT, *SPPL3*<sup>-/-</sup>, and control *HLA-A*<sup>-/-</sup>*B*<sup>-/-</sup>*C*<sup>-/-</sup> (HLA-I-deficient) and *TAPBP*<sup>-/-</sup> cells (Table S1), which were individually color barcoded and combined in a single well for optimal comparison of staining intensity (Figure S1E). In contrast to saturating W6/32 concentrations, lower W6/32 concentrations resulted in decreased binding to *SPPL3*<sup>-/-</sup> cells compared with WT cells, indicating that accessibility of the HLA-I epitope recognized by W6/32 was indeed hindered (Figures 1C and 1D). A similar result was apparent using monovalent W6/32 Fab fragments, validating this conclusion (Figure S2A). To further define SPPL3-dependent antibody accessibility to HLA-I, we performed additional titrations using 13 HLA-I-specific antibodies recognizing distinct HLA-I epitopes and one targeting a  $\beta_2m$  epitope (Figures 1C and S2A; Table S2). The binding of three antibodies (clones W6/32, TP25.99, and ROU9A6) was markedly affected by the absence of SPPL3 (Figures 1C, 1D, and S2A). By superimposing critical amino acid positions for binding of the individual antibodies onto an HLA-I structure, we defined the SPPL3-susceptible region as relatively proximal to the cellular membrane, an area that is largely conserved among HLA-I alleles (Figure 1E). Binding of several other antibodies (e.g., B1.23.2) was not affected by SPPL3, supporting that HLA-I surface expression is not targeted by SPPL3 and providing unique intra-molecular controls for further experiments. To determine whether SPPL3 differentially affected HLA-A, HLA-B, or HLA-C alleles, we recon-

stituted *HLA-A*<sup>-/-</sup>*B*<sup>-/-</sup>*C*<sup>-/-</sup> cells on a WT or *SPPL3*<sup>-/-</sup> background (Table S1) with the single original HLA-I alleles and analyzed their accessibility for antibodies. Each allele showed a difference in HLA-I accessibility between WT and *SPPL3*-deficient cells comparable to that detected by W6/32 (Figure S2B). In addition, other cell lines exhibited a similar decrease in W6/32 accessibility to HLA-I after small interfering RNA (siRNA) silencing or CRISPR-Cas9 genetic ablation of SPPL3, indicating that regulation of HLA-I accessibility is not solely restricted to HAP1 cells or their HLA-I haplotype (Figures S2C–S2E).

### SPPL3 Expression Promotes Receptor Binding to HLA-I and CD8<sup>+</sup> T Cell Activation

SPPL3 modulates antibody reactivity toward specific regions of HLA-I molecules, which raises the question of whether the HLA-I function is affected. The CD8 coreceptor, which on most T cells is essential for sensitizing responsiveness by supporting TCR docking to its cognate peptide-HLA-I complex, ligates closely to the region of HLA-I that is affected by SPPL3 (as depicted in Figures 1E and 2A) (Gao et al., 1997; Purbhoo et al., 2004; Roszkowski et al., 2003). We evaluated whether SPPL3 enhances HLA-I antigen presentation to CD8<sup>+</sup> T cells by stimulating multiple HLA-A\*02:01-restricted T cell clones specific for different tumor-expressed antigens endogenously expressed by WT and *SPPL3*<sup>-/-</sup> cells (Amir et al., 2011; van Bergen et al., 2007, 2010). All clones were more reactive to SPPL3-expressing WT cells, as determined by their IFN- $\gamma$  or granulocyte-macrophage colony-stimulating factor (GM-CSF) production (Figure 2B), indicating that SPPL3 increases functional access to HLA-I. The effect of SPPL3 on T cell function was confirmed in <sup>51</sup>Cr release assays showing reduced killing of *SPPL3*<sup>-/-</sup> cells compared with WT cells (Figures 2C and S3A).

Disturbed receptor accessibility to HLA-I may affect not only T cells but also other immune cells that express HLA-I-interacting proteins, including the inhibitory leukocyte immunoglobulin (Ig)-like receptor (LIR) and killer cell Ig-like receptor (KIR) families that suppress unwanted immune cell activation (Saverino et al., 2000; Valiante et al., 1997). The defined SPPL3-susceptible region on the HLA-I protein highly overlaps with the binding site of LIR-1, which is expressed by monocytes, B cells, and small subsets of natural killer (NK) cells and T cells (Borges et al., 1997; Colonna et al., 1997; Cosman et al., 1997) (Figure 2A). Even more pronounced than for SPPL3-affected antibodies, binding of a recombinant LIR-1 Fc fusion protein (Gonen-Gross

(C) Normalized <sup>51</sup>Cr release (specific lysis) of the indicated HAP1 cells targeted by specified T cells at different effector-target (E:T) ratios (n = 3) (see also Figure S3A).

(D) (Left) Representative titration curves of LIR-1 Fc fusion protein on indicated HAP1 cells (n = 2). (Right) Histogram of LIR-1 Fc binding at the indicated concentration (arrow).

(E) Flow cytometry staining of HLA-C\*05:01-transduced *SPPL3*<sup>-/-</sup>*HLA-A*<sup>-/-</sup>*B*<sup>-/-</sup>*C*<sup>-/-</sup> or depicted control HAP1 cells with nonsaturating concentrations of KIR2DL1 and KIR2DL2 Fc fusion proteins. Gray, unstained control.

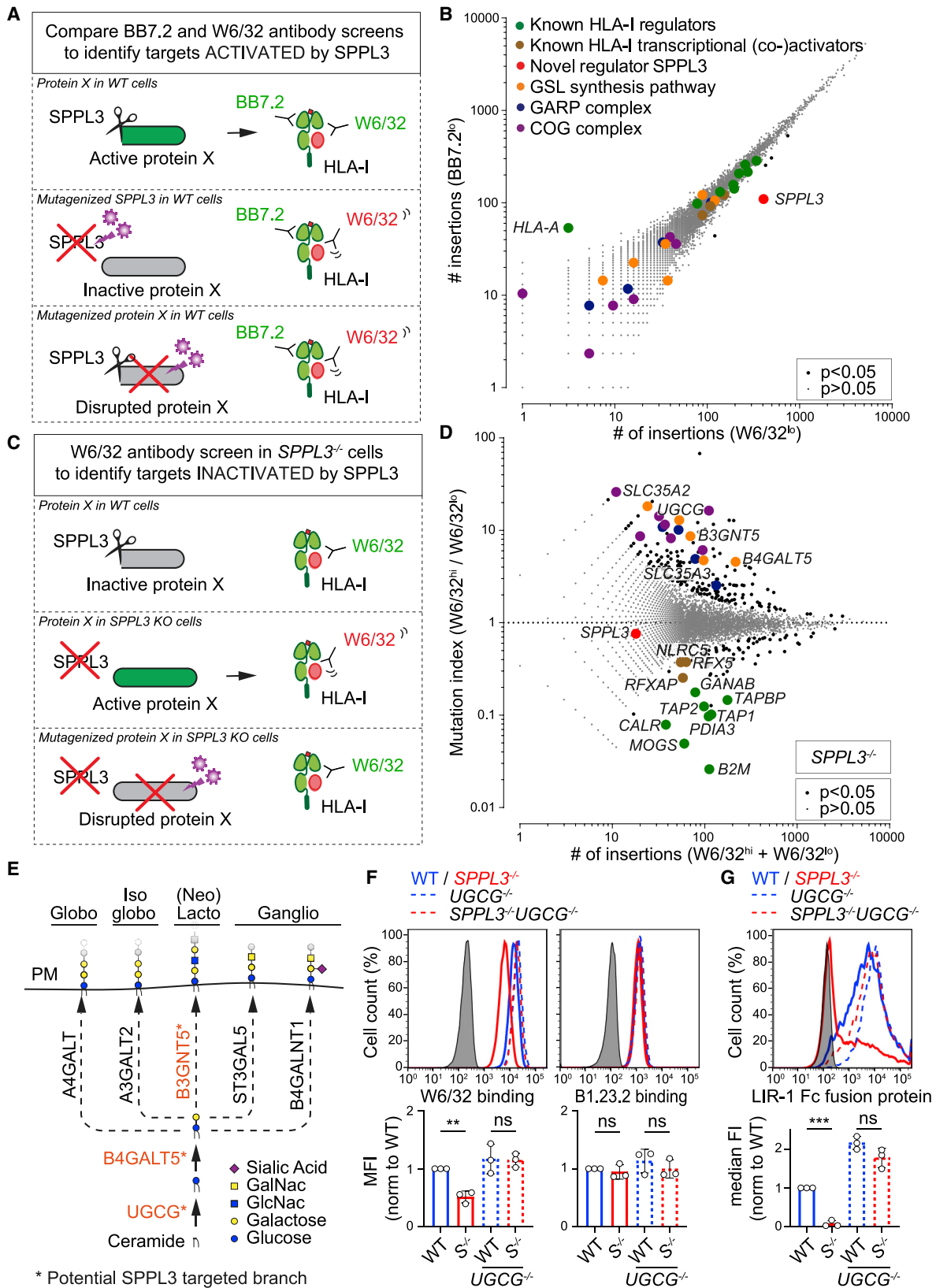
(F) Normalized quantification (median fluorescence intensity [median FI]) of HLA-I binding by the indicated fusion proteins (including data from D, E, and Figures 3G and 4E) (n = 3–6).

(G) Predicted protein structure of SPPL3 with its catalytic residues magnified.

(H) Histogram (cells from the RFP+ gate for transduced samples) and quantification (MFI normalized to the RFP- gate) of nonsaturating W6/32 stain on indicated HAP1 cells transduced with RFP-empty vector (EV), RFP-SPPL3, or catalytically inactive RFP-SPPL3 D271A (n = 5). See Figure S3C for B1.23.2 stain.

(I) IFN- $\gamma$  secretion in supernatant after coculture of the indicated T cells with RFP+ fluorescence-activated cell sorting (FACS)-sorted HAP1 *SPPL3*<sup>-/-</sup> cells transduced as in (H) or with unsorted WT or HLA-I-deficient (gray) cells (n = 2–3).

Mean  $\pm$  SD of n independent experiments is plotted in (D), (F), and (H). A representative of n experiments is shown in (B), (C), and (I). ns, not significant; \* p<0.05; \*\* p<0.01; \*\*\* p<0.001. See also Figure S3.



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et al., 2010) to HLA-I was strongly decreased on *SPPL3*<sup>-/-</sup> cells compared with WT cells, suggesting that various immune cell functions can be affected by SPPL3 (Figures 2D and 2F). Similarly, we investigated the NK cell receptors KIR2DL1 and KIR2DL2 by evaluating the binding of their recombinant Fc fusion proteins to overexpressed HLA-C\*05:01 on *HLA-A*<sup>-/-</sup>*B*<sup>-/-</sup>*C*<sup>-/-</sup> cells on a WT or *SPPL3*<sup>-/-</sup> background (Figure S3B) (Anfossi et al., 2006; Moesta et al., 2008). KIR2DL1 binding was not influenced by SPPL3 depletion, whereas KIR2DL2 binding was significantly reduced in the absence of SPPL3 (Figures 2E and 2F). Thus, the effect of SPPL3 on KIR and LIR binding to HLA-I is variable, indicating diverse, not further defined, effects on signaling toward different immune cell subsets.

### Accessibility to HLA-I Depends on SPPL3-Mediated Proteolytic Cleavage

SPPL3 contains two aspartate residues embedded in conserved YD and GxGD motifs located in transmembrane helices 6 and 7, respectively, forming the catalytic unit for proteolysis (Voss et al., 2013) (Figure 2G). To investigate whether SPPL3 catalytic activity was critical in controlling protein accessibility to HLA-I and function, we expressed WT SPPL3 or a catalytically inactive SPPL3 mutant (D271A) in *SPPL3*<sup>-/-</sup> cells (Voss et al., 2012). Flow cytometry analysis showed that only SPPL3 D271A failed to restore the accessibility of HLA-I for W6/32, indicating that SPPL3 proteolytic activity is required for antibody accessibility to HLA-I (Figures 2H and S3C). This lack of rescue was confirmed on a functional level, because expression of active SPPL3, but not inactive SPPL3, in *SPPL3*<sup>-/-</sup> cells partially restored their capacity to activate T cells (Figure 2I).

SPPL3 has previously been reported to affect protein N-glycosylation by proteolytic inactivation of glycosyltransferases in the ER and Golgi (Kuhn et al., 2015; Voss et al., 2014). Variations in the N-linked glycan of HLA-I, located at position N86 close to the SPPL3-susceptible region (Figure S3D), can affect its availability for proteins (Barbosa et al., 1987; Neefjes et al., 1990). However, liquid chromatography-mass spectrometry (LC-MS) (reverse-phase [RP] nano-LC-ESI-MS/(MS)) of HLA-I N-linked glycans revealed no differences between WT and *SPPL3*<sup>-/-</sup> cells (Figure S3E), indicating that the HLA-I N-glycan structure is not regulated by SPPL3 activity. To exclude a contribution of the HLA-I N-glycan to SPPL3-modulated HLA-I accessibility for several receptors and antibodies, we inhibited complex N-glycan formation on *SPPL3*<sup>-/-</sup> cells using the  $\alpha$ -mannosidase I and II inhibitors kifunensine and swainsonine. The decrease of complex N-glycans failed to alter the accessibility of the W6/32 epitope on *SPPL3*<sup>-/-</sup> cells (Figures S3F and S3G). This result was confirmed

in cells genetically engineered to lack complex (HLA-I) N-glycosylation through ablation of the gene encoding *GANAB* (Table S1), which resulted in lower overall HLA-I surface expression as visualized by decreased W6/32 and B1.23.2 signals (Figure S3H). Comparison of these antibody stainings between WT and *SPPL3*<sup>-/-</sup> cells showed that W6/32 accessibility to HLA-I was still impaired in the absence of SPPL3 (Figure S3H). As we ruled out a role for protein glycosylation, our finding that SPPL3 activity affects HLA-I at the cell surface suggests the involvement of at least one unknown SPPL3 target.

### SPPL3-Controlled GSLs Modulate Receptor Accessibility to HLA-I

To elucidate how SPPL3 controls protein accessibility to HLA-I, we followed two genome-wide screening strategies to specifically identify targets that are either positively or negatively regulated by SPPL3. An SPPL3-activated target affecting HLA-I would likely be a hit in the original W6/32 screen, just like SPPL3 (Figure 1B). However, the identification of such a target was complicated by the long list of significant hits. To distinguish SPPL3-activated targets from other candidates, we complemented the original screen with a new genome-wide haploid screen using a different HLA-I-specific antibody that was only mildly affected by the absence of SPPL3 (antibody BB7.2) (Figures 3A and S2A). This additional screen yielded another high-resolution snapshot of HLA-I antigen presentation (Figure S4A). A comparison of the two screens showed that SPPL3 was the only factor selectively affecting W6/32 binding, implying that no other gene was as strongly required for W6/32 accessibility to HLA-I (Figure 3B).

We then searched for potential genes negatively regulated by SPPL3 to affect HLA-I. To this end, we performed a genome-wide haploid screen in *SPPL3*<sup>-/-</sup> HAP1 cells. In these cells, which potentially lacked SPPL3-mediated suppression of the sought target, the gene trap mutagenesis of such a target or its associated pathway should improve W6/32 accessibility to HLA-I (Figure 3C). The hits from this screen converged to the GSL synthesis pathway (Figure 3D). The enzymes UGCG, B4GALT5, and B3GNT5 catalyze the synthesis of GSLs in the Golgi membrane by consecutive linkage of sugar residues derived from uridine diphosphate (UDP)-glucose, UDP-galactose, and UDP-N-acetylglucosamine (GlcNAc) donors on ceramide molecules (Figure 3E) (Allende and Proia, 2014). The latter two carbohydrate donors are transported from the cytoplasm into the Golgi by SLC35A2 and SLC35A3, respectively, which were also identified in the screen (Figure 3D) (Caffaro and Hirschberg, 2006). Other hits from the screen included proteins and

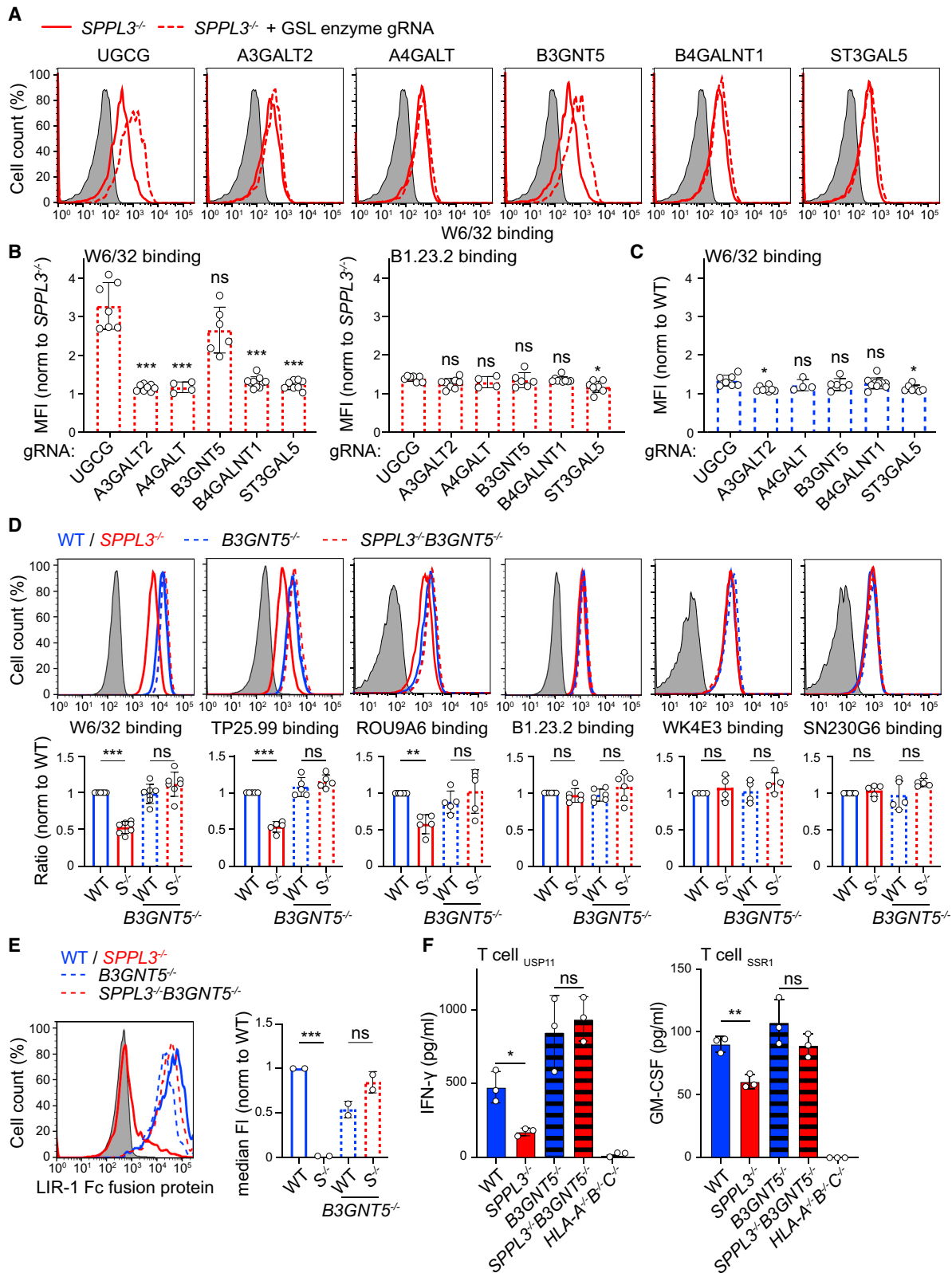
### Figure 3. SPPL3-Controlled GSLs Modulate Receptor Accessibility to HLA-I

(A–C) Schematic outlines of screening strategies to discover potential HLA-I regulators activated (A) or inactivated (C) by SPPL3. (B) Rocket plot of haploid genetic screens comparing the number of unique disruptive integrations per gene in BB7.2<sup>o</sup>- and W6/32<sup>o</sup>-sorted HAP1 populations. Two-sided FDR corrected Fisher's exact test was applied. The symbol legend is indicated (see also Figure S4A).

(D) Fish-tail plot of the haploid genetic screen in *SPPL3*<sup>-/-</sup> HAP1 cells, showing per gene the mutation index (ratio of integrations mapped in indicated populations) against the amount of integrations. Statistics and color legend as in (B) (see also Figure S4B).

(E) Schematic overview of the core enzymes involved in the synthesis of GSL subtypes. The putative SPPL3-targeted branch is shown in orange. PM, plasma membrane.

(F and G) Histograms of W6/32, B1.23.2 (F), and LIR-1 Fc (G) surface staining of the indicated HAP1 cells. Quantification (MFI, F; or median fluorescence intensity [median FI], G) is shown as mean  $\pm$  SD (n = 3 independent experiments). Gray, unstained controls; S<sup>-/-</sup>, *SPPL3*<sup>-/-</sup>. ns, not significant; \*\* p < 0.01; \*\*\* p < 0.001. See also Figure S4.



**Figure 4. B3GNT5 Function Determines HLA-I Accessibility for Its Natural Receptors**

(A) W6/32 surface staining of guide RNA (gRNA)-transduced (GFP<sup>+</sup>; gRNAs in Table S4) or control (GFP<sup>-</sup>)  $SPPL3^{-/-}$  HAP1 cells.

(B and C) Quantification showing normalized MFIs of the depicted antibody binding to  $SPPL3^{-/-}$  (B) or WT (C) cells combined data from two experiments with 2–4 gRNAs per gene (examples in A) (Figures S4C and S4D).

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complexes associated with Golgi homeostasis, such as the members of the component of oligomeric Golgi (COG) and Golgi-associated retrograde protein (GARP) complexes that facilitate GSL synthesis and trafficking (Fröhlich et al., 2015; Kingsley et al., 1986) (Figure 3D). None of these hits related to GSL metabolism emerged in the original screen with W6/32 on SPPL3-containing WT cells, strongly suggesting that in WT cells, the GSL synthesis or biosynthetic transport pathway is suppressed by SPPL3 (Figure S4B). Collectively, these observations revealed the existence of a pathway comprising GSL-mediated regulation of HLA-I access and function controlled by SPPL3.

To validate that SPPL3 reduces protein accessibility to HLA-I through manipulation of GSL synthesis, we generated GSL-deficient *SPPL3*<sup>-/-</sup> cells by additional genetic ablation of the first enzyme of the GSL synthesis pathway, UGCG (Table S1). In these *SPPL3*<sup>-/-</sup>*UGCG*<sup>-/-</sup> cells, we observed full rescue of the W6/32 HLA-I epitope accessibility without affecting SPPL3-independent B1.23.2 staining (Figure 3F), pointing toward an essential role for GSLs in shielding specific HLA-I epitopes. Importantly, GSL-deficient *SPPL3*<sup>-/-</sup> cells regained the capacity to engage the HLA-I binding receptor LIR-1 (Figure 3G), underscoring the physiological relevance of GSL-mediated epitope shielding of HLA-I.

### B3GNT5 Tunes the Capacity of HLA-I to Interact with Its Natural Receptors

The synthesis of GSLs is probably best illustrated as a chain of sugar moiety transfers catalyzed by different Golgi enzymes (Figure 3E). UGCG initiates the GSL synthesis pathway by transferring a glucose to a ceramide on the cytosolic leaflet of the Golgi membrane (Allende and Proia, 2014). After this glucosylceramide is flipped into the Golgi lumen, a galactose moiety is added by B4GALT5 or B4GALT6 to generate lactosylceramide (LacCer). This neutral GSL then serves as a substrate for various glycosyltransferases responsible for the generation of different GSL series: A4GALT (globo-series), A3GALT2 (isoglobo-series), B3GNT5 (lacto-series and nsGSLs), B4GALNT1 (gangliosides, o series), and ST3GAL5 (gangliosides, a, b, and c series) (Figure 3E) (Allende and Proia, 2014; Zhang et al., 2019). Our screening data suggested that lacto-series GSL or nsGSL production, through B3GNT5 activity, could diminish protein accessibility to HLA-I (Figures 3D and S4A). To confirm this specificity, we generated polyclonal cell lines on the *SPPL3*<sup>-/-</sup> background, each CRISPR-Cas9 targeted for one of the five branching enzymes, and then analyzed W6/32 binding by flow cytometry. W6/32 accessibility to HLA-I in *SPPL3*<sup>-/-</sup> cells was restored only by ablation of B3GNT5 or control UGCG, confirming B3GNT5 as the sole branching enzyme involved in HLA-I epitope shielding (Figures 4A, 4B, and S4C). This effect was selective for SPPL3-deficient cells, because HLA-I accessibility for W6/32 was unaffected on WT cells with corresponding genetic ablations (Figures 4C and S4D). Single-cell-derived *B3GNT5*<sup>-/-</sup>

and *SPPL3*<sup>-/-</sup>*B3GNT5*<sup>-/-</sup> cell lines were generated to corroborate a pivotal role for B3GNT5 in HLA-I epitope shielding (Table S1). As expected, additional genetic B3GNT5 depletion in *SPPL3*<sup>-/-</sup> cells restored not only W6/32 binding to its epitope but also accessibility of other SPPL3-susceptible epitopes recognized by TP25.99 and ROU9A6 (Figure 4D). Accessibility to SPPL3-independent epitopes and total HLA-I surface expression were not affected by additional B3GNT5 depletion (Figure 4D). Moreover, the lack of B3GNT5 expression in *SPPL3*<sup>-/-</sup> cells restored both binding of LIR-1 to HLA-I and their potential to activate T cells (Figures 4E and 4F). Altogether, our results suggest that active SPPL3 disrupts the B3GNT5 protein, which tunes the capacity of HLA-I to interact with its receptors.

### SPPL3 Controls the Generation of nsGSLs by Proteolytically Inactivating B3GNT5

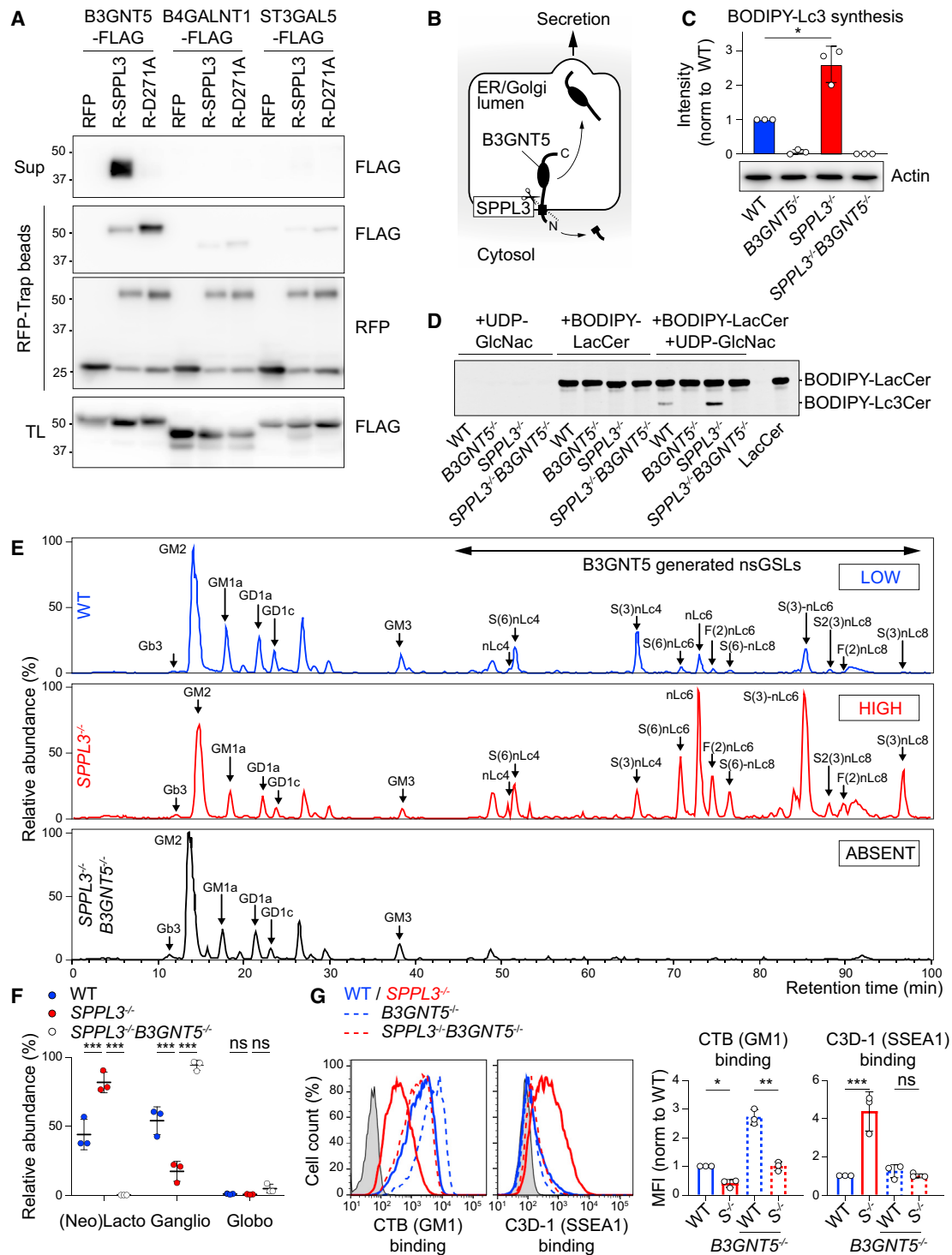
To detect a direct interaction between SPPL3 and its putative target B3GNT5, we performed coimmunoprecipitation of over-expressed epitope-tagged proteins. We coisolated B3GNT5 predominantly with the catalytically inactive SPPL3 D271A mutant, suggesting a transient interaction between SPPL3 and its substrate (Figure 5A). Cleavage of B3GNT5 by the intramembrane protease SPPL3 was confirmed in total lysate by a small reduction in the molecular weight of B3GNT5, reflecting proteolytic removal of the 1.5- to 4-kDa cytosolic tail, and by the presence of luminal B3GNT5 fragments in the supernatant (Figures 5A and 5B). Two other branching enzymes of the GSL synthesis pathway, B4GALNT1 and ST3GAL5, were poorly coisolated with SPPL3 (Figure 5A). Furthermore, cleavage products were not detected in the supernatant, indicating that B3GNT5 is a specific substrate of SPPL3 (Figure 5A).

To investigate whether SPPL3 affects B3GNT5 activity, we performed a B3GNT5 enzymatic assay. Lysates of indicated WT and genetically ablated cells were incubated with a BODIPY-conjugated analog of the B3GNT5 substrate LacCer and the donor sugar UDP-N-GlcNAc, followed by thin-layer chromatography (TLC) of extracted GSLs. The B3GNT5 product lactotriaosylceramide (BODIPY-Lc3Cer), as confirmed by LC-MS, was generated in increased amounts in *SPPL3*<sup>-/-</sup> compared with WT cell lysates (Figures 5C, 5D, and S5A). In addition, no Lc3Cer was synthesized in lysates of *B3GNT5*<sup>-/-</sup> cells, demonstrating that B3GNT5 is the sole producer of Lc3Cer in HAP1 cells. Because SPPL3 inhibits B3GNT5 activity, we next addressed the extent to which SPPL3 defines the cellular GSL profile. Glycan portions of the GSL repertoire of WT, *SPPL3*<sup>-/-</sup>, and *SPPL3*<sup>-/-</sup>*B3GNT5*<sup>-/-</sup> cells were isolated and analyzed by LC-MS. We found an extensive shift in the relative GSL abundance toward B3GNT5-produced nsGSLs, from 44% in WT cells to 82% in *SPPL3*<sup>-/-</sup> cells (Figures 5E and 5F; Table S3). The increase was most evident for complex nsGSLs containing six or more sugar residues, as determined by relative quantification of individual GSLs, suggesting that epitope shielding of HLA-I is mediated by complex nsGSLs (Figure S5B; Table

(D and E) Histograms and quantifications (MFI, D, or median fluorescence intensity, E, median FI) of surface staining of the indicated HAP1 cells using antibodies recognizing SPPL3-susceptible (W6/32, TP25.99, and ROU9A6) or SPPL3-independent (B1.23.2, WK4E3, and SN230G6) HLA-I epitopes (D; n = 4–7) or using LIR-1 Fc fusion protein (E; n = 2). *S*<sup>-/-</sup>, *SPPL3*<sup>-/-</sup>.

(F) IFN- $\gamma$  or GM-CSF secretion by depicted T cells after coculture with the indicated HAP1 cells. Representative of n = 3.

Gray histograms are unstained controls. Mean  $\pm$  SD of n independent experiments is plotted in (B)–(E). ns, not significant; \* p < 0.05; \*\* p < 0.01; \*\*\* p < 0.001. See also Figure S4.



**Figure 5. SPPL3 Controls the Generation of nsGSLs by Targeting B3GNT5**

(A) Coimmunoprecipitation of the indicated FLAG-tagged proteins with RFP-EV (RFP), RFP-SPPL3 (R-SPPL3), or RFP-SPPL3 D271A (R-D271A). Sup, supernatant; TL, total lysate. Representative of  $n = 2$ .

(B) Schematic model of B3GNT5 proteolysis by SPPL3.

(C and D) TLC of the indicated HAP1 cell lysates incubated with UDP-GlcNac and boron-dipyrromethene (BODIPY)-LacCer substrate to detect B3GNT5 activity ( $n = 3$ ). BODIPY-Lc3Cer quantification (see Figure S5A for LC-MS validation) (C) and an example chromatogram (D) are shown.

(E) Base peak chromatograms of porous graphitized carbon (PGC) LC-MS on total GSL glycan isolated from indicated HAP1 cells ( $n = 3$ ). Proposed glycan structures and their relative abundance are listed in (F), Table S3 and Figure S5B.

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S3). To validate the shift in GSL repertoire in living cells, we conducted flow cytometry-based experiments using cholera toxin subunit B, which binds the ganglioside GM1, and an antibody against the nsGSL SSEA-1 epitope. GSL-deficient *UGCG*<sup>-/-</sup> cells were negative for all probes, demonstrating probe specificity toward GSLs on our cells (Figure S5C). Compared with WT cells, *SPPL3*<sup>-/-</sup> cells expressed increased amounts of SSEA-1 nsGSLs, which were generated by B3GNT5, and decreased amounts of GM1 gangliosides (Figures 5G and S5C). Consistent with our relative quantification of individual glycans detected by LC-MS, these data demonstrate that SPPL3 dictates the composition of the GSL repertoire by inhibiting the nsGSL biosynthesis activity of B3GNT5.

### Sialic Acid Residues on nsGSL Acid Residues for HLA-I Shielding

GSLs are major constituents of membrane microdomains (Sezgin et al., 2017). A change in GSL composition may then disturb membrane protein localization, mobility, and function. We therefore investigated the mobility of HLA-I in *SPPL3*<sup>-/-</sup> cells by single-particle tracking. The mobile fraction and diffusion constant of BB7.2 Fab labeled HLA-I molecules were equal between *SPPL3*<sup>-/-</sup> and WT cells, indicating that HLA-I membrane dynamics were not detectably affected by potential alterations in membrane microdomain organization or by SPPL3 (Figure S6). This renders unlikely a scenario in which HLA-I associates with another protein in the absence of SPPL3, because this would reduce the HLA-I diffusion rate. Instead, our data suggested that decreased receptor accessibility to HLA-I was a direct consequence of interactions with nsGSLs. Such GSL-protein interactions can occur between gangliosides and hormone receptors through a charge-based linkage of GSL-derived sialic acid with positively charged amino acids (D'Angelo et al., 2013). Further analyses of the GSL signature of *SPPL3*<sup>-/-</sup> compared with WT cells revealed that the nsGSL glycan chains more frequently contain  $\alpha$ -2,3- and  $\alpha$ -2,6-linked sialic acid residues, as well as noncharged fucoses (Figures 5D and 6A; Table S3). To test the requirement of these nsGSL-localized sugar residues, we inhibited all sialyltransferase and fucosyltransferase activity and found that dose-dependent inhibition of sialylation, but not fucosylation, restored W6/32 accessibility to HLA-I in *SPPL3*<sup>-/-</sup> cells (Figures 6B–6E). The requirement for sialic acids was substantiated by the genetic ablation of CMP-sialic acid synthetase (CMAS), which also recovered W6/32 accessibility to HLA-I in *SPPL3*<sup>-/-</sup> cells (Figures 6B and 6F). Finally, the enzymatic removal of sialic acid residues at the cell surface by neuraminidase treatment diminished HLA-I shielding (Figures 6B and 6G). Thus, the B3GNT5-generated GSLs shield HLA-I through its sialic acids, likely via a direct charge-based interaction.

### Pharmacological Inhibition of GSL Synthesis in Glioma Enhances Anti-tumor Immune Activation In Vitro

After determining that nsGSL-rich target cells suppress T cell activity, we examined the effect of increased nsGSL expression or

downmodulated SPPL3 activity in tumors. Because of the complexity of identifying (large) nsGSLs, there is a limited amount of data available on their tissue expression, including tumors (Merrill and Sullards, 2017; Zhang et al., 2019). Nonetheless, elevated amounts of nsGSLs or their synthesis enzyme B3GNT5 have been observed on several tumor types, including glioma, acute myeloid leukemia (AML), and adenocarcinomas (Furukawa et al., 2015; Hakomori, 1984; Wang et al., 2012; Wikstrand et al., 1991). In addition, The Cancer Genome Atlas (TCGA) analyses demonstrated that high B3GNT5 expression in low-grade glioma correlates with decreased overall patient survival (Figure 7A). In line with our findings, the reverse held true for the B3GNT5-suppressing SPPL3 (Figure 7B). Moreover, analyses of the combined effect of B3GNT5 and SPPL3 expression showed only lower survival rates for patients with high B3GNT5 and low SPPL3 expression (Figures 7C and S7A), probably reflecting that nsGSL expression is elevated in tumors from this group. Because this indicated that gliomas may limit immune detection by exploiting the SPPL3-B3GNT5 axis, we tested the role of SPPL3 and nsGSLs in the glioblastoma cell line U-251. Overexpression of SPPL3 increased HLA-I accessibility to W6/32 without altering HLA-I expression (Figure 7D). Next, genetic depletion of GSLs (including nsGSLs) from U-251 cells resulted in a specific increase in W6/32 binding to HLA-I (Figures 7E and S7B). Moreover, in the absence of GSLs, U-251 cells were better activators of T cells (Figure 7F).

To downregulate nsGSL expression in patients, the clinically approved GSL synthesis inhibitors miglustat and eliglustat may be used (Stirnemann et al., 2017). These drugs are being used as substrate reduction therapy in Gaucher disease. We first explored whether these small-molecule drugs affect accessibility to HLA-I epitopes that are shielded in *SPPL3*<sup>-/-</sup> cells. The miglustat mimics MZ21 and MZ31, with fewer off-target effects, were also included (Ghisaidoobe et al., 2014). All GSL synthesis inhibitors fully restored W6/32 accessibility to HLA-I, despite a small proportion of GSLs remaining detectable on the cell surface (Figures 7G and S7C–S7F). Moreover, these inhibitors increased the capacity of *SPPL3*<sup>-/-</sup> cells to activate T cells (Figures 7H, 7I, and S7G). This was also observed for U-251 cells, for which HLA-I shielding was alleviated and their capacity to activate T cells was increased (Figures 7J, 7K, and S7H). Altogether, these data demonstrate that these inhibitors can boost immune responses against tumor cells that display excess nsGSLs.

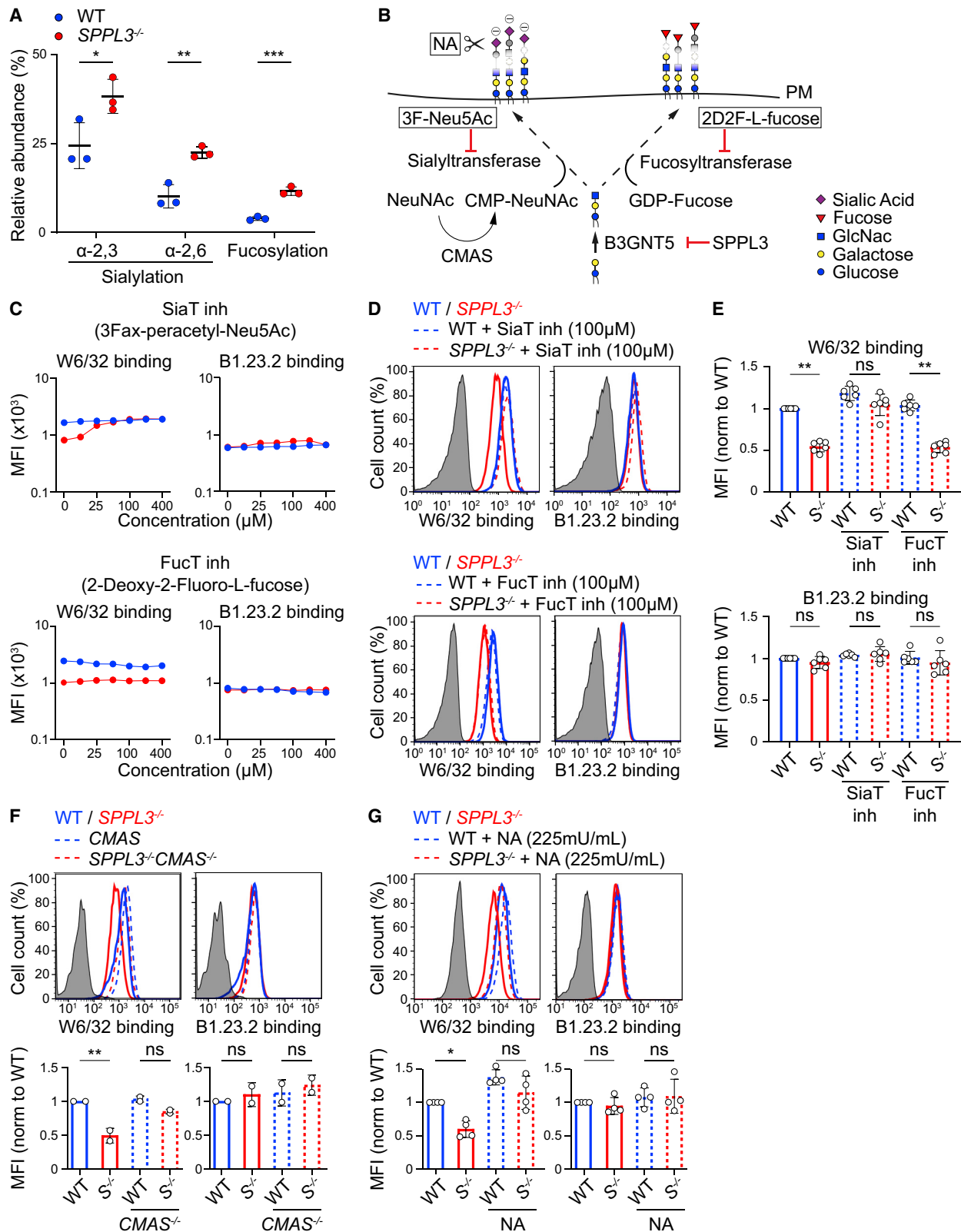
## DISCUSSION

The process of HLA-I antigen presentation has been a topic of long-standing interest, giving rise to a detailed understanding of this complex pathway. We here describe an additional element for the equation of successful antigen presentation, namely, the SPPL3-B3GNT5 pathway responsible for the production of a subset of GSLs. GSLs are present on every cell, yet their functional roles in the cell membrane remain largely

(F) Quantified relative abundance of the three major GSL types of the indicated cells.

(G) Histograms and normalized MFI of cholera toxin B (CTB) and C3D-1 binding to indicated HAP1 cells (n = 3). Gray, unstained control; *S*<sup>-/-</sup>, *SPPL3*<sup>-/-</sup> (see Figure S5C).

Mean  $\pm$  SD of n independent experiments is plotted in (C), (F), and (G). ns, not significant; \* p < 0.05; \*\* p < 0.01; \*\*\* p < 0.001. See also Figure S5.



**Figure 6. Sialic Acid Residues on nsGSLs Are Required for HLA-I Shielding**

(A) Quantified relative abundance of sialylated and fucosylated nsGSLs in WT or *SPPL3*<sup>-/-</sup> HAP1 cells. Data from Figure 5E were reused.

(B) Schematic view of targetable steps in nsGSL sialylation and fucosylation. NA, neuraminidase.

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unknown. By iteratively conducting sensitive genome-wide screens, we uncovered a role for a subset of GSLs in immunity controlled by the aspartyl protease SPPL3. These nsGSLs shield HLA-I molecules, limiting their interactions with several immune cell receptors and decreasing CD8<sup>+</sup> T cell responses. We identified SPPL3 as a switch controlling the expression of nsGSLs through proteolytic inhibition of the nsGSL-synthesizing enzyme B3GNT5. Altogether, our study reveals a layer of immune regulation that acts late in the HLA-I antigen presentation pathway by shielding critical HLA-I epitopes at the cell surface.

Understanding nsGSL function at a molecular level and in physiological and pathophysiological settings is challenging given that their isolation, their analytical dissection, and in particular their experimental manipulation have been extraordinarily demanding to date (Merrill and Sullards, 2017; Zhang et al., 2019). Hence, no validated methods are available to study nsGSL-protein interactions, restricting options to directly probe the nsGSL-HLA-I interaction. Our current data indicate that the interaction between nsGSLs and HLA-I molecules must be transient, because a high antibody dose can overcome limited accessibility to HLA-I. In addition, we show that this interplay is independent of carbohydrate-carbohydrate interactions (D'Angelo et al., 2013) between nsGSLs and HLA-I N-glycans. However, the profound shielding of large HLA-I patches by nsGSLs can be explained by the ability of nsGSLs, in contrast to other GSL subtypes, to carry huge glycan chains of up to 60 sugar residues (Miller-Podraza et al., 1993, 1997). These long carbohydrate trees may reach HLA-I domains involved in the interaction with receptors such as LIR-1. The nsGSLs may sterically compete with proteins for accessibility to HLA-I or restrict protein accessibility by altering HLA-I orientation toward the cellular membrane (Mitra et al., 2004). In addition, our data do not exclude a direct interaction between the GSL ceramide and the HLA-I transmembrane domain (Contreras et al., 2012), which could contribute to the positioning the nsGSL glycan chain close to HLA-I. Finally, we show that sialic acid residues on GSLs are essential for HLA-I shielding. The negatively charged sialylated nsGSLs may establish ionic interactions with HLA-I, which has abundant positively charged patches at its molecular surface (Li et al., 2012). Similar GSL-protein interactions have been found between sialic acids on short GSL glycans and exposed positively charged amino acid residues close to the plasma membrane (D'Angelo et al., 2013). This shows that sialylated nsGSLs may shield cell surface receptors other than HLA-I and possibly affect their cognate interactions. This assumption is supported by SPPL3 being identified in genome-wide CRISPR-Cas9 screens for surface detection of butyrophilin (BTN) molecules by a functional V $\gamma$ 9V $\delta$ 2<sup>+</sup>  $\gamma\delta$  TCR or CD47 and CD59 by antibodies (Davis et al., 2015; Logtenberg et al., 2019; Rigau et al., 2020). In these cases, the underlying molecular mechanism of the SPPL3 ef-

fect was not resolved, yet these cases suggest that the SPPL3-B3GNT5 pathway constitutes a mechanism to fine-tune communication between cells, including a functional tumor cell-T cell interaction, as we report here.

Various malignant cells exhibit alterations in their GSL surface repertoire, to which several specific functions have been attributed. Some GSLs can serve as signaling molecules to control cellular processes such as apoptosis and proliferation, whereas other GSL species can confer anti-cancer drug resistance by inhibiting proteins that facilitate their membrane transport (Liu et al., 2013; Ogretmen and Hannun, 2004). We here propose that changes in the tumor GSL repertoire, in particular increments of sialic-acid-containing nsGSLs, limit HLA-I signaling to T cells as a means to evade immune surveillance. In support of this hypothesis, our *in vitro* data show that GSLs diminish the capacity of CD8<sup>+</sup> T cells to respond to glioma, a tumor type with high expression of nsGSLs (Furukawa et al., 2015). Furthermore, HLA-I-related NK cell activation against tumors lacking SPPL3 may be restricted according to recent genome-wide CRISPR-Cas9 screens (Pech et al., 2019). Thus, nsGSL upregulation by tumors such as glioma might allow T cell escape while marginalizing NK cell recognition. In addition to *in vitro* experimentation, analyses involving glioma patients revealed worst overall survival when the SPPL3 and B3GNT5 expression signature of the tumor suggests high nsGSL synthesis. Such correlation with patient outcome may have been influenced by covariates, which potentially include nsGSL-mediated shielding of other immune or nonimmune receptors or membrane turnover (Catalaa et al., 2006; Righi et al., 2009). Other tumor types, including AML, colorectal carcinoma, adenocarcinoma, and ductal carcinoma *in situ* (DCIS), also overexpress B3GNT5 and its product nsGSLs (Hakomori, 1984; Potapenko et al., 2015; Wang et al., 2012; Wikstrand et al., 1991), suggesting that nsGSL overexpression is a general strategy for tumor survival. Furthermore, pathogens such as cytomegalovirus, respiratory syncytial virus, and HIV alter the GSL composition of the host cell, potentially inducing immune evasion through HLA-I shielding (Fantini et al., 2000; Moore et al., 2008; Radsak and Wiegandt, 1984). Except for low-resolution data concerning cytomegalovirus-induced nsGSL expression upon infection (Andrews et al., 1989; Radsak and Wiegandt, 1984), little is known about which viral infections influence complex nsGSL expression.

In this study, we have presented GSLs as highly relevant molecules affecting the efficiency of immune responses. nsGSLs and their molecular switch SPPL3 represent an unexplored avenue for therapeutic intervention in cancer, infection, and autoimmune diseases. Two small-molecule drugs inhibiting GSL synthesis are registered: miglustat (Zavesca) and eliglustat (Cerdelga). These structurally different UGCG inhibitors (Platt et al., 1994; Shayman, 2010) have been approved for the treatment of patients with lysosomal storage disorders, such as

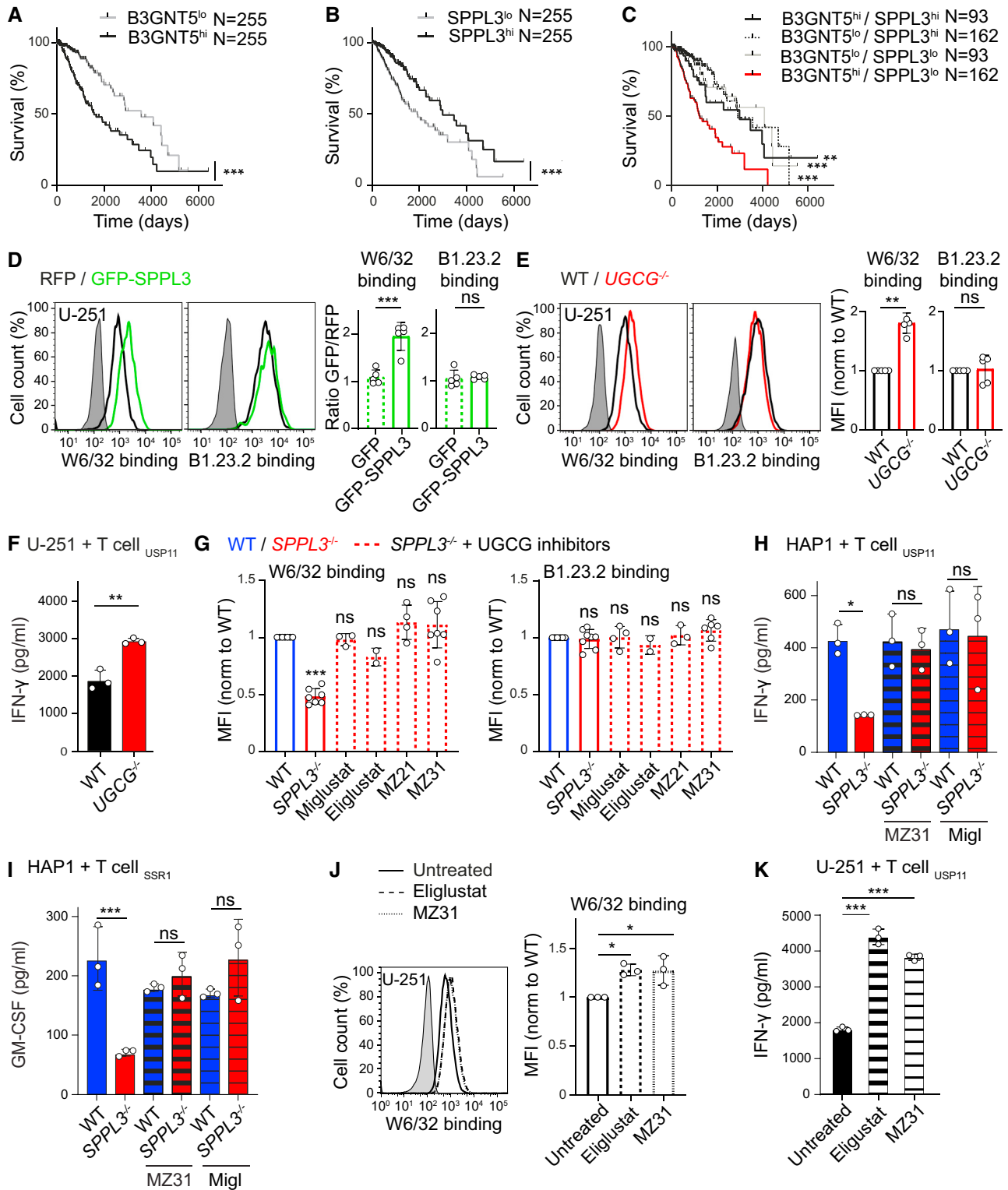
(C) Antibody staining of the indicated HAP1 cells cultured with a serial dilution of sialyltransferase (SiaT) or fucosyltransferase (FucT) inhibitors.

(D and E) Histograms (D) and normalized MFI (E) of the depicted antibody binding to HAP1 cells precultured with 100  $\mu$ M inhibitors as in (C) (n = 6).

(F) Histograms and quantification of the depicted antibody staining of the indicated HAP1 cells (n = 2).

(G) Histograms and quantification of W6/32 and B1.23.2 binding to NA-treated HAP1 cells (n = 4).

Gray histograms are unstained controls. S<sup>-/-</sup>, SPPL3<sup>-/-</sup>. Mean  $\pm$  SD of n independent experiments is plotted in (A) and (E)-(G). ns, not significant; \* p<0.05; \*\* p<0.01; \*\*\* p<0.001.



**Figure 7. Pharmacological Inhibition of GSL Synthesis in Glioma Enhances Anti-tumor Immune Responses**

(A–C) TCGA-derived Kaplan-Meier curves showing the survival of patients with tumors expressing low or high B3GNT5 (A), low or high SPPL3 (B), or any of the four combinations thereof (C) (see Figure S7A).

(D) Histograms of W6/32 and B1.23.2 binding to U-251 glioblastoma cells overexpressing GFP-SPPL3 or RFP-EV, combined in a single well. Quantification (MFI GFP+ cells/MFI RFP+ cells) includes a GFP-EV control (n = 5).

(E) Histograms and normalized MFI of the depicted antibody staining of WT and UGCG<sup>-/-</sup> U-251 cells (n = 4–5) (see Figure S7B).

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type I Gaucher disease and Niemann-Pick disease type C (Lachmann, 2003; Wraith and Imrie, 2009). Therefore, therapeutic application can be extended efficiently to include immune enhancement against tumors or pathogen-infected cells. GSL synthesis inhibition may even be combined successfully with existing immunotherapies, such as PD-1 blockade, because of the potential synergy between enhanced tumor cell immunogenicity and simultaneous T cell activation.

## STAR★METHODS

Detailed methods are provided in the online version of this paper and include the following:

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- QUANTIFICATION AND STATISTICAL ANALYSIS

## SUPPLEMENTAL INFORMATION

Supplemental Information can be found online at <https://doi.org/10.1016/j.immuni.2020.11.003>.

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## AUTHOR CONTRIBUTIONS

Conceptualization and design, M.L.M.J. and R.M.S. Data acquisition, analysis, and interpretation, M.L.M.J., M.R., A.A.d.W., T.Z., B.C., R.Plutzer, V.A.B., A.X., T.V., S.B., X.K., C.G., L.J., E.S., S.H., R.Plomp, and R.M.S. Resources and discussion, A.M., S.F., F.H.J.C., M.H.M.H., M.G., A.H., and H.O. Supervision and conceptual discussion, J.B.H., M.W., T.R.B., J.N., and R.M.S. Writing, M.L.M.J. and R.M.S. Editing, J.B.H., M.R., A.A.d.W., T.Z., M.W., and J.N.

## DECLARATION OF INTERESTS

T.R.B. is a cofounder and SAB member of Haplogen GmbH and a cofounder and director of Scenic Biotech BV.

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(F) IFN- $\gamma$  secretion by the indicated T cells against WT or UGCG<sup>-/-</sup> U-251 cells (n = 3).

(G) Normalized MFI of the depicted antibody staining of the indicated HAP1 cells precultured with specified UGCG inhibitors (n = 2–7) (see Figures S7C–S7F).

(H and I) IFN- $\gamma$  or GM-CSF secretion by the indicated T cells cocultured with WT or SPPL3<sup>-/-</sup> HAP1 cells that were precultured with or without the specified UGCG inhibitor (n = 3) (see Figure S7G for more T cell clones).

(J) Histogram and normalized MFI of W6/32 binding to WT U-251 cells precultured with the indicated UGCG inhibitors (n = 3) (see Figure S7H).

(K) IFN- $\gamma$  secretion by the indicated T cells after coculture with the depicted inhibitor-pretreated U-251 cells (n = 3).

Gray histograms are unstained controls. Mean  $\pm$  SD of n independent experiments is plotted in (D), (E), (G), and (J). A representative of n experiments is shown in (F), (H), (I), and (K). ns, not significant; \* p<0.05; \*\* p<0.01; \*\*\* p<0.001. See also Figure S7.

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STAR★METHODS

KEY RESOURCES TABLE

REAGENT or RESOURCE	SOURCE	IDENTIFIER
<b>Antibodies</b>		
Mouse monoclonal anti-HLA-ABC-PerCP-eFluor710 (W6/32)	Thermo Fisher Scientific	Cat# 46-9983-42; RRID:AB_10804486
Mouse monoclonal anti-HLA-BC-APC (B1.23.2)	Thermo Fisher Scientific	Cat# 17-5935-42; RRID:AB_11151509
Mouse monoclonal anti-HLA-A2-APC (BB7.2)	Thermo Fisher Scientific	Cat# 17-9876-42; RRID:AB_11149299
Mouse monoclonal anti-CD15-FITC (C3D-1)	Millipore	Cat# FCMAB182F; RRID:AB_11214339
Goat anti-mouse IgG (H+L) - Alexa Fluor 647	Thermo Fisher Scientific	Cat# A28181; RRID:AB_2536165
Mouse APC anti-human IgM (MHM-88)	Biologend	Cat# 314510; RRID:AB_493011
Mouse monoclonal anti-human IgG (MH161-1)	Sanquin home made	Hybridoma; RRID:AB_1286031
Rabbit polyclonal anti-RFP	Netherlands Cancer Institute home made	<a href="#">Rocha et al., 2009</a>
Mouse monoclonal anti-HLA-ABC (W6/32)	Dr. J. Neefjes (NKI, the Netherlands)	Hybridoma; RRID:AB_964524
Mouse monoclonal anti-HLA-BC (HC10)	Dr. J. Neefjes (NKI, the Netherlands)	Hybridoma; RRID:AB_2728622
Mouse monoclonal anti-HLA-A (HCA2)	Dr. J. Neefjes (NKI, the Netherlands)	Hybridoma
Mouse monoclonal anti-HLA-A2 (BB7.2)	Dr. M. Heemskerk (LUMC, the Netherlands)	Hybridoma; RRID:AB_627935
Mouse monoclonal anti-FLAG (M2)	Sigma-Aldrich	Cat# F3165; RRID:AB_259529
Mouse monoclonal anti-b-actin (AC-15)	Sigma-Aldrich	Cat# A5441; RRID:AB_476744
Mouse monoclonal anti-NGFR(CD271)-PE/Cy7 (ME20.4)	Biologend	Cat# 345109; RRID:AB_11204073
Goat anti-rabbit IgG (H+L)-HRP	Thermo Fisher Scientific	Cat# G-21234; RRID:AB_2536530
Mouse monoclonal anti-FLAG M2-HRP antibody	Sigma-Aldrich	Cat# A8592; RRID:AB_439702
APC AffiniPure F(ab') <sub>2</sub> Fragment Goat Anti-Human IgG, Fcγ fragment specific	Jackson ImmunoResearch	Cat# 109-136-098; RRID:AB_2337693
HLA-I/B2M specific antibodies, see <a href="#">Table S3</a>		N/A
<b>Chemicals, Peptides, and Recombinant Proteins</b>		
LIR-1 Fc fusion protein (human IgG1)	Dr. O. Mandelboim (Hebrew University Hadassah Medical School, Israel)	<a href="#">Gonen-Gross et al., 2010</a>
Recombinant human KIR2DL1/CD158a Fc chimera protein	R&D systems	Cat# 1844-KR
Recombinant human KIR2DL2/CD158b1 Fc chimera protein	R&D systems	Cat# 3015-KR
Alexa Fluor® 555 NHS Ester	Thermo Fisher Scientific	Cat# A20009
Alexa Fluor® 647 NHS Ester	Thermo Fisher Scientific	Cat# A37573
CFSE	Invitrogen	Cat# C1157
Alexa Fluor 350 NHS Ester	Thermo Fisher Scientific	Cat# A10168
Violet proliferation dye 450	BD Horizon	Cat# 562158
DyLight 650 NHS ester	Thermo Fisher Scientific	Cat# 62265
Cholera Toxin B Subunit from Vibrio cholerae-FITC	Sigma	Cat# C1655; CAS: 131096-89-4
Animal-Free Recombinant Human IFN-γ	Peprtech	Cat# AF-300-02
Sialyltransferase inhibitor (3Fax-peracetyl Neu5Ac)	Sigma	Cat# 566224; CAS: 117405-58-0

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REAGENT or RESOURCE	SOURCE	IDENTIFIER
Fucosyltransferase inhibitor (2-Deoxy-2-fluoro-L-fucose)	Carbosynth	MD06089; CAS: 70763-62-1
Neuraminidase from <i>Clostridium perfringens</i>	Sigma-Aldrich	N2876; CAS: 9001-67-6
Endoglycoceramidase I	New England Biolabs	Cat# P0773S
Miglustat	Dr. H. Overkleef (Leiden University, the Netherlands)	<a href="#">Ghisaidoobe et al., 2014</a>
Eliglustat	Bio-Connect	Cat# HY-14885A
MZ21	Dr. H. Overkleef (Leiden University, the Netherlands)	<a href="#">Ghisaidoobe et al., 2014</a>
MZ31	Dr. H. Overkleef (Leiden University, the Netherlands)	<a href="#">Ghisaidoobe et al., 2014</a>
Kifunensine (mannosidase inhibitor)	Santa Cruz	Cat# sc-201364; CAS: 109944-15-2
Swainsonine	Sigma-Aldrich	Cat# S8195; CAS: 72741-87-8
Brefeldin A	Sigma-Aldrich	Cat#B7651; CAS: 20350-15-6
BODIPY FL C5-Lactosylceramide complexed to BSA	Thermo Fisher Scientific	Cat# B34402
UDP-N-acetyl-D-glucosamine disodium salt	Santa Cruz	Cat# sc-286851
Endoglycoceramidase I (derived from <i>Rhodococcus triatomea</i> and expressed in <i>Escherichia coli</i> )	New England Biolabs	Cat# P0773S
Endoglycosidase H from <i>Streptomyces plicatus</i>	Sigma-Aldrich	Cat# A0810; CAS: 37278-88-9
Critical Commercial Assays		
IFN- $\gamma$ ELISA Kit	Sanquin	Cat# M9333
GMCSF-ELISA Kit	Biologend	Cat# 432002
Deposited Data		
Raw sequence data	NCBI Sequence Read Archive	PRJNA665349 (SAMN16252402)
Experimental Models: Cell Lines		
Human: HAP1	Dr. T. Brummelkamp (NKI, the Netherlands)	
Human: MeJuSo	Dr. J. Neefjes (LUMC, the Netherlands)	Authenticated: 19 ZE 000486 (2019)
Human: U-251	Laboratory of Dr. H. Versteeg (LUMC, the Netherlands)	N/A
Human: SW620	Dr. T. de Gruijl (Amsterdam UMC, the Netherlands)	N/A
Human: HEK293T	ATCC	Cat# CRL-3216; RRID:CVCL_0063
Human: Phoenix Ampho	Dr. J. Neefjes (LUMC, the Netherlands)	N/A
Murine: BB7.2 cell line (hybridoma)	Dr. M. Heemskerk (LUMC, the Netherlands)	N/A
Murine: W6/32 cell line (hybridoma)	Dr. J. Neefjes (LUMC, the Netherlands)	N/A
Human: CD8+ T cell clone reactive to USP11 peptide	Dr. M. Heemskerk (LUMC, the Netherlands)	<a href="#">Amir et al., 2011</a>
Human: CD8+ T cell clone reactive to FDPS peptide	Dr. M. Heemskerk (LUMC, the Netherlands)	<a href="#">Amir et al., 2011</a>
Human: CD8+ T cell clone reactive to VPS13B peptide	Dr. M. Heemskerk (LUMC, the Netherlands)	<a href="#">Amir et al., 2011</a>
Human: CD8+ T cell clone reactive to SSR1 peptide	Dr. M. Griffioen (LUMC, the Netherlands)	<a href="#">Van Bergen et al., 2010</a>
Human: CD8+ T cell clone reactive to ADIR peptide	Dr. M. Griffioen (LUMC, the Netherlands)	<a href="#">van Bergen et al., 2007</a>

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REAGENT or RESOURCE	SOURCE	IDENTIFIER
<b>Continued</b>		
<b>Oligonucleotides</b>		
siGENOME Human SPPL3 siRNA	Dharmacon	D-006042-02,03,04
siGENOME Human B2M siRNA (SMARTpool)	Dharmacon	M-004366-00
siGENOME Non-Targeting siRNA Pool#1	Dharmacon	D-001206-13
gRNA sequences used for CRISPR/Cas9 mediated genome editing, see <a href="#">Table S4</a>	This paper	N/A
PCR and sequencing primers for KO validation, see <a href="#">Table S5</a> .	This paper	N/A
<b>Recombinant DNA</b>		
pX330	Addgene	Plasmid #42230
TIA-2A-blast	Dr. T. Brummelkamp (NK1, the Netherlands)	<a href="#">Blomen et al., 2015</a>
LentiCRISPR_v2	Addgene	Plasmid #52961
pL-CRISPR.EFS.GFP	Addgene	Plasmid #57818
pMXs-puro retroviral expression vector	Cell Biolabs	Cat# RTV-012
pMXs-puro-GFP	This paper	N/A
pMXs-puro-GFP-SPPL3	This paper	<a href="#">Voss et al., 2012</a>
pMXs-puro-GFP-SPPL3DA	This paper	<a href="#">Voss et al., 2012</a>
pMXs-puro-FLAG-N	This paper	N/A
pMXs-puro-FLAG-C	This paper	N/A
pMXs-puro-FLAG-B3GNT5	This paper	IMAGE:202800754
pMXs-puro-B3GNT5-FLAG	This paper	IMAGE:202800754
pMXs-puro-RFP	This paper	N/A
pMXs-puro-RFP-SPPL3	This paper	<a href="#">Voss et al., 2012</a>
pMXs-puro-RFP-SPPL3DA	This paper	<a href="#">Voss et al., 2012</a>
pMXs-puro-FLAG-B4GALNT1	This paper	IMAGE:202800771
pMXs-puro-FLAG-ST3GAL5	This paper	IMAGE:202759803
puc2CL6IN-HLA-C*05:01 (with mutated signal peptide from HLA-A*02:01 [M4V])	Dr. A. Halenius (University Medical Center Freiburg)	N/A
pLZRS-HLA-A*02:01-IRES-ΔNGFR	Dr. M. Griffioen (LUMC, the Netherlands)	<a href="#">(Griffioen et al., 2012; Van Bergen et al., 2010)</a>
pLZRS-HLA-B*40:01-IRES-ΔNGFR	Dr. M. Griffioen (LUMC, the Netherlands)	<a href="#">(Griffioen et al., 2012; Van Bergen et al., 2010)</a>
pLZRS-HLA-C*03:03-IRES-ΔNGFR	Dr. M. Griffioen (LUMC, the Netherlands)	<a href="#">(Griffioen et al., 2012; Van Bergen et al., 2010)</a>
<b>Software and Algorithms</b>		
Phyre2	<a href="http://www.sbg.bio.ic.ac.uk/">http://www.sbg.bio.ic.ac.uk/</a>	<a href="#">Kelley et al., 2015</a>
PyMOL v2.0	<a href="http://pymol.org">http://pymol.org</a>	Schrödinger, LCC
Compass Data Analysis v4.0 and v5.0 software	<a href="http://www.bruker.com">http://www.bruker.com</a>	Bruker Daltonics
FlowJo Single Cell Analysis v10 software	<a href="http://www.flowjo.com">http://www.flowjo.com</a>	FlowJo, LLC
Fiji – plugin ThunderSTORM		<a href="#">Ovesný et al., 2014</a>
Glycoworkbench		<a href="#">Ceroni et al., 2008</a>
Glycomod		<a href="#">Cooper et al., 2001</a>
TCGA survival and expression data	<a href="http://oncoLnc.org">http://oncoLnc.org</a>	<a href="#">Anaya, 2016</a>
Graphpad Prism	<a href="http://www.graphpad.com">http://www.graphpad.com</a>	N/A
<b>Other</b>		
RFP-Trap_A antibody	ChromoTek	Cat# rta-20, RRID:AB_2631362
Acclaim PepMap100 C18 column, 100 μm × 2cm, C18 particle size 5 μm, pore size 100Å.	Dionex/Thermo Fisher Scientific	Cat# 164199

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REAGENT or RESOURCE	SOURCE	IDENTIFIER
Acclaim PepMap RSLC nanocolumn, 75 $\mu$ m $\times$ 15cm, C18 particle size 2 $\mu$ m, pore size 100 $\text{\AA}$	Dionex/Thermo Fisher Scientific	Cat# TF164534
$^{51}\text{Cr}$ , sodium chromate in normal saline (pH 8-10)	Perkin-Elmer	Cat# NEZ030
Hypercarb KAPPA Column, 30 $\times$ 0.32 mm, particle size 5 $\mu$ m, pore size 250 $\text{\AA}$ .	Thermo Scientific	Cat# 03170480
Hypercarb KAPPA Column, 100 mm $\times$ 75 $\mu$ m, particle size 3 $\mu$ m, pore size 250 $\text{\AA}$ .	Thermo Scientific	Cat# 03170480

**RESOURCE AVAILABILITY****Lead Contact**

Further information and requests for resources and reagents should be directed to and will be fulfilled by the Lead Contact, Robbert Spaapen, Ph.D ([r.spaapen@sanquin.nl](mailto:r.spaapen@sanquin.nl)).

**Materials Availability**

Plasmids and genetically ablated cell lines generated in this study will be available from the lead contact with a completed Material Transfer Agreement (MTA).

**Data and Code availability**

The published article includes all data generated or analyzed during this study, except for the processed screen results, which are accessible in an interactive database (<https://phenosaurus.nki.nl/>). The accession number for the raw sequence data of the screens reported in this paper is NCBI Sequence Read Archive: PRJNA665349. The bioSample accession number for these data in NCBI Sequence Read Archive: SAMN16252402.

**EXPERIMENTAL MODEL AND SUBJECT DETAILS****Mammalian cell lines and T cell clones**

HAP1 (HLA-A\*02:01, HLA-B\*40:01 and HLA-Cw\*03:04, near-haploid, derived from the male chronic myeloid leukemia KBM7 cell line with a fibroblast-like phenotype), MelJuSo (HLA-A\*01:01, B\*08:01 and C\*07:01, female melanoma authenticated at Eurofins), SW620 (HLA-A\*24:02, A\*02:01, B\*07, B\*15 and C\*07:04, male colon, derived from metastatic site: lymph node) (kindly provided by Dr. T. de Gruijl (Amsterdam UMC, the Netherlands)) and U-251 (HLA-A\*02:01, male glioblastoma) (kindly provided by Dr. H. Versteeg (LUMC)) cell lines were cultured in IMDM (GIBCO and Lonza) supplemented with 10% FCS and antibiotics (PenStrep; Invitrogen) at 37°C and 5% CO<sub>2</sub>. HEK293T (ATCC Cat# CRL-3216, RRID: CVCL\_0063) and its derivative Phoenix amphi (female embryonic kidney) cells were cultured in similarly supplemented DMEM (GIBCO). The HLA-A\*02:01-restricted CD8<sup>+</sup> T cell clones reactive against antigens derived from the tumor-expressed proteins USP11, FDPS, VPS13B, ADIR, and SSR1 that are all endogenously expressed by HAP1 cells were previously described (Amir et al., 2011; van Bergen et al., 2007, 2010) and expanded using a feeder cell-cytokine mixture in medium with human serum (Sanquin) as described (Oostvogels et al., 2014).

**METHOD DETAILS****Haploid genetic screening**

Genome-wide haploid genetic screening was performed in either early passage WT or CRISPR/Cas9 generated *SPPL3*<sup>-/-</sup> HAP1 cells using directly conjugated W6/32 or BB7.2 antibodies. Retroviral mutagenesis was performed on  $\sim 100 \times 10^6$  cells using GT-GFP or GT-BFP plasmids as previously described (Brockmann et al., 2017).  $2 \times 10^9$  expanded mutagenized cells were fixed, stained and sorted into two separate populations based on the fluorescent intensity of the respective HLA-I antibody staining (Brockmann et al., 2017). Gene trap integration sites were amplified using a LAM-PCR with a biotinylated primer on genomic DNA isolated from sorted cells. Biotinylated products were captured on streptavidin-coated beads followed by a single-stranded DNA linker ligation and a subsequent amplification step using two primers to generate fragments that include a genomic region flanking the insertion site in addition to adaptors for deep sequencing. Deep sequencing reads were aligned to the human genome (hg19) and intersected with protein coding genes to obtain the numbers of unique disruptive integrations mapped per gene in both populations either lowly or highly recognized by the respective HLA-I antibody. Enrichment of mutations in genes was assessed using a Fisher's exact test corrected for false discovery (Benjamini-Hochberg). The approach is described in detail in Brockmann et al. (2017).

### Plasmids

pMXs-puro vector (Cell Biolabs) was equipped with a novel multiple cloning site with or without N- or C-terminal tags (RFP, GFP and FLAG). SPPL3 and its inactive mutant SPPL3 D271A (kind gift from Dr. R. Flührer) (Voss et al., 2012) were recloned into pMXs-puro-RFP or -GFP using XhoI/BamHI restriction sites. B3GNT5 and B4GALNT1 were PCR amplified from IMAGE:202800754 and IMAGE:202800771 and cloned into pMXs-puro-FLAG-C or -N by EcoRI/BamHI restriction sites. ST3GAL5 was amplified from IMAGE:202759803 and cloned into pMXs-puro-FLAG-C using EcoRI/BclI digestion into an EcoRI/BamHI digested plasmid. The pLZRS-based retroviral vectors containing HLA-A\*02:01/B\*40:01/C\*03:03-IRES-ΔNGFR were described before (Griffioen et al., 2012; Van Bergen et al., 2010). Generation of retroviral supernatants and transduction of cells were performed as described (Spaapen et al., 2008). HLA-C\*05:01 (IMGT/HLA database) with a mutated signal peptide from HLA-A\*02:01 (M4V) was purchased as a gBlock gene fragment (Integrated DNA Technologies, Inc.) and used as a template for amplification. The PCR product was cloned into the puc2CL6IN lentiviral vector using NheI and BamHI. gRNAs (Table S4) were cloned into the pX330 expression vector or the lentiviral vectors lentiCRISPR\_v2 or pLCRISPR.efs.GFP (Addgene) as described (Heckl et al., 2014; Sanjana et al., 2014).

### Genome editing

SPPL3<sup>-/-</sup> and TAPBP<sup>-/-</sup> HAP1 cells were created by in frame integration of a blasticidin-resistance gene after cotransfection of pX330 with TIA-2Ablast (using Extremegene HP, Sigma) as described for other targets (Blomen et al., 2015; de Waard et al., 2020). Clones growing after blasticidin selection (10 μg/mL, Life Technologies) were sequence verified for specific genome editing by Sanger sequencing (primers in Table S5). Lentiviral constructs containing gRNAs targeting SPPL3, UGCG, CMAS and the five core GSL-enzymes were cotransfected into HEK293T with the packaging enzymes psPAX2, pVSVg, pAdVantage using polyethylenimine (PEI; Polyscience) for virus production. Filtered viral supernatants were used for transduction by spinoculation in the presence of 8 μg/mL protamine sulfate. Cells were selected using puromycin (0.25 μg/mL; GIBCO), blasticidin (10 μg/mL; GIBCO), geneticin (550 μg/mL; Formedium) or gated on based on the coexpression of GFP or RFP. Polyclonal gene-deficient populations without selection (core GSL-enzymes for HAP1 cells) or after puromycin selection (SPPL3 and CMAS for HAP1 cells and UGCG for U-251 cells) were used for flow cytometry or T cell assays, or genetically ablated clones were made by limiting dilution and sequence verified. HLA-A<sup>-/-</sup>HLA-B<sup>-/-</sup>HLA-C<sup>-/-</sup> cells were generated by pX330 transfection followed by single cell sort by flow cytometry using W6/32 and sequence verification of clones (de Waard et al., 2020).

### Sanger sequencing

Constructs were sequenced using BigDye Terminator Kit (Applied Biosystems). Genomic DNA isolated from cell lines using DirectPCR (Cell) lysis reagent (Viagen) supplemented with Proteinase K (Sigma) was amplified using primers mentioned in Table S5 and directly sequenced using BigDye Terminator Kit (Applied Biosystems). Sequences were analyzed using Snapgene (GSL Biotech). Sequence decomposition to assess the size of deletions or insertions of genome-edited diploid cells was performed using Tide (Brinkman et al., 2014).

### siRNA transfections

Gene silencing was performed in a 96F well plate using 5 μL siRNA (500nM stock) mixed with 0.1 μL DharmaFECT1 #1 (Dharmacon) in 4.9 μL IMDM. The mixture was incubated for 20min on a shaker at RT and mixed with 4700 cells/200ul. After three days cells were analyzed using flow cytometry. siGENOME human SPPL3 (D-006042, Dharmacon) and β<sub>2</sub>m (M-004366, Dharmacon) siRNAs were used to silence SPPL3 and β<sub>2</sub>m respectively. Non-targeting siRNA (siCTRL, D-001206-13-20, Dharmacon) was used as a negative control.

### Inhibitors and enzymes

UGCG inhibitors MZ31 (used concentration: 2 μM), MZ21 (2 μM), miglustat (100 μM) were produced as previously described (Ghisai-doobe et al., 2014) and eliglustat (200nM) was obtained from Bio-Connect. Swainsonine (20 μg/mL) and kifunensine (25 μM) were obtained from Sigma. N-glycosylation inhibitor activity was confirmed by incubating W6/32 immunoprecipitated HLA-I molecules with Endoglycosidase H (Sigma) in a 20 μl reaction mixture (50 μM Sodiumcitrate (pH5.5), 0.2% SDS, 2 μl Endoglycosidase H and protease inhibitors) for 18h at 37°C followed by HLA-I detection on immunoblot. Sialyltransferase inhibitor (3Fax-peracetyl Neu5Ac, 100 μM) was obtained from Sigma and fucosyltransferase inhibitor (2-Deoxy-2-fluoro-L-fucose, 100 μM) was obtained from Carbosynth. Cells were cultured for 2 or 3 days at 37°C with inhibitors and analyzed by flow cytometry. Cells were incubated with neuraminidase (N2876, Sigma, 225mU/mL) for 1h at 37°C.

### Fab fragment production and labeling

W6/32 Fab fragments were prepared by 1h pepsin (1 μg/μL, pH3.5) treatment at 37°C in a pH3.5 buffer containing citric acid (0.07M) and sodium citrate (0.03M), followed by reduction using DTT (2.5mM, Sigma-Aldrich). BB7.2 Fab fragments were prepared using a papain:antibody ratio of 1:100 (16 μg/mL papain) for 20h at 37°C. All Fab fragments were purified by gel filtration (Superdex 200, 10/300 GL, GE Healthcare Life Sciences). Monomeric Fab fragments were either conjugated to Alexa Fluor® 555 NHS Ester (AF555; ThermoFisher Scientific) or Alexa Fluor® 647 NHS Ester (AF647; ThermoFisher Scientific) according to the manufacturer's labeling protocol. To remove unconjugated fluorophores, the labeled Fab fragments were further purified by gel filtration (Superdex 75, 10/300 GL, GE Healthcare Life Sciences). Fractions containing monomeric fluorophore-conjugated Fab fragments

were concentrated to a protein concentration of 0.2–0.5 mg/mL using Amicon Ultra-4 centrifugal filters (10 kDa cut off, Merck Millipore) and then stored in 50% glycerol at  $-20^{\circ}\text{C}$ . The protein to dye ratios were determined by spectrophotometry at 280 nm and the corresponding absorption maximum of the dyes at 555 nm and 650 nm. The protein to dye ratio of the randomly conjugated Fab fragments were 1.0 (BB7.2 Fab-AF555), 0.94 (W6/32 Fab-AF647) and 0.4 (W6/32 Fab-AF555).

### Flow cytometry using antibodies

Trypsinized cells were incubated with specific antibodies diluted in PBS for 30 min at  $4^{\circ}\text{C}$  and washed up to five times in PBS (with or without 0.5% BSA) before incubation with secondary antibodies if required (30 min at  $4^{\circ}\text{C}$ ). All stains for which no titrations are shown were performed using non-saturating antibody concentrations. Stained cells were fixed in PBS containing 1% formaldehyde (Merck). DAPI (1  $\mu\text{M}$ , Sigma-Aldrich) was used to exclude dead cells from analyses. Cells were fluorescently barcoded using the fluorescent dyes CFSE (125 nM, Invitrogen), Alexa Fluor 350 NHS Ester (40  $\mu\text{M}$ , Thermo Fisher Scientific) or Violet proliferation dye 450 (2.5  $\mu\text{M}$ , BD Horizon) diluted in PBS. Cells were incubated with a fluorescent dye for 15 min (vortexed every 5 min) and washed three times in ice-cold complete IMDM (see above). Barcoded cells were mixed prior to plating in 96V-bottom wells for antibody staining. Stained samples (barcoded or not) were analyzed or sorted on BD flow cytometers (Canto II, Fortessa, LSR II or ARIA II). Flow cytometry data was analyzed using FlowJo (Tree Star, Inc).

### Flow cytometry using other proteins

Fusion proteins LIR-1 Fc (kindly provided by Ofer Mandelboim), KIR2DL1-Fc and KIR2DL2-Fc (R&D Biosystems) were reconstituted in PBS (100  $\mu\text{g}/\text{mL}$ ) and stored at  $-80^{\circ}\text{C}$ . Cholera toxin B subunit (CTB)-FITC (Sigma) was reconstituted in sterile water (500  $\mu\text{g}/\text{mL}$ ) and used at 10  $\mu\text{g}/\text{mL}$ . Trypsinized cells were washed in PBS with 3% FCS or 0.5% BSA and stained in 40  $\mu\text{L}$  (fusion) protein dilution for 30 min to 2 h on ice. For fusion proteins, cells were washed twice and incubated on ice in 40  $\mu\text{L}$  secondary antibody APC AffiniPure F(ab')<sub>2</sub> Fragment Goat Anti-Human IgG, Fc $\gamma$  fragment specific (Jackson ImmunoResearch) (KIR2DL1, KIR2DL2) or mouse anti-human IgG (MH161-1, Sanquin) in-house conjugated to DL650 (Thermo Fisher Scientific) for 30–45 min. After two washes cells were resuspended in PBS/3%FCS containing DAPI and analyzed on BD flow cytometers.

### T cell assays

Target cells were cocultured with T cells in a 1:1 ratio for 18 h as previously described (Spaapen et al., 2008). Cytokine content of cell free supernatants was determined using standard sandwich ELISA according to the manufacturer's instructions (IFN- $\gamma$  and GM-CSF; Sanquin and Biolegend). Some target cells were precultured for 2 days in the presence or absence of UGCG inhibitors. For cytotoxicity experiments, target cells were loaded with 100  $\mu\text{Ci}$  <sup>51</sup>Chromium (Perkin-Elmer) for 90 min at  $37^{\circ}\text{C}$  and then washed twice with PBS. 4000 target cells were then cocultured with T cells at indicated effector target (E:T) ratios for 4–6 h in round-bottom plates after which 30  $\mu\text{L}$  of the supernatant was harvested and analyzed for radioactivity using a gamma counter (Wallac). Medium and 0.1% Triton X-100 served as spontaneous and maximal release controls. The percentage of specific lysis was calculated as  $[(\text{experimental cpm} - \text{spontaneous cpm}) / (\text{maximal cpm} - \text{spontaneous cpm})] \times 100\%$ .

### Immunoprecipitation

The medium of transiently transfected HEK293T cells was filtered through a puradisc 30 syringe filter (0.2  $\mu\text{m}$ , FP 30/0.2) to remove cell material. Before sample buffer was added. HEK293T or CRISPR/Cas9 edited HAP1 cells (confluent 6 cm dish) were lysed for 20 min in lysis buffer containing 0.8% NP-40, 10% glycerol, 150 mM NaCl, 50 mM Tris-HCl pH 8.0, 1 mM EDTA, 5 mM MgCl<sub>2</sub> and protease inhibitors (Roche Diagnostics, EDTA free). Lysate was centrifuged for 20 min at 12,000 rpm and the supernatant was incubated with RFP-Trap beads (Chromotek) or antibody coated Protein-G Sepharose beads for 2 h. Beads were washed four times in lysis buffer before addition of Laemmli Sample Buffer (containing 5%  $\beta$ -mercaptoethanol) followed by 5 min incubation at  $95^{\circ}\text{C}$ . Coimmunoprecipitated proteins were separated by SDS-PAGE for western blotting and detected by antibody staining.

### SDS-PAGE and western blotting

Samples were separated by SDS-PAGE (10% or 12% acrylamide gel) and transferred to a PVDF membrane (Immobilon-P, 0.45  $\mu\text{m}$ , Millipore) at 300 mA for 3 h. The membranes were blocked in PBS/5% Milk (Skim milk powder, Oxoid) and incubated with a primary antibody for 1 h diluted in PBS/0.1% Tween/5% Milk, washed thrice for 10 min in PBS/0.1% Tween and incubated with the secondary antibody for 45 min diluted in PBS/0.1% Tween/5% Milk and washed thrice again in PBS/0.1% Tween. The filter was incubated with ECL reagent (SuperSignal West Dura Extended Duration Substrate, Thermo Fisher Scientific) and the signal was detected using the Chemidoc XRS+ imager (Bio-Rad) or Amersham Imager 600.

### BFA assay

To determine HLA-I turnover, HAP1 cells were seeded in 96w plates (80% confluent) and cultured in the presence of Brefeldin A (0.5  $\mu\text{g}/\text{mL}$  BFA, Sigma Aldrich) at  $37^{\circ}\text{C}$  for indicated times. BFA containing medium was removed and the cells were washed in cold PBS, trypsinized and harvested in PBS/0.5% BSA. Cells were kept on ice and stained for flow cytometry using non-saturating amounts of W6/32 and B1.23.2 antibodies (see flow cytometry section).

### Crystal structures

Structural prediction software Phyre2 was used to create a model of SPPL3 using the primary consensus sequence CCDS9208.1 (Kelley et al., 2015). Models of HLA-A\*02:01 were made using the crystal structure 3MRG courtesy of the RCSB PDB (Reiser et al., 2014; Winn et al., 2011). The structure 3QZW was used in conjunction with the CCP4 program ARIAIMOL (Berman et al., 2000) to determine the hydrogen bonding contacts between human HLA-A\*24:02 and human CD8 alpha-alpha dimer (Shi et al., 2011). A similar method was used to assess the contacts between LIR-1 and HLA-A\*02:01 using the structure 1P7Q (Berman et al., 2000; Willcox et al., 2003). All figures have been produced using the PyMOL molecular graphics software (Version 2.0 Schrödinger, LLC).

### B3GNT5 activity assay

$5 \times 10^6$  HAP1 cells were harvested by trypsinization and cells were lysed in lysis buffer (2% Triton X-100, 50mM sodium cacodylate pH7.4, 10mM  $MnCl_2$ , and protease inhibitors (Roche Diagnostics, EDTA free)) by incubating on ice for 30min. Nuclei were precipitated by centrifugation at 15,000xg. Equal volumes of lysis buffer containing  $2\mu M$  BODIPY-C5-Lactosylceramide complexed to BSA (Thermo Fisher Scientific) and/or 1mM UDP-N-acetyl-D-glucosamine (Santa Cruz) were added to post-nuclear supernatants containing equal amounts of protein in  $50\mu L$  lysis buffer. Samples were incubated at  $37^\circ C$  for 4h and subjected to lipid extraction using Bligh-Dyer method (Bligh and Dyer, 1959).  $100\mu L$  2% NaCl,  $250\mu L$  chloroform, and  $500\mu L$  methanol were added to the reactions and samples were vortexed. Phase separation was induced by addition of  $250\mu L$  0.45% NaCl and  $250\mu L$  chloroform and lower phases were collected. Upper phases were re-extracted twice more and collected lower phases were dried under a nitrogen flow. Dried lipids were resuspended in chloroform:methanol (2:1 v/v) solution and spotted on TLC plates. TLC plates were developed in chloroform:methanol:water (60:25:4 v/v/v) and imaged using Typhoon FLA9500 (GE Healthcare) scanner equipped with a 473nm laser and BPB1 filter (530DF30). Identity of BODIPY-Lc3Cer was confirmed by MS/MS. Structures were assigned based on MS/MS fragmentation pattern in negative mode following nomenclature from Domon and Costello (1988).

### GSL extraction and purification by RP-SPE

GSLs were extracted from  $1 \times 10^7$  HAP1 WT, *SPPL3*<sup>-/-</sup> and *SPPL3*<sup>-/-</sup> *B3GNT5*<sup>-/-</sup> cell lines in triplicate in glass vials equipped with a Teflon-lined screw cap. Cells were washed three times in 1mL of water followed by centrifugation at 2000xg for 30min. The supernatant was removed and replaced by  $300\mu L$  of 2-propanol. The samples were vortexed for 5min and incubated for 15min at  $75^\circ C$ . A volume of  $350\mu L$  of MTBE (Sigma-Aldrich) was added to the samples followed by 15min sonication.  $200\mu L$  of water was added to the cell pellets and incubated for 4h with shaking at room temperature. The upper phase containing GSLs was collected after centrifugation at 2700xg for 20min. Then,  $400\mu L$  of MTBE was added, followed by sonication and centrifugation. The upper phase was collected and pooled to the previous sample. The process of adding MTBE, sonication, centrifugation and removing upper phase was repeated another two times. The combined upper phases were dried under vacuum in an Eppendorf Concentrator 5301 (Eppendorf) at  $30^\circ C$ . Before purification of the GSLs using RP-solid phase extraction (SPE), the samples were dissolved in  $200\mu L$  methanol and vortexed for 10min, followed by addition of  $400\mu L$  water. TC18-RP-cartridges were prewashed with 2mL of chloroform/methanol (2:1, v/v), 2mL of methanol followed by equilibration with 2mL methanol/water (1:2, v/v). The extracted GSLs were loaded to the cartridge for 3 times and washed with 3mL methanol/water (1:2, v/v). The GSLs were eluted from the column with 2mL methanol and 2mL chloroform/methanol (2:1, v/v). The samples containing the eluate were evaporated under nitrogen for 1h and dried under vacuum in an Eppendorf Concentrator at  $30^\circ C$ . The collection and dry of GSLs eluate were performed in glass tube.

### GSL glycan release by EGCCase I and purification

To release the glycans from the GSLs, a mixture of  $2\mu L$  Endoglycoceramidase I (EGCase I recombinant clone derived from *Rhodococcus triatomea* and expressed in *Escherichia coli* (12mU, NEB)),  $4\mu L$  10x EGCCase I buffer (500mM HEPES, 1M NaCl, 20mM DTT and 0.1% Brij 35, pH5.2, NEB) and  $34\mu L$  water (pH5.2) was added to each sample and incubated for 16h at  $37^\circ C$ . The released glycans were collected and applied to a TC18-RP-cartridges (Waters) which was preconditioned with 2mL of methanol and 2mL of water. The sample vials were washed with  $200\mu L$  of water and residual glycans were loaded to the cartridge. Then,  $500\mu L$  of water was added to the cartridge to wash the glycans from the column. The flow-through and wash fractions were pooled and dried in an Eppendorf Concentrator at  $30^\circ C$ .

### Reduction, desalting and carbon SPE cleanup of GSL glycans

The reduction was carried out as described previously with slight modifications (Jensen et al., 2012). In brief, GSL glycans were reduced to alditols in  $20\mu L$  of sodium borohydride (500mM, Sigma-Aldrich) in potassium hydroxide (50mM, Sigma-Aldrich) for 2h at  $50^\circ C$ . Subsequently,  $2\mu L$  of 100% glacial acetic acid was added to neutralize the solution and quench the reaction. The desalting of GSL glycans was performed on cation exchange columns (Sigma-Aldrich) which consist of  $60\mu L$  of AG50W-X8 resin beads deposited onto reversed phase  $\mu C18$  ZipTips (Perfect Pure, Millipore) as previously described (Jensen et al., 2012). Glycan alditols were eluted with  $50\mu L$  of water twice. The combined flow-through and eluate were pooled and dried under vacuum in an Eppendorf Concentrator at  $30^\circ C$ . The carbon SPE clean-up was performed and the purified glycan alditols were resuspended in  $10\mu L$  water for porous graphitized carbon (PGC) LC-ESI-MS/MS analysis.

### Analysis of GSL glycans using PGC LC-ESI-MS/MS

Porous graphitized carbon (PGC) LC-ESI-MS/MS analysis of GSL glycan alditols was performed on a Dionex Ultimate 3000 nano-LC system equipped with a Hypercarb PGC trap column (5  $\mu\text{m}$  Hypercarb Kappa, 32  $\mu\text{m}$  x 30mm, Thermo Fisher Scientific) and a Hypercarb PGC nano-column (3  $\mu\text{m}$  Hypercarb Kappa, 75  $\mu\text{m}$  x 100mm, Thermo Fisher Scientific) coupled to an amaZon speed ion trap mass spectrometer (Bruker Daltonics). Mobile phase A consisted of 10mM ammonium bicarbonate (Sigma-Aldrich). Mobile phase B was 60% (v/v) acetonitrile (Biosolve) / 10mM ammonium bicarbonate (Sigma-Aldrich). To analyze glycans, 2  $\mu\text{L}$  injections were performed and separation was achieved with a gradient of B (1%–71% at 0.7%/min) followed by a 10min wash step using 95% of B at a flow rate of 0.6  $\mu\text{L}/\text{min}$ . MS scans from  $m/z$  340 to 1700 were recorded in enhanced mode using negative ion mode. MS/MS spectra were recorded selecting the top 3 highest intensity peaks. Glycan structures were assigned based on glycan composition obtained from accurate mass, relative PGC elution position, MS/MS fragmentation pattern in negative-ion mode and general glycobiochemical knowledge (Karlsson et al., 2004), with help of Glycoworkbench (Ceroni et al., 2008) and Glycomod (Cooper et al., 2001) software tools. Extracted ion chromatograms were used to integrate area under the curve (AUC) for each individual glycan isomer using Compass Data Analysis software v.5.0. The most abundant peaks in the glycan profile were manually picked and integrated. Relative quantitation of individual glycans was performed on the total area of all included glycans within one sample normalizing it to 100%.

### HLA-I glycan analysis, in-gel tryptic digestion

The glycopeptide generation and analysis were performed as described previously with slight modifications (Plomp et al., 2014). BB7.2 immunoprecipitated samples were loaded on SDS-PAGE. Bands containing HLA-I were excised and cut into pieces. The gel pieces were washed with 25mM ammonium bicarbonate, dehydrated with acetonitrile (ACN) and reduced in-gel for 30min at 55°C with 100  $\mu\text{L}$  10mM DTT in 25mM ammonium bicarbonate solution. Thereafter they were dehydrated in ACN followed by cysteine alkylation for 20min with 100  $\mu\text{L}$  of a 55mM iodoacetamide (Sigma-Aldrich) in 25mM ammonium bicarbonate solution in the dark for 45min. This was repeated twice, and the gel pieces were subsequently dried in a centrifugal vacuum concentrator at 30°C for 10min. Enzymatic digestion of trypsin was performed by adding 50  $\mu\text{L}$  of 25mM ammonium bicarbonate containing 0.6  $\mu\text{g}$  of trypsin (sequencing grade modified trypsin, Promega) to the dried gel particles. The samples were kept on ice for 1h and were subsequently incubated overnight at 37°C. The solution surrounding the gel pieces was collected and stored at –20°C. 20  $\mu\text{L}$  of 25mM ammonium bicarbonate was added to the remaining gel pieces, and incubated at 37°C for another hour. The solution was again collected and added to the first fraction prior to freezing.

### Glycopeptide analysis by reverse-phase (RP) nanoLC-ESI-MS(/MS)

Glycopeptides were analyzed by RP nanoLC-ESI-MS(/MS) on an Ultimate 3000 RSLCnano system (Dionex / Thermo Fisher Scientific) coupled to an HCTultra-ESI-ion trap-MS (Bruker Daltonics). 5  $\mu\text{L}$  sample was injected and concentrated on a trap column (Acclaim PepMap100 C18 column, 100  $\mu\text{m}$  x 2cm, C18 particle size 5  $\mu\text{m}$ , pore size 100Å, Dionex / Thermo Fisher Scientific) before separation on an Acclaim PepMap RSLC nanocolumn (75  $\mu\text{m}$  x 15cm, C18 particle size 2  $\mu\text{m}$ , pore size 100Å, Dionex / Thermo Fisher Scientific). A flow rate of 700nL/min was applied. Solvent A consisted of 0.1% formic acid in water; solvent B, 0.1% formic acid in 95% ACN and 5% water. A linear gradient was applied with the following conditions:  $t = 0\text{min}$ , 3% solvent B;  $t = 5\text{min}$ , 3% solvent B;  $t = 20\text{min}$ , 27% solvent B;  $t = 21\text{min}$ , 70% solvent B;  $t = 23\text{min}$ , 70% solvent B;  $t = 24\text{min}$ , 3% solvent B;  $t = 43\text{min}$ , 3% solvent B. Samples were ionized in positive ion mode with an online nanospray source (4500V) using fused-silica capillaries and a Distal Coated SilicaTip Emitter (New Objective) with an internal diameter of 20  $\mu\text{m}$  (10  $\mu\text{m}$  at the tip) and a length of 5cm. Solvent evaporation was performed at 220°C with a nitrogen flow of 3L/min. For the detection of glycopeptides, the MS ion detection window was set at  $m/z$  500–1800, and the MS/MS detection window at  $m/z$  140–2200, with automated selection of the three highest peaks in the spectrum for MS/MS analysis. The LC-MS/MS results were analyzed using DataAnalysis 4.0 software (Bruker Daltonics) and screened manually for the masses of common oxonium fragment ions ( $m/z$  366.1, [1 hexose + 1 GlcNAc + H]<sup>+</sup>;  $m/z$  657.2, [1 hexose + 1 GlcNAc + 1 N-acetylneuraminic acid + H]<sup>+</sup>;  $m/z$  528.2, [2 hexoses + 1 GlcNAc + H]<sup>+</sup>), which are characteristic for fragmentation spectra of glycopeptides. Glycopeptide MS/MS spectra were further analyzed manually to derive the oligosaccharide structure and the mass of the peptide moiety.

### Preparation of fibronectin-coated glass slides for single particle tracking experiments

Glass slides (24mm x 50mm #1.5 borosilicate, VWR) were immersed in a 1:1 mixture of concentrated sulfuric acid (Sigma) and 30% hydrogen peroxide (Sigma) for at least 30min, rinsed with deionized water, air-dried and glued with picodent twinsil extrahart (Picodent) to the bottom of 8-well LabTek chambers (Nunc). Slides were coated with 20 $\mu\text{g}/\text{mL}$  fibronectin (Sigma-Aldrich) in PBS for 1–2h at 37°C and rinsed with 1 X PBS.

### Single particle tracking of HLA-I molecules on HAP1 WT and SPPL3<sup>-/-</sup> cells

0.2 x 10<sup>6</sup> HAP1 WT and SPPL3<sup>-/-</sup> cells were stained with a single molecule dilution of the BB7.2-Fab-AF555 on ice for 30min and washed twice in imaging buffer (HBSS, GIBCO, 1% FCS, 2mM MgCl<sub>2</sub> and CaCl<sub>2</sub>). Cells were kept on ice or seeded onto fibronectin-coated glass slides for imaging at room temperature (23–27°C) and in TIRF mode. AF555 was excited with a 532 nm laser (Obis) with a power density of 0.8kW/cm<sup>2</sup> and the emission channel was cleaned up with a TRITC filter (605/52) installed within the fast emission filter wheel (Sutter Instrument Company). We recorded single HLA-I (BB7.2 Fab-AF555) trajectories over 500 frames with an illumination time of 16ms and a total time lag of 20.6ms between two adjacent images (100 X 100 pixel ROI size). Microscopy images were

processed and analyzed with the open-source image processing package Fiji (Schindelin et al., 2012). XY localization, intensity and positional accuracy of single fluorescence emitters was calculated with the Fiji plugin ThunderSTORM (Ovesný et al., 2014). After determining the localization of every fluorophore in the image stack we combined these localizations to trajectories based on a published approach (Gao and Kilfoil, 2009) with custom-made algorithms written in MATLAB (MathWorks). We calculated the mean square displacement (MSD) describing the average of the square displacements between two points of the trajectory according to  $MSD(t_{lag}) = \langle (r \cdot (t + t_{lag}) - r(t))^2 \rangle$ . The first three MSD values as a function of time lag ( $t_{lag}$ ) were used to calculate the diffusion coefficient (D) for each trajectory according by fitting  $MSD = 4 \cdot D \cdot t_{lag} + 4 \cdot \sigma_{xy}^2$ , with  $\sigma_{xy}$  representing the localization precision (Wieser et al., 2007). Multiple fractions (i.e., a fast and a slow moving fraction of molecules) were discriminated by analyzing the step-size distributions of square displacements for several time lags (Schütz et al., 1997). By assuming free Brownian motion of one mobile fraction, the cumulative probability for finding a square displacement smaller than  $r^2$  is given by  $P = 1 - \exp(-r^2 / 4Dt_{lag})$ ; two different fractions  $\alpha$  and  $(1-\alpha)$  with diffusion coefficients D1 (e.g., fast) and D2 (e.g., slow/immobile) can be distinguished by fitting the bi-exponential function  $P = 1 - \alpha \cdot \exp(-r^2 / 4D_1 t_{lag}) - (1 - \alpha) \cdot \exp(-r^2 / 4D_2 t_{lag})$ . We calculated the fraction of mobile and slow/immobile molecules, the diffusion rate of mobile molecules and the diffusion rate of slow/immobile molecules of all HLA-I trajectories present on a single cell (Bramshuber et al., 2018).

### QUANTIFICATION AND STATISTICAL ANALYSIS

All error bars correspond to the standard deviation of the mean. For all figures, the number of independent experiments (n) are described in the legend. For quantifications of flow cytometry data we plotted data from median fluorescent intensities (fusion proteins, no normal distribution) or mean fluorescent intensities (antibody and CTB stainings) as clarified in the legends. Data from genome-wide screens were analyzed using two-sided Fisher's exact test followed by FDR (Benjamini-Hochberg) correction of the p value. Other statistical evaluations were done by a Student's t test (analysis of two data groups), one-way ANOVA (three groups or more), two-way ANOVA (two variables), Mann-Whitney U test (non-parametric analyses) or Log-rank (survival analyses) with Prism-Graphpad software (<http://www.graphpad.com>). TCGA survival and expression data were retrieved from [OncoLnc.org](http://OncoLnc.org) (Anaya, 2016). We subdivided patients with the 50% lowest SPPL3 or B3GNT5 expressing tumors versus the 50% highest expressing tumors. \*p < 0.05, \*\*p < 0.01, \*\*\*p < 0.001 and ns (= not significant). EC<sub>50</sub> values of titrations were calculated using non-linear four parameter fit modeling with Prism software.

**Update**

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## Correction

# The SPPL3-defined glycosphingolipid repertoire orchestrates HLA class I-mediated immune responses

Marlieke L.M. Jongsma, Antonius A. de Waard, Matthijs Raaben, Tao Zhang, Birol Cabukusta, René Platzer, Vincent A. Blomen, Anastasia Xagara, Tamara Verkerk, Sophie Bliss, Xiangrui Kong, Carolin Gerke, Lennert Janssen, Elmer Stickel, Stephanie Holst, Rosina Plomp, Arend Mulder, Soldano Ferrone, Frans H.J. Claas, Mirjam H.M. Heemskerk, Marieke Griffioen, Anne Halenius, Hermen Overkleeft, Johannes B. Huppa, Manfred Wuhrer, Thijn R. Brummelkamp, Jacques Neefjes, and Robbert M. Spaapen\*

\*Correspondence: [r.spaapen@sanquin.nl](mailto:r.spaapen@sanquin.nl)

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In our article entitled “The SPPL3-defined glycosphingolipid repertoire orchestrates HLA class I-mediated immune responses,” we employed a glioblastoma cell line in some experiments to which we referred as U373. Short tandem repeat profiling-based authentication revealed that we had actually used the glioblastoma cell line U-251 instead of U373. The U-251 and U373 cell lines were derived from two different glioblastoma patients. An error occurring decades ago caused U-251 to be incorrectly distributed by multiple cell line repositories as “U373,” resulting in a misreporting of the cell line’s name that still occasionally occurs today (<https://www.ncbi.nlm.nih.gov/pmc/articles/PMC4303149/>). We have now corrected the name of the cell line we used to U-251. The misidentification does not affect the results, conclusions and implications discussed. The authors apologize for any confusion that may have been caused.

