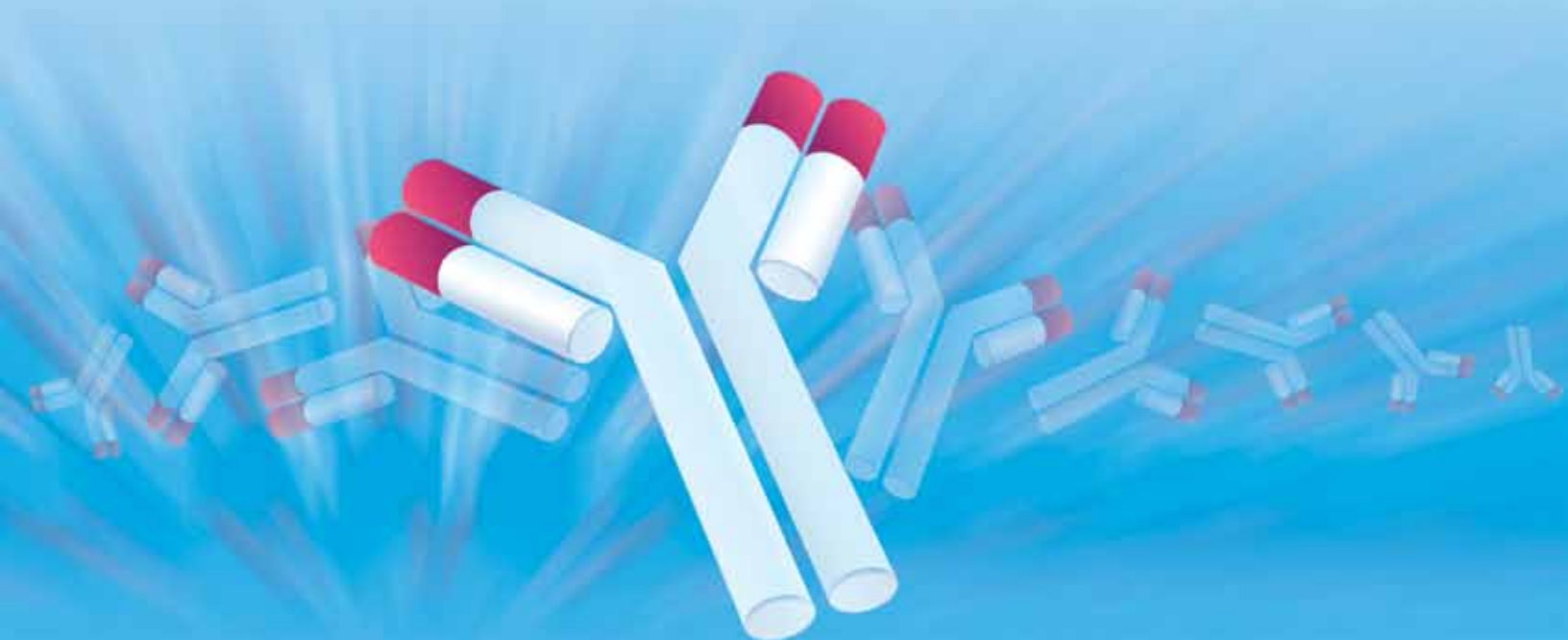


# Monoclonal Antibody Characterization

Achieving Higher Throughput and Productivity



Now sold under the  
Thermo Scientific brand

**Thermo**  
SCIENTIFIC



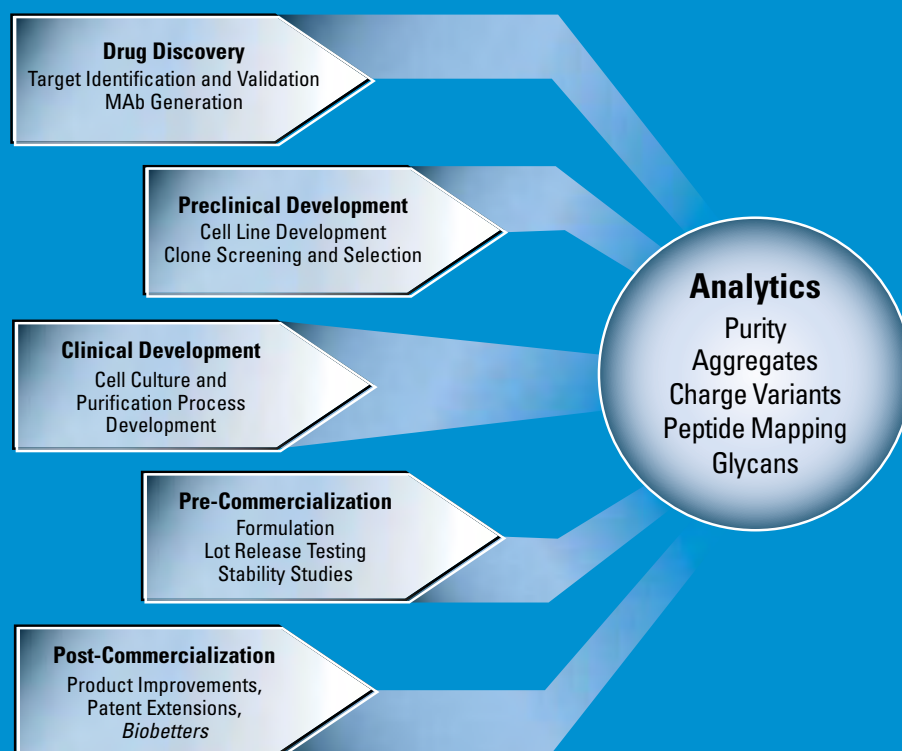
Passion. Power. Productivity.

# Dionex Solutions to Accelerate Monoclonal Antibody R&D and Characterization

## The throughput and productivity challenge

- Increasing number of MAb candidates entering the clinical pipeline
- Advances in automation in upstream processes like cell culture and purification process development, and formulation screening
- Strict Quality-by-Design (QbD) guideline requiring enhanced antibody characterization

All result in a large increase in sample requests for characterization.



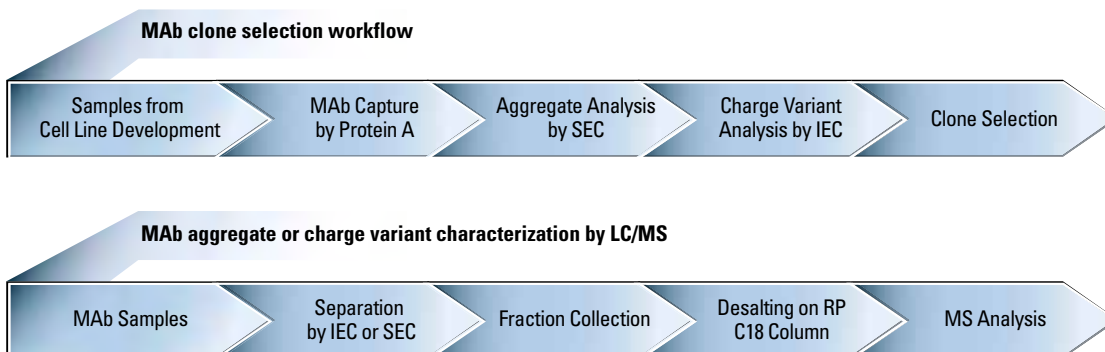
## The analytical solutions

Dionex provides best-in-class analytical solutions for MAb therapeutics, improving throughput and productivity, thus reducing time to market.

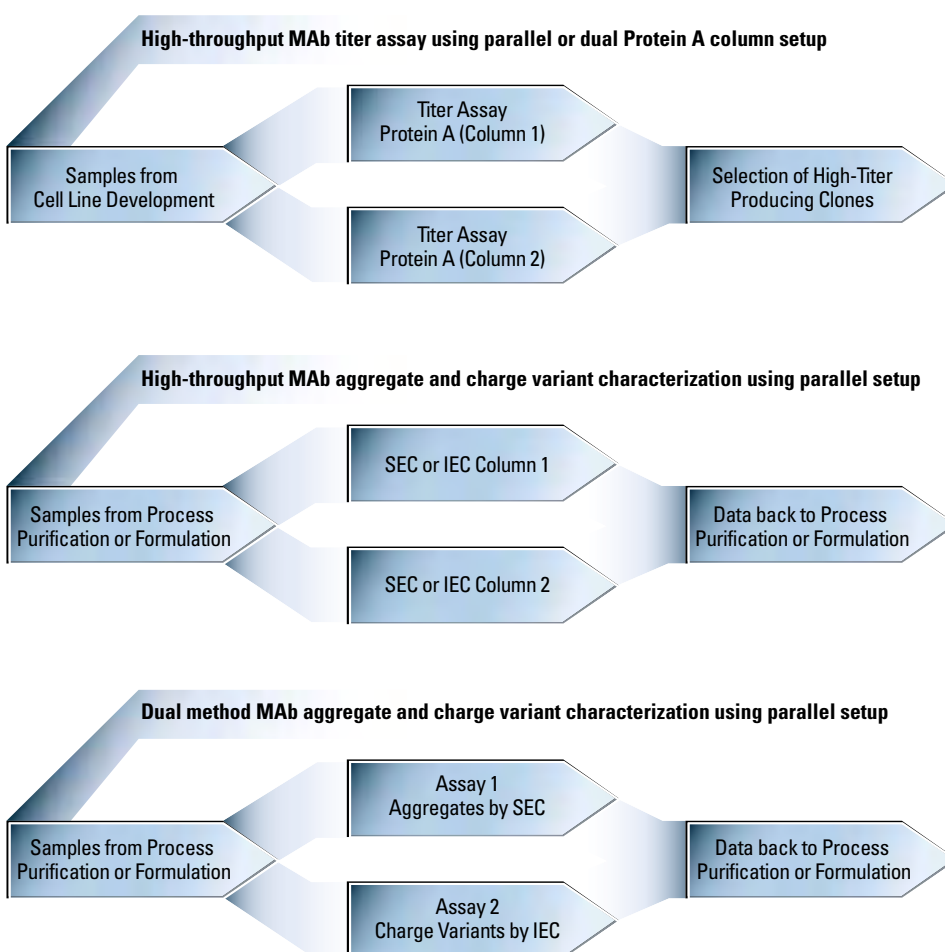
- Best-in-class columns for high-resolution and high-throughput characterization using ion-exchange chromatography (IEC) and size-exclusion chromatography (SEC)
- MAb characterization platforms to increase sample throughput and streamline multistep workflows
- Powerful and flexible Chromeleon® Chromatography Data System software
- Innovative UltiMate® 3000 RSLC system for UHPLC solutions for fast MAb characterization and peptide mapping
- High performance MAb glycan analysis

# Whatever Your Workflow, You Can Achieve Higher Throughput and Productivity

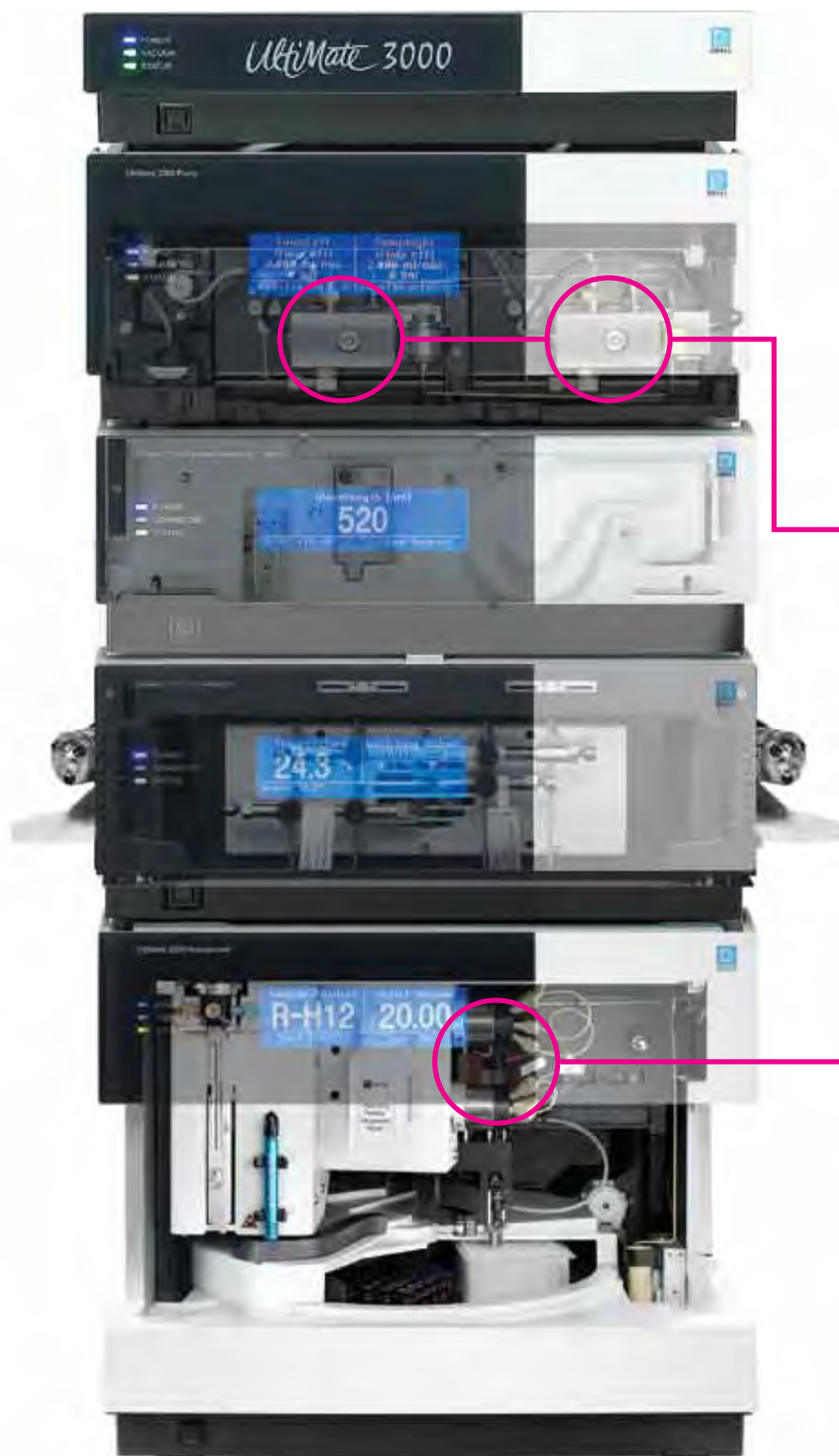
## Multistep MAb characterization workflows



## High-throughput MAb characterization workflows



# The Dionex MAb Characterization Platform—



## UltiMate 3000 MAb fully biocompatible platform

- Titanium pumps; PEEK™ fluidics, valves and injection needle
- Full compatibility with all biological buffers
- Prevents iron poisoning of columns
- Maintains protein modifications
- Less system corrosion and maintenance

## Dual gradient pump design: 2-in-1 system

- Multistep Automation: allows automation of two or more methods like Protein A capture, SEC aggregate, and IEC charge variant analysis
- Tandem Analyses: shortens run times by utilizing the power of off-line column regeneration
- Parallel LC: double throughput for productivity, cost, and space saving

## Autosampler with fraction collection and re-injection

- Dual valve design allows injection, fraction collection, and re-injection
- Optimized for automated workflows like:
  - Automated multistep workflow automation
  - Protein purification
  - Sample fractionation and desalting prior to MS detection
  - Sample derivatization like digestion or neutralization between multistep separations

*UltiMate 3000 MAb Analysis Platform with dual ternary gradient pumps and autosampler with integrated fraction collector.*

# For Your Throughput and Productivity Needs

## Chromeleon Chromatography Data System (CDS) software

Features of Chromeleon CDS software:

- Automated peak integration
- Automated control of fraction collection into autosampler
- Automated method template and sequence generation for multi-dimensional workflows
- Automatic rejection of samples based on set criteria e.g., automated rejection of clones with low antibody titer
- Dilutions or variable injection volumes for the next steps
- One-click report generation
- Validation report templates and sequences for increased automation of method validation



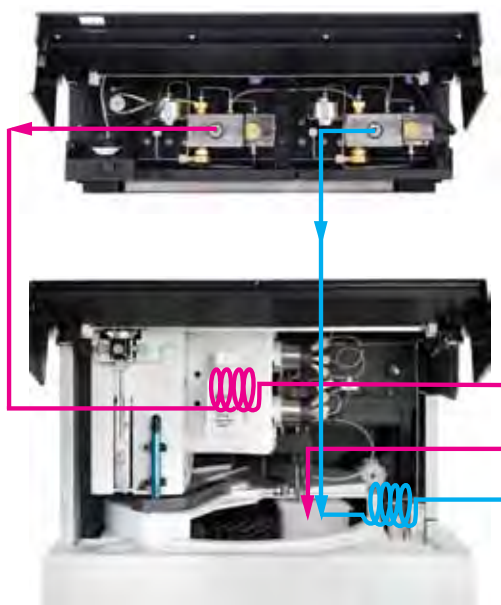
*Fully integrated solutions for R&D, analytical method development and QA/QC.*



# Boost Productivity with Automated Multistep MAb Capture and Characterization

**Automate your multistep MAb capture and characterization—  
reduce hands-on time and increase productivity**

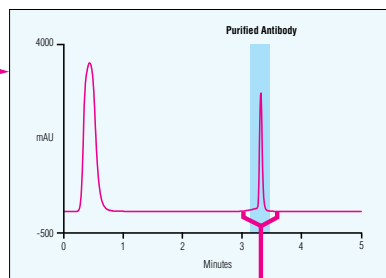
*Dual Ternary Gradient Pump. 2-in-1 Pump Design.*



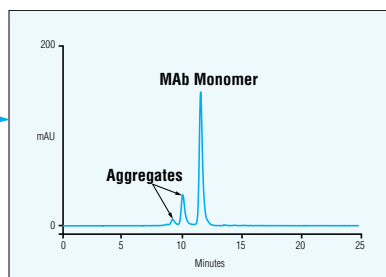
*Autosampler with Injection, Fraction Collection,  
and Re-Injection. Two sampling technologies in one.*

*The Multistep MAb Analysis Platform allows the purification and analysis of hundreds of MAb samples. Cell culture fluid samples containing antibodies are injected onto a Protein A column for antibody recovery, then the purified antibody samples are automatically injected, first onto the SEC column, and then onto the IEC column—fully automated.*

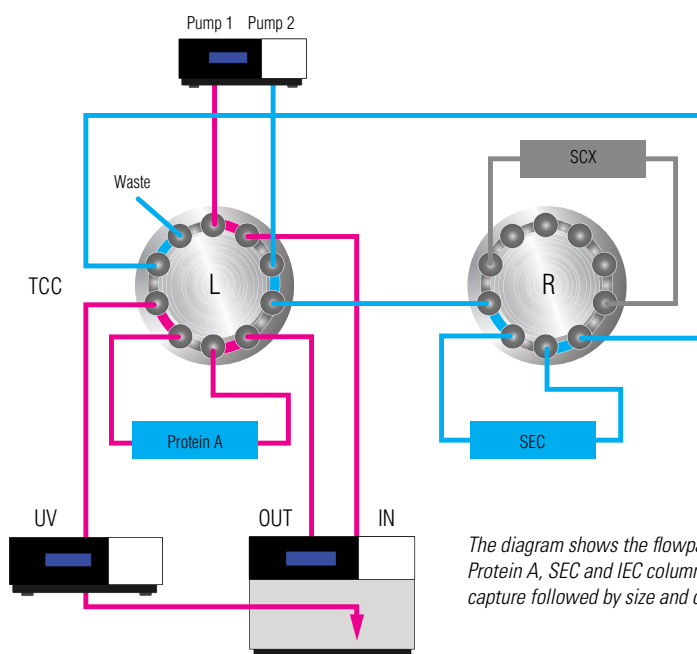
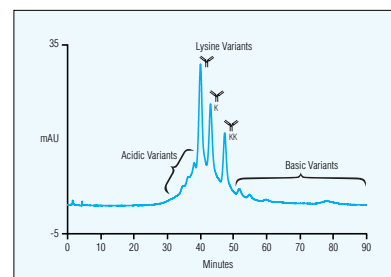
*Step 1: Protein A capture of IgG (MAb) and  
automated fraction collection.*



*Step 2: SEC aggregate analysis.*



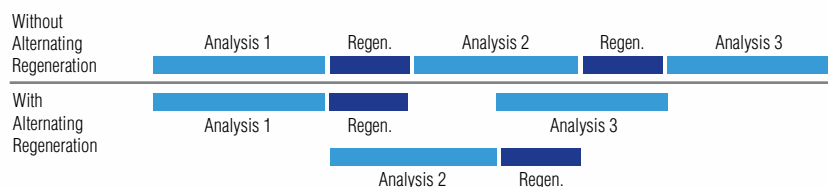
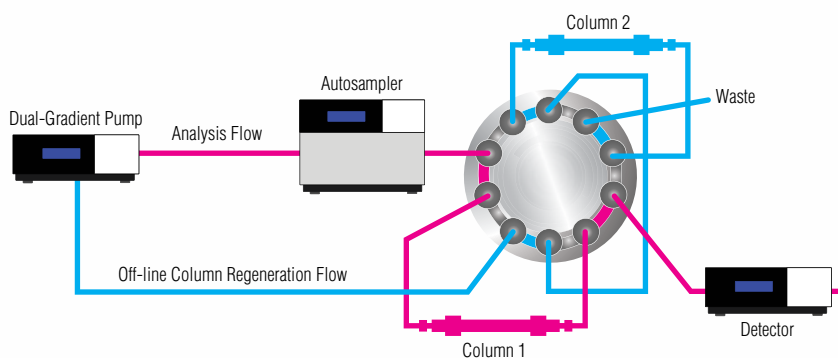
*Step 3: IEC charge variant analysis.*



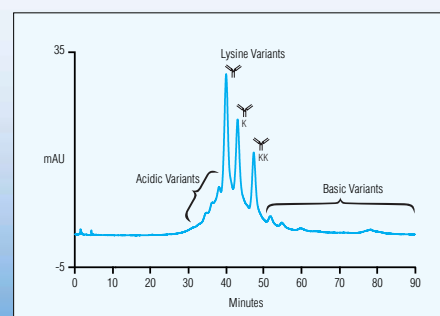
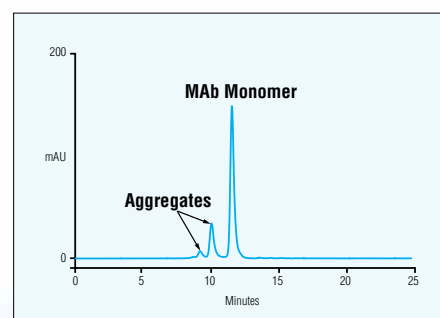
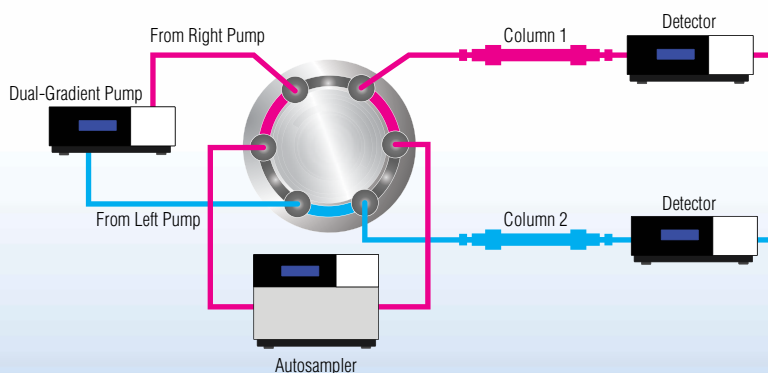
*The diagram shows the flowpath configuration of the  
Protein A, SEC and IEC columns for fully automated MAb  
capture followed by size and charge characterization.*

# Accelerate Product Development by Increasing Sample Throughput

Analyze over a 1000 MAb samples a day by utilizing the power of off-line column regeneration



With the same samples perform multiple separation steps including SEC and IEC in a parallel configuration



# Monoclonal Antibody Characterization Using

---

## Charge Characterization with IEC

- Glycosylation
- Sialylation
- C-Terminal Lysine
- Deamidation
- Glutamine cyclization
- Maleuric acid adduct
- Oxidation
- Cysteinylation
- Disulfide related
- Succinimide
- Isomerization

## Column

ProPac® WCX-10 (4 × 250 mm)

ProPac WCX-10HT (4 × 50 mm)

## NEW!

MABPac™ SCX-10 (4 × 250 mm)

MABPac SCX-10 (4 × 150 mm)

MABPac SCX-10HT (4 × 50 mm)

MABPac SCX-10 for LC/MS (2 × 250 mm)

---

## Hydrophobicity Characterization with HIC

- Isomerization
- Succinimide
- Oxidation
- Amidation
- Aggregation
- Clipping

## Column

ProPac HIC (4.6 × 250 mm)

ProPac HIC (4.6 × 100 mm)

ProPac HIC-10 (2.1 × 100 mm)

ProPac HIC-10 (7.8 × 75 mm)

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## Size Characterization with SEC

- Monomers, aggregates, and fragments
- Under non-denaturing conditions using both high- and low-salt mobile phases and volatile eluents for LC/MS

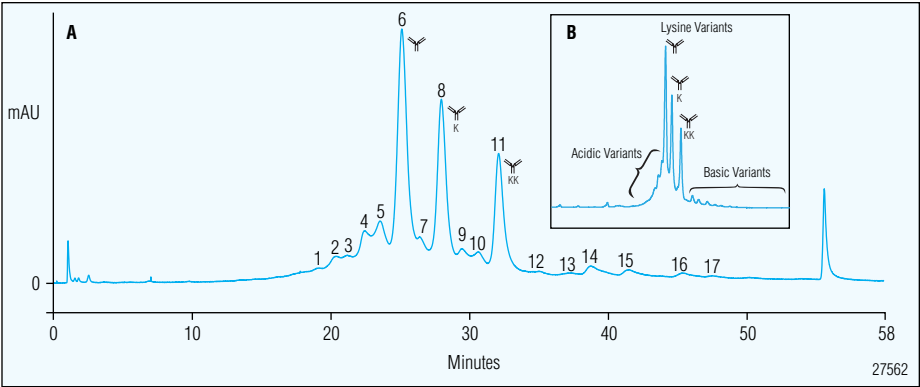
## Column

MABPac SEC-1, 5 µm, 300 Å (4.0 × 300 mm)

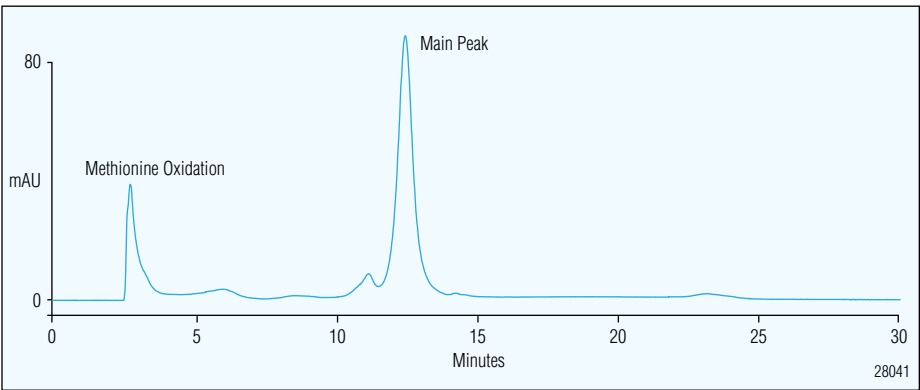
MABPac SEC-1, 5 µm, 300 Å (4.0 × 150 mm)



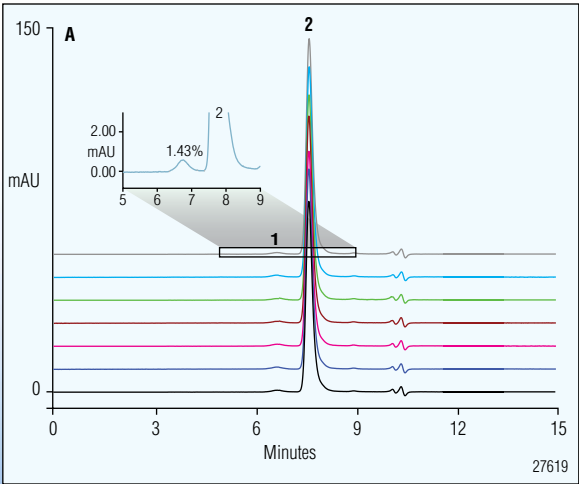
# High-Resolution Columns



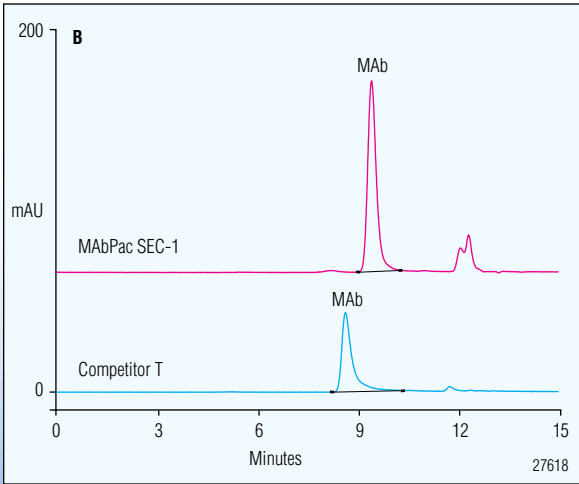
A. MAbPac SCX-10 column is the next generation IEC column for MAb charge characterization from Dionex.  
B. ProPac WCX-10 column (inset) is the industry gold standard for charge variant characterization.



ProPac HIC-10 column - Separation of populations of MAb variants using hydrophobic interaction chromatography (e.g., methionine oxidation monitoring).



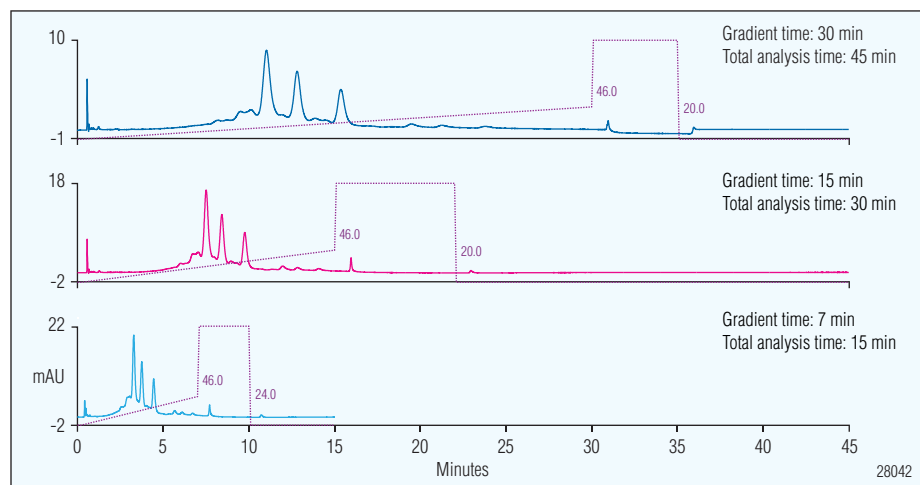
A. MAb analysis using the MAbPac SEC-1 column demonstrating excellent ruggedness.



B. MAb analysis in volatile buffer for LC/MS—MAb Pac SEC-1 vs the competitor's column.

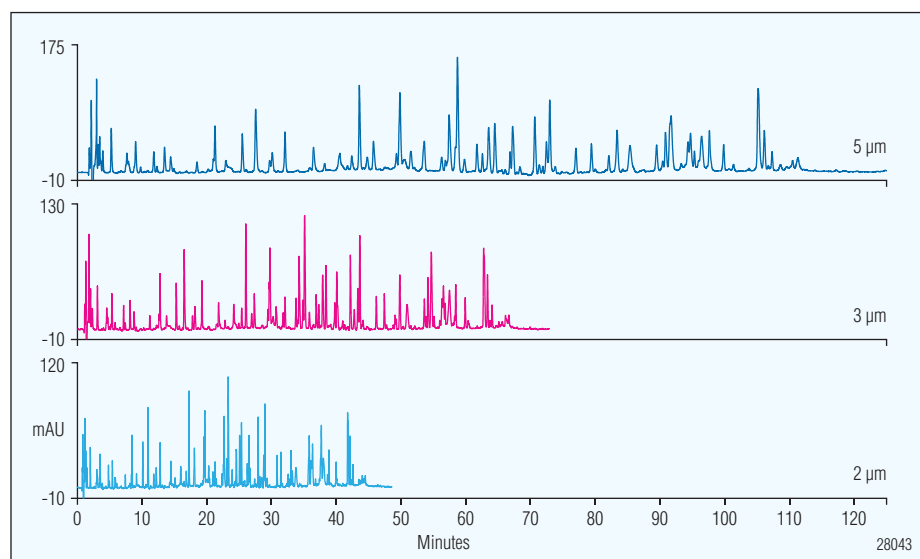
# Fast MAb Characterization and Peptide Mapping

## Method acceleration using the new MAbPac SCX-10 column, 4 × 150 mm



Elution profiles of a MAb sample from a 4 × 150 mm MAbPac SCX-10 column with different gradient elution times. The gradient time and total analysis time is indicated in the chromatogram.

## Fast MAb peptide mapping—application of UHPLC and reduced particle size



Performing peptide mapping with smaller particle columns allows the user to achieve identical resolution in less time. Here a method transfer was performed using 2.1 × 100 mm columns packed with 5, 3, and 2 μm particles respectively on the UltiMate 3000 Rapid Separation LC (RSLC).

# The Preferred and Proven Platform for MAb Glycan Characterization

## MAb glycosylation—why measure it?

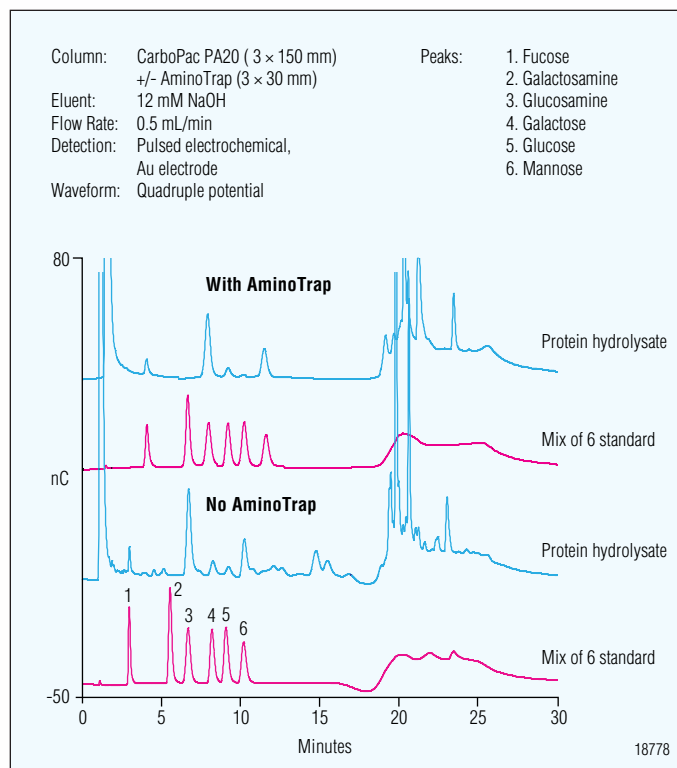
- Increasing relevance—biosimilars
- Glycosylation can affect:
  - Biological activity
  - Pharmacokinetics and clearance *in vivo*
  - Stability
  - Immunogenicity
- Analysis of glycosylation is important to:
  - Meet regulatory requirement
  - Ensure product consistency
  - Develop new generation drugs with modified glycosylation

## Dionex solution

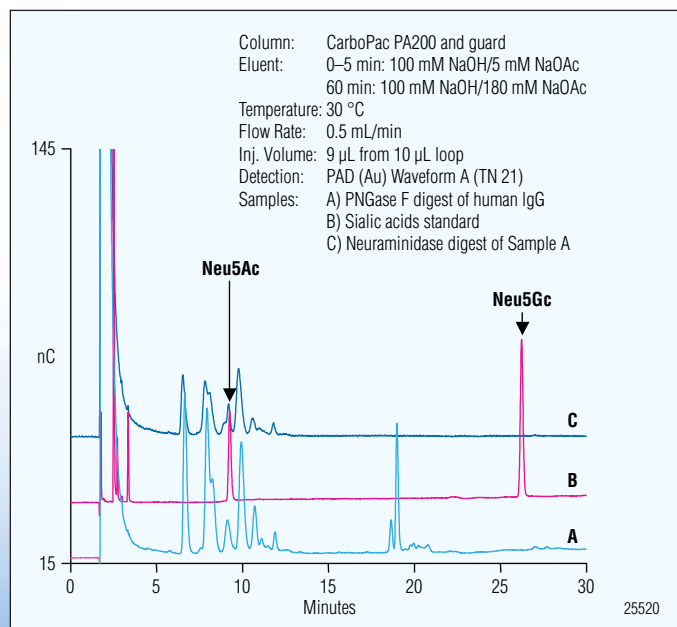
- High Performance Anion-Exchange with Pulsed Amperometric Detection (HPAE-PAD)
  - The workhorse in carbohydrate analysis
  - High Sensitivity—0.1 to 1 pmole detection limits
  - Label-Free—No derivatization necessary



ICS-5000—New capillary system for carbohydrate analysis.



Profiling MAb hydrolysate on the CarboPac® PA20 column, with and without an AminoTrap™ precolumn.



Monitoring release of sialic acids from human IgG N-linked oligosaccharides by HPAE-PAD.

## Selected Peer Reviewed Publications

1. Lyubarskaya, Y.; Houde, D.; Woodard, J. Murphy, D.; Mhatre, R. Analysis of Recombinant Monoclonal Antibody Isoforms by Electrospray Ionization Mass Spectrometry as a Strategy for Streamlining Characterization of Recombinant Monoclonal Antibody Charge Heterogeneity. *Anal. Biochem.* **2006**, 348, 24–39.
2. Vlasak, J.; Ionescu, R. Heterogeneity of Monoclonal Antibodies Revealed by Charge-Sensitive Methods. *Curr. Pharm. Biotechnol.* **2008**, 9, 468–481.
3. Valliere-Douglass, J.; Wallace, A.; Balland, A. Separation of Populations of Antibody Variants by Fine Tuning of Hydrophobic Interaction Chromatography Operating Conditions. *J. Chromatogr., A* **2008**, 1214, 81–89.
4. Decrop, W.; Gendeh, G.; Swart, R. Development of an Automated Method for Monoclonal Antibodies Purification and Analysis. *Chromatography Today*, Jun 2009, 8–10.
5. Farnan, D.; Moreno, G.T. Multiproduct High-Resolution Monoclonal Antibody Charge Variant Separations by pH Gradient Ion-Exchange Chromatography. *Anal. Chem.* **2009**, 81(21), 8846–8857.
6. Farnan, D.; Moreno, D.T.; Stults, J.; Becker, A.; Tremintin, G.; van Gils, M. Interlaced Size Exclusion Liquid Chromatography of Monoclonal Antibodies. *J. Chromatogr., A* **2009**, 1216(51), 8904–9.
7. Rea, J.C.; Moreno, G.T.; Lou, Y.; Parikh, R.; Farnan, D. High-Throughput Multi-Product Liquid Chromatography for Characterization of Monoclonal Antibodies. *BioPharm International* **2010**, 23, 44–51.
8. Grey, C.; Edebrink, P.; Krook, M.; Jacobsson, S.P. Development of a High Performance Anion Exchange Chromatography Analysis for Mapping of Oligosaccharides. *J. Chromatogr., B Anal. Technol. Biomed. Life Sci.* **2009**, 877, 1827–1832.



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