

High Performance Liquid Chromatography (HPLC) is a suitable technique for analysis of harmful food additives and contaminants. COSMOSIL ODS and special columns offer improved separation for a large variety of these substances.

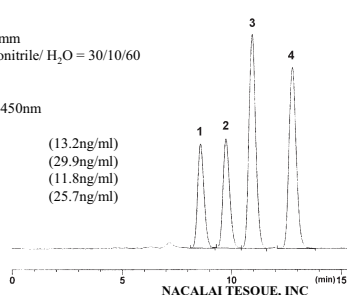
(1) Aflatoxins

COSMOSIL Application Data

Column: 5C₁₈-AR-II
 Column size: 4.6mm I.D.-150mm
 Mobile phase: Methanol/ Acetonitrile/ H₂O = 30/10/60
 Flow rate: 0.8 ml/min
 Temperature: 40°C
 Detection: Ex.365nm Em.450nm

Sample: 1; Aflatoxin G2 (13.2ng/ml)
 2; Aflatoxin G1 (29.9ng/ml)
 3; Aflatoxin B2 (11.8ng/ml)
 4; Aflatoxin B1 (25.7ng/ml)

Inj.Vol: 20 μl



Data courtesy of a customer

AP-1148

Aflatoxins are naturally occurring mycotoxins. They can colonize and contaminate grains before harvesting or during storage. Aflatoxins are among the most carcinogenic substances known. COSMOSIL 5C₁₈-AR-II can be used to separate aflatoxins.



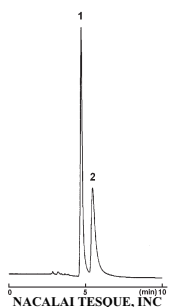
(2) 2- and 4-methylimidazole (2-MI and 4-MI)

COSMOSIL Application Data

Column: HILIC
 Column size: 4.6mm I.D.-250mm
 Mobile phase: Acetonitrile/ 50mmol/l Ammonium Acetate = 90/10
 Flow rate: 1.0 ml/min
 Temperature: 30°C
 Detection: UV220nm

Sample: 1; 4-Methylimidazole (0.25mg/ml)
 2; 2-Methylimidazole (0.25mg/ml)

Inj.Vol.: 1.0μl



AP-1203

2- and 4-methylimidazole (2-MI and 4-MI) are by-products formed during the manufacturing of caramel coloring used in popular soft drinks and foods. These two chemicals are selected by the National Cancer Institute for a long-term study because of the high potential for human exposure. The gas chromatography and reversed-phase HPLC columns with ion-pairing reagent have been traditionally used, but COSMOSIL HILIC offers complete separation of 2- and 4-methylimidazole without ion-pairing reagent.



(3) Melamines

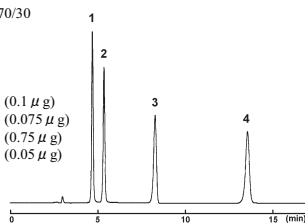
After recent melamine contamination in pet food and dairy products, determination of melamine has become a high priority. COSMOSIL HILIC offers excellent separation of melamine and cyanuric acid. They form a crystalline complex that is more toxic than either one alone.

• Melamines

COSMOSIL Application Data

Column: HILIC
 Column size: 4.6mm I.D.-250mm
 Mobile phase: Acetonitrile/ 10mmol/l Ammonium Acetate = 70/30
 Flow rate: 1.0 ml/min
 Temperature: 30°C
 Detection: UV225nm

Sample: 1; Melamine (0.1 μg)
 2; Ammeline (0.075 μg)
 3; Cyanuric Acid (0.75 μg)
 4; Ammelide (0.05 μg)



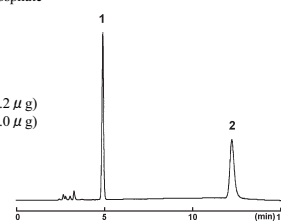
AP-1076

• Oxalic Acids

COSMOSIL Application Data

Column: HILIC
 Column size: 4.6mm I.D.-250mm
 Mobile phase: Acetonitrile/ 10mmol/l Phosphate buffer(pH7.0) = 50/50
 Flow rate: 1.0 ml/min
 Temperature: 30°C
 Detection: UV210nm

Sample: 1; Oxamic Acid (0.2 μg)
 2; Oxalic Acid (1.0 μg)

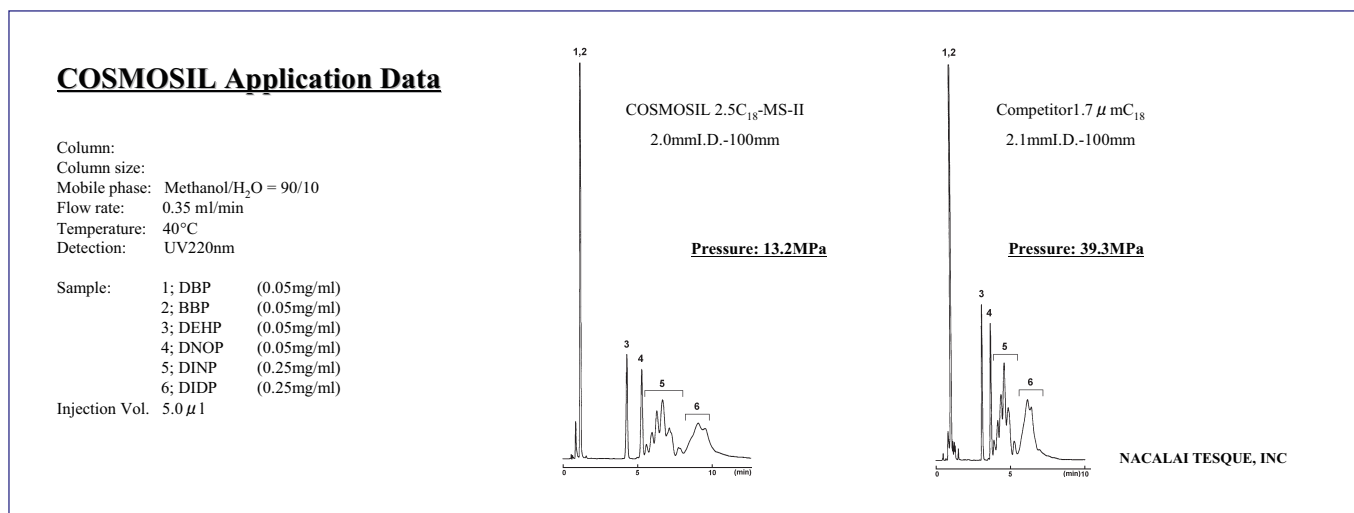


AP-0307

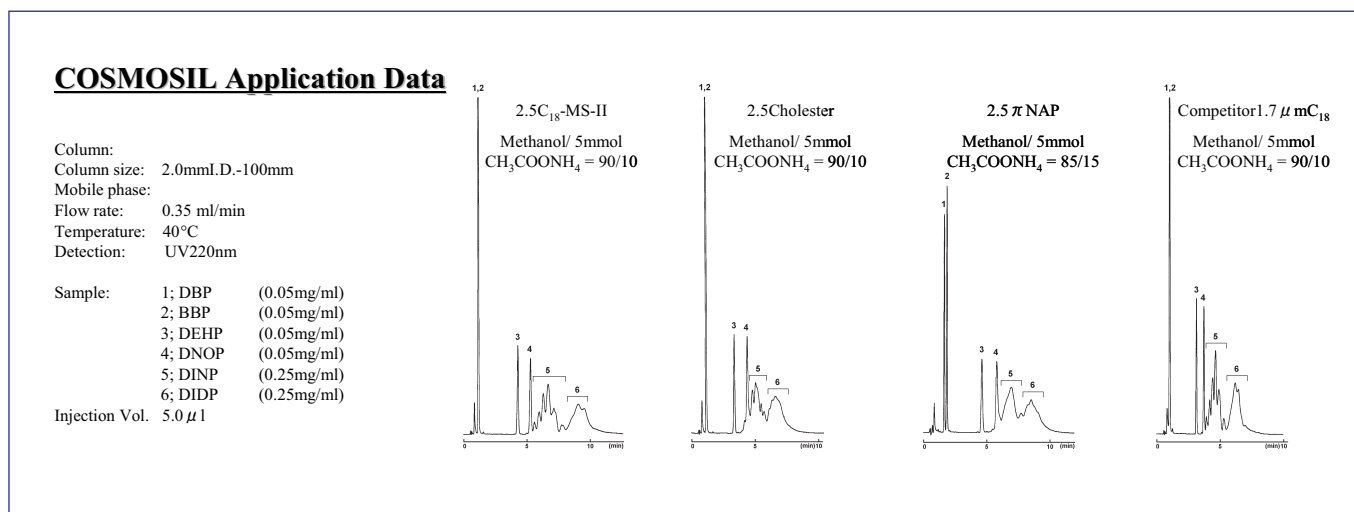
(4) DEHP (Bis(2-ethylhexyl)phthalate)

A few years ago, news reported that DEHP (Bis(2-ethylhexyl)phthalate), a plasticizer, has contaminated the food supply. If taken into the human body in large amounts, DEHP can cause cancer in addition to the risk of liver function damages.

COSMOSIL 2.5C₁₈-MS-II produces an equivalent chromatogram compared with a competitor's 1.7µm column. 2.5C₁₈-MS-II has longer retention time. More importantly, 2.5C₁₈-MS-II operates about 1/3 the pressure of competitor's 1.7 µm column, putting it within the range of conventional HPLC equipment.



COSMOSIL 2.5πNAP enables separation of DBP(Dibutyl Phthalate)(Sample 1) and BBP (Butyl Benzyl Phtharate)(Sample 2) that are difficult to separate with C₁₈ columns.



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