

# Fast Screening of Chiral Stationary Phases for Chiral Separations on Zirconia

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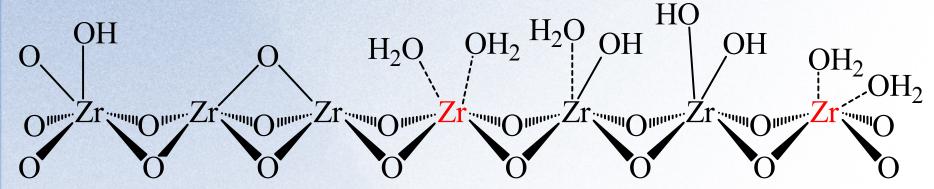


## Goal-To Make Zirconia Based Chiral Stationary Phases for Fast Chiral Selector Screening

- Why Zirconia?
- Synthetic Approach
- Zirconia-based vs Silica-based CSPs
- Chromatographic Comparison of Different Anchors
- Stability Study
- Example Separations on Zirconia Based CSPs
- Use for Fast Chiral Selector Screening
- Conclusions Zirconia Based CSPs Have Comparable Chromatographic Performance Compared to Silica Based CSPs. Fast Chiral Selector Screening is Possible.



#### Surface Chemistry of Zirconia

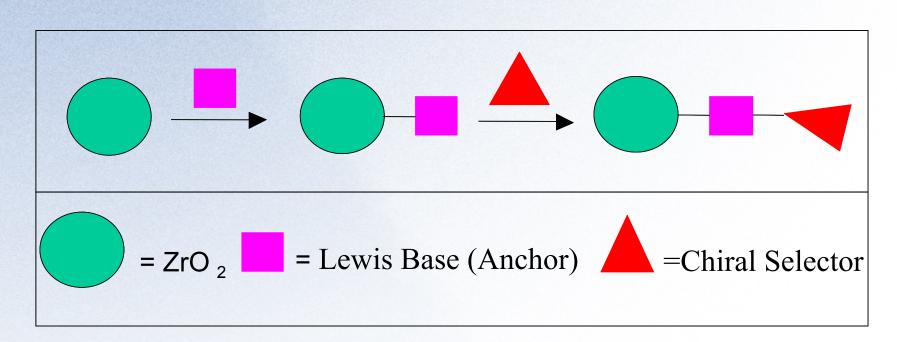


RPO<sub>3</sub><sup>2</sup>- or Catechol

Lewis Acid: 
$$Zr^{4+}$$
:  $H_2O + R-COO^- = Zr^{4+}$ :  $OOC-R + H_2O$ 



#### New Way to Attach Chiral Selectors to Zirconia Surface





#### **Three Anchors Studied**

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1) APPA (Aminopropylphosphonic acid)

3) ASPA (Aspartic acid)

2) DHNP (3,4-Dihydroxynorephedrine)

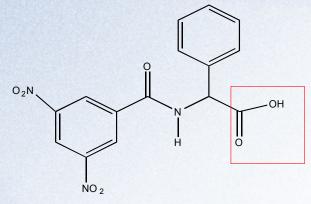
Anchors should have two function groups: (1) A group anchoring to zirconia surface, and (2) A group bonding to Chiral selector.



## **Chiral Selectors in This Study**

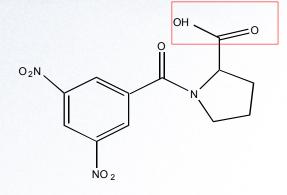
**DNB-LEU** (3,5-dinitrobenzoylLeucine)

NAP-VAL (Naphthoylvaline)



**DNB-PG** (3,5-dinitrobenzoylphenylglycine)

2-NAP-LEU (2-Naphthoyl-leucine),

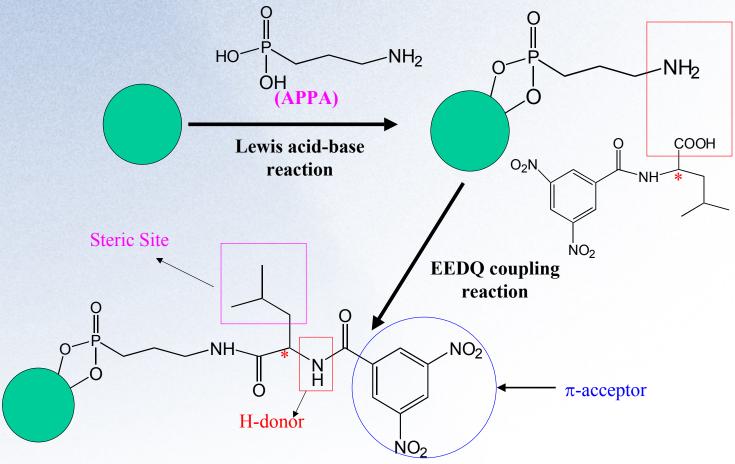


**DNB-PRO** (3,5-dinitrobenzoylproline)

NAP (naproxen)



## **Example of Lewis Acid-Base Modified Zirconia CSPs**





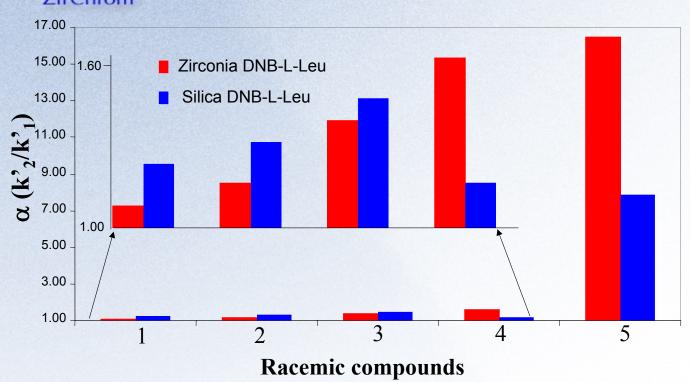
## List of Zirconia and Silica CSPs Studied

Column	CSP	Anchor
<b>Z</b> 1	DNB-Leu	APPA
Z2	DNB-Leu	Aspartic acid
<b>Z</b> 3	DNB-Leu	DHNP
Z4	DNB-PG	APPA
<b>Z</b> 5	DNB-PG	Aspartic acid
<b>Z</b> 6	DNB-PG	DHNP
<b>Z</b> 7	DNB-Pro	DHNP
<b>Z</b> 8	NAP-Leu	APPA
<b>Z</b> 9	NAP-Val	DHNP
Z10	Naproxen	APPA
R1	DNB-PG	
R2	DNB-Leu	

Z1-Z10 zirconia based CSPs, R1, R2-commercialized silica based CSPs



## Direct Comparison of DNB-L-LEU Zirconia and Silica Based CSPs

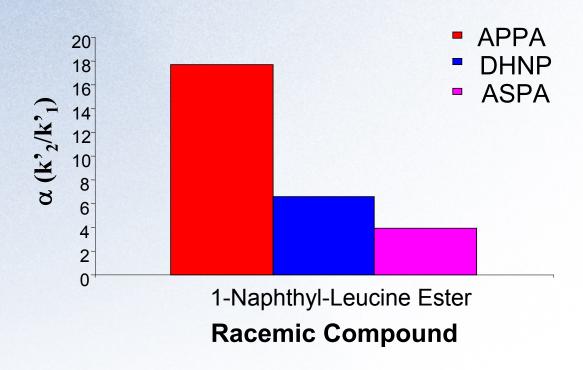


- 1 trans-stilbene oxide
- 2 1,1'-bi-2-naphthol
- 3 trifluoranthyl ethanol
- 4 napropamide
- 5 1-naphthyl leucine ester

Much better separations for napropamide and 1-naphthyl leucine ester are obtained on zirconia-based CSPs.



# Chromatographic Comparison of Differently Anchored Zirconia-based DNB-L-LEU

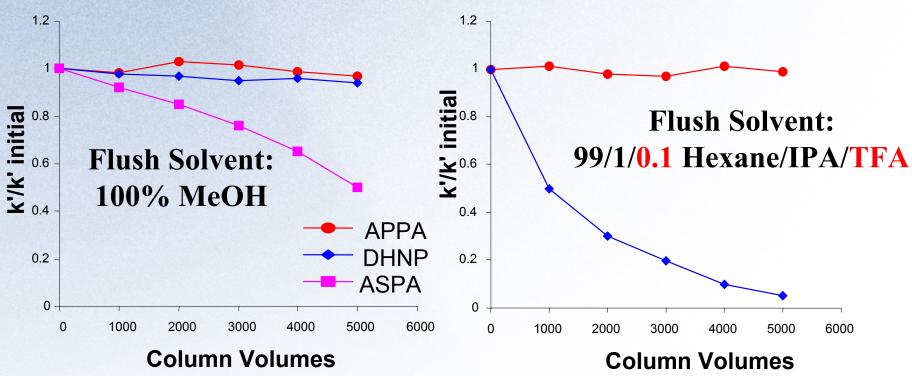


Different anchors show different selectivity.



## Stability Comparison of Differently Anchored Zirconia-Based DNB-L-





Test solute: trifluoranthryl ethanol. Note that the retention factor ratio is for the less retained isomer.



## Novel Chiral Selector Screening Method

- Attached a CSP to a bare zirconia column insitu.
- Flush the column to remove unbound CSP.
- Screen chiral target compound.
- Strip off CSP using basic conditions.
- Repeat Cycle until desired resolution is achieved using different CSPs.



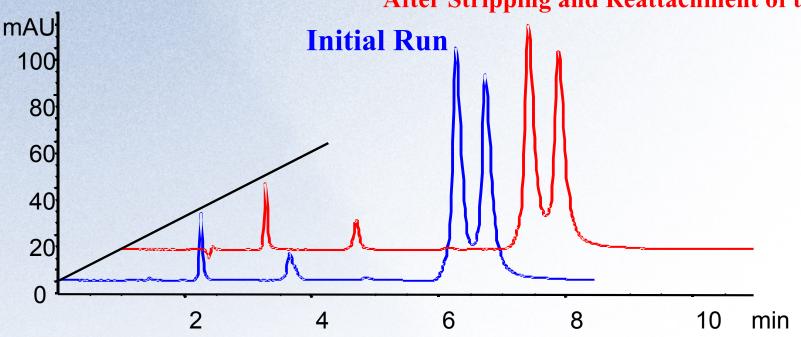
## **Example Attachment and Detachment Cycle**

- Pass a solution of 20 mM N-(4-nitrobenzoyl)-L-glutamic acid (CSP) in tetrahydrofuran for 10 minutes at a column temperature of 60°C and a flow rate of 1 mL/min.
- Flushed column with 100% THF for 10 minutes at 2 mL/min at ambient temperature.
- Separate a racemate solution of (±)-2,2,2-trifluoro-1-(9-anthyl)ethanol.
- Strip the CSP by flushing the column with a 50 mM solution of tetramethylammonium hydroxide solution (pH 12) for 20 minutes at 60°C using a flow rate of 1 mL/min.
- Repeat procedure using the same CSP



### **Proof of Concept**

**After Stripping and Reattachment of the CSP** 



Comparison between the initial and final separation of  $(\pm)$ -2,2,2-trifluoro-1-(9-anthyl)ethanol leucine ester during a single CSP screening cycle.

Chromatographic conditions: mobile phase: 99/1 hexane/IPA; flow rate: 1 ml/min; temperature: 30 °C, solute concentration = 1mg/mL, 5 microliter injection.



#### **Conclusions**

- Flexible attachment chemistry.
- APPA is the best anchor in terms of column stability.
- Zirconia based CSPs have comparable chromatographic performance compared to the commercial silica based CSPs for a wide range of chiral compounds.
- Phase II testing will involve *new anchors and different CSPs* with more optimization of screening techniques.
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