



Microfluidic Systems for Radiochemistry Synthesis and Functionalization of Polymer Monoliths for Chromatographic Separation of U and Eu

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DE LA RECHERCHE À L'INDUSTRIE

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**Marion Losno, René Brennetot,
Clarisse Mariet ¹**

¹ Den – Service d'Etudes Analytiques et de
Réactivité des Surfaces (SEARS), CEA,
Université Paris-Saclay, F-91191, Gif sur Yvette,
France

Ivan Ferrante, Stéphanie Descroix ²

² MMBM Group, Institut Curie Research Center,
CNRS UMR 168, Paris, France

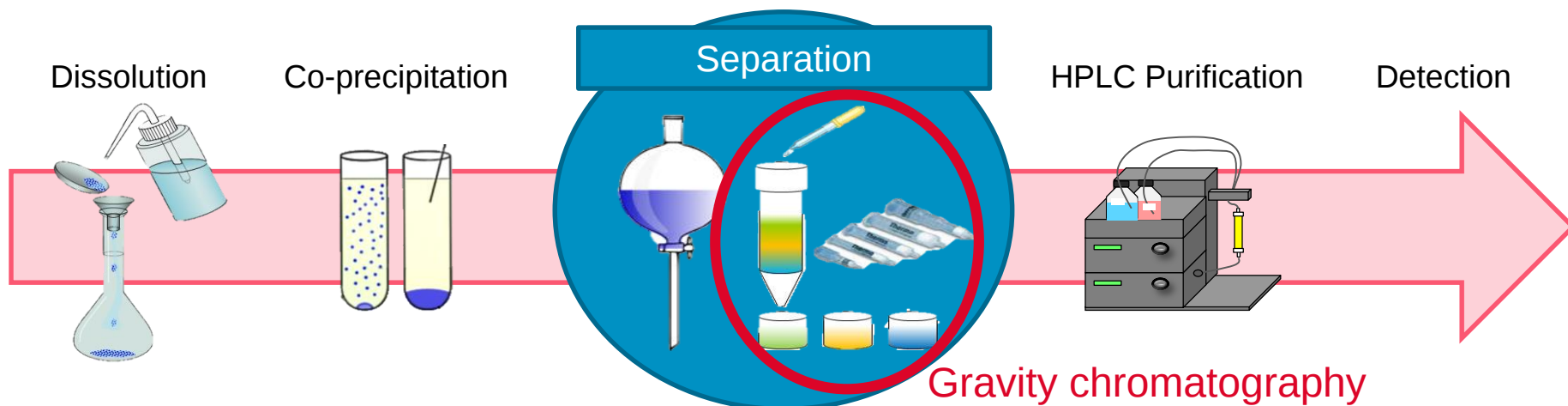


MICROFLUIDIC SYSTEMS FOR RADIOCHEMISTRY: SYNTHESIS AND FUNCTIONALIZATION OF POLYMER MONOLITHS FOR CHROMATOGRAPHIC SEPARATION OF U AND EU

ATALANTE 2016 | Marion Losno

June 9th, 2016





■ Many constraints have to be taken into account such as

- Glove box handling
- Concentrated irradiant samples
- Waste management

😊 Benefits of miniaturization

■ A little amount of sample needed for the analysis

- Decrease of the volume handled by the operators
- Decrease of the dose received
- Decrease of the amount of waste produced -> reduces analysis cost



■ Benefits related to plastic devices

- Easy to produce (thermoformable)
- Disposable : no cross-contamination

■ Benefits related to lab-on-CDs

- No need for external pumping device or connection
- Very limited instrumentation (only a rotor) -> limited cost of maintenance
- Easy automation and multiplexed analysis



 How to develop this type of technology ?

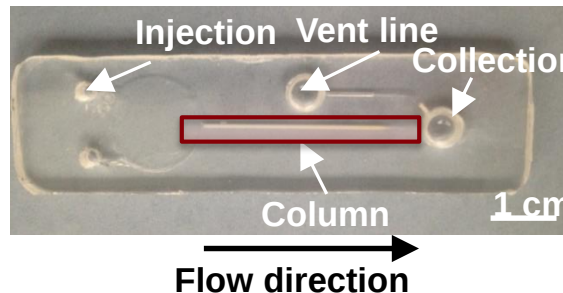
■ Step 1 : Lab-on-CD configuration *

- CD platform made of PEEK including 4 microsystems holders



■ Step 2 : microsystem design *

- Use of Cyclo Olefin Copolymer (cheap, easy to structure and suited physico-chemical properties)



■ Step 3 : implementation of a monolithic anion-exchange stationary phase for radionuclides separation

* Bruchet, A. et al. (2013). "Centrifugal microfluidic platform for radiochemistry: potentialities for the chemical analysis of nuclear spent fuels." *Talanta* **116**: 488-494.

■ Constraints due to the analysis

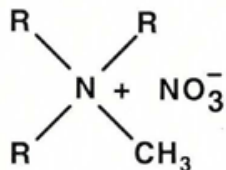
■ Nitric acid resistance

- Corresponding to the dissolution medium of radiochemical samples

When $[\text{HNO}_3] > 4\text{mol.L}^{-1}$ *

- Formation of $[\text{UO}_2(\text{NO}_3)_x]^{(2-x)}$ ($1 \leq x \leq 3$) **anionic complex**
- **Cationic form** of lanthanides (Ln^{3+})

- Functionalization by an anion exchanger will enable the separation of the two components in $[\text{HNO}_3] > 4 \text{ mol.L}^{-1}$



Quaternary ammonium was chosen for its efficiency at macro scale

* Chiarizia, R., R. Gatrone and E. Horwitz (1995). "Am (III) and Eu (III) extraction by Aliquat-336 and benzyl substituted quaternary ammonium salts from nitrate and thiocyanate solutions." Solvent Extraction and Ion Exchange **13**(4): 615-645.

- Stationnary phase constraints
 - Easy to synthesize
 - Localization of the phase
- } ☺ Photochemistry

Stationnary phase = polymer + grafting of a molecule of interest

- Choice of organic methacrylate monolith [1]

- Photosynthetizable
- Localized
- Chemical resistance
- Mechanical resistance
- Low flow resistance

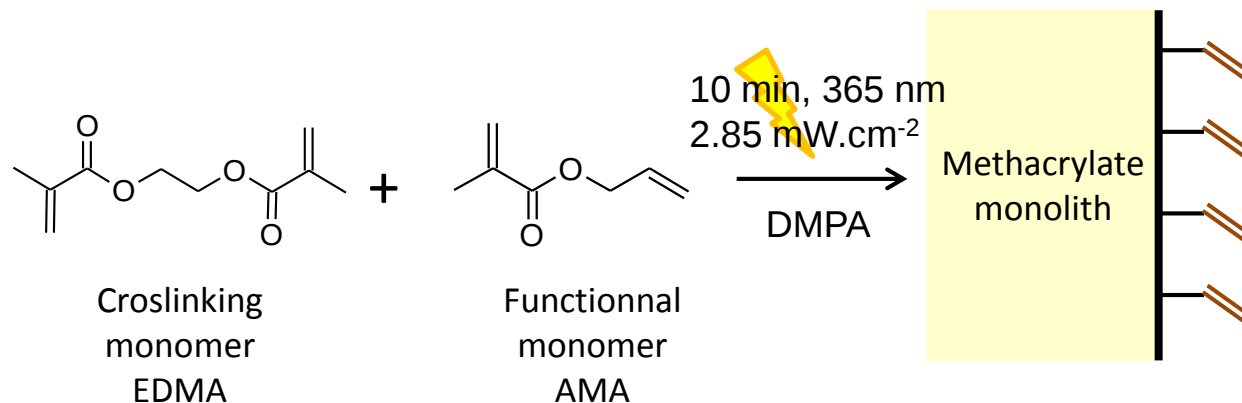
- Choice of thiolene « click chemistry » [2]

- Photochemical reaction
- Localized
- Chemical resistance
- Versatility

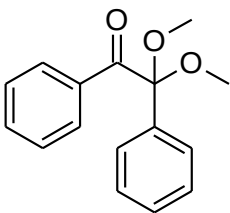
[1] Svec, F. and J. M. J. Fréchet (1996). "New designs of macroporous polymers and supports: From separation to biocatalysis." *Science* **273**(5272): 205-211.

[2] Kolb, H. C., M. G. Finn and K. B. Sharpless (2001). "Click chemistry: Diverse chemical function from a few good reactions." *Angewandte Chemie International Edition* **40**(11): 2004-2021.

ORGANIC MONOLITH SYNTHESIS OPTIMIZATION AT MACROSCALE



- Monomers : 24 wt% Ethylene glycol dimethacrylate (EDMA) and 16 wt% Allylmethacrylate (AMA)
- Porogenic solvent : 24 wt% 1,4-butanediol, 35 wt% 1-propanol, 1 wt% water
- Photoinitiator : 1 wt% DMPA



Globule size $\approx 1 \mu\text{m}$

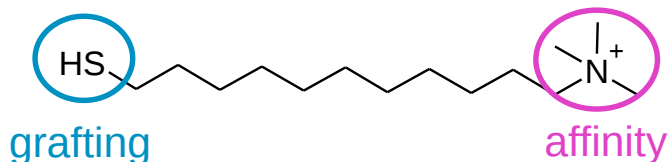
Permeability $\approx 10^{-14} \text{ m}^2$

Density $\approx 0.4 \text{ g.mL}^{-1}$

Total porosity $\approx 60 \%$

Choice of molecules

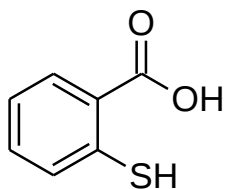
- Molecule to be grafted : ammonium thiol



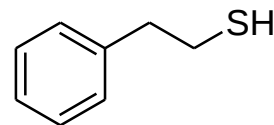
- ☺ Commercially available
- ☹ Price
- ☹ Characterization

→ Study of the reaction through different molecules

Thiosalicylic acid

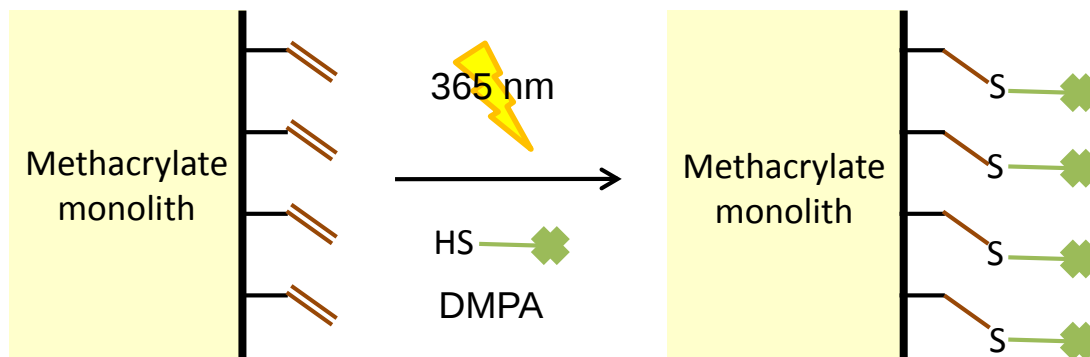


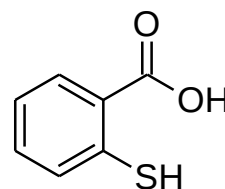
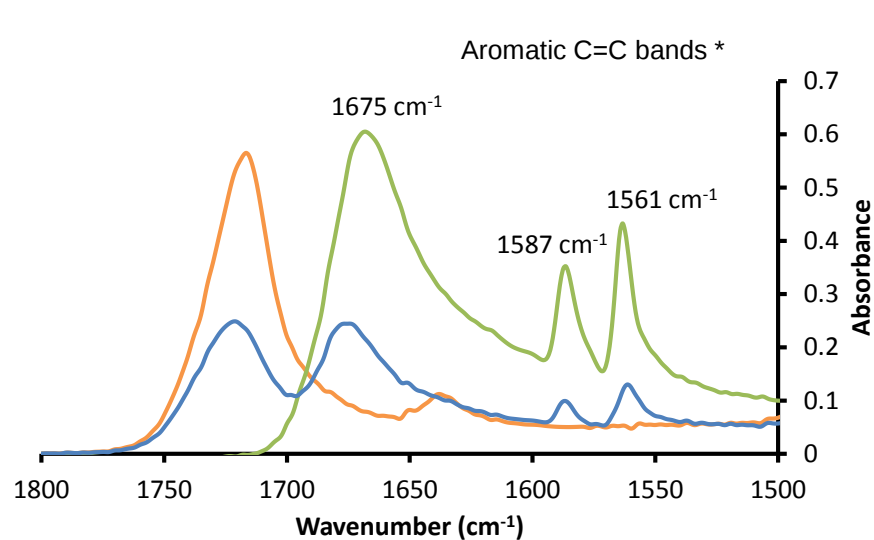
- ☺ Commercially available
- ☺ Price
- ☺ Characterization



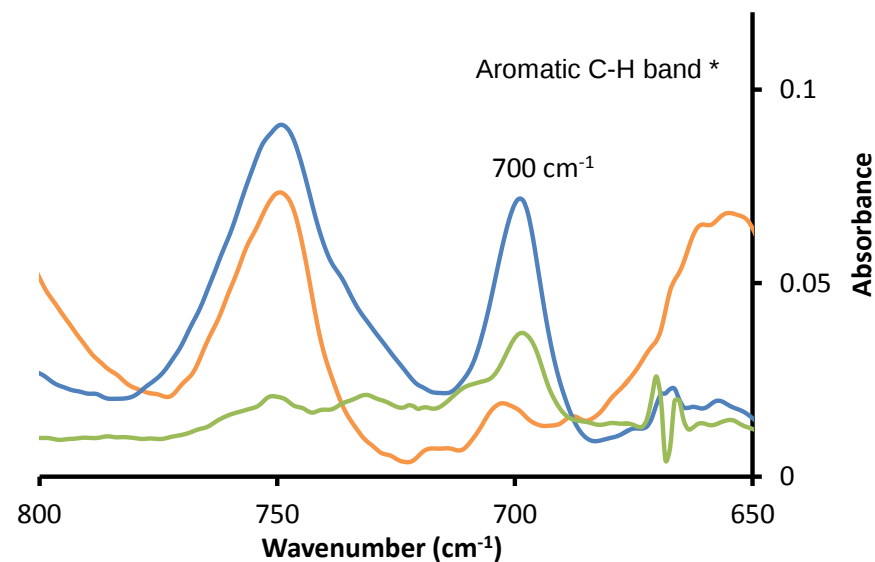
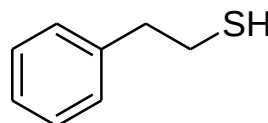
2-phenylethanethiol

Procedure





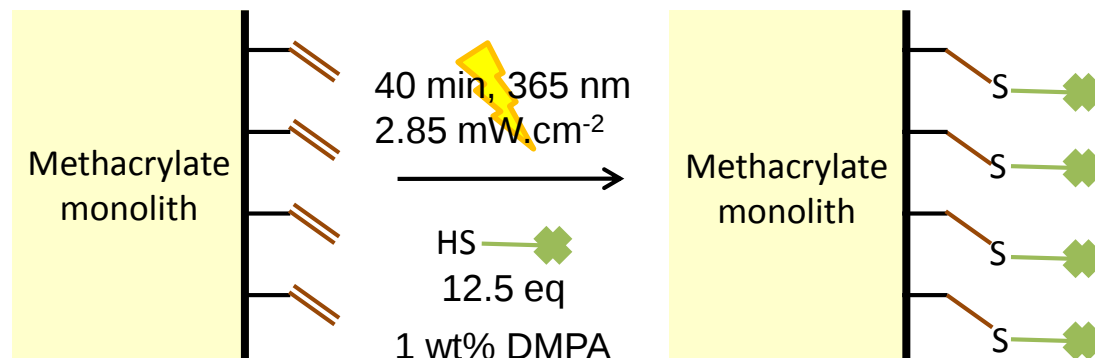
Blank monolith
Thiol
Grafted monolith



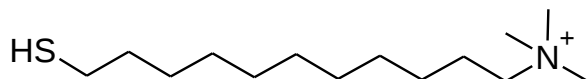
* Skoog, D. A., D. M. West and F. J. Holler (1997). *Chimie analytique*. Bruxelles, Boeck and Larcier.

* Silverstein, Basler and Morill (1998). *Identification spectrométrique de composés organiques*, Boeck and Larcier.

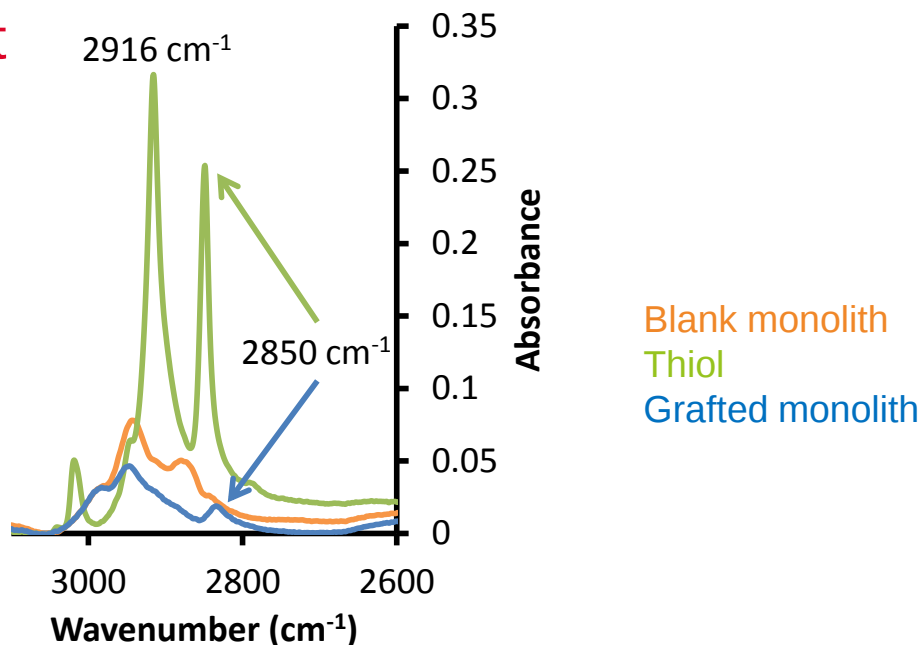
Optimized conditions



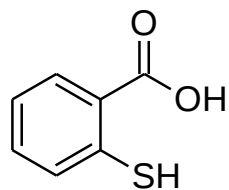
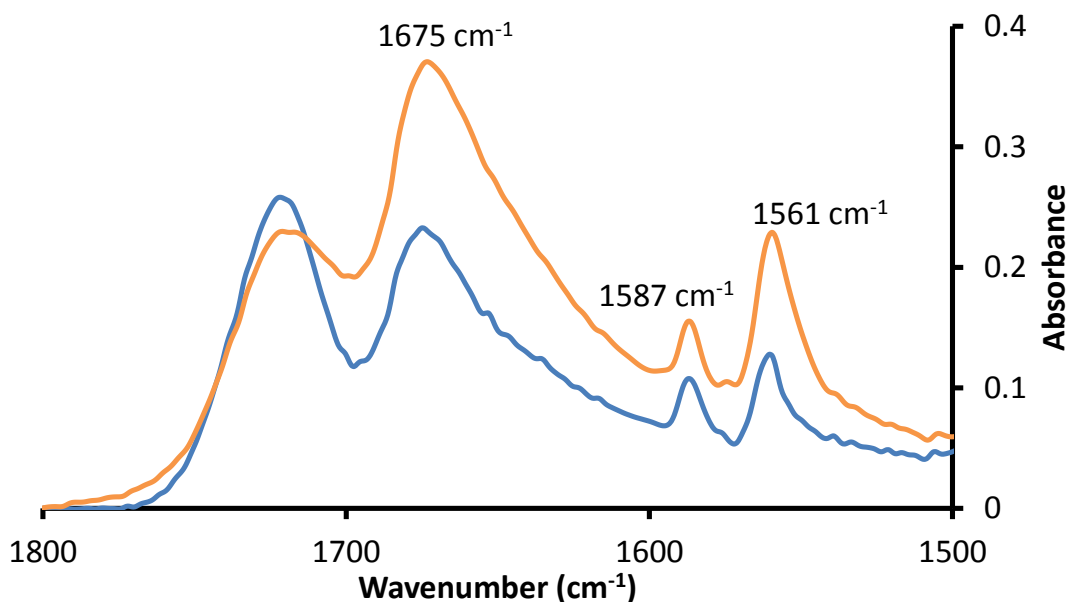
Molecule of interest



Versatile functionalization



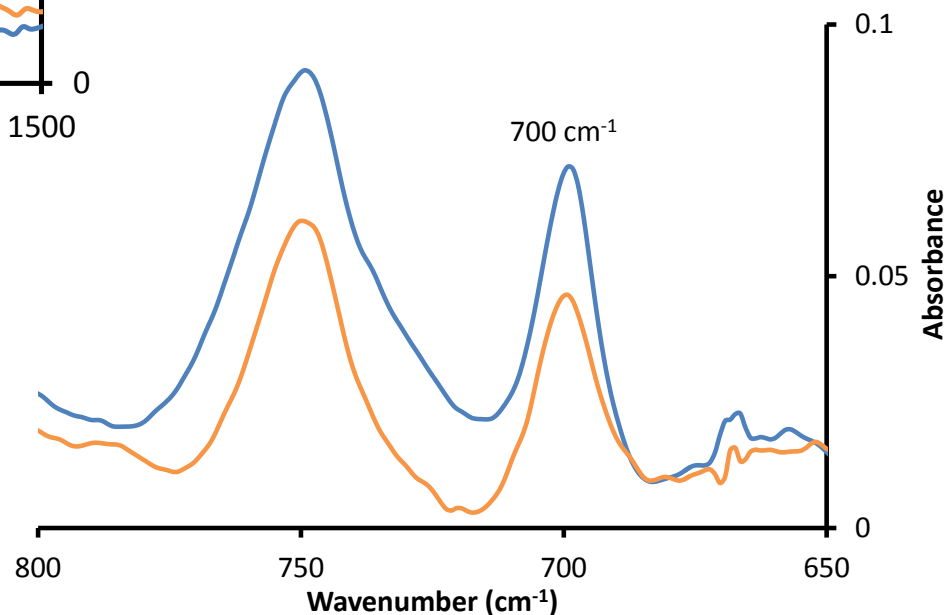
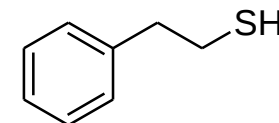
CHEMICAL RESISTANCE OF THE FORMED C-S BOND



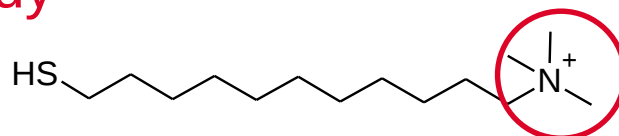
The C-S bond is robust in nitric acid medium

Grafted monolith

Grafted monolith after 24 h in a 8 mol.L⁻¹ nitric acid aqueous solution

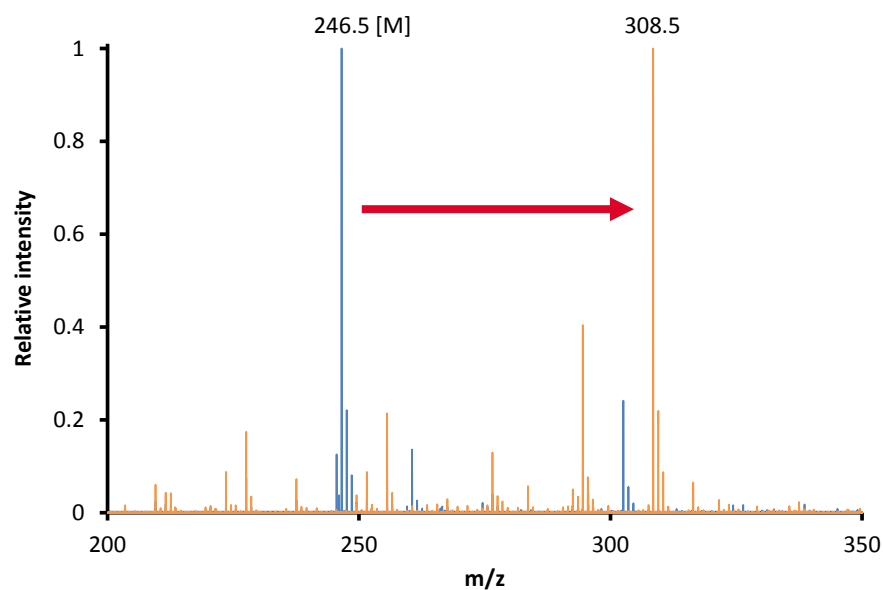


ESI-MS Study

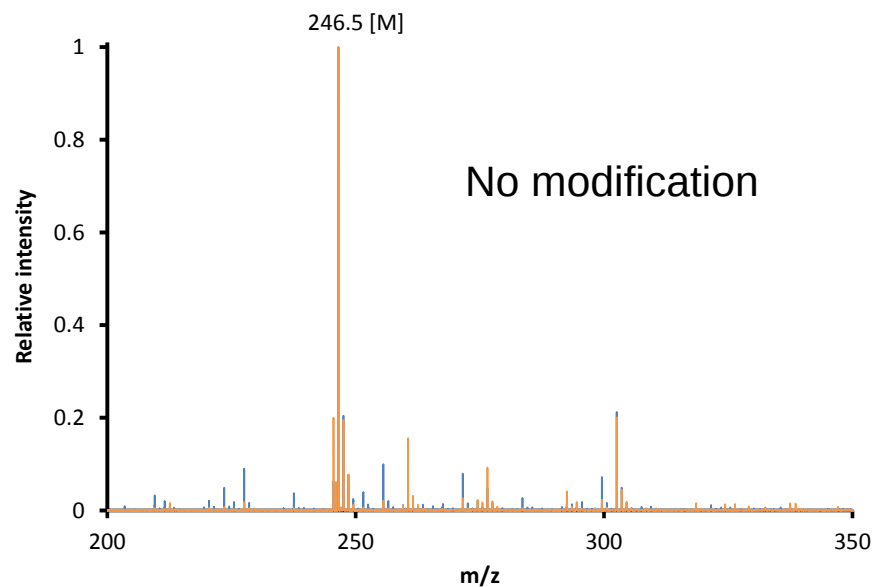


Free molecule

Free molecule after 24 h
in nitric acid solution



$[HNO_3] = 8 \text{ mol.L}^{-1}$



$[HNO_3] = 5 \text{ mol.L}^{-1}$

* Losno, M. et al (2016). "Photochemical synthesis and versatile functionalization method of a robust porous poly(ethylene glycol methacrylate-co-allyl methacrylate) monolith dedicated to radiochemical separation in a centrifugal microfluidic platform." *Micromachines* **7**(3): 45.

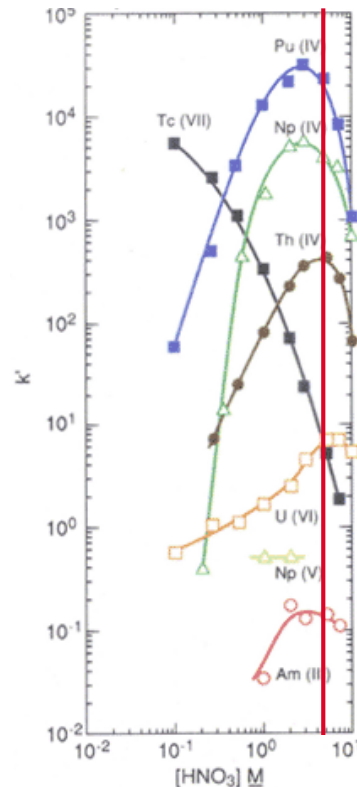
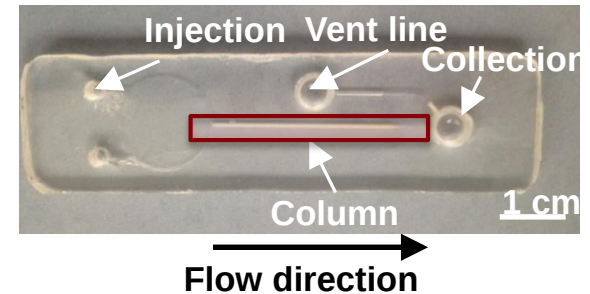
* Losno, M. et al (2016). "Microsystems for Anion Exchange Separation of Radionuclides in Nitric Acid Media" *Procedia chemistry*

Summary of the presented results

- A stationary phase for radiochemical analysis in hard acidic medium was developed
- A robust and versatile method of functionalization was proposed
- The C-S bond formed via thiol-ene chemistry is strong enough to be used in strong nitric acid medium
- The ammonium thiol presents sufficient resistance to be used in $[\text{HNO}_3] = 5 \text{ mol.L}^{-1}$ medium

Outlook

- Transfert of the optimized monolith in the COC microsystem
- Radiochemical analysis in 5 M nitric acid medium ?



5 mol.L⁻¹

An anion exchange stationary phase : TEVA resin

THANKS FOR YOUR ATTENTION



Commissariat à l'énergie atomique et aux énergies alternatives
Centre de Saclay | 91191 Gif-sur-Yvette Cedex
DEN/DANS/DPC/SEARS/LANIE – Bât. 391 – PC 33
T. +33 (0)1 69 08 83 85 | F. +33 (0)1 69 08 54 11

Etablissement public à caractère industriel et commercial | RCS Paris B 775 685 019

Direction de l'énergie nucléaire
Direction déléguée aux activités nucléaires de Saclay
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