### ผลของความแรงตัวทำละลายต่อพฤติกรรม โครมา โทรกราฟีของสารประกอบ ฟีนอลิกบางชนิดในรีเวิร์สเฟสไฮเพอร์ฟอร์มานซ์ลิควิด โครมา โทกราฟี



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วิทยานิพนธ์นี้เป็นส่วนหนึ่งของการศึกษาตามหลักสูตรปริญญาวิทยาศาสตรมหาบัณฑิต ภาควิชาเคมี บัณฑิตวิทยาลัย จุฬาลงกรณ์มหาวิทยาลัย พ.ศ. 2538

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# OF CERTAIN PHENOLIC COMPOUNDS IN REVERSED PHASE HIGH PERFORMANCE LIQUID CHROMATOGRAPHY

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สมศักดิ์ ศิริไชย : ผลของความแรงตัวทำละลายต่อพฤติกรรมโครมาโทกราฟีของสารประกอบฟีนอ ลิกบางชนิดในฟีนอลิกบางชนิดในรีเวิร์สเฟสไฮเพอร์ฟอร์มานซ์ลิควิดโครมาโทกราฟี (EFFECT OF SOLVENT STRENGTH ON THE CHROMATOGRAPHIC BEHAVIOR OF CERTAIN PHENOLIC COMPOUNDS IN REVERSED PHASE HIGH PERFORMANCE LIQUID CHROMATOGRAPHY) อ.ที่ปรึกษา : รศ.คร. วิทยา เรืองพรวิสุทธิ์, อ.ที่ปรึกษาร่วม : ผศ.คร.สุรพจน์ วงศ์ใหญ่ , 157 หน้า. ISBN 974-632-892-1

ในงานวิจัยครั้งนี้ ได้ศึกษาผลของความแตกต่างของความแรงระหว่างตัวทำละลายสำหรับสารตัวอย่าง และตัวชะต่อพฤติกรรมทางโครมาโทกราฟีของกลุ่มสารพวกฟีนอล คือ พาราเซตามอล ฟีนอล เมธิลพาราเบน เอธิลพาราเบน และโพรพิลพาราเบน บนคอลัมน์ชนิครีเวิร์สเฟสไฮเพอร์ฟอร์มานซ์ลิควิคโครมาโทกราฟีที่ช่วง อุณหภูมิ 20° ถึง 35 ° ซ. คอลัมน์ที่ใช้ในการศึกษาได้แก่ LiChrosorb RP-8 และ LiChrosorb RP-18 โดยมี ขนาคอนุภาค 10 ไมโครเมตร ซึ่งบรรจุอยู่ในคอลัมน์ขนาคเส้นผ่าศูนย์กลางภายใน 4.6 มม. และยาว 25 ซม. โดยที่องค์ประกอบของตัวชะ คือ เมธานอลและน้ำ ในอัตราส่วน 60 ต่อ 40 (โดยปริมาตร)

จากการศึกษาพบว่า จำนวนเพลตตามทฤษฎี (the number of theoretical plates , N) เพิ่มขึ้น เมื่อ เปอร์เซ็นต์ของน้ำในตัวทำละลายสารตัวอย่างเพิ่มขึ้น และมีค่ามากที่สุดเมื่อความเข้มข้นของน้ำในตัวทำละลายมาก กว่า 40 เปอร์เซ็นต์ และพบว่า ผลที่เกิดขึ้นเนื่องจากความแรงของตัวทำละลายจะเพิ่มขึ้นเมื่ออุณหภูมิเพิ่มขึ้น จากการทดลองพบว่า ปริมาณน้ำในตัวทำละลายสารตัวอย่างทำให้จำนวนเพลตตามทฤษฎีเพิ่มขึ้น โดยมีความ สัมพันธ์ระหว่าง N กับ polarity index  $(P_i)$  ของตัวทำละลาย เป็นฟังก์ชันโพลีโนเมียล องศาสอง ตามสมการ  $N = A_0 + A_1 P_i + A_2 P_i^2$  ของทุกอุณหภูมิที่ศึกษา สำหรับรีเทนชันไทม์ (retention time) ของสารที่ศึกษาจะไม่ เปลี่ยนแปลงตามความแรงของตัวทำละลาย พร้อมกันนี้ได้อธิบายผลอันเกิดจากความแรงของตัวทำละลายนี้ในรูป แบบของการเกิด peak distortion และ peak compression

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SOMSAK SIRICHAI: EFFECT OF SOLVENT STRENGTH ON THE CHROMATOGRAPHIC BEHAVIOR OF CERTAIN PHENOLIC COMPOUNDS IN REVERSED PHASE HIGH-PERFORMANCE LIQUID CHROMATOGRAPHY. THESIS ADVISOR: ASSOC.PROF.VITHAYA RUANGPORNVISUTI, Dr.rer.nat., ASSIST.PROF.SURAPOTE WONGYAI, Dr.rer.nat. 157 pp. ISBN 974-632-892-1

The effects resulting from differences between sample solvent and mobile phase composition on chromatographic behavior of paracetamol, phenol, methylparaben, ethylparaben and propylparaben at 20° to 35°C on reversed-phase high-performance liquid chromatography were investigated. Two columns used in this study were LiChrosorb RP-8 (10 µm, 4.6 mm I.D. x 25 cm.) and LiChrosorb RP-18 (10 μm , 4.6 mm I.D. x 25 cm.) with methanol/water (60:40 , v/v) as mobile phase. The number of theoretical plates (N) for the studied compounds increased as the % (v/v) water in dissolving solvent incerased and reached maximum value after 40% (v/v) water concentration. The effect of sample solvent strength was enhanced at higher temperature. At 0 to 40% (v/v) water in dissolving solvent, the number of theoretical plates varied with the solvent polarity index (P<sub>i</sub>) as a second-degree polynomial function, i.e.,  $N = A_0 + A_1P_i + A_2P_i^2$ at all temperature studied. The retention time of each compound was not affected by the The proposed mechanisms of peak distortion and peak compression occuring solvent strength. from this effect are discussed.

ภาควิชา	เคมี	ลายมือชื่อนิสิต	สมศักดิ์ ศิริโชย
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	RP-18 at temperatures 25° and 35°C and polarity index (P <sub>i</sub> ) in the injection
	slovent
4.47	The relationship between number of theoretical plates (N) of ethylparaben in
	RP-18 at temperatures 25° and 35°C and polarity index (P <sub>i</sub> ) in the injection
	slovent
4.48	The relationship between number of theoretical plates (N) of propylparaben in
	RP-18 at temperatures 25° and 35°C and polarity index (P <sub>i</sub> ) in the injection
	slovent
A1	Chromatograms of standard mixtures of phenols dissolved in 2% methanol (a),
	10% methanol (b), 20% methanol (c), 30% methanol (d), 40% methanol (e),
	50% methanol (f), 60% methanol (g), 70% methanol (h), 80% methanol (i),
	90% methanol (j) and pure methanol (k) on reversed-phase column.
	Peak identification and chromatographic conditions as given in Figure 4.984

A2	Chromatograms of standard mixtures of phenols dissolved in 2% methanol (a),
	10% methanol (b), 20% methanol (c), 30% methanol (d), 40% methanol (e),
	50% methanol (f), 60% methanol (g), 70% methanol (h), 80% methanol (i),
	90% methanol (j) and pure methanol (k) on reversed-phase column.
	Peak identification and chromatographic conditions as given in Figure 4.1090
A3	Chromatograms of standard mixtures of phenols dissolved in 2% methanol (a),
	10% methanol (b), 20% methanol (c), 30% methanol (d), 40% methanol (e),
	50% methanol (f), 60% methanol (g), 70% methanol (h), 80% methanol (i),
	90% methanol (j) and pure methanol (k) on reversed-phase column.
	Peak identification and chromatographic conditions as given in Figure 4.1196
A4	Chromatograms of phenol where phenol dissolved in 2% methanol (a),
	10% methanol (b), 20% methanol (c), 30% methanol (d), 40% methanol (e),
	50% methanol (f), 60% methanol (g), 70% methanol (h), 80% methanol (i),
	90% methanol (j) and pure methanol (k) on reversed-phase column.
	Chromatographic conditions as given in Figure 4.12101
A5	Chromatograms of standard mixtures of phenols dissolved in 2% methanol (a),
	10% methanol (b), 20% methanol (c), 30% methanol (d), 40% methanol (e),
	50% methanol (f), 60% methanol (g), 70% methanol (h), 80% methanol (i),
	90% methanol (j) and pure methanol (k) on reversed-phase column.
	Peak identification and chromatographic conditions as given in Figure 4.13105

A6	Chromatograms of phenol where phenol dissolved in 2% methanol (a),
	10% methanol (b), 20% methanol (c), 30% methanol (d), 40% methanol (e),
	50% methanol (f), 60% methanol (g), 70% methanol (h), 80% methanol (i),
	90% methanol (j) and pure methanol (k) on reversed-phase column.
	Chromatographic conditions as given in Figure 4.14110
A7	Chromatograms of standard mixtures of phenols dissolved in 2% methanol (a),
	10% methanol (b), 20% methanol (c), 30% methanol (d), 40% methanol (e),
	50% methanol (f), 60% methanol (g), 70% methanol (h), 80% methanol (i),
	90% methanol (j) and pure methanol (k) on reversed-phase column.
	Peak identification and chromatographic conditions as given in Figure 4.15114
A8	Chromatograms of standard mixtures of phenols dissolved in 2% methanol (a),
	10% methanol (b), 20% methanol (c), 30% methanol (d), 40% methanol (e),
	50% methanol (f), 60% methanol (g), 70% methanol (h), 80% methanol (i),
	90% methanol (j) and pure methanol (k) on reversed-phase column.
	Peak identification and chromatographic conditions as given in Figure 4.16120
A9	Chromatograms of standard mixtures of phenols dissolved in 2% methanol (a),
	10% methanol (b), 20% methanol (c), 30% methanol (d), 40% methanol (e),
	50% methanol (f), 60% methanol (g), 70% methanol (h), 80% methanol (i),
	90% methanol (j) and pure methanol (k) on reversed-phase column.
	Peak identification and chromatographic conditions as given in Figure 4.17126

A10	Chromatograms of standard mixtures of phenols dissolved in 2% methanol (a),
	10% methanol (b), 20% methanol (c), 30% methanol (d), 40% methanol (e),
	50% methanol (f), 60% methanol (g), 70% methanol (h), 80% methanol (i),
	90% methanol (j) and pure methanol (k) on reversed-phase column.
	Peak identification and chromatographic conditions as given in Figure 4.18