Structure-function characterization of three human antibodies targeting the vaccinia virus adhesion molecule D8

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ABSTRACT

Vaccinia virus (VACV) envelope protein D8 is one of three glycosaminoglycan adhesion molecules and binds to the linear polysaccharide chondroitin sulfate (CS). D8 is also a target for neutralizing antibody responses that are elicited by the smallpox vaccine, which has enabled the first eradication of a human viral pathogen and is a useful model for studying antibody responses. However, to date, VACV epitopes targeted by human antibodies have not been characterized at atomic resolution. Here, we characterized the binding properties of several human anti-D8 antibodies and determined the crystal structures of three VACV-MAb variants, VACV-66, VACV-138, and VACV-304, separately bound to D8. While all these antibodies bound D8 with high affinity and were moderately neutralizing in the presence of complement, VACV-138 and VACV-304 also fully blocked D8 binding to CS-A, the low affinity ligand for D8. VACV-138 also abrogated D8 binding to the high-affinity ligand CS-E, but we observed residual CS-E binding was observed in the presence of VACV-304. Analysis of the VACV-138 and VACV-304 binding sites along the CS binding crevice of D8, combined with different efficiencies of blocking D8 adhesion to CS-A and CS-E allowed us to propose that D8 has a high and low affinity CS binding region within its central crevice. The crevice is amenable to protein engineering to further enhance both specificity and affinity of binding to CS-E. Finally, a wild-type D8 tetramer specifically bound to structures within the developing glomeruli of the kidney, which express CS-E. We propose that through structure-based protein engineering, an improved D8 tetramer
could be used as a potential diagnostic tool to detect expression of CS-E, which is a possible biomarker for ovarian cancer.

Smallpox is caused by infection with variola virus and was a major health threat until successful global vaccination efforts led to its complete eradication from the general human population (1). This eradication of an infectious agent was achieved by immunizing with Vaccinia virus (VACV), a related orthopoxvirus with low virulence and the active ingredient in the smallpox vaccine (2,3). Immunization with VACV leads to the robust production of highly neutralizing antibodies targeting several VACV envelope proteins, including A27, A33, B5, D8, H3, L1, and others, which are expressed on different viral envelopes (3,4). A27, D8, H3 and L1 are expressed on the outer membrane of the intracellular mature virion (IMV), while A33 and B5 are found in the more fragile extracellular enveloped virion (EEV), which has an additional host cell derived envelope. Both the IMV and the EEV particles are infectious virions, with IMV being more abundant and mainly responsible for spread between hosts, and EEV being involved in cell-to-cell spread within the host after infection with the IMV (5). As a result, a potent antibody-mediated immune response against VACV targets both infectious virions.

Among the targeted envelope proteins, A27, D8, and H3 are glycosaminoglycan (GAG) adhesion molecules. GAGs are linear, mostly sulfated polysaccharides with repeating disaccharide units. The precise sulfation of each GAG is regulated by specific sulfotransferases, leading to the production of different isoforms. GAGs are covalently attached to various core proteins to form the proteoglycan (PG). They are ubiquitously expressed on cell surfaces and in extracellular matrices (6). In addition to providing structural integrity, GAGs participate in various physiological processes such as cell adhesion, cell proliferation, migration, neurite outgrowth, microbial adhesion and infection, inflammation and angiogenesis (7-11). Many pathogens encode GAG-binding proteins as a major route of entry (12). While A27 and H3 (13-15) bind to heparan sulfate (HS), D8 binds to chondroitin sulfate (CS) (16). Chondroitin sulfate (β-GlcA-β-(1-3)GalNAc-sulfate) is the most abundant GAG and has 4 major isotypes (chondroitin-4-sulfate, CS-A; chondroitin-6-sulfate, CS-C; chondroitin-2,6-sulfate, CS-D; chondroitin-4,6-disulfate, CS-E) in mammals (17).

We have determined previously the structure of VACV D8 protein, which revealed an N-terminally located carbonic anhydrase (CAH) domain (residues 1-235), followed by a disordered stalk region (residues 236-267) that is required for higher oligomeric assembly of the D8 hexamer in solution, and for embedding D8 in the virial envelope (18,19). We have further identified CS-E as the optimal ligand for D8 and identified a positively charged crevice as the CS binding site (19). Computational docking suggested that the crevice is capable of binding to at least an octasaccharide of CS-E (18).

We also previously characterized the binding pattern for a panel of D8-reactive murine antibodies, all of which targeted the CAH domain rather than the stalk region, and can be grouped into 4 different specificity groups based on cross-blocking studies (18,20,21). One of these antibodies, LA5, is moderately neutralizing in the presence of complement and prevents D8 binding to CS-E, since LA5 binds above the CS-E binding crevice (18,19).

However, to date, VACV epitopes that are targeted by human antibodies have not been characterized to atomic resolution using X-ray crystallography, and in total only the structures of eight murine antibodies bound to their VACV antigens A27, A33, D8, and L1 have been determined (19,22-25).

Previously, we reported isolation of a panel of 21 anti-D8 human monoclonal antibodies (MAb) and assessed their neutralizing and protective capabilities (26). In this study, we have characterized the epitopes for some of these human D8 antibody and determined the first crystal structures of human antibodies bound to D8, using three clones designated VACV-66, VACV-138 and VACV-304. Similarly to the murine antibodies, human antibodies bound to the CAH domain of D8 and were moderately neutralizing in the presence of complement. Two antibodies, VACV-138 and VACV-304, also fully blocked D8 binding to CS-A, while VACV-304 only partially blocked D8 binding to the high affinity ligand CS-E, revealing a high- and low-affinity CS binding region within the central D8 crevice.

RESULTS

Human anti-D8 MAbs identify a novel D8 epitope. We analyzed four human anti-D8 MAbs VACV-66, VACV-138, VACV-249, and VACV-304 for their ability to block each other’s binding to recombinant D8 using a real-time binding assay (biolayer interferometry, BLI). VACV-66 was the
only antibody that did not compete with any other MAb for D8 binding, suggesting it bound a distinct epitope on D8 (Fig 1A). VACV-138, VACV-249, and VACV-304 all competed with each other for binding to D8, suggesting that their epitopes on D8 were overlapping. However, a previous study suggested that crossblocking of VACV-249 with VACV-304 depends on the order of antibodies used in the assay, suggesting the overlap is only partial (26). Next, we used ELISA to analyze the ability of VACV-66 and VACV-304, as representatives of the two major human MAb groups, to block binding of murine anti-D8 MAbs in a previously characterized antibody panel (21). We included representative murine MAbs from specificity groups I, II and IV to ask whether the human MAbs recognized common or unique epitopes on D8 (Fig 1B). While VACV-304 blocked binding of mouse CC7.1 (group II), human VACV-66 appeared to recognize a unique epitope on D8, notwithstanding that there was slight cross-blocking with the murine group IV antibodies HG12 and LA5 (Fig 1). LA5 had been shown previously to block D8 binding to its ligand CS-E (18,19), raising the possibility that some human MAbs may also prevent D8 binding to CS-E.

**Human and murine anti-D8 MAbs possess moderate neutralizing potency**- We next tested the ability of anti-D8 MAbs to neutralize IMV in a flow cytometry-based assay. While the murine control anti-L1 MAb M12B9 neutralized IMV in the absence of complement, most anti-D8 MAbs required the presence of complement to neutralize IMV, regardless of whether they were of human or murine origin (Fig 2). Murine MAb JF11 was the only anti-D8 MAb that did not neutralize, while neutralization mediated by MAb JE11 also was weak. Compared to L1, D8 appeared to be a target only for antibodies with a moderate level of neutralizing potency, likely because the GAG adhesion molecules A27 and H3 can compensate for the function of D8.

**Human anti-D8 MAbs bind with high affinity**- Next, we determined the binding affinity of human anti-D8 MAbs for recombinant D8. While VACV-66, VACV-138 and VACV-249 bound D8 with relatively low affinity ($K_D = 0.35$ to $1.9 \text{ nM}$), VACV-304 bound with much higher affinity ($K_D=16\text{pM}$, Fig 3). MAb VACV-249 had a comparably low association rate ($k_{on}= 2 \times 10^4 1/\text{Ms}$), while VACV-66, VACV-138 and VACV-304 bound similarly fast ($k_{on}= 6.5 \text{ to } 7.2 \times 10^5 1/\text{Ms}$). Surprisingly, the high affinity of VACV-304 was a result of extremely slow dissociation, which was roughly 100-times slower compared to VACV-66 ($k_{off}= 1 \times 10^{-5}$ vs. $1 \times 10^{-3} 1/\text{s}$). This finding suggested that VACV-304 formed a stable complex with D8 that was characterized by a long half-life in solution. As the kinetic measurements were performed with immobilized intact IgG, only the 1:1 binding interaction between a single Fab and D8 was measured. Increases in avidity due to antibody bivalency were not addressed in this experimental design. As a result, binding of the antibody to D8 embedded in the virion membrane during natural infection is likely even higher.

**Human antibodies can interfere with D8 adhesion to CS** - We next asked if any of the antibodies could prevent binding of D8 to its host ligand chondroitin sulfate-E (CS-E), similarly to the murine antibody LA5 (18), and whether the ability to block CS binding correlated with their neutralization capacity. MAbs were pre-bound to oligomeric D8, since monomeric D8 does not bind to CS-A. The D8-MAb complexes were assessed for their binding ability to CS-A and CS-E using GAG microarrays (Fig 4). While VACV-66 did not impair D8 binding to CS-A or CS-E, both VACV-138 and VACV-304 fully abrogated D8 binding to CS-A. Although binding to CS-E was greatly reduced when vv304 bound to D8, a low level of binding (25% of maximum binding) was consistently observed. As CS-E is the high-affinity D8 ligand, and CS-A is only bound by oligomeric D8, VACV-304 likely binds at one of the extremities of the crevice. This has been confirmed by structural studies discussed below. VACV-304 would then leave the remainder of the D8 crevice uncovered and available to bind CS-E. CS-A, on the other hand, would not be able to bind to only a partially accessible D8 crevice due to its lower binding affinity compared to CS-E. Although the MAbs do not show great differences in differences in extent of IMV neutralization (~20 -50%), all of the MAbs that prevented CS adhesion (VACV-138, VACV-304 and LA5) neutralize to slightly higher levels compared to MAbs VACV-66, JE11, and JF11 that did not block CS adhesion (Figs 2 and 3 (18)). VACV-249 was not included in the CS microarray but since its binding site on D8 partially overlaps with that of VACV-138 and VACV-304, it is likely that VACV-249 also could impair D8 binding to CS-E.
Structural characterization of the D8 epitopes targeted by human MAbs - To identify and characterize the D8 epitopes that are targeted by the human MAbs, we determined the crystal structures of VACV-66, VACV-138 and VACV-304 bound to the CAH domain of D8, to resolutions of 2.23 Å, 2.9 Å, or 2.9 Å respectively (Table 1).

VACV-66-D8 complex - VACV-66 binds on the side of D8, away from the CS-binding crevice, explaining the inability of VACV-66 to interfere with D8 attachment to CS-E (Fig 5). The VACV-66 binding site on D8 is rotated 90° compared to the CS-E binding crevice and is adjacent to that of the murine group II, III, and IV antibodies, correlating with a low degree of cross-blocking with representative antibodies from these groups (Fig 1).

The Fab binds to a total of 21 discontinuous D8 residues. The heavy chain (HC) buries a total of 1683 Å², while the light chain (LC) buries a total of 313 Å² (Table S1). The HC dominates the overall interactions and binds to 20 D8 residues (Gln3, Gln4, Asn18, Arg20, Leu21, Phe56, Pro58, Tyr61, Lys109, His110, Asp111, Asp112, Ile115, Leu170, Asp179, Val181, Asp228, Thr229, Glu230, and Tyr232) while the LC contacts three D8 residues (Asn18, Ala19, and Arg20). All three HC CDRs are involved in the interaction, forming an intricate H-bond network of 21 H-bonds and one salt bridge, while the LC forms a cation-pi interaction between Tyr33L and Arg20 of D8 (Fig 6 and Table S2). VACV-66 exhibited a relatively fast dissociation from D8, possibly as a result of the lack of strong electrostatic, or hydrophobic interactions with D8.

VACV-66 is the only one of the three antibodies studied that did not engage in salt-bridges with the antigen (Table S2). The VACV-66 binding site on D8 also is rather shallow and broad, with only minimal participation of the LC in antigen binding.

VACV-138-D8 complex - VACV-138 blocks D8 adhesion to CS and binds directly over the CS binding crevice (Fig 5). VACV-138 was not included in the murine antibody cross-blocking study, but shares several contact residues with murine LA5, which also interferes with CS-E adhesion. Compared to LA5 however, the footprint of VACV-138 is not located centrally above the binding crevice but rather is shifted along the crevice. VACV-138 engages D8 in a slightly tilted manner compared to the more centrally placed murine LA5 (19). The VACV-138 footprint is delineated by Lys48 and Glu105 at one end, and Thr39 and Asn175 at the other end, thereby sitting over both sides of the crevice.

The Fab binds to a total of 21 discontinuous D8 residues and the HC buries a total of 1465 Å², while the LC buries a total of 435 Å². As a result, the LC is slightly more involved in D8 binding compared to MAb VACV-66. Nevertheless, the HC dominates the overall interactions and binds to 17 D8 residues while the LC contacts five D8 residues (Table S1). All three HC CDRs bind D8 with a total of 15 H-bonds and three salt bridges, while the LC only forms one polar interaction using L3 (Fig 6 and Table S2). The additional electrostatic interactions compared to VACV-66 are consistent with the reduced dissociation from D8 and an overall higher binding affinity.

VACV-304-D8 complex - Similar to VACV-138, VACV-304 also blocks D8 adhesion to CS and binds over the CS binding crevice (Fig 5). However, in contrast to VACV-138, VACV-304 binds closer to the end of the crevice, delineated by Lys48 and Lys98 on one end and Asn145 and Glu105 on the other end. Likely due to the slightly altered footprint compared to VACV-138 and the loss of interaction with D8 residue Asn175, the CS binding crevice is more accessible upon VACV-304 binding and thus residual binding to the high affinity ligand CS-E was observed (Fig 4). In addition, slight cross-blocking with LA5 was observed, since VACV-304 and LA5 share the D8 contact residues Thr39, Arg44, and Asn145, which are the peripheral binding residues of LA5 (19).

The Fab binds to a total of 21 discontinuous D8 residues, and the HC buries a total of 1010 Å², while the LC buries a total of 977 Å². The HC binds to 14 D8 residues while the LC contacts ten D8 residues (Table S1). All three HC CDRs bind D8 with a total of 8 H-bonds, while the LC forms 5 H-bonds and 1 salt bridge using L2 and L3 (Fig 6 and Table S2). In contrast to VACV-66 and VACV-138, the LC of VACV-304 is far more involved in D8 binding, as judged by the buried surface area and the number of contact residues. This finding is consistent with the highest observed binding affinity among the tested human MAbs.

In summary, the Fab binding affinity correlates well with the number of electrostatic interactions, as well as the involvement of the LC in D8 binding. VACV-66, which has the lowest affinity, has the least involvement of the LC in binding and lacks salt-bridges, while VACV-304 has the highest affinity, establishes several salt-bridges and also the most contacts using the LC.
The affinity differences are driven mostly by the off-rate, suggesting a locking mechanism when both LC and HC bind to D8, leading to a reduced off-rate. VACV-304 has a 100-fold reduced off-rate compared to VACV-66.

**Antibody binding to D8 mutants** - We generated a panel of ten D8 mutants in which either one (Gln3, Tyr85, Tyr101, Glu105, Glu106, Asp112, Asn1675, and Arg220) or two (Arg44/Arg220 and Lys48/Arg220) residues were replaced with alanine to assess the effect of a particular amino acid side chain in antibody binding (S1 Fig). The D8 double mutants Lys48/Arg220 and Arg44/Arg220 had been shown previously to abrogate D8 binding to CS-E (18).

**MAb VACV-66** - None of the mutations tested significantly affected VACV-66 binding. Of the ten mutants, Gln3 and Asp112 are contact residues but did not appear to be critical for binding. In the structure, Gln3 forms a H-bond with Tyr55H of VACV-66, but since the carboxylate of Asp179 also contacts Tyr55H, the interaction with Gln3 appears to be dispensable. Asp112 forms two potential H-bonds with Y60H but since the mutation has no effect, loss of these interactions is likely compensated for by the other contact residues. Several D8 mutants appear to increase antibody dissociation slightly, but since they are located outside the binding interface they likely have an allosteric rather than a direct effect on the antibody binding.

**MAb VACV-138** - Binding of VACV-138 to the D8 mutants Arg44Ala/Arg220A was abrogated, since Arg44 is a centrally located contact residue that binds to Asp52H through two salt-bridges. The Lys48Ala/Arg220A mutant only showed slightly increased dissociation, suggesting contact of VACV-138 with Lys48, which lies at the periphery of the epitope. This finding was somewhat surprising, given that Lys48 engages both Thr28H with an H-bond and Asp31H using a salt-bridge. Arg220, which is present in both mutants, is not part of the epitope, and as such the Arg220A single-position mutant did not exhibit reduced binding to VACV-138. The Glu105Ala mutation also demonstrated no effect on binding since it is located at the periphery of the epitope as well (Fig 5).

**MAb VACV-304** - Since VACV-304 binds D8 slightly offset along the CS binding crevice compared to VACV-138, VACV-304 binding to the mutant Arg44Ala/Arg220Ala was not impaired, in contrast to the binding of VACV-138. Arg44 is not a contact residue for VACV-304, although VACV-304 and VACV-138 bind at a similar D8 site. This finding is in contrast to that for VACV-138, for which the binding to the two aforementioned mutants was reversed. Lys48 is a LC contact and appeared important for D8 binding, since it engaged both Asp50L with a salt-bridge and also forms extensive non-polar VdW interactions with Phe31L.

**The D8 crevice has a low- and high-affinity binding region for CS-E** - As VACV-304 binds to one extremity of the CS-E binding crevice, residual binding to the high affinity ligands CS-E was observed consistently (Fig 4). At least half of the binding crevice was not covered upon VACV-304 binding (Fig 7). As CS-A binding was not observed in the presence of VACV-304, we postulate that there is an affinity threshold necessary for binding, which is not reached by CS-A but is by CS-E, correlating with the finding that CS-A is a lower affinity ligand for D8. It should be noted that monomeric D8 does not bind to CS-A but only to CS-E (18). In addition, LA5 binds to the opposite extremity of the binding crevice and fully blocks CS-E binding. Therefore, we speculate that the binding crevice in D8 has a high- and low-affinity CS binding region. A detailed view at the D8/CS-E computationally docked model correlates well with a high-affinity CS binding region located at the C-terminal end of the D8 crevice, which starts with residues Arg220 and His176 and ends in the middle of the crevice around Lys108 and Arg44 (S2 Fig). This region is covered by the antibody LA5 but is accessible when VACV-304 is bound. In this model, more polar interactions with CS-E are predicted in the high affinity region compared to the low affinity region, where only a few interactions are predicted. In conclusion, the low-affinity CS binding region is amenable to protein engineering to increase the specificity and affinity towards CS-E. Such a protein would be useful in generating a highly specific and high-affinity CS-E binding reagent for detecting changes in CS-E expression in various tissues, which would correlate with cancer progression, since CS-E has been identified as a potential biomarker for ovarian cancer (27).

**Immunofluorescence microscopy using D8 tetramers** - As monomeric D8 specifically binds to the glycosaminoglycan CS-E, we asked whether we
could use D8 in microscopy to identify CS-E expression in tissues and generated a D8 tetramer conjugated to Alexa Fluor 488 (AF488). Since the CS-E specific antibody GD3G7 was reported to bind to developing glomeruli of the kidney (27), we assessed the staining of D8 on kidney sections isolated from one week old mice. Glomeruli were stained in green using phalloidin, a toxin with high affinity for F-actin enriched in these structures and D8 control tetramer, which cannot bind to CS-E, in red (Fig 8). While D8 does indeed bind to specific structures in glomeruli, it only stains localized regions, suggesting a highly specific binding pattern. The D8 control tetramer showed no binding to the glomerulus, in contrast to wildtype D8. However, there was a low-grade immunoreactivity of this reagent to adjacent regions. This altered binding behavior likely is the result of the four introduced mutations (Lys41Ala, Arg44Ala, Lys48Ala, Arg220Ala) that greatly reduce the positive charge of D8 and may favor hydrophobic interactions to unknown structures outside the glomerulus.

**DISCUSSION**

Many microbes exist that express glycosaminoglycan (GAG) binding proteins to adhere to their host cells for subsequent cell entry and infection (28,29). These adhesion molecules have evolved to acquire unique binding specificities. Vaccinia virus (VACV) expresses three envelope proteins, H3, A27 and D8 that are glycosaminoglycan adhesion molecules. While H3 and A27 bind to heparin and HS, D8 binds chondroitin sulfate (13-16). Antibodies that target A27, H3 and D8 can block viral adhesion to the host cell and protect from infection, especially if multiple viral envelope proteins are targeted simultaneously (26,30). It is difficult to correlate MAb binding affinity and the ability of a single anti D8 MAb to block viral cell adhesion with neutralization potency, because VACV expresses several antigens that can functionally compensate for each other. This is especially true for the three GAG adhesion molecules A27, H3, and D8. Blocking viral adhesion via one antigen is likely not sufficient to prevent adhesion and subsequent infection. However, many antibodies targeting several different antigens are elicited during the cause of viral infection, and together these antibodies confer protection. As a result, the smallpox vaccine is considered the gold standard of vaccines (31).

Since GAGs are the most abundant heteropolysaccharides in the body and virtually expressed on every cell, they are ideal receptors for viral adhesion. They consist of long unbranched chains with repeating disaccharide units. Most GAGs, including chondroitin sulfate (CS) are sulfated. The precise sulfation of CS is regulated by specific sulfotransferases, leading to the production of different isoforms. All GAGs are attached to specific serine residues of a core-protein to form the proteoglycan (PG) (32). They are ubiquitously expressed on cell surfaces and in extracellular matrices. In addition to providing structural integrity, GAGs participates in various physiological processes such as cell adhesion, cell proliferation, migration, neurite outgrowth, microbial adhesion and infection, inflammation and angiogenesis (7-11). The upregulation of individual GAG-specific sulfotransferases during cellular stress results in the expression of different GAG isoforms and can be a biomarker for cellular changes leading to disease or a method to monitor disease progression. Chondroitin sulfate (GlcAβ1-3GalNAcβ1-4)n is the most abundant GAG and has 3 major isoforms (chondroitin-4-sulfate, CS-A; chondroitin-6-sulfate, CS-C; chondroitin-4,6-O-disulfate, CS-E) (17). The CS sulfotransferase GalNAc4S-6ST, which converts CS-A into CS-E is upregulated in ovarian cancer and results in an enrichment of CS-E on the cancer cell surface, while CS-C and CS-E are enriched in colon carcinomas (33), (34). CS-E has been proposed to be involved in ovarian cancer tumorigenesis as it mediates binding of the vascular endothelial growth factor (VEGF) (27,33). More recently, the malaria protein VAR2CSA was shown to bind a form of CS-A that is both expressed in the placenta and various cancers, especially on cancer-associated proteoglycans including CD44 and CSPG but not on healthy tissues (35). These findings suggest that detection of defined isoforms of CS can be a useful diagnostic in monitoring cancer progression. Several anti CS-specific antibodies exist that target CS-A. Antibodies against different CS isoforms (CS-A, CS-C, and CS-E) exist but often their specificity toward the individual sulfation patterns is limited, or their epitope is rather small (1-2 disaccharides), leading to antibody cross-reactivity with other similar sulfated isoforms (36-39). The CS-E specific antibody GD3G7 also binds moderately to oversulfated dermatan sulfate (DS) from hagfish (IdoA-GalNAc4S6S) and shark skin DS (40). Therefore, the antibody also recognizes IdoA, a component of dermatan sulfate and is not.
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exclusively CS specific. A new CS-E specific antibody has recently been reported that can distinguish between iduronic and glucuronic acid and is, similar to D8, specific for CS (and not DS) (39). However, as the GAGs were mostly obtained commercially and purified from natural source, heterogeneity in the CS sulfation patterns is to be expected, i.e. squid cartilage from Sigma is labeled as containing ~61% CS-E, among other CS structures. An additional limitation is that no structural information of any antibody bound to any CS exists so the antibody recognition of and interaction with a particular GAG molecular species is unknown. Some antibodies, such as the CS-E specific antibody from (36) are very specific but also moderately binds to the CS-E disaccharide. This could lead to off-target binding, e.g the CS-E specific antibody could stain tissues or cellular structures that are enriched for CS-A or CS-C but contain one or two CS-E disaccharides to which it binds, falsely suggesting that this cell expresses largely CS-E. In summary, the different CS binding antibodies have different limitations and a more specific CS binding reagent that recognizes longer stretches, such as minimally a hexamer is needed. This specific binding reagent would allow a more careful distinction of cell surface expressed CS isoforms and their role in disease onset or progression, ultimately leading to better diagnostics needed for biomarker discovery.

Since D8 specifically binds to CS-E using a longitudinal binding crevice capable of binding at least 4 disaccharide units (18), we propose that the D8 protein is a suitable scaffold for the design of a highly specific CS-E binding protein as a potential diagnostic tool to monitor cancer progression that is linked to CS-E upregulation, such as ovarian cancer.

**EXPERIMENTAL PROCEDURES**

**Human antibody generation-** Human anti-D8 MAbs VACV-66, VACV-138, VACV-249 and VACV-304 were generated as reported previously (26). Hybridoma cells secreting VACV-specific MAbs were generated previously (26) and grown in serum-free medium (Gibco). MAbs were purified from culture supernatants using a HiTrap MabSelect Sure column (GE Healthcare). Nucleotide sequences of variable gene segments were determined by Sanger sequencing from cloned cDNA generated by reverse transcription PCR of mRNA, using variable gene-specific primers, as reported previously (41).

**Murine antibody generation-** Murine anti-D8 MAbs JE11, JF11 and LA5, as well as anti-L1 MAbs M12B9 and 39D4, were reported previously (19,20,23).

**Competition-binding ELISA-** The ELISA assay was performed as previously reported (21). Briefly, recombinant D8 (residues 1-263) was coated on 96-well flat bottom plates, washed with PBS + 0.05% Tween 20 and blocked by adding 10% fetal bovine serum. The antibody of interest was added to the D8 bound wells, washed off and the ability of a second antibody to bind simultaneously to D8 was assessed. For this, the second antibody was HRP-conjugated, and its binding to D8 was assessed by measuring optical density at 490 nm. Total cross-blocking results in no signal, while a distinct binding site on D8 would result in maximum signal of the second antibody.

**Antibody-mediated neutralization of virus-** In vitro antibody neutralization was performed using a flow cytometry-based assay, as reported previously (23,26,42). Briefly, Vero E6 cells (ATCC, CRL-1586) were seeded in 96-well plates (1x10^5 cells/well) and allowed to adhere for 5 h. Cells were then infected with 1.25x10^4 PFU of purified VACV-GFP and MAbs (final MAbs concentration of 20 µg/mL) for 12 h at 37°C and 5% CO₂ in 50 µL in the presence (2% final concentration) or absence of sterile baby rabbit complement (CEDARLANE).

**D8 constructs and tetramer generation-** The VACV (Acam2000) D8 ectodomain (residues 1-263), as well as its carbonic anhydrase domain (CAH, residues 1-235), were cloned separately into the pET22b vector to generate hexameric (residues 1-263) or monomeric (residues 1-235) D8 protein forms. For D8 tetramer generation, we added a C-terminal 15 amino acid glycine-serine linker (GGGGS)₃ followed by a biotin protein ligase (birA) tag and a hexa-histidine tag to the CAH construct. A CS-E binding deficient D8 quadruple mutant (K41A, R44A, K48A, R220A) was generated as a specificity control for immunofluorescence experiments. The resulting expression plasmids were transformed into CodonPlus BL21 cells (Agilent), and D8 expression and purification was performed as reported (19). Briefly, expression was induced with 1 mM IPTG at 37 °C for 4 h at a cell density of 0.6 (OD₆₀₀). Cells were pelleted by centrifugation (10
min x 4000 g), resuspended in 50 mM Tris pH8.0, and sheared with a microfluidizer (3 rounds at 1400 bars, Microfluidics). Crude lysate was clarified by centrifugation (1 h x 50,000 g), and D8 was purified using ion metal affinity chromatography (IMAC) on 1 mL HisTrap cartridges (GE Healthcare). After washing the column with 20 cv of 50 mM Tris pH8.0, 300 mM NaCl and 20 mM imidazole, followed by an additional wash step with 20 cv 50 mM imidazole containing wash buffer, D8 was eluted with 50 mM Tris pH 8.0, 300 mM NaCl and 250 mM imidazole. D8 was dialyzed against 10 mM Tris pH 8.0, 200 mM NaCl prior to size-exclusion chromatography (SEC). For tetramer preparation, birA-tagged D8 was expressed and purified in a similar manner and biotinylated using the biotin protein ligase, according to manufacturer’s suggestions (Avidity). Free biotin was removed using 10 kDa MWCO filtration devices, and D8 was tetramerized by incubation with Alexa fluor conjugated streptavidin using molar excess of D8. (5 D8 monomers per 1 streptavidin tetramer). D8 tetramer was purified from excess D8 monomer using SEC.

Glycosaminoglycan microarray assay- CS GAGs enriched in CS-A, CS-C, CS-D, and CS-E (Seikagaku Corp., Tokyo, Japan), dermatan sulfate (DS; Sigma-Aldrich, St. Louis, MO), hyaluronic acid (HA; Sigma-Aldrich, St. Louis, MO), heparin (Hep; Neoparin, Alameda, CA), heparan sulfate (HS; Sigma-Aldrich, St. Louis, MO), or chondroitin sulfate (CS; Sigma-Aldrich, St. Louis, MO) were printed on poly-DL-lysine-coated glass surfaces, as described previously (43,44). Antibodies were pre-incubated at a 10-fold molar excess with hexameric D8 before assessing D8 binding to CS-A and CS-E. Experiments were conducted as previously published (18).

Real-time binding studies using biolayer interferometry (BLI)- Intact IgGs were loaded onto anti-human Fc sensor tips (at a 20 nM concentration) and binding to increasing concentrations of monomeric D8 (0.625 to 40 nM) was measured, as previously reported (18). For qualitative assessment of antibody binding to variant D8 protein, a single antibody was loaded on the sensor tip and binding to wildtype D8 or seven variant D8 proteins was measured simultaneously using a saturating concentration of monomeric D8 (5 µM). Only a complete abrogation of binding was considered as significant, since overall binding amplitudes differed between individual D8 variant proteins.

Fab preparation- Purified IgGs for MAbs VACV-66, VACV-138, and VACV-304 (1 mg/mL) were digested with 2% (w/w) activated papain (Sigma #P3125) for 4 h at 37 ºC in in 100 mM Tris/HCl pH, 1 mM EDTA, 10 mM cysteine. The papain digestion was stopped by adding 20 mM iodoacetamide (IAA). Digestion mixtures were dialyzed against PBS for subsequent protein A purification to remove undigested IgG and Fc. The protein A flow-through containing Fabs was concentrated and purified by size exclusion chromatography (SEC) on a Superdex S200 GL10/300 column (GE Healthcare), using 50 mM Hepes, pH7.5, 150mM NaCl as running buffer.

Fab-D8 complex preparation- 20% molar excess of monomeric D8 protein was used to prepare the D8:Fab complex with Fab made from MAbs VACV-66, VACV-138, or VACV-304. Complexes were prepared at low concentration (~0.2 mg/mL) and incubated on ice for 5 minutes, concentrated, and separated from unbound D8 using SEC on a Superdex 200 100/300 GL column (GE). Fractions corresponding to the D8:Fab complexes (MW~82 kDa) were pooled and concentrated to ~10 mg/mL for subsequent crystallization.

Crystallization and structure determination- Initially, suitable crystallization conditions were determined using a liquid handling robot (Phenix, Art Robbins ltd.) and commercially available screens. Quality diffracting crystals then were grown manually using the sitting drop vapour diffusion method by mixing 0.5 µL protein with 0.5 µL precipitate; 20% PEG 3350, 200 mM sodium malonate pH 5.0 (D8/VACV-66 and D8/VACV-138), or 20% PEG 400, 200 mM sodium citrate tribasic (D8/VACV-304). Crystals were flash-cooled at 100 K in mother liquor containing 20% glycerol. Data collection for the D8/VACV-66 crystal included modification from a regular phi scan (SSRL BL11-1, Kappa =30°; 0.25 oscillations), due to a very long axis (c=444 Å). Diffraction data was integrated using XDS (45). All diffraction data were scaled with Scala and phase determination by molecular replacement (MR) was carried out using Phaser (46). MR template for the Fab VACV-66 was PDB code 4FQQ (highest sequence similarity), separated into variable and constant domains, while PDB 4E9O (D8) was used
as a template for D8. MR search models for VACV-138 and VACV-304 were PDB ID 1RZ7 and PDB ID 3LMJ, respectively. MR solutions first were refined in ccp4 using rigid body refinement, followed by iterative rounds of model building in coot (47) and restrained refinement using TLS refinement in REFMAC5, as part of the CCP4 suite (48-50). Refinement was carried out to a final R_{cryst} and R_{free} of 21.8% and 25.7% (D8/VACV-66), 25.0% and 26.9% (D8/VACV-138), or 24.7% and 29.7% (D8/VACV-304). Data collection and refinement statistics are presented in Table 1.

Structural analyses and presentation- The quality of the models was examined with the program Molprobity (51). Buried surface areas (BSA) and atomic contacts were calculated using PDBePISA (http://www.ebi.ac.uk/msd-srv/prot_int/pistart.html). Figures were prepared using PyMOL.

Accession numbers- The coordinates and structure factors of the D8:Fab complexes have been deposited in the Protein Data Bank (www.rcsb.org) with codes 5USH, 6B9J, and 5USL. The genetic features of the human anti-D8 antibodies were reported previously (26).

Immunofluorescence- One week old mice were sacrificed by CO2 inhalation, followed by immediate decapitation. Kidneys were removed and frozen in Tissue-Tek optimum cutting temperature medium (Sakura Finetek). Eight micron cryostat sections were fixed with methanol-free 4% formaldehyde (EMS Diasum) for 10 minutes at room temperature. Following 2 washes with PBS (Gibco), slides were blocked with 5% normal donkey serum, 0.3% Triton X-100 in PBS for 1 hour and incubated with 156 ng/ml of D8 or control tetramer that is unable to bind to CS-E (D8 coupled to Alexa Fluor 488 streptavidin diluted in 1% BSA, 0.3% Triton X-100 in PBS. Following an overnight incubation at 4C slides were washed with PBS and counterstained with Alexa Fluor 647 phalloidin (Life Technologies, 1:100) and Hoechst (Life Technologies, 10 µg/ml). Slides were washed and mounted with Prolong Gold using #1.5 coverslips. Microscopy was performed using a Zeiss LSM780 confocal laser scanning microscope using 63x 1.4NA objective. Imaging was done using 405, 488 and 633 nm laser lines in a sequential mode to eliminate cross-talk between channels. Acquired Z-stacks were projected to a single plane and contrast was adjusted using Fiji (52). All images were acquired and processed in an identical way. Experiments were repeated 3 times with 2 different batches of tetramers, yielding similar results.

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Conflict of interest: The authors declare that they have no conflict of interest with the contents of this article.

Author contributions: MHM conducted most of the biochemical and structural experiments, analyzed the results and drafted the paper. AS assisted with biochemical and structural experiment. IMG generated human antibodies. GM performed the GAG microarray. ZM performed immunofluorescence microscopy using the D8 tetramers. MH assisted with crystallization experiments, JW generated the D8 tetramers, XM and YX generated the murine antibodies, TK performed antibody crossblocking ELISA and antibody neutralization experiments. TD assisted with data collection, KL conceived and supervised the immunofluorescence microscopy experiment and edited the manuscript. SC supervised and conceived antibody neutralization experiments and edited the manuscript. BP conceived and supervised the antibody crossblocking experiment and edited the manuscript. LCHW conceived and supervised the GAG microarray experiment and edited the paper. JEC conceived and supervised the biochemical and structural experiments, supervised the overall project and wrote the manuscript.
REFERENCES


FOOTNOTES
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The abbreviations used are: MAb, monoclonal antibody; Fab, fragment antigen binding; VACV, vaccinia virus; CS, chondroitin sulfate; GAG, glycosaminoglycan; HC, heavy chain; LC, light chain
TABLE 1. Data collection and refinement statistics

<table>
<thead>
<tr>
<th>Data collection</th>
<th>D8/VACV-66</th>
<th>D8/VACV-138</th>
<th>D8/VACV-304</th>
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<td>90, 90, 90</td>
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<td></td>
<td></td>
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<tr>
<td>No. of unique reflections</td>
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<td>R_pim (%)</td>
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<td>11.2 [57.2]</td>
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<td>Multiplicity</td>
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<td>6.2 [6.4]</td>
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<td>8.3 [1.4]</td>
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<td>99.4 [99.8]</td>
<td>94.8 [97.1]</td>
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Refinement statistics

| No. atoms | 10,202 | 9,990 | 10,022 |
| Protein   | 10,028 | 9,923 | 10,022 |
| Waters    | 174    | 48    | 0      |
| Ramachandran plot (%) |          |        |        |
| Favored   | 97.0   | 94.7  | 95.2   |
| Allowed   | 99.6   | 99.8  | 99.8   |
| R.m.s. deviations |          |        |        |
| Bonds (Å) | 0.007  | 0.007 | 0.004  |
| Angles (°) | 1.23   | 1.19  | 0.85   |
| B-factors (Å²) |        |        |        |
| Protein   | 52.0   | 50.5  | 56.6   |
| Waters    | 44.7   | 63.3  | /      |
| R factor (%) | 21.8  | 24.1  | 24.7   |
| R_free (%) | 25.7   | 26.8  | 29.7   |

FIGURE LEGENDS

FIGURE 1. Cross-Blocking of human and mouse D8 Abs reveal a novel epitope. (A) Human antibody cross-blocking reveals two specificity groups (columns labeled 1 and 2). (B) Overlap between mouse and human MAbs identify a separate epitope for VACV-66. Black cells correspond to Abs that fully block D8 binding of each other and are thus assumed to share overlapping epitopes. Empty cells correspond to Abs that did not affect the D8 binding of each other. Relative strength of cross-blocking is indicated in shades of grey, since epitopes may overlap partially and allow simultaneous binding of both antibodies. Columns 1-4 indicate the four different specificity groups.

FIGURE 2. Human anti-D8 MAbs are moderately neutralizing. VACV IMV neutralization activity of purified human (VACV-249, VACV-304, and VACV-66) and mouse (JE11, JF11, and LA5) anti-D8 MAbs in the absence (black bars) or presence (red bars) of complement. Murine anti-L1 MAb (M12B9) was used as positive control and anti-L1 MAb 39D4 as control for a weakly neutralizing MAb.

FIGURE 3. Real-time binding kinetics using biolayer interferometry. MAbs were immobilized on anti-human Fc capture antibody sensors and the 1:1 binding of each Fab to monomeric D8 was measured by
dipping into increasing concentrations (colored curves) of D8. Kinetic measurements were unaffected by antibody avidity. All antibodies bind with high affinity.

FIGURE 4. Interference of D8 binding to CS-A or CS-E by human MAbs. Hexameric D8 was incubated with MAbs VACV-66, VACV-138, VACV-304, or alone prior to binding to GAG microarrays. Antibody blocking was assessed by comparing the amount of D8 and D8-MAb complex bound to the microarray. Blocking experiments were performed in triplicate (+/- SEM, error bars). While VACV-66 does not block D8 binding to CS-A or CS-E, VACV-138 blocks binding to both. VACV-304 fully blocks binding to CS-A while slight binding to CS-E was observed (~25% of maximum binding).

FIGURE 5. Structural overview of human MAb-D8 complexes. (A) Binding of VACV-66, VACV-138, or VACV-304 to D8. (B) MAb footprint on D8 is colored by chain on the D8 protein otherwise shown in grey. Heavy chain footprint is shown in orange and light chain in green. The CS-binding site is indicated with a black dot (A) and a yellow line (B). Both VACV-138 and VACV-304 bind above the CS binding crevice, while VACV-66 binds on the side of D8 away from the crevice.

FIGURE 6. Interactions between MAbs and D8. Binding of VACV-66 (A), VACV-138 (B), or VACV-304 (C) to D8. D8 is shown in grey and the MAb CDRs are colored as follows: L1 (green), L2 (light green), L3 (dark green), H1 (yellow), H2 (cyan), and H3 (orange). Potential H-bonds are indicated with blue dotted lines.

FIGURE 7. Footprint of CS-blocking MAbs on D8. Schematic representation of MAb binding sites (circles) and illustration of a high- (H) and low- (L) affinity binding region for CS within the central binding crevice (blue rectangle) of D8.

FIGURE 8. D8 tetramers specifically detect chondroitin sulfate E in tissues. Kidney tissue samples were flash-frozen, sectioned, fixed in 4% PFA and reacted with 156 ng/ml of either D8 or D8 control tetramers coupled to Alexa Fluor 488. Samples were counterstained with phalloidin to reveal dense F-actin in the glomerulus and with the DNA stain Hoechst. The D8 tetramer immunoreactivity was localized within the glomeruli. Negative control with the mutated D8 tetramer revealed absence of binding in the glomerulus but low-grade immunoreactivity in adjacent regions. Bar = 20 µm.
FIGURES

Figure 1

Figure 2
Figure 3

<table>
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<th>$K_D$ (nM)</th>
<th>$k_{on}$ (1/Ms)</th>
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Figure 4
Figure 5
Human antibody responses to vaccinia virus D8

Figure 6

Figure 7
Figure 8
Structure-function characterization of three human antibodies targeting the vaccinia virus adhesion molecule D8

Michael H Matho, Andrew Schlossman, Iuliia M Gilchuk, Greg Miller, Zbigniew Mikulski, Matthias Hupfer, Jing Wang, Aruna Bitra, Xianghzi Meng, Yan Xiang, Tom Kaeever, Tzanko Doukov, Klaus Ley, Shane Crotty, Bjoern Peters, Linda C Hsieh-Wilson, James E Crowe Jr. and Dirk M Zajonc

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