## Kremasil

# GLP-1 agonist purification toolbox

Methodology to maximize purity and throughput in Liraglutide and Semaglutide purification

### Nouryon

## A toolbox to reach your purification goals

GLP-1 agonists are a family of peptides that have become popular in the early treatment of diabetes type ll. Achieving the right purity for these peptides is however quite challenging.

A set of purification methods have been developed using Kromasil stationary phases for removal of impurities in peptide crudes that can be combined in a systematic way to improve purity from crude to 99.5%. This set is called the **GLP-1 agonist purification toolbox**.

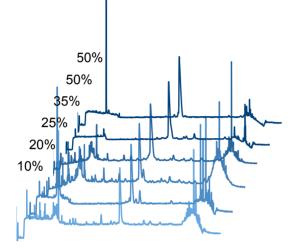
With a set of well adapted purification methods and routines in place, high efficiency and long column lifetime can be maintained to maximize throughput and productivity.

### Liraglutide and Semaglutide

Liraglutide and Semaglutide are a couple of the main GLP-1 agonists commercially produced on the market. They have a couple of amino acid substitutions compared to human GLP-1 to slow down peptidase degradation and are both acetylated with a hydrocarbon side chain.

However, if inadequately handled during purification, they easily degrade or aggregate. Most of these issues are avoided when using the toolbox methods.

Structure representation of Liraglutide (pdb id 4apd) using Mol\* Viewer (D. Sehnal et al (2021) Nucleic Acids Research. doi: 10.1093/nar/gkab314) and RCSB PDB.



### **Reaching for purity**

Crudes of Liraglutide are typically anything in the range from 10 to 50% in purity when entering downstream processing. The goal of the toolbox is to offer purification methods that will bring purity up to 99.5%.

Typical crude purity profiles for Liraglutide

### **Tools selection**

Operating the major factors affecting chromatography efficiency in HPLC, many different combinations of the following properties were screened:

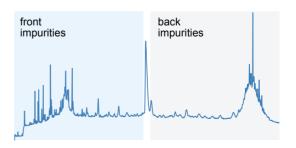
- Kromasil stationary phase
- Mobile phase organic solvent
- Mobile phase buffer

The screening compilation resulted in set of methods combinations with focus on the relative location of the impurities removed from the crude: back or front. The most efficient combinations have been compiled in the table below.

### Toolbox methodology

The methodical approach is to select two methods from the toolbox (tables on next page) that will target front and back impurities from the crude, respectively. In some cases this can be achieved in a single step.

> In a typical crude, impurities can be grouped as front and back impurities



Targeted impurities	Stationary phases	Organic solvents	Buffer salts			
front	Kromasil C4	acetonitrile	potassium phosphate			
	Kromasit C4	acetomitrite	trifluoroacetic acid			
	Kromasil C8	acetonitrile	potassium phosphate			
	Kromasit Co	propanol	ammonium acetate			
	Knonso sil Dhonul	acetonitrile	ammonium acetate			
	Kromasil Phenyl	acetomitrite	potassium phosphate			
	Kromasil diC4	2-propanol	ammonium acetate			
		acetonitrile	potassium phosphate			
		ethanol	ammonium acetate			
back	Kromasil C4	acetonitrile	potassium phosphate			
	Kromasil C8	acetonitrile	potassium phosphate			
			ammonium acetate			
	Kromasil Phenyl	acetonitrile	ammonium carbonate			
			potassium phosphate			
	Kromasil diC4	acetonitrile	potassium phosphate			

### Toolbox combinations for the purification of Semaglutide

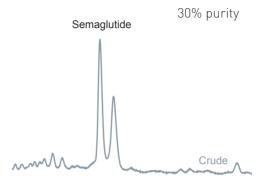
### Toolbox combinations for the purification of Liraglutide

Targeted impurities	Stationary phases	Organic solvents	Buffer salts			
front		ethanol	ammonium formate			
	Kromasil C4	ethanot	ammonium acetate			
		acetonitrile	ammonium acetate			
	Kromasil C8	ethanol	ammonium acetate			
	Kromasil Phenyl	acetonitrile	ammonium carbonate			
	Kromasil diC4	acetonitrile	ammonium carbonate			
	KI OMASIL UIC4	acetominie	ammonium acetate			
back	Kromasil C4	ethanol	ammonium carbonate			
		ethanot	citric acid			
		2-propanol	ammonium acetate			
		acetonitrile	ammonium acetate			
		acetonitrite	ammonium carbonate			
	Kromasil C8	ethanol	ammonium acetate			
	RI UTTASIL CO	acetonitrile	ammonium carbonate			
	Kromasil Phenyl	ethanol	ammonium acetate			
	Kromasil diC4	acetonitrile	ammonium carbonate			
	Riomasit ulo4	acetomente	ammonium acetate			

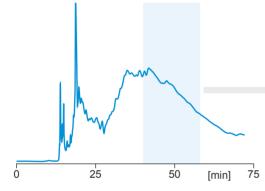
### Examples with Semaglutide

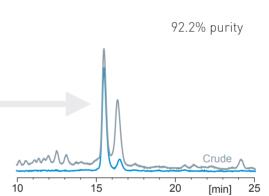
In this example, starting with a 30% purity crude of Semaglutide, the two-steps method uses Kromasil C4 in both steps, but at different pH, obtaining 92.2% purity after the first step and then 99.5% after the second step.

Starting crude with focus around the product peak. Other impurities are present further away, before and after the focus area.



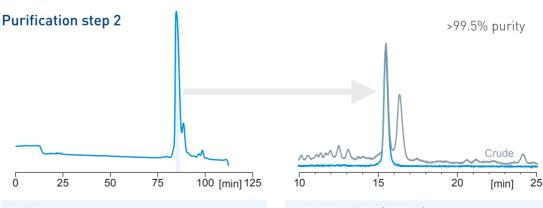
Purification step 1





Conditions

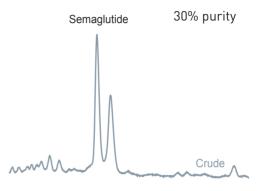
Stationary phase: Kromasil 100-10-C4 Mobile phase organic solvent: acetonitrile Mobile phase buffer: trifluoroacetic acid (TFA)

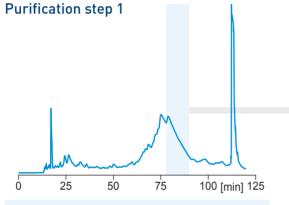


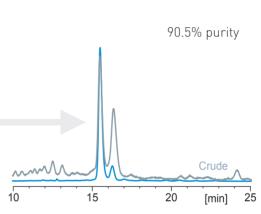
### Conditions

Stationary phase: Kromasil 100-10-C4 Mobile phase organic solvent: acetonitrile Mobile phase buffer: ammonium acetate Analytical conditions (all above) Stationary phase: Kromasil 100-3.5-C4 Mobile phase organic solvent: acetonitrile Mobile phase salt: trifluoroacetic acid Example with Semaglutide using phosphate buffers at different pH on the same C4 packing media. Also starting with the 30% purity crude, the first steps at low pH reaches a 90.5% purity. The second step is run at neutral pH achieving the final 99.5% purity.

Starting crude with focus around the product peak. Other impurities are present further away, before and after the focus area.

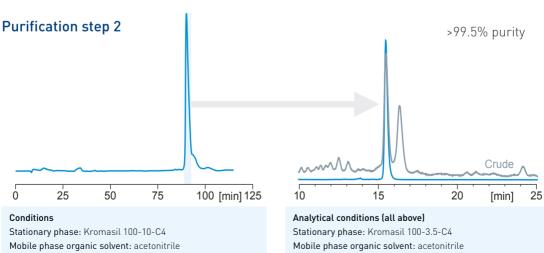






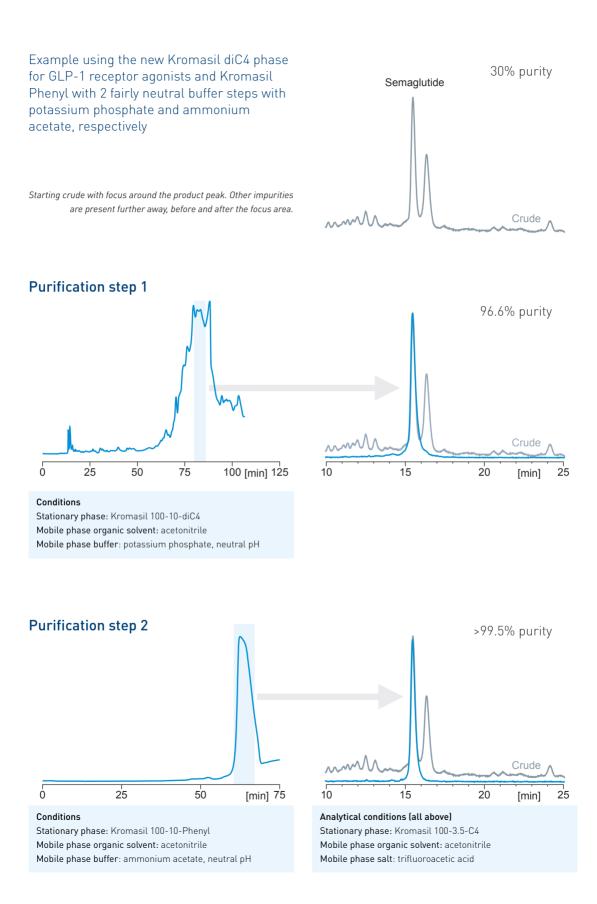
#### Conditions

Stationary phase: Kromasil 100-10-C4 Mobile phase organic solvent: acetonitrile Mobile phase salt: potassium phosphate, low pH

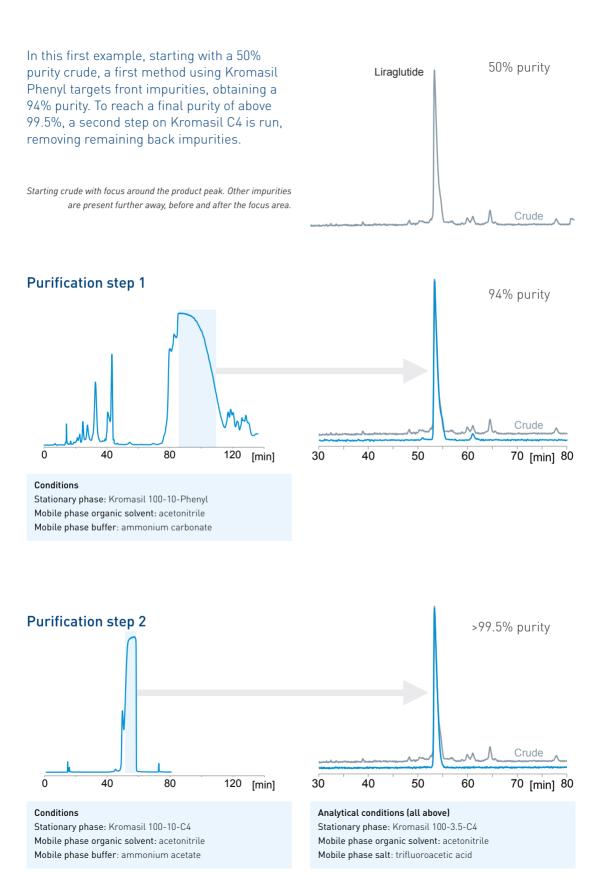


Mobile phase salt: potassium phosphate, neutral pH

Mobile phase salt: trifluoroacetic acid

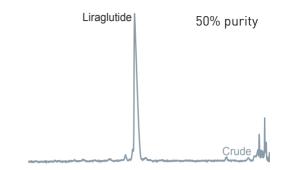


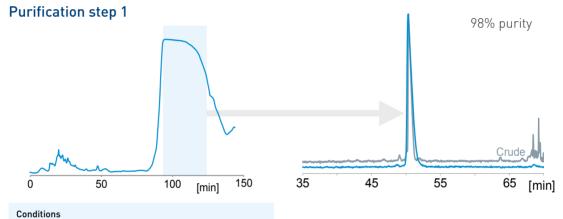
### Examples with Liraglutide



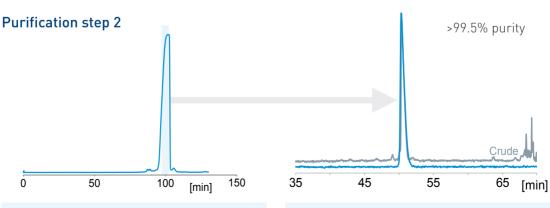
In this second example, starting with another 50% purity crude, a first method using Kromasil C8 targeting both front and back impurities, reaching a 98% purity. To achieve the final purity of above 99.5%, the second step is run on Kromasil C4, removing the remaining impurities.

Starting crude with focus around the product peak. Other impurities are present further away, before and after the focus area.



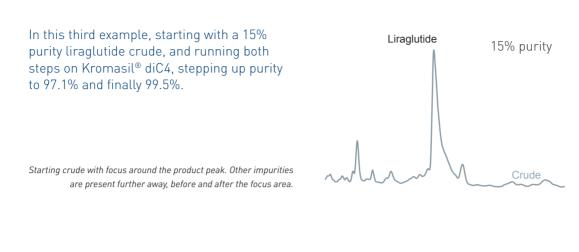


Stationary phase: Kromasil 100-10-C8 Mobile phase organic solvent: ethanol Mobile phase buffer: ammonium acetate

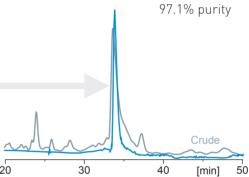


### Conditions

Stationary phase: Kromasil 100-10-C4 Mobile phase organic solvent: acetonitrile Mobile phase buffer: ammonium acetate Analytical conditions (all above) Stationary phase: Kromasil 100-3.5-C4 Mobile phase organic solvent: acetonitrile Mobile phase salt: trifluoroacetic acid

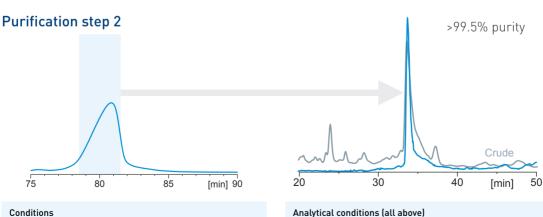


Purification step 1



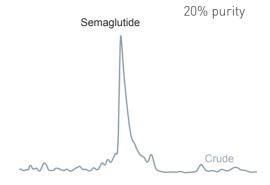
#### Conditions

Stationary phase: Kromasil 100-10-diC4 Mobile phase organic solvent: acetonitrile Mobile phase buffer: ammonium carbonate, neutral pH



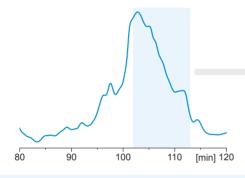
#### Conditions

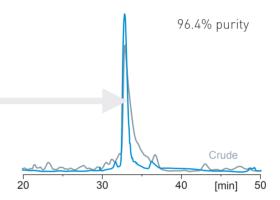
Stationary phase: Kromasil 100-10-diC4 Mobile phase organic solvent: acetonitrile Mobile phase buffer: ammonium acetate, mild basic pH Analytical conditions (all above) Stationary phase: Kromasil 100-3.5-C4 Mobile phase organic solvent: acetonitrile Mobile phase buffer: potassium phosphate, 10 mM pH 7.5 This fourth example is similar to the previous one but with a different crude with 20% liraglutide purity. Both steps are also run on Kromasil<sup>®</sup> diC4, stepping up purity to 96.4% and, finally, virtually 100%.



Starting crude with focus around the product peak. Other impurities are present further away, before and after the focus area.

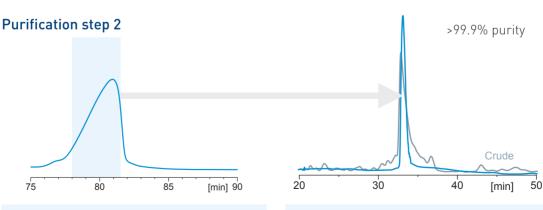






#### Conditions

Stationary phase: Kromasil 100-10-diC4 Mobile phase organic solvent: acetonitrile Mobile phase buffer: ammonium carbonate, neutral pH



### Conditions

Stationary phase: Kromasil 100-10-diC4 Mobile phase organic solvent: acetonitrile Mobile phase buffer: ammonium acetate, mild basic pH Analytical conditions (all above) Stationary phase: Kromasil 100-3.5-C4 Mobile phase organic solvent: acetonitrile Mobile phase buffer: potassium phosphate, 10 mM pH 7.5

### Doing it right

While peptide crudes can be challenging when repeatedly injected on an HPLC column, having a regeneration routine can be beneficial to maintain a good column lifetime and efficiency and thereby process productivity.

### Troubleshooting back-pressure increase

When purifying demanding peptide crudes like Liraglutide, the column quickly gets fouled by absorbed impurities with repeated injections, even when performing a traditional 2 column volume wash between injections. We propose here to perform an additional wash under mild conditions after a sequence of injections to quickly and fully recover column capacity and efficiency, optimizing productivity and throughput.

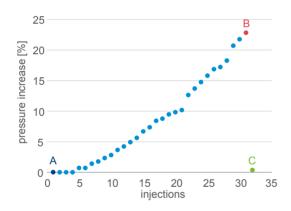
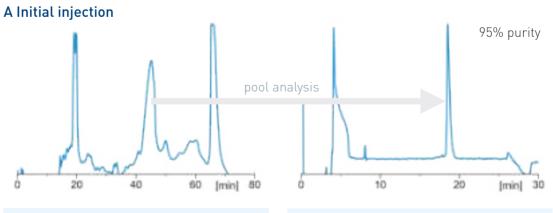
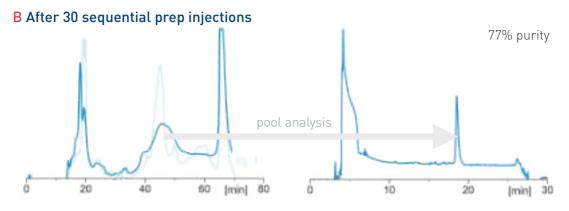


Figure showing the back-pressure over the column increasing between 30 consecutive crude injections from A to B. C shows restored normal back-pressure after the additional column wash. Chromatograms illustrating the status at these three points are shown in the following figures.

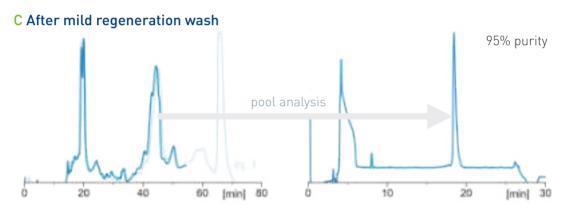


Prep conditions (in A, B and C) Stationary phase: Kromasil 100-10-C4 Mobile phase organic solvent: acetonitrile Mobile phase buffer: ammonium acetate

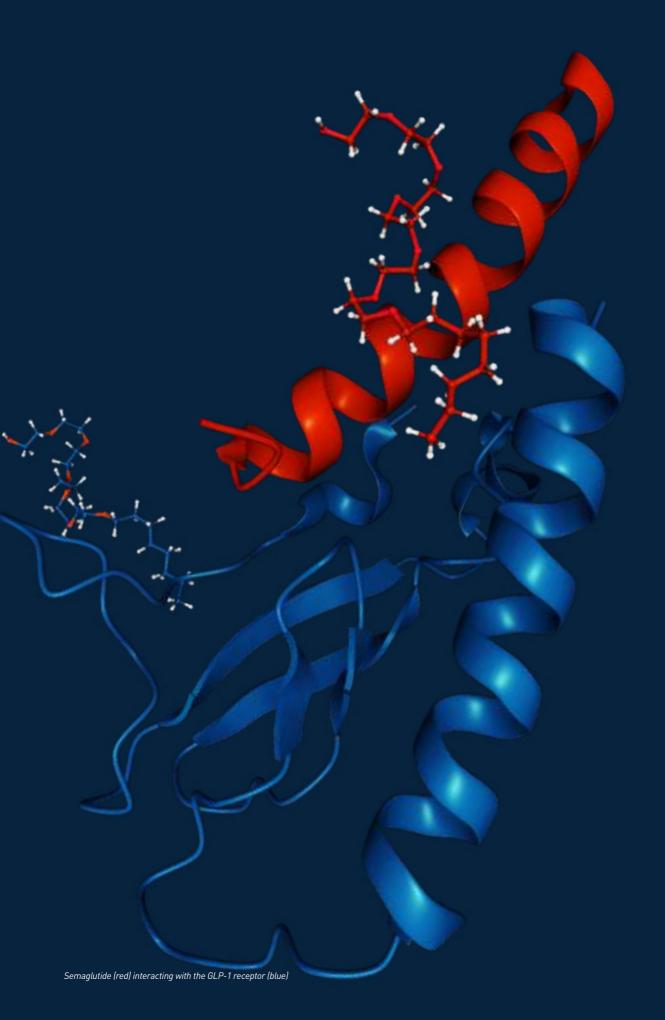
Analytical conditions (in A, B and C) Stationary phase: Kromasil 100-3.5-C4 Mobile phase organic solvent: acetonitrile Mobile phase salt: trifluoroacetic acid



After 30 preparative injections, column efficiency is impaired resulting in lower purity in extracted fractions. In the preparative chromatogram, the initial peak shape is shown in light blue as reference.



After the regeneration wash, peak shape and thereby efficiency and purity has been practically restored. The mild wash allows for a quick turn around to proceed with new injections. In the preparative chromatogram, the initial peak shape is shown in light blue as reference.



### Availability of Kromasil® Classic

					Particle size, [µm]						
Family	Phase	1.8	2.5	3	3.5	5	7	10	13	16	25
60 Å	SIL					•	•	٠	•	٠	
60 Å	CN							•		•	
60 Å	Diol							٠			
60 Å	HILIC-D							•			
100 Å	SIL	•	•		•	•	•	•	•	•	
100 Å	C1										
100 Å	C4	•	•		•		•	•	•	•	
100 Å	diC4								•		
100 Å	C8	•	•		•		•	•	•	•	
100 Å	C18		•		•		•		•		
100 Å	C18(w)							•			
100 Å	NH2				•		•		•		
100 Å	Phenyl							•		•	
300 Å	SIL					•		•		•	
300 Å	C4							•		•	
300 Å	diC4							•		•	
300 Å	C8					•		•		•	
300 Å	C18							•		•	

• : standard product, available in bulk quantities

• : analytical product, only available in slurry-packed columns

• : bare silica product in analytical particle sizes available in bulk for contracted OEM producers

www.kromasil.com/classic

### Contact us

Are you also working on some GLP-1 agonist peptides and is interested in how this toolbox can reduce your development time and boost your productivity? Don't hesitate to contact us and reach out to our team of skilled application scientists for more details:

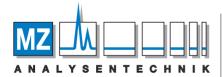
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Not only will you benefit from our patented silica technology, but you gain a strong partner with a reliable track record in the eld of silica products. For the past 70 years, we have pioneered new types of silica. Our long experience in the field of silica chemistry is the secret behind the development of Kromasil® stationary phases, and the success of our Kromasil® team. Kromasil® chromatography phases are available in bulk and in high-pressure slurry-packed columns.

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