There is strong interest in the production of bispecific monoclonal antibodies that can simultaneously bind two distinct targets or epitopes to achieve novel mechanisms of action and efficacy. Regeneron’s bispecific technology, based upon a standard IgG, consists of a heterodimer of two different heavy chains, and a common light chain. Coexpression of two heavy chains leads to the formation of two parental IgG impurities, the removal of which is facilitated by a dipeptide substitution in the Fc portion of one of the heavy chains that ablates Fc Protein A binding. Therefore, the affinity capture (Protein A) step of the purification process must perform both bulk capture and high resolution of these mAb impurities, a task current commercially available resins are not designed for. Resolution can be further impaired by the ability of Protein A to bind some antibodies in the variable region of the heavy chain (VH). This article details development of a novel Protein A resin. This resin combines an alkali stable ligand with a base matrix exhibiting excellent mass transfer properties to allow high capacity single step capture and resolution of bispecific antibodies (bsAbs) with high yields. The developed resin, named MabSelect SuRe™ pcc, is implemented in GMP production processes for several bsAbs. © 2018 The Authors Biotechnology Progress published by Wiley Periodicals, Inc. on behalf of American Institute of Chemical Engineers

Keywords: bispecific antibody, protein A chromatography, MabSelect SuRe™, affinity chromatography, z-domain

Introduction

Bispecific antibodies (bsAbs) are antibody-derived proteins with the ability to bind to two different epitopes on the same or different antigens. As such, they combine specificities of two antibodies: an attractive therapeutic concept, especially in the fields of immuno-oncology and immune disease. A variety of novel molecular mechanisms have been proposed, including: recruitment of immune cells to target cells, simultaneous inhibition of two cell surface receptors, simultaneous blocking of two ligands, and receptor cross-linking. The combination of two binding specificities on a single molecule could also be attractive without a mechanistic necessity, as it would avoid the complicated and costly development of combination therapies.

There is no one common molecular structure of a bsAb. On the contrary, over the past two decades over 60 different formats of bsAbs have been proposed, with varying size, valency, flexibility, half-life, biodistribution, ease of manufacture, options for Fc-mediated effector functions, and immunogenic potential. Generally, formats can be divided into two major classes, those bearing an Fc region and those lacking an Fc region. Bispecifics from both classes have achieved marketing approval. Catumaxomab, an Fc-containing bsAb for the treatment of patients with
malignant ascites was approved in 2009. Blinatumomomab was approved in 2014 for the treatment of Philadelphia chromosome-negative precursor B cell ALL. Blinatumomomab, a bispecific T-cell engager (BiTE) is an example of a format lacking an Fc region. It is based on the two single-chain variable fragments joined via a flexible linker.

Regeneron’s bsAb format was designed as a fully human IgG antibody. It consists of a heterodimer of two different heavy chains, which confer binding specificity, and a common light chain. Therefore, light-heavy chain pairing issues are addressed by choosing heavy chains that can retain their different specificities but employ identical light chains. Coexpression of two different heavy chains does however lead to the formation of two parental IgG impurities. The removal of these impurities is facilitated by a dipeptide substitution in the Fc portion of one of the heavy chains (then named Fc*) that ablates Protein A binding (Figure 1). This allows the isolation of the bispecific dimer via selective elution from a Protein A column. The Fc*-containing parental will flow through the column, while the bispecific can be separated from the Protein A binding parental by using the decreased avidity with which the bispecific binds to Protein A. The two amino acid residues substituted are taken from the equivalent region in the IgG3 isotype; therefore, no new non-human potentially immunogenic sites are introduced.

While the exact residues replaced can vary depending on the heavy chains derived from the V H3 gene family have been shown to exhibit the formation of either heavy chains (then named Fc*) that ablates Protein A binding (Figure 1). This allows the isolation of the bispecific dimer via selective elution from a Protein A column. The Fc*-containing parental will flow through the column, while the bispecific can be separated from the Protein A binding parental by using the decreased avidity with which the bispecific binds to Protein A. The two amino acid residues substituted are taken from the equivalent region in the IgG3 isotype; therefore, no new non-human potentially immunogenic sites are introduced. While the exact residues replaced can vary depending on the IgG isotype and individual antibody, in all cases the substitutions, replace the histidine residue critical for Fc-Protein A binding.

The drawback of this approach is that, assuming that Fc* and the unmodified heavy chain (Fc) are produced in equal amounts and formation of the bispecific is thermodynamically equivalent to formation of the FcFc and Fc*Fc* parental impurities, the theoretical expression ratio will be 1:2:1 (FcFc:bsAb:Fc*Fc*). This means cell culture productivity is intrinsically half that of a standard monoclonal antibody. The advantage over other Fc-containing bispecific formats is that the fully human nature of the antibody minimizes immunogenicity while maximizing molecule stability and pharmacokinetics. An additional advantage over similar full length antibody formats, such as the knobs-into-holes design, is that if parental antibodies or other variants do form with these techniques, their removal can be highly challenging. In contrast, the star substitution eases purification development by providing a well characterized, platformable mechanism of separation of parental antibody impurities.

The classical interaction between Staphylococcal Protein A (SpA) and the Fc region has been studied in detail using X-ray crystallography and exploited by the Fc* substitution, which ablates the interaction. However, a subset of antibodies bind Protein A in the variable region of the heavy chain (Figure 2). In particular, IgGs that contain heavy chains derived from the V H3 gene family have been shown to exhibit this behavior. Nearly half of human V H3 germline genes belong to this family. Because of this, on SpA-based resins, many bsAbs exhibit poor resolution of the two binding species as well as retention of the non-binding Fc*Fc* parental antibody. This is because V H binding reduces the avidity difference between the bispecific and the FcFc parental, and Fc*Fc* parental is also retained by V H binding. This phenomenon can be resolved through the use of the alkali stabilized MabSelect SuRe ligand. This is because while binding studies between SpA and antibodies have shown that all five domains of SpA (E, D, A, B, and C) bind antibodies via the Fc region, only domains D and E exhibit significant Fab binding.

As the MabSelect SuRe ligand is a tetramer of the Z-domain, a protein-engineered version of the native SpA B-domain, it would be expected to lack significant Fab binding, and this has been demonstrated in multiple studies.

This finding is also supported by previous work where it was shown that use of MabSelect SuRe allows for flow-through removal of the Fc*Fc* parental antibody during column loading. However, commercially available resins functionalized with the MabSelect SuRe ligand did not provide adequate separation of the bispecific from the FcFc parental. This necessitated insertion of another Protein A step after primary capture to remove this SpA binding impurity. Resolution was achieved via the use of a commercially available resin (POROS™ MabCapture™ A, Thermo Scientific) exhibiting superior mass transfer and resolution characteristics based on (i) large through pores that give an element of perfusive flow in the base matrix, (ii) a large mean pore diameter of 1,100 Angstroms, and (iii) a smaller average particle size of 45 μm (c.f. to MabSelect SuRe resin average particle size of 85 μm).

Although this resin used SpA, the Fab-binding was minimized via the use of chaotropic salts in the elution
buffer. A salt tolerant multimodal resin was then used in a positive mode as both a polishing step, and to achieve buffer exchange. The resultant process was broadly applicable, with bispecific purities (Eq. 1) >95% and a robust operating space. However, the requirement of an extra chromatographic step was compounded by the increased cost of Protein A resins compared to nonproteinaceous ligands.\textsuperscript{34,35} This led to the desire to create a resin that could combine the mass transfer benefits of the POROS MabCapture A base matrix with the MabSelect SuRe ligand to allow single step affinity chromatography for all star substitution-based bsAbs.

### Materials and Methods

**Materials**

All bsAbs (C, D, and F) used in this study were expressed in CHO cells and produced at Regeneron Pharmaceuticals. Commercially available chromatographic resins used were obtained from their manufacturers: MabSelect SuRe and MabSelect SuRe pcc (GE Healthcare), and POROS MabCapture A (Thermo Scientific). All chemicals used were supplied by J.T. Baker. Developmental resins were supplied by GE Healthcare Custom Design Resin Group.

**Equipment**

Laboratory scale chromatographic separations were performed using an AKTA\textsuperscript{TM} avant 25 chromatographic system (GE Healthcare) and 1.0 cm inner diameter (ID) Omnitit\textsuperscript{®} Benchmark chromatography columns (Omnitit Ltd.). Pilot- and production-scale chromatography was conducted on a AKTAPROCESS\textsuperscript{™} chromatography skid using either 20 cm ID INDEXTM chromatography columns or ReadyToProcess\textsuperscript{™} MabSelect SuRe pcc 20 columns (cat#29–1875-64; GE Healthcare), with the exception of pilot scale pressure-flow data that used an Axichrom\textsuperscript{™} 200 (GE Healthcare). UPLC analysis leveraged an ACQUITY\textsuperscript{®} UPLC system from Waters Corporation. All material was derived from cell culture using either a 500 L or 2,000 L single use bioreactor (Thermo Scientific).

### Pilot-scale preparation of affinity capture pools for chromatographic development

When clarified cell culture fluid was not used directly, load material for affinity resolving development was produced by affinity capture chromatography using 20 ± 1 cm bed height MabSelect SuRe columns to remove the Fc*Fc* parental.

After equilibration with 2 CV of 20 mM sodium phosphate pH 7.2, the columns were loaded to 10–40 g FcFc + bsAb per L of packed bed with clarified cell culture fluid. Columns were washed and all bound protein eluted in 3 CVs using a proprietary buffer system at pH 3.0. After elution a 2 CV column strip is performed to regenerate the column. The entire elution peak containing FcFc and bispecific was collected and neutralized to pH 7.0 ± 0.2 with 2 M Tris base.

### Resin prototype screening using gradient elution methodology

Chromatography resins screened were packed to 20 ± 1.0 cm bed height in 1.0 cm ID columns. Columns were equilibrated with 2 CVs of 20 mM sodium phosphate, pH 7.2 before load application of bsAb C to 10 g FcFc + bsAb/L. Load material had been previously subjected to affinity capture as described above, and was 77% bispecific purity as per Eq. 1. Following loading, columns were washed with a proprietary wash buffer system and eluted in 40 mM acetate, 500 mM calcium chloride, pH 6–3 over 20 CV. All steps were performed at a 4 min residence time. UNICORN\textsuperscript{™} 6.1 software (GE Healthcare) was used for chromatographic analysis including calculation of chromatographic peak resolution (Rs see Eq. 2) assuming Gaussian peaks using the width at half height method (resolution algorithm 3).

### Affinity resolving chromatography using isocratic elution approach

All affinity resolving chromatography runs were performed using 20 ± 1 cm bed height columns. Columns were equilibrated with 2 CVs of 20 mM sodium phosphate, pH 7.2 before load application. Following loading, columns were washed with a proprietary wash buffer system and eluted using 40 mM acetate, 500 mM calcium chloride at a specified pH and elution volume. Pool collection began 0.5 CV into the elution volume and was terminated at the end of the elution step. Following elution, columns were stripped with 2 CV of buffer. Column clean in place (CIP) was performed with 0.5 N NaOH at a 15 min contact time. Statistical design of experiments (DoE), analysis, and modeling were performed using JMP\textsuperscript{®} 11.1.1 (SAS Institute Inc). Small-scale pools were neutralized to pH 7–8 using 2 M Tris base. Pilot- and production-scale pools were buffer exchanged by standard flat sheet ultrafiltration techniques into low conductivity buffer, subjected to viral inactivation by low pH hold, neutralized, and filtered before process- and product-related impurity measurement. Yield was calculated over the affinity resolving step only.

### Process- and product-related impurity measurement

Soluble aggregate quantification was achieved by two ACQUITY UPLC PrST SEC Columns, 200 Å, 1.7 μm, 4.6 × 150 mm\textsuperscript{-2} cat#186005225 in series in a 10 mM sodium phosphate, 500 mM sodium chloride, pH 7.0 mobile phase. Bispecific purity was measured using three prepacked
POROS A 20 μm columns (2.1 × 30 mm², 0.1 mL) cat#2-1001-00 in series and an isocratic elution buffer system. Bispecific and FcFc titers were measured using a POROS A 20 μm column (2.1 × 30 mm², 0.1 mL) cat#2–1001-00, and Fc*Fc* titers were measured by loading the Protein A flow-through over a POROS G 20 μm column (2.1 × 30 mm², 0.1 mL) cat#2-1002-00 (Thermo Scientific). HCP quantification was performed using a commercially available ELISA kit cat#F550 (Cygnus Technologies). Leached affinity ligand was quantified using the commercially available ELISA kit cat#F400 (Cygnus Technologies).

Results and Discussion

Base matrix selection and screening

The aim of this study was to couple the MabSelect SuRe ligand to a base matrix capable of resolving the bispecific from the FcFc parental, while maintaining acceptable flow properties for large-scale commercial protein production for pharmaceutical applications. Three agarose-based prototype resins with a range of mean particle sizes and mean pore diameters were provided (Table 1). For base matrix evaluation, ligand density was fixed at an equivalent level to the commercially available MabSelect SuRe. MabSelect SuRe resin was used as a control.

Bispecific antibody C (bsAb C) was used as a model bsAb for resin screening. Table 2 details attributes of this and other bsAbs used in this study. Resins were loaded to 10 g FcFc + bsAb/L resin with load bispecific purity (Eq. 1) of 77%. For the purposes of screening the prototype resins, elution was achieved with a pH 6-3 pH gradient over 20 column volumes (CV) rather than by isocratic step elution. This allowed measurement of the chromatographic peak resolution obtained between bsAb C and FcFc peaks as calculated by Eq. 2. Resolutions varied from 1.06 to 1.66 between the four resins studied (Figure 3).

All three prototype resins demonstrated improved resolution (Rs = 1.07 – 1.66) from MabSelect SuRe control (Rs = 1.06). As might be expected for separation of a large protein such as an antibody, resolution seems to be related to both average particle size and mean pore diameter. Specifically, the Capto ImpAct base matrix combined the second smallest (50 μm) particle size with a larger mean pore diameter and was found to provide the best resolution of tested resins. Based on these data, Capto ImpAct was chosen as the base matrix for the new resin.

Influence of ligand density upon dynamic binding capacity and resolution

The Capto ImpAct base matrix was functionalized with MabSelect SuRe ligand at a range of ligand densities (2–11 mg ligand per mL of resin) and the dynamic binding capacity (g FcFc + bsAb/L) at 5% breakthrough of bsAb (DBC5%) determined at a four minute residence time with bsAb C (see Figure 4). It should be noted the bsAb breaks through before the FcFc parental via a displacement mechanism likely caused by the avidity difference between the two molecules (data not shown). A higher ligand density on the chromatography medium directly translates into an increased binding capacity, but diminishing results are observed at higher ligand densities. Increasing Protein A ligand density leading to a situation of diminishing returns for binding capacity due to increased steric hindrance on the ligand surface is studied in a previous work.34

Following this initial univariate screening experiment, a 31-run D-optimal custom DoE study was performed with the aim of determining an optimal ligand density that could give the most robust separation between the bispecific and the parental antibodies (FcFc and Fc*Fc*). An isocratic elution was chosen to simplify possible plant fit and technology transfer while also reducing buffer consumption and processing time. Other factors known to be critical to the bispecific separation from a previous study (eluion pH and length [CV], and column loading) were also included as factors (Table 3).31 This allowed possible interactions between ligand density and other process parameters to be determined in order to make the most informed ligand density choice. Residence time for all chromatographic steps, including loading, wash, and elution, was held constant at 4 min (300 cm/h) as this was previously observed not to be a significant factor for bispecific yield or bispecific purity with isocratic elution. Yield and pool bispecific purity were studied as responses. The load material for the study was bsAb C that had been previously subjected to standard positive mode affinity chromatography using MabSelect SuRe resin. This resulted in a load bispecific purity for the DoE of 77%, with all parental antibody present being FcFc.

Good models were obtained for both bispecific yield (R² = 0.97, R²_adj = 0.96, RMSE = 4.2%, P < 0.001) and bispecific purity (R² = 0.97, R²_adj = 0.97, RMSE = 1.4%, P < 0.001) using a standard least squares fit algorithm. Using a 95% confidence interval, ligand density was found to be a significant parameter for both yield and purity (P < 0.0001 for both responses), and several interactions and quadratic effects were also identified (Table 4).

The model predicted that attainable bispecific purities are optimized at increased ligand densities (Figure 5). A recent work observed competitive binding of antibody species on ProA resins despite the tight binding affinity.36 The speed of this displacement was shown to vary between different commercial SpA based resins, attributed to ligand attachment chemistry. We believe this competitive binding mechanism may explain this increase in purity with higher ligand densities. A higher ligand density may result in more frequent multiple binding events of the same antibody per ligand for the FcFc parental, which would not be possible with the bispecific because one of the two Fc binding sites has been ablated. Thus, a higher ligand density enhances the mechanism of separation via facilitating multiple binding to the FcFc parental.

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**Table 1. Properties of Chromatographic Base Matrices Studied**

| Base Matrix               | Average Particle Size (d50V) | Relative Mean Pore Diameter |
|--------------------------|-----------------------------|-----------------------------|
| MabSelect SuRe (control) | 85                          | Large                       |
| MabSelect Xtra™           | 75                          | Extra large                 |
| Capto ImpRes             | 40                          | Medium                      |
| Capto ImpAct             | 50                          | Large                       |

**Table 2. Attributes of the bsAbs Studied**

| BsAb | Isoelectric Point | Subtype | Member of V γ3 Gene Family? |
|------|-------------------|---------|-----------------------------|
| C    | 9.2               | IgG1    | Y                           |
| D    | 8.8               | IgG4    | Y                           |
| F    | 9.3               | IgG4    | Y                           |

DBC5%
Optimizing the model for both yield and purity suggested conditions predicted to give 100% purity with >99% bispecific yield, as well as the highest DBC 5%, at the highest ligand density studied (11 mg/mL). Therefore, a MabSelect SuRe ligand density of 11 mg/mL was targeted for the new resin, using the Capto ImpAct base matrix, and this new resin was named MabSelect SuRe pcc.

Sweet spot analysis was conducted using the models for bispecific yield and purity obtained. CaCl₂ concentration in the elution buffer was held constant at 500 mM and elution volume fixed at 4 CVs. As shown in (Figure 6), the desirable operating window, or sweet spot, is indicated by the white area when imposing constraints of bispecific yield >80% (blue contour lines) and bispecific purities >95% (red contour lines).

In a previous work, similar studies were performed to evaluate both MabSelect SuRe and POROS MabCapture A resins for ability to separate bsAb C from the parental antibodies (FcFc and Fc*Fc*) in Protein A pool (i.e., affinity resolving chromatography step). Here, load material had also been previously subjected to standard positive mode affinity chromatography using MabSelect SuRe resin, resulting in 64% bspecific purity. The MabSelect SuRe pcc resin seems to exhibit a larger sweet spot, indicating increased process robustness, with the same bsAb when compared with the two resins used in the previous study. Further data

Table 3. Factors Studied using Custom D-Optimal DoEs Study to Determine Optimal Ligand Density for Prototype Resin

| Factor                  | Range Studied          | Levels |
|-------------------------|------------------------|--------|
| Elution pH              | 4.3–4.9                | 3      |
| Loading                 | 50–100% of determined  | 3      |
|                         | DBC for ligand density |        |
| Ligand Density          | 2–11 mg/mL             | 4      |
| Step Elution Length     | 4–6 CV                 | 3      |
gained directly comparing MabSelect SuRe pcc to MabSelect SuRe and POROS MabCapture A resin using other Regeneron bsAbs further supports this claim (data not shown). Moreover, the use of the MabSelect SuRe ligand, as compared with the SpA used by POROS MabCapture, alleviates the need for a prior affinity capture step to resolve the Fc*Fc* parental antibodies (FcFc and Fc*Fc*) decreasing cost of goods and increasing productivity of the purification process.

As studied in the previous work, the unit operation is sensitive to column loading and therefore also the bispecific purity of the load material.31 Thus, column loading, and the effect of resin cycling upon resin degradation, need to be carefully controlled as part of the manufacturing strategy. Furthermore, the reproducibility of the upstream process to provide material of consistent bispecific purity must be assessed as part of the development strategy.

**Alkaline stability and pressure flow characteristics**

The MabSelect SuRe ligand was originally developed to withstand stronger alkaline conditions than SpA, thereby facilitating CIP protocols with cleaning agents such as sodium hydroxide.37 As the developed MabSelect SuRe pcc resin utilizes this affinity ligand, it was expected to demonstrate similar alkaline stability to other commercially available resins produced with this ligand (e.g., MabSelect SuRe and MabSelect SuRe LX, [GE Healthcare]).

Alkaline stability was assessed at small-scale (20 cm BH, 1.0 cm I.D.). Dynamic binding capacity was measured at a six minute residence time using bsAb D before (naive resin) and after soaking in 0.1–0.5 M sodium hydroxide for up to 12 h. MabSelect SuRe pcc was found to retain >95% binding capacity after a 12 h exposure to up to 0.3 N sodium hydroxide. Furthermore, 90% binding capacity was retained with 12 h exposure of up to 0.5 M sodium hydroxide (Figure 7). Assuming a 15 min exposure time per CIP step, with CIP performed every four cycles, 12 h would correspond to ~200 production cycles.

Pressure flow characteristics of the new resin were assessed at small-scale (20 cm BH, 1.0 cm I.D.). Dynamic binding capacity was measured at a six minute residence time using bsAb D before (naive resin) and after soaking in 0.1–0.5 M sodium hydroxide for up to 12 h. MabSelect SuRe pcc was found to retain >95% binding capacity after a 12 h exposure to up to 0.3 N sodium hydroxide. Furthermore, ~90% binding capacity was retained with 12 h exposure of up to 0.5 M sodium hydroxide (Figure 7). Assuming a 15 min exposure time per CIP step, with CIP performed every four cycles, 12 h would correspond to ~200 production cycles.

Pressure flow characteristics of the new resin were assessed at small-scale (20 cm BH, 1.0 cm I.D.) and production-scale (20.0 cm BH, 35.7 cm I.D.). These column diameters were deliberately chosen for evaluation as more representative of production batches, which would require larger columns with less wall support.38 Pressure drop over packed columns was measured at a range of linear velocities at 22°C using water as the mobile phase. The nonlinear profile observed is characteristic of compressible media (Figure 8). Based on these data a maximum linear velocity of 200 cm/h was recommended with 20 cm bed height columns to avoid pressure limitations with scale-up.
Process performance and scale-up

The performance of MabSelect SuRe pcc was assessed in the context of a complete downstream process for clinical manufacturing. As part of determining operating conditions for this process, a custom D-optimal DoE (N = 12) was performed using a small-scale column (20 cm BH, 1.0 cm I.D.) and clarified cell culture fluid containing bsAb F, a bsAbs, of the IgG4 subtype (Table 2). Elution pH (range of 0.6 pH units), elution length (4–6 CV), and column loading (35–65 g FcFc1bsAb/L resin), and all primary interactions and quadratic effects were considered as factors, based on the prior study (Table 4). It should be noted that 4–6 CV are used for elution to maximize yield, as the use of the higher elution pH to allow selective elution of the bispecific causes elution peak broadening. Residence time was held constant for all chromatographic steps at 6 min, the maximal recommended flow at production-scale due to loss of wall support in larger columns. Good models were obtained for both responses of yield (R² = 0.99, R²adj = 0.97, RMSE = 4.4%, P < 0.001) and bispecific purity (R² = 0.98, R²adj = 0.97, RMSE = 0.62%, P < 0.001) using a least squares fit algorithm. Using the model, a sweet spot analysis was performed, with the desirable area defined as >80% yield and >95% bispecific purity (%; Figure 9). Based on this and further analysis of the predictive models, isocratic elution over 5 CV at pH 4.4 ± 0.1 was shown to predict robust operation,
with >95% bispecific purity and >80% yields over a wide range of column loadings (35–65 g/L) throughout the entire pH range.

The purification scheme developed for bsAb F was scaled up and process performance and product quality analyzed at pilot-scale (500 L bioreactor) and during GMP production (2 kL bioreactor, N = 2). Load material was cell culture fluid clarified by direct depth filtration and all runs were performed at a constant residence time of 6 min. The results are shown in Table 5 and show a high level of reproducibility across scale and site. The goals of >80% yield and >95% bispecific purity were met for both scales of production. High molecular weight (HMW) species, leached Protein A bispecific purity were met for both scales of production. The authors thank Michelle LaFond, Mark Czeterko, Anthony DeBiase, Shey Ng, and Matthew Klaas’ group for carrying out antibody productions, Benjamin Adams for column packing, and Michael Perrone, Audrey Rodriguez, and Kathir Muthusamy for impurity quantification. They acknowledge Scott Staton for invaluable help with figures.

### Conclusions

A previous study detailed an approach for purification of a fully human bsAbs format. This process was limited by the need for two Protein A resins when applied to bsAbs of the Vh3 gene family. A novel Protein A resin, MabSelect SuRe pcc, has been developed that combines an alkali stabilized, non Vh-region binding ligand with a base matrix exhibiting excellent mass transfer properties. Thus, high capacity single step capture and resolution of native format bsAbs with high yield is achieved, and demonstrated at pilot- and production-scale. This allows the consolidation of the two Protein A columns used in the prior approach, increasing productivity while decreasing cost of goods. A downstream processing platform has therefore been established to facilitate the manufacturing of fully human bsAbs.

Although MabSelect SuRe pcc was developed for the purification of bsAbs, the high DBC (>60 g/L) at moderate residence times, and the alkaline stability (0.5 N NaOH) compare favorably with other commercially available Protein A resins. The resin may be especially appropriate for use in continuous capture by periodic counter-current chromatography (PCC). The ability of PCC to use shorter column bed heights could allow use of shorter residence times without pressure restrictions, important because of the relatively small average particle size (50 μm) of the MabSelect SuRe pcc resin.

### Conflicts of Interest

The authors are currently or have been employees of either Regeneron Pharmaceuticals Inc. or GE Healthcare.

### Acknowledgments

The authors thank Michelle LaFond, Mark Czeterko, Anthony DeBiase, Shey Ng, and Matthew Klaas’ group for carrying out antibody productions, Benjamin Adams for column packing, and Michael Perrone, Audrey Rodriguez, and Kathir Muthusamy for impurity quantification. They acknowledge Scott Staton for invaluable help with figures.

### Tables

**Table 5. Quantitative Purification Performance of Single Step Affinity Resolving Chromatography using MabSelect SuRe pcc at Pilot- and Production-Scale**

| Parameter                        | Pilot Scale | Production Scale | Production Scale |
|----------------------------------|-------------|------------------|------------------|
|                                  | Batch 1     | Batch 2          | Batch 2          |
| Average Protein A (ppm)          | 2.9         | 0.8              | 0.5              |
| Output                           | 98.7        | 98.1             | 98.5             |
| Final Bispecific Purity (%)      | 92.3        | 92.4             | 83.0             |
| Yield (%)                        | 113         | 229              | 210              |
| HCP (ppm)                        | 6.0         | 4.3              | 5.2              |
| HMW (%)                          | 60          | 56               | 53               |
| Column Loading (g FcFc + bsAb/L resin) | 20         | 20               | 20               |
| Bioreactor Size (L)              | 500         | 2,000            | 2,000            |
| MabSelect SuRe pcc CV (L)        | 6.3         | 20               | 20               |
| Column Loading                   | 6.3         | 20               | 20               |

### Equations

**Equation 1:** Definition of bispecific purity

$$\text{Bispecific Purity} = \frac{[\text{Bispecific}]}{([\text{Bispecific}]+[\text{FcFc}]+[\text{Fc}+\text{Fc*}])} \times 100$$

**Equation 2:** Equation used to determine peak resolution (Rs) using peak widths at half height ($W_{1/2}$) and their retention times ($t_R$)

$$Rs = 1.8 \left( \frac{t_{R2} - t_{R1}}{W_{1/21} + W_{1/22}} \right)$$

### Notation

- BH = bed height
- bsAb = bispecific antibody
- CIP = clean in place
- CV = column volume
- DBC = dynamic binding capacity
- DoE = statistical design of experiments
- Fc*Fc* = parental antibody containing star substitution on both heavy chains
- FcFc = parental antibody containing native heavy chains
- FcFc* = bispecific antibody containing a single chain with the star substitution
- HCP = CHO host cell protein
- ID = inner diameter
- Rs = chromatographic peak resolution
- SpA = Staphylococcal Protein A

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