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APPENDICES

Appendix A Investigation of Characteristic Peaks of FT-IR Spectrum of Undoped and Doped Poly(p-phenylene)

FT-IR spectrometer (Nicolet/Nexus 670) was used to characterize functional groups of undoped and doped poly(p-phenylene) at 50:1 mole ratio of dopant to moomer unit (uPPP and 50:1 dPPP) and operated in the absorption mode in a wavenumber range of 4000-400 cm⁻¹ with 32 scans and ±4 cm⁻¹ resolution. Optical grade potassium bromide (KBr) was used as a background material and dried before being mixed with PPP at a ratio of 20:1.

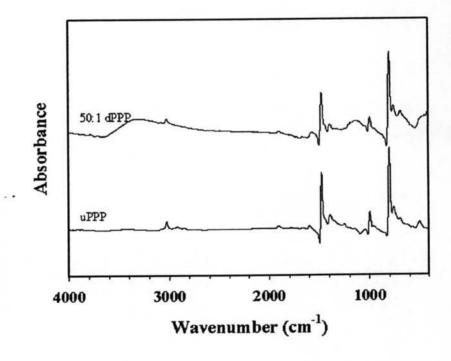


Figure A1 The FT-IR spectra of uPPP and 50:1 dPPP.

The FT-IR spectrum of PPP synthesized by Kovacic's direct route (Kovacic et al., 1963) through oxidative cationic polymerization was shown in Figure A-1. The characteristic absorption peak situated at 805 cm⁻¹ was corresponding to para aromatic substitution. The less intense absorption peaks occurred at 763 and 695 cm⁻¹ were assignable to mono substitution of benzene rings. Moreover, there were additional bands of secondary intensity of para band located at 999, 1396, and 1479 cm⁻¹ (Kovacic et al., 1963). The several precise assignments of characteristic FT-IR spectrum for PPP were summarized in Table A-1 (Kvarnstrom et al., 1997; Lacaze et al. 1997).

The doping of PPP caused the occurrence of new absorption peaks at 1545 and 1180 cm⁻¹that were not appeared in the undoped state of PPP. The doping induced IR characteristic peaks were not due to specific vibrations of dopant molecules or between the molecules and the polymer chain, because the similar peaks appeared for K-, AsF₅- and FeCl₃-doping, therefore they are intrinsic vibration of the polymer chain in doping state (Yli-Lahti et al., 1985; Shacklette et al., 1980; Yaniger et al., 1984).

Table A1 Typical IR assignments of uPPP

Wavenumber (cm ⁻¹)	Characteristic bands	References Dale, 1957 and Castiglioni et al., 1989	
3026±0.6 [3029]	C-H stretching of aromatic		
1599±0.6 [1600]	Quinoid structure	Pham et al., 1990	
1396±2.0, 1479±0.4 [1390, 1480]	C-C stretching of aromatic	Pham et al., 1990	
999±0.5, 1250±4.0 [1000, 1255]	C-H in-plane vibration of para disubstituted phenyl rings	Soubiran et al., 1998	
805±0.6 [817-805]	C-H out of plane vibration of para disubstituted phenyl rings	Froyer et al., 1985	
763±2.0 [760]	C-H out of plane vibration mono substituted phenyl rings	Kvarnstrom et al. 1985	
695±1.0 [692]	C-H out of plane vibration of mono substituted phenyl rings	Rakovic et al., 1991	

Appendix B Thermal Property of uPPP and 50:1 dPPP

The thermal property of uPPP and 50:1 dPPP was examined by a thermal gravimetric technique. Thermal Gravimetric Analyzer (duPont/TGA 2950) was used in operating mode: heating rate of 10 °C/min and temperature scan from 30-800 °C under air atmosphere. There was single transition step of both uPPP and dPPP decomposition under air atmosphere. Single transition of uPPP and dPPP decomposition under air atmosphere was noticed as shown in Figure B-1. It was obviously observed that uPPP is thermally and thermooxidatively stable. There was no significant decomposition appeared below 400 °C. It meant that at temperature up to 400 °C uPPP demonstrated good resistance to thermal degradation and air oxidation (Kovacic *et al.*, 1963). At approximately 569 °C PPP backbone began to decompose responsible for the loss of benzene.

dPPP proposed less thermal stability with gradually decomposed when temperature increased, than the undoped one, because doping induced the defect in the polymer chain. However, dPPP still withstand the heat until most of dPPP molecule decomposed at around 480 °C. Moreover, after decomposition dPPP had higher amount of residue than uPPP due to the addition of ferric chloride dopant.

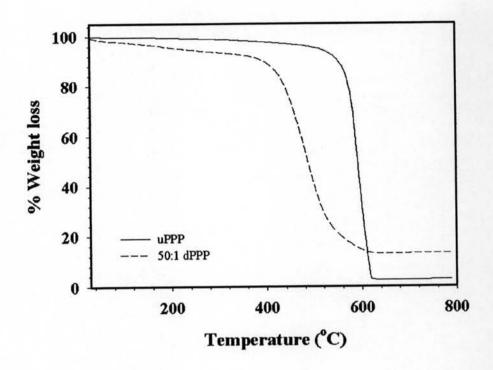


Figure B1 TGA thermogram of uPPP and 50:1 dPPP.

Table B1 Decomposition of uPPP and 50:1 dPPP

Sample	T _d (°C)	% Weight loss	% Residue
uPPP	569±4	98.04±0.83	1.96±0.83
50:1 dPPP	480±6	86.95±1.10	13.05±1.10

Appendix C Identification of Crystallinity of uPPP and 50:1 dPPP and Structure of Zeolite

X-ray Diffractrometer (Phillips, Rigaku) was used in order to identify the crystallinity of uPPP and 50:1 dPPP and ensure the structure of zeolite. Both uPPP and dPPP powder were compressed into pellet form with 10 mm in diameter and 0.15-0.25 mm in thickness. Zeolite powder was packed into a glass plate. The XRD patterns showed that uPPP had quite high crystalline accompanying with the d-spacing value of 4.53, 3.63 and 3.24 Å. The most intense d-spacing was 4.53 Å which closely corresponded to the length of a phenyl unit (Kovacic et al., 1963; Marvel et al., 1959). Therefore, it was postulated that the rings were very nearly coplanar. However, the pattern of uPPP did show the existence of some amorphous regions.

Some peaks presenting in the uPPP pattern were disappeared or reduced in intensity after doping with FeCl₃ which meant FeCl₃ dopant can decease the crystallinity of PPP. It might be due to the distortion of the polymer chain after being doped.

By using XRD technique, the structure and crystallinity ZSM-5 was confirmed. XRD patterns and peak positions matched those observed for the characteristic structure of MFI. The peak intensities and low background lines demonstrated the high crystallinity of the sample.

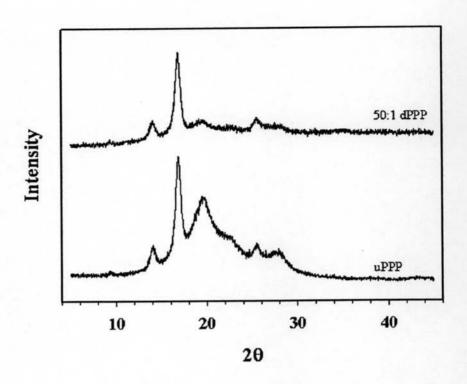


Figure C1 XRD pattern of uPPP and 50:1 dPPP.

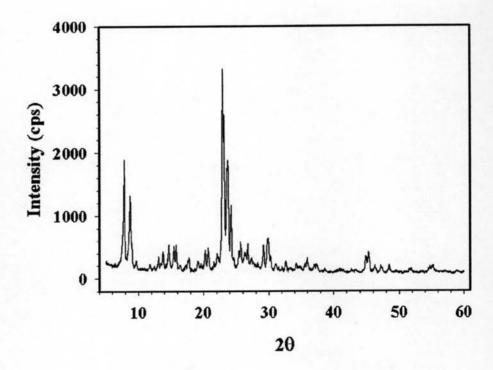


Figure C2 XRD pattern of ZSM-5 with SiO₂/Al₂O₃ of 23.

Appendix D Determination of Particle Size and Particle Size Distribution of PPP and Zeolite Powder

Particle Size Analyser (Malvern/Mastersizer X Version 2.15) was employed to examine the particle size and particle size distribution of materials. Synthesized PPP was ground and subsequently sieved with 54 μm sieve before being analyzed.

Table D1 Mean particle diameter of materials

		Mean p	article diame	eter (µm)	
Materials -	1	2	3	Avg.	SD
PPP	33.74	33.41	33.69	33.61	0.18
ZSM-5(23)	5.46	5.45	5.47	5.46	0.01

Table D2 Raw data of particle size of uPPP

Parti	cle size			uF	PP		
	ter (µm)		1		2		3
Size low	Size high	In %	Under %	In %	Under %	In %	Under %
0.20	0.48	0.11	0.11	0.11	0.11	0.11	0.11
0.48	0.59	0.33	0.43	0.32	0.43	0.34	0.40
0.59	0.71	0.45	0.88	0.45	0.88	0.47	0.91
0.71	0.86	0.45	1.33	0.44	1.32	0.46	1.37
0.86	1.04	0.34	1.67	0.33	1.65	0.35	1.72
1.04	1.26	0.19	1.86	0.19	1.84	0.20	1.92
1.26	1.52	0.10	1.97	0.10	1.94	0.11	2.03
1.52	1.84	0.12	2.09	0.12	2.06	0.13	2.16
1.84	2.23	0.27	2.36	0.26	2.33	0.27	2.43
2.23	2.70	0.48	2.84	0.48	2.81	0.50	2.93
2.70	3.27	0.68	3.52	0.68	3.49	0.70	3.63
3.27	3.95	0.90	4.42	0.90	4.40	0.93	4.56
3.95	4.79	1.18	5.60	1.18	5.58	1.21	5.77
4.79	5.79	1.51	7.11	1.52	7.09	1.55	7.31
5.79	7.01	1.91	9.03	1.92	9.01	1.95	9.26
7.01	8.48	2.36	11.39	2.36	11.37	2.39	11.66
8.48	10.27	2.86	14.24	2.87	14.25	2.90	14.56
10.27	12.43	3.45	17.70	3.48	17.73	3.51	18.07
12.43	15.05	4.23	21.92	4.29	22.02	4.31	22.38
15.05	18.21	5.32	27.25	5.40	27.42	5.41	27.80
18.21	22.04	6.80	34.06	6.89	34.31	6.87	43.67
22.04	26.68	8.58	42.63	8.66	42.97	8.57	43.23
26.68	32.29	10.34	52.98	10.41	53.38	10.22	53.46
32.29	39.08	11.52	64.49	11.57	64.95	11.32	64.77
39.08	47.30	11.51	75.99	11.53	76.47	11.30	76.06
47.30	57.25	9.83	85.82	9.81	86.28	9.67	85.72
57.25	69.30	7.10	92.93	7.04	93.32	7.00	92.73
69.30	83.87	4.25	97.17	4.15	97.47	4.23	96.96
83.87	101.52	2.06	99.23	1.94	99.42	2.11	99.08
101.52	122.87	0.73	99.97	0.58	100.00	0.82	99.90
122.87	148.72	0.03	100.00	0.00	100.00	0.10	100.00
148.72	180.00	0.00	100.00	0.00	100.00	0.00	100.00

Table D3 Raw data of particle size of ZSM-5(23)

Parti	cle size			ZSM-5(23)					
diameter (µm)			1		2	3			
Size low	Size high	In %	Under %	In %	Under %	In %	Under %		
0.05	0.12	0.00	0.00	0.00	0.00	0.00	0.00		
0.12	0.15	0.00	0.00	0.00	0.00	0.00	0.00		
0.15	0.19	0.00	0.00	0.00	0.00	0.00	0.00		
0.19	0.23	0.00	0.00	0.00	0.00	0.00	0.00		
0.23	0.28	0.00	0.00	0.00	0.00	0.00	0.00		
0.28	0.35	0.00	0.00	0.00	0.00	0.00	0.00		
0.35	0.43	0.00	0.00	0.00	0.00	0.00	0.00		
0.43	0.53	0.05	0.05	0.05	0.06	0.06	0.06		
0.53	0.65	0.30	0.35	0.30	0.36	0.31	0.37		
0.65	0.81	0.68	1.03	0.70	1.06	0.71	1.08		
0.81	1.00	1.23	2.26	1.25	2.31	1.28	2.35		
1.00	1.23	1.92	4.18	1.96	4.27	1.99	4.34		
1.23	1.51	2.70	6.88	2.72	6.99	2.73	7.07		
1.51	1.86	3.37	10.25	3.36	10.34	3.33	10.40		
1.86	2.30	3.84	14.09	3.79	14.14	3.74	14.14		
2.30	2.83	4.43	18.52	4.41	18.54	4.40	18.53		
2.83	3.49	6.12	24.66	6.17	24.73	6.25	24.79		
3.49	4.30	9.35	34.00	9.47	34.19	9.60	34.38		
4.30	5.29	14.17	48.19	14.25	48.45	14.25	48.64		
5.29	6.52	18.51	66.67	18.37	66.80	18.05	66.67		
6.52	8.04	17.38	84.04	17.17	83.96	16.84	83.49		
8.04	9.91	10.69	94.71	10.66	94.60	10.74	94.21		
9.91	12.21	4.27	99.01	4.35	98.98	4.60	98.85		
12.21	15.04	0.98	100.00	1.02	100.00	1.14	100.00		
15.04	18.54	0.00	100.00	0.00	100.00	0.00	100.00		
18.54	22.84	0.00	100.00	0.00	100.00	0.00	100.00		
22.84	28.15	0.00	100.00	0.00	100.00	0.00	100.00		
28.15	34.69	0.00	100.00	0.00	100.00	0.00	100.00		
34.69	42.75	0.00	100.00	0.00	100.00	0.00	100.00		
42.75	52.68	0.00	100.00	0.00	100.00	0.00	100.00		
52.68	64.92	0.00	100.00	0.00	100.00	0.00	100.00		
64.92	80.00	0.00	100.00	0.00	100.00	0.00	100.00		

Appendix E Density Measurement

Density of PPP and zeolite can be measured by using pycnometer at controlled temperature. Blank pycnometer was firstly weighed and then water was added into the pycnometer and weight of pycnometer with water was measured again. The density of water can be calculated through equation (E1)

$$\rho_{w} = \frac{(a-b)}{25} \tag{E1}$$

where ρ_w is the density of water (g.cm⁻³), a is the weight of pycnometer with water (g), b is the weight of blank pycnometer (g).

The blank pycnometer was weighed again and then sample powder was introduced into the pycnometer and the total weight was measured. After that the water was added into the same pycnometer and again the total weight was measured. In this step, the volume of added water will be obtained from equation (E2) and finally the density of the sample can be calculated through equation (E3)

$$A = \frac{(e - d)}{\rho_{w}} \tag{E2}$$

where A is the volume of added water (cm³), e is the weight of pycnometer with sample and added water (g), d is the weight of pycnometer with sample (g).

$$\rho_m = \frac{(d-b)}{25-A} \tag{E3}$$

where ρ_m is the density of material (g.cm⁻³).

Table E1 Density of materials

Materials	Density(g.cm ⁻³)	
PPP	1.3273±0.0011	
ZSM-5(23)	1.9739±0.0004	

Appendix F Morphology

Scanning Electron Microscope (SEM) (JEOL/JSM 5200) was utilized to examine the morphology of materials. Samples were coated with gold before being characterized in order to suppress the charge up phenomena. The morphology of PPP (Figure F-1) shows rough surface and irregular shapes.

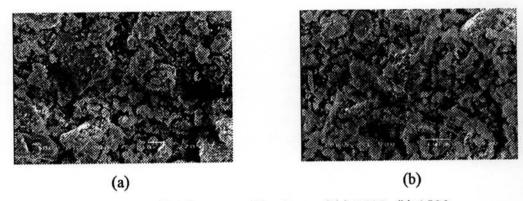


Figure F1 Morphology of PPP at magnifications of (a) 1000; (b) 1500.

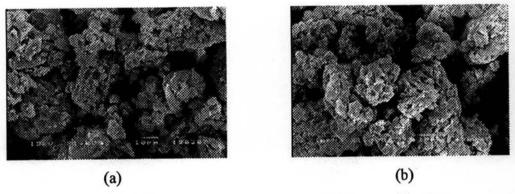


Figure F2 Morphology of dPPP at magnifications of (a) 1000; (b) 1500.

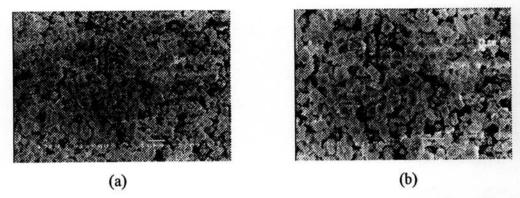


Figure F3 Morphology of ZSM-5(23) at magnifications of (a) 1000; (b) 1500.

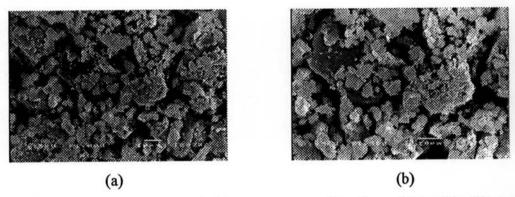


Figure F4 Morphology of dPPP(90)/NaZ23 at magnifications of (a) 1000; (b) 1500.

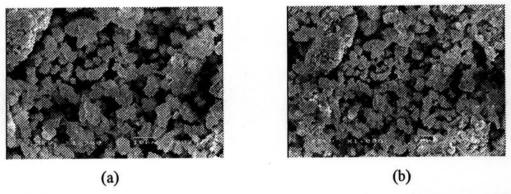


Figure F5 Morphology of dPPP(80)/NaZ23 at magnifications of (a) 1000; (b) 1500.

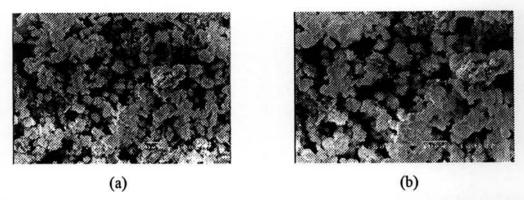


Figure F6 Morphology of dPPP(70)/NaZ23 at magnifications of (a) 1000; (b) 1500.

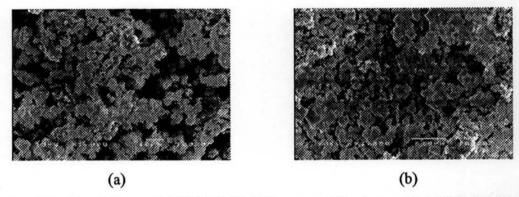


Figure F7 Morphology of dPPP(70)/NaZ23 at magnifications of (a) 1000; (b) 2000.

Appendix G Identification of Si/Al and Cation Exchange Level of Zeolite

An Atomic Absorption Spectrometer (AAS) (Varian/SpectrAA 300) was used to determine the amount of Si and Al containing in zeolite and the amount of desired cation after ion exchange processes. The known weight zeolite was dissolved with hydrofluoric acid (HF) and hydrochloric acid (HCl). After the volume of solution was adjusted, the solution was filtered before being tested. Standard, blank and sample solutions were prepared for testing of each cation and different lamps were used for different cations.

The Si/Al ratio of zeolite can be calculated via equation (G1)

$$Si/Al \ ratio = \frac{mole \ of \ Si/g \ of \ zeolite}{mole \ of \ Al/g \ of \ zeolite}$$
 (G1)

Table G1 Si/Al ratios of zeolite

Zeolite -	Cation content	Si/Al	
	Si	Al	SIAI
ZSM-5(23)	1.3849×10^{-2}	1.0930×10^{-3}	12.67

Before performing cation exchange process, ZSM-5 zeolite contains the NH₄⁺ cation. After the process, cation containing in the ZSM-5 was changed and the actual amount of balancing cation was then determined. Cation exchange level can be calculated on the assumption that 100% cation exchange possesses identical mole number of incoming cation and Al as followed equation (G-2)

Cation exchange level =
$$\frac{\text{mole of cation/g of zeolite}}{\text{mole of Al/g of zeolite}}$$
 (G-2)

Table G2 The cation exchange level of NH₄ZSM-5(23) with Na⁺ and K⁺

Sample	Cation	Times	[Al ₃ ⁺] (mmol/g zeolite)	[cation] (mmol/g zeolite)	% Exchange
ZSM-5(23)	Na ⁺	1 st	1.0978	1.0674	97.23
	K ⁺	1 st	1.0978	1.0576	96.34

Appendix H Surface Area and Pore Volume of Zeolite

Surface area and pore volume of zeolite were determined by employing a Surface Area Analyzer (SAA) (Quantachrome/Autosorb-1). Dried zeolite was weighed and out gassed at 250 °C over night before being performed adsorption and desorption with He and N₂ gases.

Table H1 Surface area and pore volume of zeolite

Zeolite	BET Surface area (m²/g)	Pore volume (cm ³ /g)	
NH ₄ ZSM-5(23)	290.1±0.85	0.1819±0.0031	
HZSM-5(23)	332.6±6.51	0.2075±0.0006	
NaZSM-5(23)	283.1±8.63	0.1759±0.0112	
KZSM-5(23)	273.3±1.34	0.1663±0.0074	

It was well-known that if the zeolite contains different cation, the surface area and pore volume of the zeolite are also different due to the various types of cation whose size of cation are not the same. From table G-1, the surface area and pore volume of ammonium cation form existing in ZSM-5 are 290.1m²/g and 0.1819 cm³/g, respectively. After calcination without passing cation exchanged process, ammonium cation became H⁺ because of the removal of NH₃ during calcination process, therefore the surface are and pore volume of zeolite increase up to 332.6 m²/g and 0.2075 cm³/g, respectively. Due to K⁺ possesses the larger cation size than Na⁺, thus ZSM-5 containing K⁺ have less surface area and pore volume than NaZSM-5.

Appendix I Correction Factor (K) Measurement

A two point probe meter connected with a source power supplier (Keithley/Model 6517A) was employed to determine the electrical conductivity of materials. A constant voltage is applied and a current is simultaneously measured.

According to the geometric effects of the probe, the geometrical correction factor was taken, depending on the configuration and probe tip spacing

$$K = \frac{w}{I} \tag{I1}$$

where K is geometric correction factor, w is width of probe tip spacing (cm), l is length between probes (cm).

The constant K can be determined by using standard materials whose specific resistivity values are known. In our case, silicon wafer chips were used as the standard materials. The resistance was measured by using our custom made two-point probe, obtained by applying various voltages and simultaneously measuring currents. The geometric correction factor was calculated via equation (I2)

$$K = \frac{\rho}{R \times t} = \frac{I \times \rho}{V \times t} \tag{12}$$

where ρ is resistivity of standard silicon wafer (Ω .cm), which was calibrated by using four point probe at Thailand Microelectronic Centre (TMEC), R is resistance of film (Ω .), t is film thickness (cm), I is measured current (A) and V is applied voltage (V).

Table I1 The correction factor of several two point probes.

Probe	Correction Factor (K)							
	1	2	3	Avg.	SD			
1	3.05E-05	2.785E-05	7.445E-06	2.195E-05	1.902E-06			
2	7.422E-05	9.872E-06	4.200E-05	4.205E-05	4.550E-05			
3	1.506E-05	4.847E-05	3.180E-05	3.177E-05	2.362E-05			
4	8.063E-06	8.213E-06	8.260E-06	8.178E-06	1.028E-07			
5	2.385E-05	2.221E-05	2.755E-05	2.454E-05	2.735E-06			
6	3.501E-05	3.453E-05	3.488E-05	3.481E-05	2.495E-07			

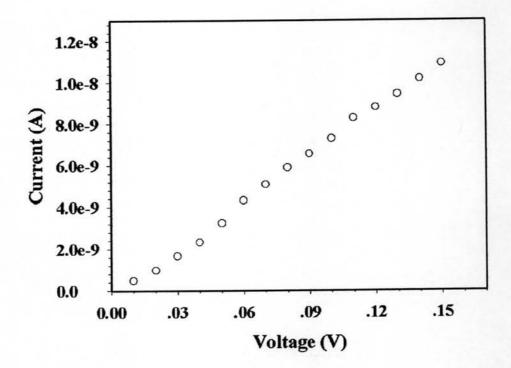


Figure I1 Voltage vs. current data of the probe calibration with Si-wafer whose sheet resistivity of 107.373 Ω /sq, at 24-25 °C, 50-60 %RH.

Table 12 Voltage-current data of the probe 1 calibration with Si-wafer whose sheet resistivity of 107.373 Ω /sq, at 24-25 °C, 50-60 %RH

Appli	ed Voltag	ge (V)		Current (A)	
1	2	3	1	2	3
0.01	0.01	0.01	5.80E-10	4.48E-10	4.25E-10
0.02	0.02	0.02	1.16E-09	8.97E-10	8.51E-10
0.03	0.03	0.03	1.70E-09	1.60E-09	1.64E-09
0.04	0.04	0.04	2.24E-09	2.30E-09	2.43E-09
0.05	0.05	0.05	3.28E-09	3.17E-09	3.25E-09
0.06	0.06	0.06	4.23E-09	4.32E-09	4.39E-09
0.07	0.07	0.07	5.06E-09	5.08E-09	5.09E-09
0.08	0.08	0.08	6.13E-09	5.96E-09	5.60E-09
0.09	0.09	0.09	6.63E-09	6.61E-09	6.42E-09
0.10	0.10	0.10	7.28E-09	7.28E-09	7.34E-09
0.11	0.11	0.11	8.33E-09	8.25E-09	8.25E-09
0.12	0.12	0.12	8.90E-09	8.75E-09	8.71E-09
0.13	0.13	0.13	9.48E-09	9.36E-09	9.42E-09
0.14	0.14	0.14	1.02E-08	1.01E-08	1.02E-08
0.15	0.15	0.15	1.10E-08	1.07E-08	1.10E-08

Table I3 Voltage-current data of the probe 2 calibration with Si-wafer whose sheet resistivity of 107.373 Ω /sq, at 24-25 °C, 50-60 %RH

Appli	ied Voltag	ge (V)		Current (A)	
1	2	3	1	2	3
0.05	0.05	0.05	3.64E-08	3.49E-08	3.40E-08
0.10	0.10	0.10	7.17E-08	7.10E-08	6.66E-08
0.15	0.15	0.15	1.00E-07	9.98E-08	1.00E-07
0.20	0.20	0.20	1.33E-07	1.27E-07	1.22E-07
0.25	0.25	0.25	1.62E-07	1.64E-07	1.50E-07
0.30	0.30	0.30	1.93E-07	1.94E-07	1.89E-07
0.35	0.35	0.35	2.38E-07	2.40E-07	2.39E-07
0.40	0.40	0.40	2.87E-07	2.79E-07	2.78E-07
0.45	0.45	0.45	3.23E-07	3.23E-07	3.12E-07
0.50	0.50	0.50	3.63E-07	3.65E-07	3.53E-07
0.55	0.55	0.55	4.01E-07	3.95E-07	3.85E-07
0.60	0.60	0.60	4.35E-07	4.34E-07	4.30E-07
0.65	0.65	0.65	4.85E-07	4.78E-07	4.73E-07
0.70	0.70	0.70	5.04E-07	4.99E-07	4.94E-07

Table I4 Voltage-current data of the probe 3 calibration with Si-wafer whose sheet resistivity of 107.373 Ω /sq, at 24-25 °C, 50-60 %RH

Appli	ied Voltag	ge (V)		Current (A)	
1	2	3	1	2	3
0.05	0.05	0.05	5.91E-09	5.89E-09	6.01E-09
0.10	0.10	0.10	1.45E-08	1.42E-08	1.40E-08
0.15	0.15	0.15	2.36E-08	2.31E-08	2.29E-08
0.20	0.20	0.20	3.16E-08	3.02E-08	2.98E-08
0.25	0.25	0.25	3.72E-08	3.63E-08	3.59E-08
0.30	0.30	0.30	4.38E-08	4.22E-08	4.19E-08
0.35	0.35	0.35	4.94E-08	4.80E-08	4.81E-08
0.40	0.40	0.40	5.61E-08	5.51E-08	5.48E-08
0.45	0.45	0.45	6.14E-08	6.08E-08	6.08E-08
0.50	0.50	0.50	6.80E-08	6.76E-08	6.76E-08
0.55	0.55	0.55	7.54E-08	7.38E-08	7.36E-08
0.60	0.60	0.60	8.14E-08	8.22E-08	8.12E-08
0.65	0.65	0.65	9.09E-08	9.18E-08	9.12E-08
0.70	0.70	0.70	1.04E-07	1.03E-07	1.03E-07

Table I5 Voltage-current data of the probe 4 calibration with Si-wafer whose sheet resistivity of 107.373 Ω /sq, at 24-25 °C, 50-60 %RH

Appli	ied Voltag	ge (V)		Current (A)	
1	2	3	1	2	3
0.01	0.01	0.01	8.74E-10	8.41E-10	8.68E-10
0.02	0.02	0.02	1.09E-09	1.06E-09	1.05E-09
0.03	0.03	0.03	1.33E-09	1.40E-09	1.33E-09
0.04	0.04	0.04	2.54E-09	2.58E-09	2.54E-09
0.05	0.05	0.05	3.07E-09	3.07E-09	3.26E-09
0.06	0.06	0.06	4.83E-09	4.87E-09	5.22E-09
0.07	0.07	0.07	5.32E-09	5.43E-09	5.28E-09
0.08	0.08	0.08	6.14E-09	6.39E-09	6.62E-09
0.09	0.09	0.09	7.83E-09	8.03E-09	7.63E-09
0.10	0.10	0.10	8.62E-09	8.75E-09	8.24E-09
0.11	0.11	0.11	9.58E-09	8.91E-09	8.85E-09
0.12	0.12	0.12	9.52E-09	9.69E-09	9.65E-09
0.13	0.13	0.13	1.11E-08	1.15E-08	1.26E-08
0.14	0.14	0.14	1.35E-08	1.34E-08	1.38E-08
0.15	0.15	0.15	1.39E-08	1.42E-08	1.43E-08

Table I6 Voltage-current data of the probe 5 calibration with Si-wafer whose sheet resistivity of 107.373 Ω /sq, at 24-25 °C, 50-60 %RH

Appl	ied Voltag	ge (V)		Current (A)	
1	2	3	1	2	3
0.005	0.005	0.005	7.03E-10	5.68E-10	9.56E-10
0.01	0.01	0.01	2.33E-09	2.22E-09	2.52E-09
0.02	0.02	0.02	3.98E-09	4.32E-09	6.68E-09
0.03	0.03	0.03	7.16E-09	6.70E-09	6.68E-09
0.04	0.04	0.04	9.36E-09	8.15E-09	9.38E-09
0.05	0.05	0.05	1.44E-08	1.31E-08	1.53E-08

Table I7 Voltage-current data of the probe 6 calibration with Si-wafer whose sheet resistivity of 107.373 Ω /sq, at 24-25 °C, 50-60 %RH

Appl	Applied Voltage (V)			Current (A)	
1	2	3	1	2	3
0.005	0.005	0.005	1.20E-09	1.24E-09	1.31E-09
0.010	0.010	0.010	3.85E-09	3.61E-09	3.65E-09
0.015	0.015	0.015	4.75E-09	4.85E-09	5.01E-09
0.020	0.020	0.020	5.97E-09	6.03E-09	5.82E-09
0.025	0.025	0.025	7.07E-09	7.13E-09	7.37E-09
0.030	0.030	0.030	9.63E-09	9.69E-09	9.72E-09
0.035	0.035	0.035	1.24E-08	1.23E-08	1.29E-08
0.040	0.040	0.040	1.42E-08	1.39E-08	1.42E-08
0.045	0.045	0.045	1.72E-08	1.59E-08	1.47E-08

Appendix J Conductivity Measurement

The electrical conductivity (σ) can be measured by using the two-point probe mater connected with a voltage supplier (Keithley, 6517A) which its constant voltage can be changed and the current is reported. The conductivity measurement was performed under atmospheric pressure, 40-60% RH and at 25-27 °C. The responsive current linearly proportional to the applied voltage is called the linear ohmic regime which can be identified by plotting the applied voltage against with the current. The voltage and the current in the regime were converted to the electrical conductivity by following equation (J1)

$$\sigma = \frac{1}{\rho} = \frac{1}{R_s x t} = \frac{I}{K x V x t} \tag{J1}$$

where σ is specific conductivity (S/cm), ρ is specific resistivity (Ω .cm), R_s is sheet resistance (Ω /sq), t is thickness of sample pellet (cm), V is applied voltage (voltage drop) (V), I is measured current (A), and K is geometric correction factor of the two-point probe meter.

In this measurement, the probe 2 was used; whose K is 4.205 x10⁻⁵ and the thickness of the samples was measured by using a thickness gauge.

Table J1 The specific conductivity (S/cm) of uPPP, dPPP with various doping level and NaZSM-5(23) zeolite

Sample	Specific conductivity (S/cm					
uPPP	(1.171±0.073)E-05					
10:1 dPPP	(1.260±0.248)E-04					
20:1 dPPP	(2.214±1.140)E-02					
50:1 dPPP	(8.701±4.246)E-01					
100:1 dPPP	(1.481±0.733)E+00					
NaZSM-5(23)	(8.670±2.543)E-03					

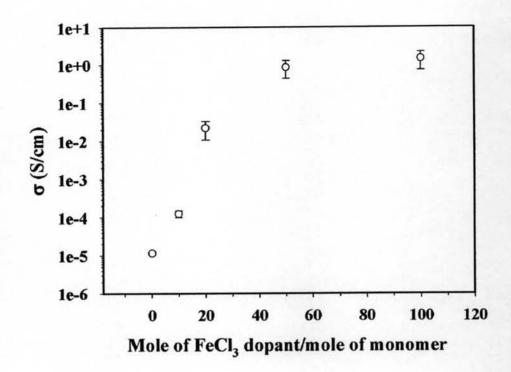


Figure J1 The specific conductivity of dPPP at various mole ratios between dopant and monomer at 24-25 °C, 50-60 %RH.

Table J2 Voltage-current data in linear regime of uPPP at 24-25 °C, 50-60 %RH

App	Applied voltage (V)			sure curren	t (A)	Con	ductivity (S	/cm)
1	2	3	1	2	3	1	2	3
0.2	0.2	0.2	1.50E-12	1.57E-12	1.63E-12	8.59E-06	9.02E-06	9.37E-06
0.4	0.4	0.4	3.66E-12	3.58E-12	3.55E-12	1.05E-05	1.03E-05	1.02E-05
0.6	0.6	0.6	5.86E-12	6.20E-12	5.98E-12	1.12E-05	1.19E-05	1.14E-05
0.8	0.8	0.8	8.58E-12	8.22E-12	8.22E-12	1.23E-05	1.18E-05	1.18E-05
1.0	1.0	1.0	1.05E-11	1.05E-11	1.04E-11	1.21E-05	1.21E-05	1.19E-05
1.2	1.2	1.2	1.25E-11	1.26E-11	1.24E-11	1.19E-05	1.20E-05	1.18E-05
1.4	1.4	1.4	1.42E-11	1.42E-11	1.42E-11	1.17E-05	1.17E-05	1.17E-05
1.6	1.6	1.6	1.57E-11	1.58E-11	1.58E-11	1.13E-05	1.13E-05	1.13E-05
1.8	1.8	1.8	1.78E-11	1.78E-11	1.75E-11	1.13E-05	1.13E-05	1.12E-05
2.0	2.0	2.0	1.92E-11	1.88E-11	1.88E-11	1.10E-05	1.08E-05	1.08E-05

Table J3 Voltage-current data in linear regime of 10:1 dPPP at 24-25 °C, 50-60 %RH

App	lied voltage	e (V)	Mea	sure curren	it (A)	Con	ductivity (S	/cm)
1	2	3	1	2	3	1	2	3
0.1	0.1	0.1	9.30E-12	1.04E-11	1.08E-11	1.31E-04	1.47E-04	1.53E-04
0.2	0.2	0.2	1.87E-11	1.82E-11	1.82E-11	1.32E-04	1.29E-04	1.29E-04
0.3	0.3	0.3	2.87E-11	2.83E-11	2.82E-11	1.35E-04	1.33E-04	1.33E-04
0.4	0.4	0.4	3.75E-11	3.72E-11	3.72E-11	1.32E-04	1.31E-04	1.31E-04
0.5	0.5	0.5	4.62E-11	4.55E-11	4.51E-11	1.31E-04	1.29E-04	1.28E-04
0.6	0.6	0.6	5.45E-11	5.39E-11	5.38E-11	1.29E-04	1.27E-04	1.27E-04
0.7	0.7	0.7	6.28E-11	6.29E-11	6.20E-11	1.27E-04	1.27E-04	1.25E-04
0.8	0.8	0.8	7.13E-11	7.13E-11	7.07E-11	1.26E-04	1.26E-04	1.25E-04
0.9	0.9	0.9	7.99E-11	7.88E-11	7.90E-11	1.26E-04	1.24E-04	1.24E-04
1.0	1.0	1.0	8.82E-11	8.82E-11	8.75E-11	1.25E-04	1.25E-04	1.24E-04
1.1	1.1	1.1	9.71E-11	9.59E-11	9.55E-11	1.37E-04	1.23E-04	1.23E-04

Table J4 Voltage-current data in linear regime of 20:1 dPPP at 24-25 °C, 50-60 %RH

App	lied voltage	e (V)	Meas	ured curre	nt (A)	Con	ductivity (S.	/cm)
1	2	3	1	2	3	1	2	3
1	1	1	2.41E-09	2.23E-09	1.81E-09	3.51E-03	3.25E-03	2.63E-03
2	2	2	8.85E-09	8.47E-09	7.99E-09	6.44E-03	6.17E-03	5.82E-03
3	3	3	1.60E-08	1.58E-08	1.55E-08	7.76E-03	7.66E-03	7.55E-03
4	4	4	2.47E-08	2.44E-08	2.43E-08	9.01E-03	8.89E-03	8.85E-03
5	5	5	3.46E-08	3.44E-08	3.42E-08	1.01E-02	1.00E-02	9.97E-03
6	6	6	4.51E-08	4.48E-08	4.47E-08	1.10E-02	1.09E-02	1.08E-02
7	7	7	5.55E-08	5.53E-08	5.50E-08	1.16E-02	1.15E-02	1.14E-02
8	8	8	6.62E-08	6.58E-08	6.55E-08	1.21E-02	1.20E-02	1.19E-02
9	9	9	7.68E-08	7.68E-08	7.68E-08	1.24E-02	1.24E-02	1.24E-02
10	10	10	8.88E-08	8.94E-08	8.85E-08	1.29E-02	1.30E-02	1.29E-02

Table J5 Voltage-current data in linear regime of 50:1 dPPP at 24-25 °C, 50-60 %RH

App	lied voltage	e (V)	Meas	ured curre	nt (A)	Con	ductivity (S	/cm)
1	2	3	1	2	3	1	2	3
0.1	0.1	0.1	1.11E-08	1.09E-08	1.08E-08	0.111813	0.109453	0.108562
0.2	0.2	0.2	3.37E-08	3.28E-08	3.14E-08	0.169619	0.164998	0.158225
0.3	0.3	0.3	5.37E-08	5.44E-08	5.27E-08	0.180072	0.182382	0.176912
0.4	0.4	0.4	7.51E-08	7.29E-08	6.91E-08	0.188950	0.183487	0.173976
0.5	0.5	0.5	9.07E-08	8.98E-08	8.87E-08	0.182579	0.180773	0.178435
0.6	0.6	0.6	1.11E-07	1.10E-07	1.06E-07	0.186147	0.184265	0.178023
0.7	0.7	0.7	1.24E-07	1.19E-07	1.19E-07	0.177664	0.171037	0.170959
0.8	0.8	0.8	1.41E-07	1.43E-07	1.42E-07	0.176984	0.179755	0.178662
0.9	0.9	0.9	1.55E-07	1.59E-07	1.54E-07	0.173118	0.177641	0.171850
1.0	1.0	1.0	1.70E-07	1.64E-07	1.57E-07	0.171437	0.164798	0.157886

Table J6 Voltage-current data in linear regime of 100:1 dPPP at 24-25 °C, 50-60 %RH

App	lied voltage	e (V)	Meas	ured curre	nt (A)	Con	ductivity (S	1.232245 1.254015 1.335709 1.413778	
1	2	3	1	2	3	1	2	3	
0.1	0.1	0.1	1.02E-07	1.03E-07	1.05E-07	0.972079	0.983163	1.000698	
0.2	0.2	0.2	2.51E-07	2.58E-07	2.63E-07	1.196080	1.232245	1.254015	
0.3	0.3	0.3	4.13E-07	4.2E-07	4.44E-07	1.313635	1.335709	1.413778	
0.4	0.4	0.4	6.38E-07	6.36E-07	6.41E-07	1.522853	1.517735	1.529188	
0.5	0.5	0.5	8.77E-07	8.97E-07	9.06E-07	1.673749	1.712993	1.729517	
0.6	0.6	0.6	1.20E-06	1.25E-06	1.27E-06	1.905577	1.990477	2.026039	
0.7	0.7	0.7	1.63E-06	1.65E-06	1.69E-06	2.216643	2.256428	2.306414	
0.8	0.8	0.8	1.98E-06	1.93E-06	1.95E-06	2.364017	2.307162	2.326751	
0.9	0.9	0.9	2.27E-06	2.3E-06	2.35E-06	2.406210	2.436363	2.490092	
1.0	1.0	1.0	2.66E-06	2.63E-06	2.61E-06	2.542574	2.515294	2.494274	

Table J7 Voltage-current data in linear regime of NaZSM-5(23) at 24-25 °C, 50-60 %RH

Apr	lied voltage	(V)	Meas	ured curre	nt (A)	Con	ductivity (S	/cm)
1	2	3	1	2	3	1	2	3
0.5	0.5	0.5	3.77E-09	2.13E-09	3.77E-09	5.64E-03	1.79E-03	3.23E-03
1.0	1.0	1.0	1.37E-08	5.46E-09	1.37E-08	1.03E-02	2.29E-03	5.26E-03
1.5	1.5	1.5	2.25E-08	9.99E-09	2.25E-08	1.12E-02	2.80E-03	5.93E-03
2.0	2.0	2.0	3.14E-08	1.78E-08	3.14E-08	1.17E-02	3.74E-03	6.74E-03
2.5	2.5	2.5	4.02E-08	2.98E-08	4.02E-08	1.20E-02	4.99E-03	7.66E-03
3.0	3.0	3.0	4.78E-08	3.9E-08	4.78E-08	1.19E-02	5.46E-03	7.92E-03
3.5	3.5	3.5	5.55E-08	4.36E-08	5.55E-08	1.18E-02	5.23E-03	7.76E-03
4.0	4.0	4.0	6.34E-08	5.31E-08	6.34E-08	1.18E-02	5.57E-03	7.97E-03
4.5	4.5	4.5	7.14E-08	5.9E-08	7.14E-08	1.18E-02	5.51E-03	7.93E-03
5.0	5.0	5.0	8.01E-08	8.07E-08	8.01E-08	1.20E-02	6.78E-03	8.81E-03
5.5	5.5	5.5	8.78E-08	1.06E-07	8.78E-08	1.19E-02	8.12E-03	9.66E-03
6.0	6.0	6.0	9.67E-08	1.15E-07	9.67E-08	1.20E-02	8.06E-03	9.67E-03
6.5	6.5	6.5	1.04E-07	1.32E-07	1.04E-07	1.20E-02	8.50E-03	9.94E-03
7.0	7.0	7.0	1.12E-07	1.44E-07	1.12E-07	1.19E-02	8.60E-03	9.99E-03
7.5	7.5	7.5	1.17E-07	1.43E-07	1.17E-07	1.17E-02	8.02E-03	9.51E-03
8.0	8.0	8.0	1.23E-07	1.62E-07	1.23E-07	1.15E-02	8.50E-03	9.75E-03
8.5	8.5	8.5	1.31E-07	1.62E-07	1.31E-07	1.15E-02	8.01E-03	9.44E-03
9.0	9.0	9.0	1.41E-07	1.76E-07	1.41E-07	1.17E-02	8.19E-03	9.64E-03
9.5	9.5	9.5	1.47E-07	1.92E-07	1.47E-07	1.16E-02	8.48E-03	9.78E-03
10.0	10.0	10.0	1.53E-07	1.91E-07	1.53E-07	1.15E-02	8.02E-03	9.43E-03

Appendix K Electrical Conductivity Sensitivity Measurement

The electrical conductivity of dPPP and dPPP/zeolite composite pellets under both N₂ and NH₃ gas atmosphere were carried out by using two point probe, under pressure 1 atm, at 29±1 °C. The electrical conductivity response of the sample is defined as the difference between the equilibrium conductivity of sample under exposed to NH₃ and that of samples under N₂ exposure.

$$\Delta \sigma = \sigma_{NH_3} - \sigma_{N_2 \text{ inotial}} \tag{K1}$$

The electrical conductivity sensitivity is calculated from the electrical conductivity response divided by equilibrium conductivity under exposed to N_2 .

$$Sensitivity = \Delta \sigma / \sigma_{N_2 \text{ initial}}$$
 (K2)

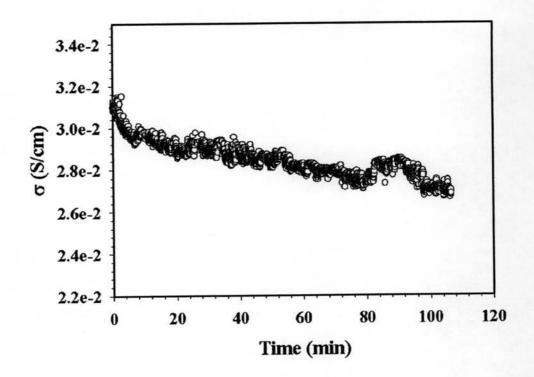


Figure K1 Specific conductivity of 50:1 dPPP when exposed to 0.15625%v NH₃.

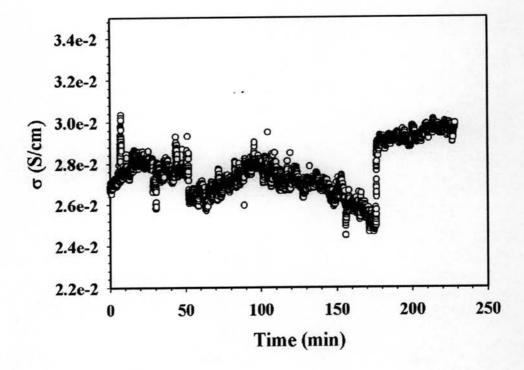


Figure K2 Specific conductivity of 50:1 dPPP after evacuating 0.15625%v NH₃ and exposed to N₂.

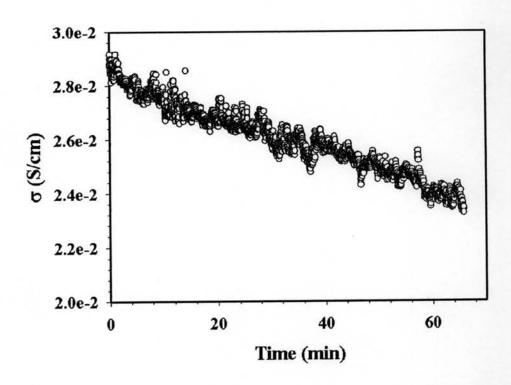


Figure K3 Specific conductivity of 50:1 dPPP when exposed to 0.3125%v NH₃.

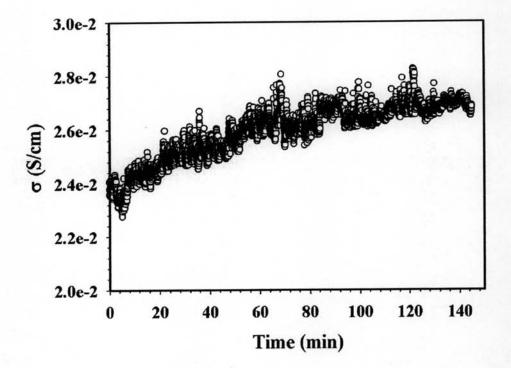


Figure K4 Specific conductivity of 50:1 dPPP after evacuating 0.3125%v NH $_3$ and exposed to N_2 .

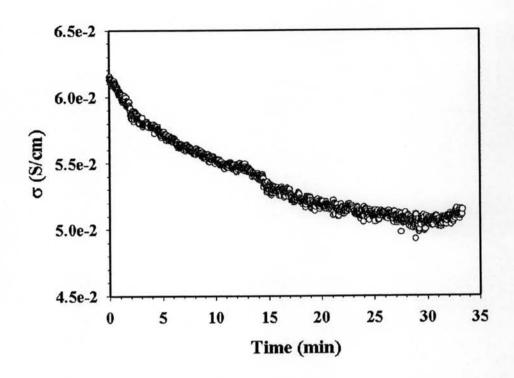


Figure K5 Specific conductivity of 50:1 dPPP when exposed to 0.625%v NH₃.

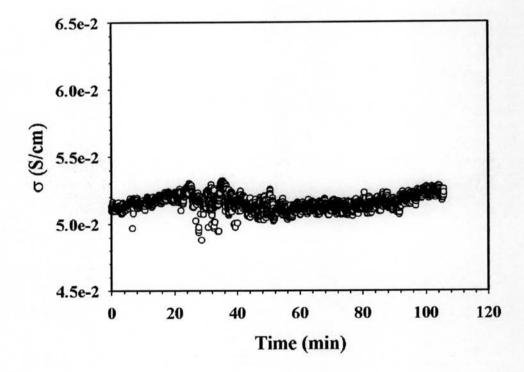


Figure K6 Specific conductivity of 50:1 dPPP after evacuating 0.625%v NH₃ and exposed to N₂.

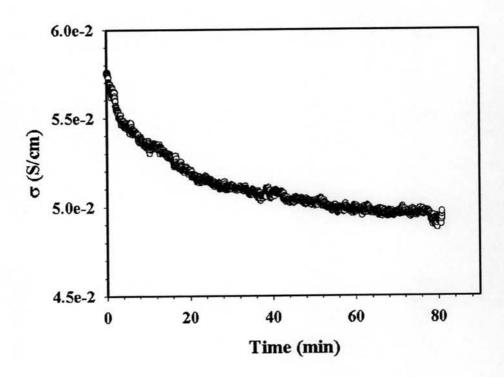


Figure K7 Specific conductivity of 50:1 dPPP when exposed to 1.25%v NH₃.

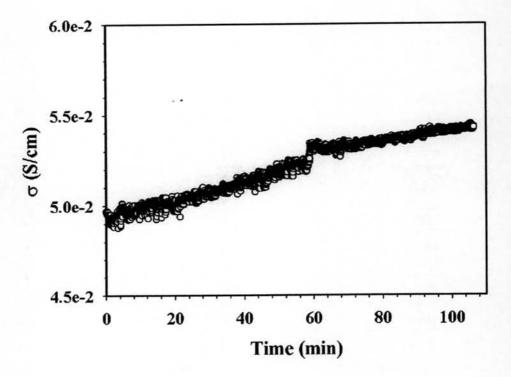


Figure K8 Specific conductivity of 50:1 dPPP after evacuating 1.25%v NH₃ and exposed to N_2 .

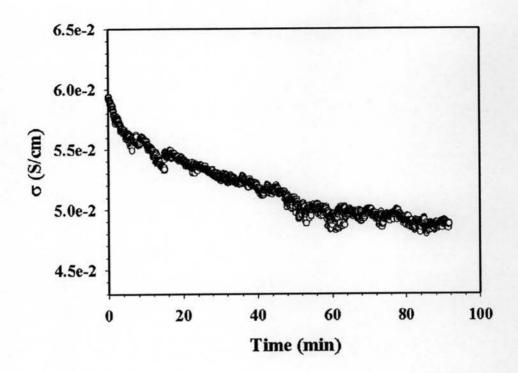


Figure K9 Specific conductivity of 50:1 dPPP when exposed to 2.5%v NH₃.

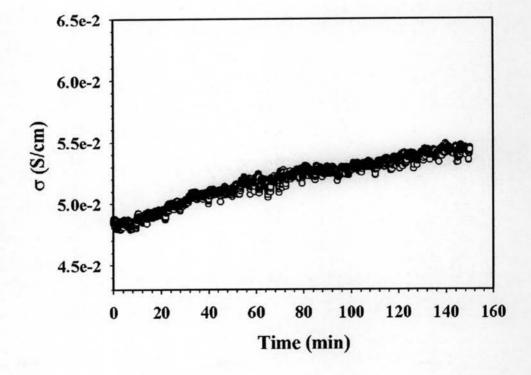


Figure K10 Specific conductivity of 50:1 dPPP after evacuating 2.5%v NH₃ and exposed to N₂.

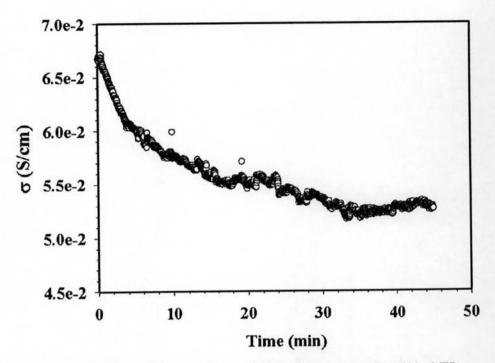


Figure K11 Specific conductivity of 50:1 dPPP when exposed to 5%v NH₃.

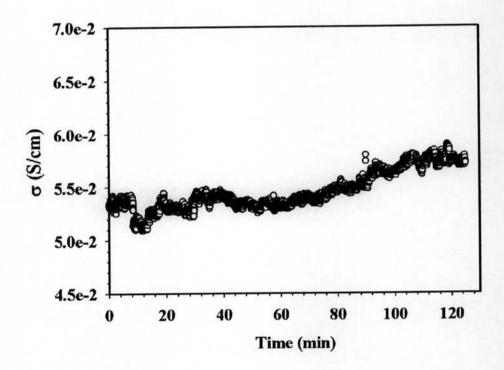


Figure K12 Specific conductivity of 50:1 dPPP after evacuating 5%v NH₃ and exposed to N₂.

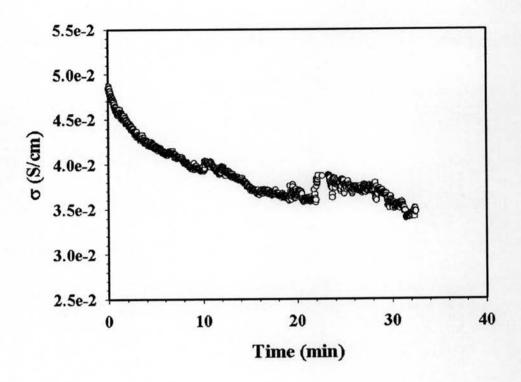


Figure K13 Specific conductivity of 50:1 dPPP when exposed to 10%v NH₃.

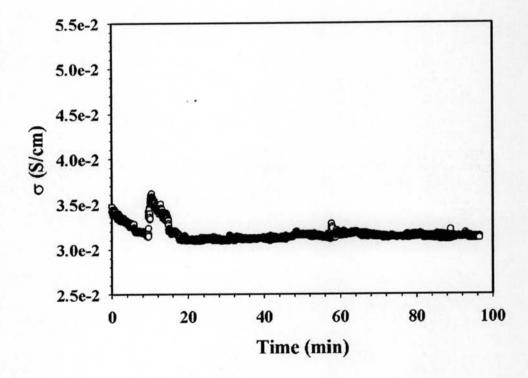


Figure K14 Specific conductivity of 50:1 dPPP after evacuating 10%v NH₃ and exposed to N₂.

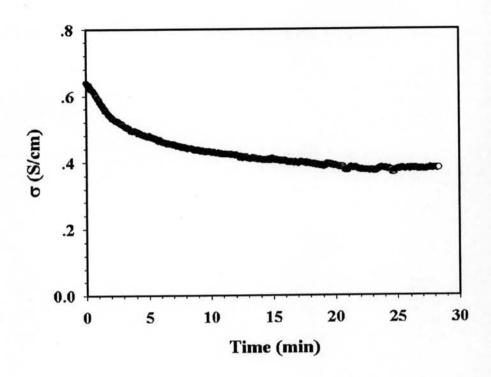


Figure K15 Specific conductivity of dPPP(90)/NaZ23 when exposed to 5%v NH₃.

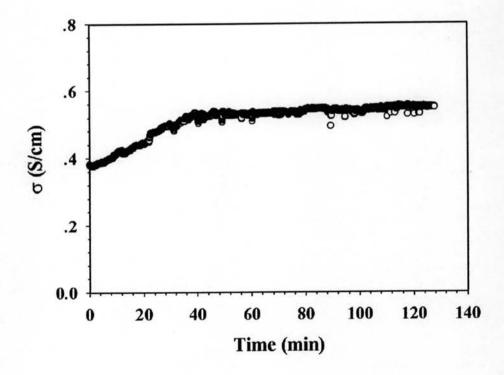


Figure K16 Specific conductivity of dPPP(90)/NaZ23 after evacuating 5%v NH₃ and exposed to N₂.

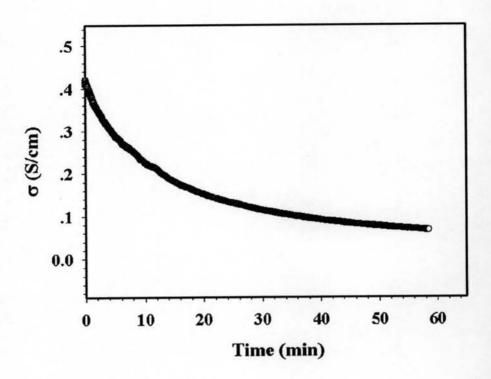


Figure K17 Specific conductivity of dPPP(80)/NaZ23 when exposed to 5%v NH₃.

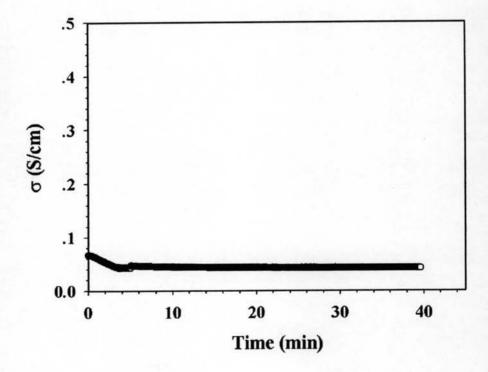


Figure K18 Specific conductivity of dPPP(80)/NaZ23 after evacuating 5%v NH₃ and exposed to N₂.

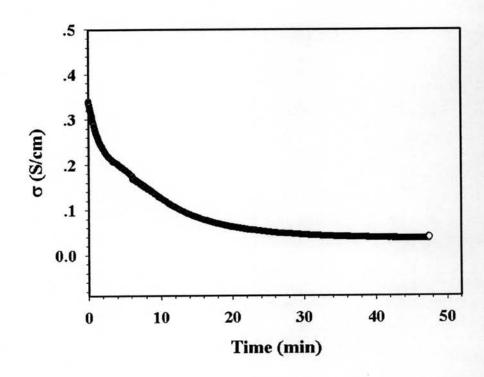


Figure K19 Specific conductivity of dPPP(70)/NaZ23 when exposed to 5%v NH₃.

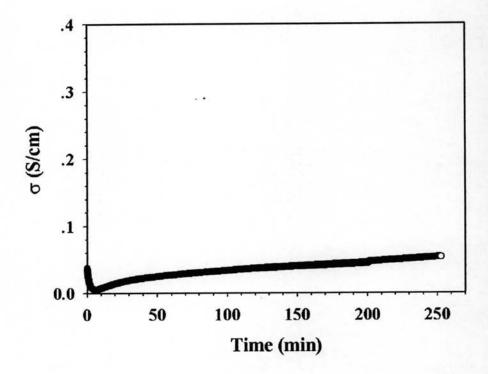


Figure K20 Specific conductivity of dPPP(70)/NaZ23 after evacuating 5%v NH₃ and exposed to N₂.

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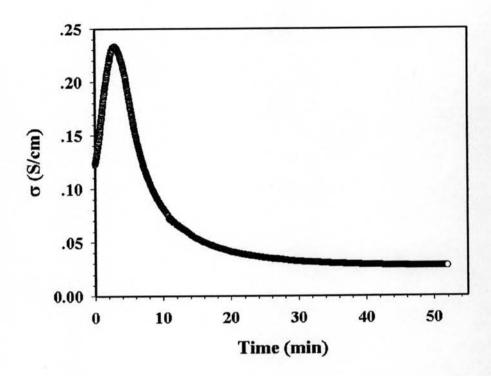


Figure K21 Specific conductivity of dPPP(60)/NaZ23 when exposed to 5%v NH₃.

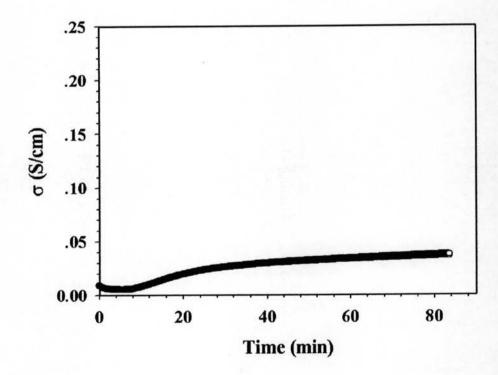


Figure K22 Specific conductivity of dPPP(60)/NaZ23 after evacuating 5%v NH₃ and exposed to N₂.

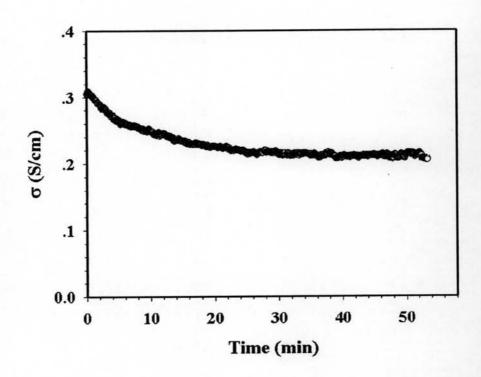


Figure K23 ecific conductivity of dPPP(90)/NaZ23 when exposed to 1.25%v NH₃.

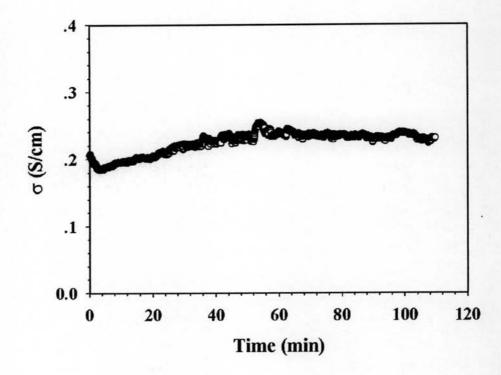


Figure K24 Specific conductivity of dPPP(90)/NaZ23 after evacuating 1.25%v NH₃ and exposed to N₂.

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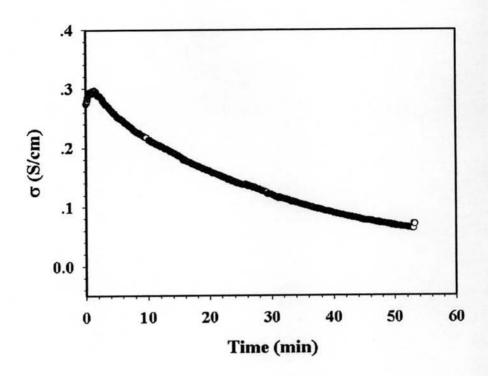


Figure K25 Specific conductivity of dPPP(80)/NaZ23 when exposed to 1.25%v NH₃.

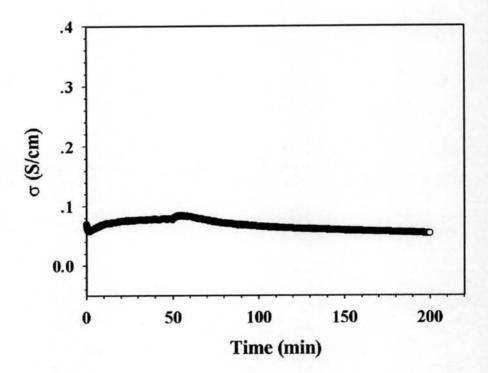


Figure K26 Specific conductivity of dPPP(80)/NaZ23 after evacuating 1.25%v NH₃ and exposed to N₂.

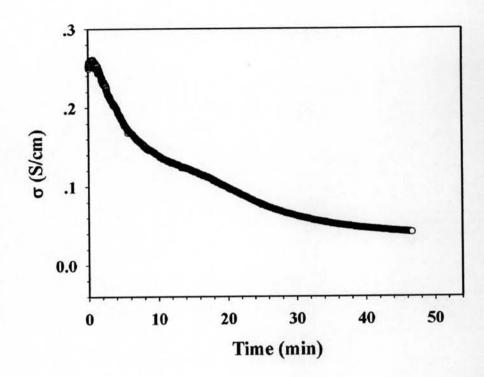


Figure K27 Specific conductivity of dPPP(70)/NaZ23 when exposed to 1.25%v NH₃.

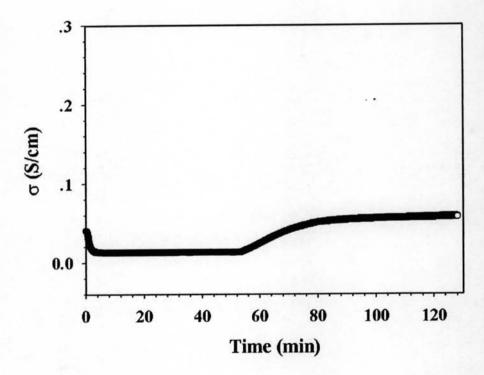


Figure K28 Specific conductivity of dPPP(70)/NaZ23 after evacuating $1.25\%v~NH_3$ and exposed to N_2 .

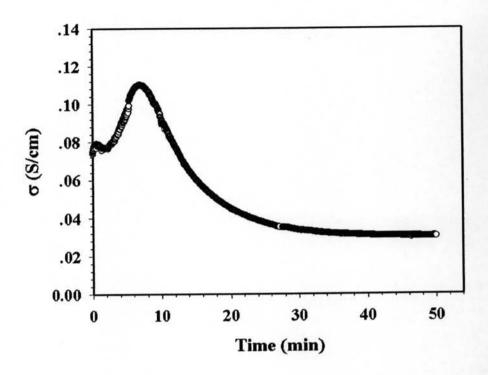


Figure K29 Specific conductivity of dPPP(60)/NaZ23 when exposed to 1.25%v NH₃.

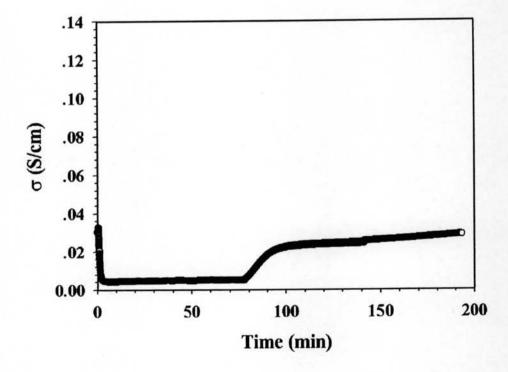


Figure K30 Specific conductivity of dPPP(60)/NaZ23 after evacuating 1.25%v NH₃ and exposed to N₂.

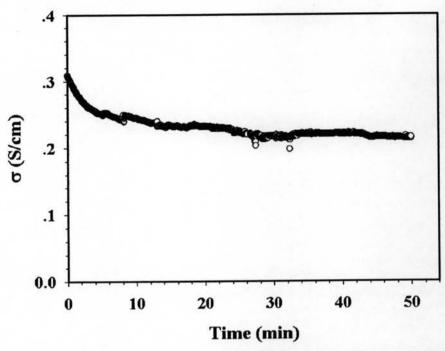


Figure K31 Specific conductivity of dPPP(90)/NaZ23 when exposed to 0.625%v NH₃.

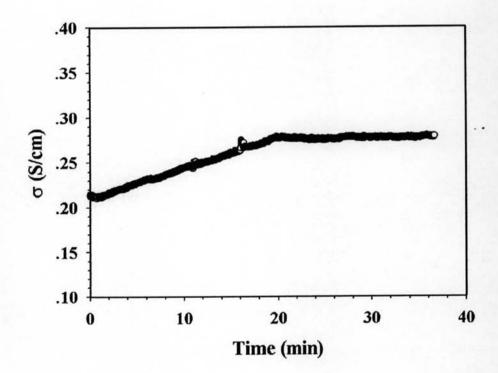


Figure K32 Specific conductivity of dPPP(90)/NaZ23 after evacuating 0.625%v NH₃ and exposed to N₂.

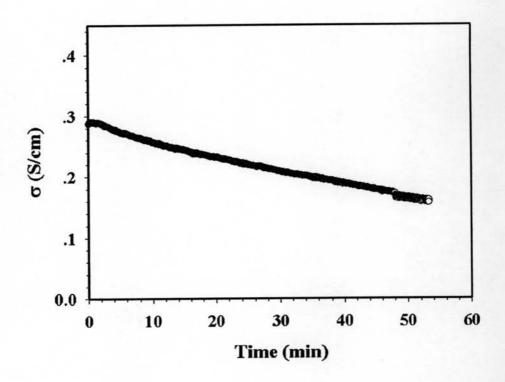


Figure K33 Specific conductivity of dPPP(80)/NaZ23 when exposed to 0.625%v NH₃.

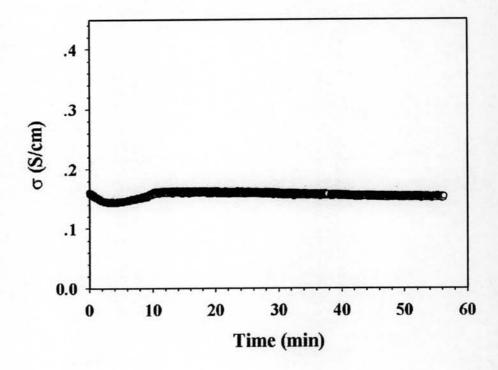


Figure K34 Specific conductivity of dPPP(80)/NaZ23 after evacuating 0.625%v NH₃ and exposed to N₂.

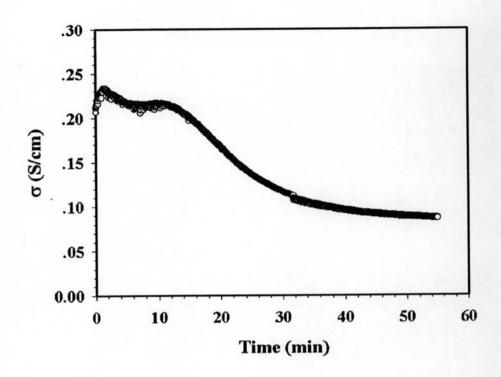


Figure K35 Specific conductivity of dPPP(70)/NaZ23 when exposed to 0.625%v NH₃.

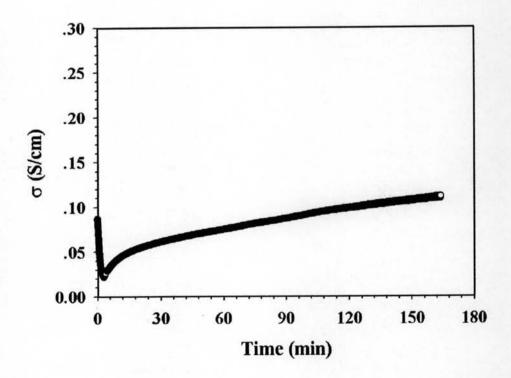


Figure K36 Specific conductivity of dPPP(70)/NaZ23 after evacuating 0.625%v NH₃ and exposed to N₂.

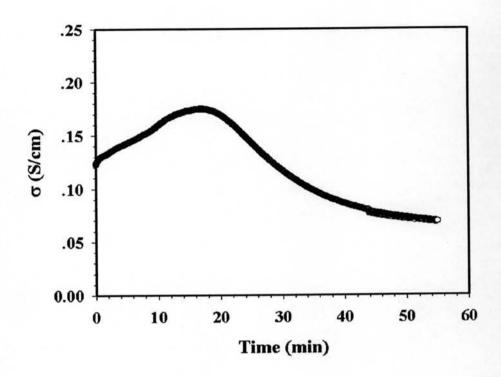


Figure K37 Specific conductivity of dPPP(60)/NaZ23 when exposed to 0.625%v NH₃.

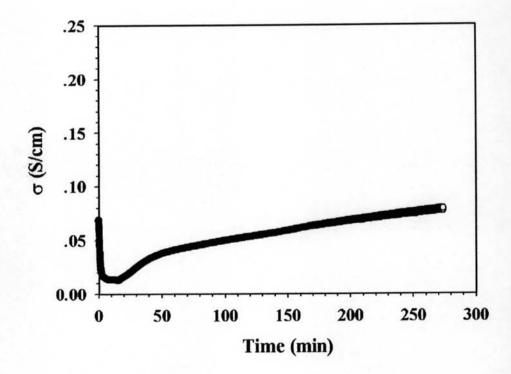


Figure K38 Specific conductivity of dPPP(60)/NaZ23 after evacuating 0.625%v NH₃ and exposed to N₂.

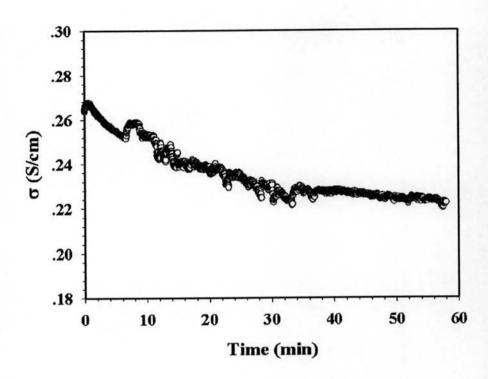


Figure K39 Specific conductivity of dPPP(90)/KZ23 when exposed to 0.625%v NH₃.

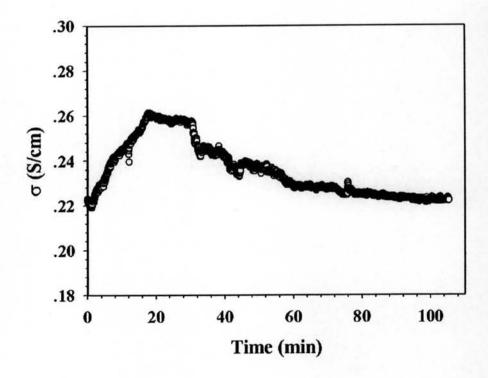


Figure K40 Specific conductivity of dPPP(90)/KZ23 after evacuating 0.625%v NH₃ and exposed to N₂.

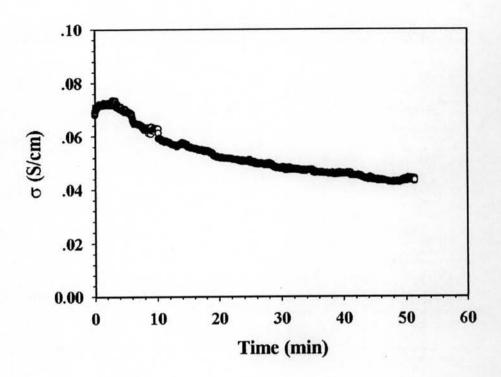


Figure K41 Specific conductivity of dPPP(90)/NH₄Z23 when exposed to 0.625%v NH₃.

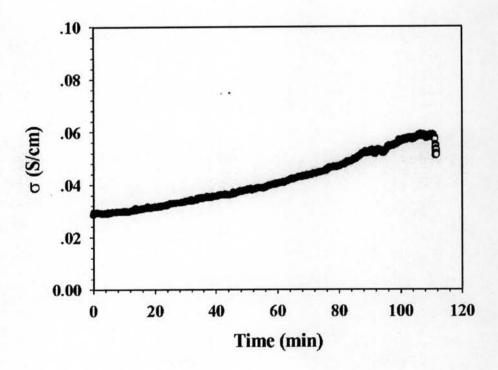


Figure K42 Specific conductivity of dPPP(90)/NH₄Z23 after evacuating 0.625%v NH₃ and exposed to N₂.

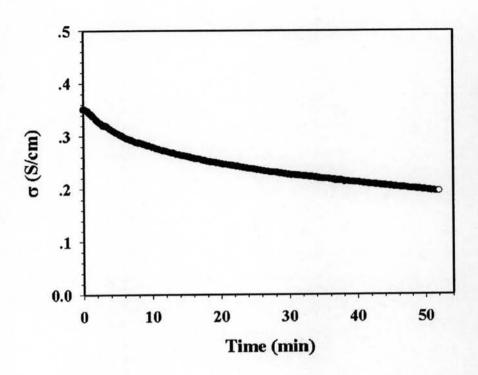


Figure K43 Specific conductivity of dPPP(90)/HZ23 when exposed to 0.625%v NH₃.

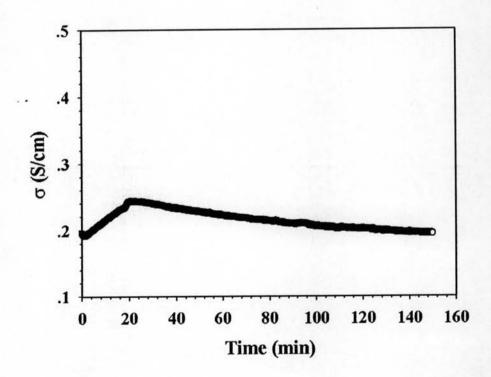


Figure K44 Specific conductivity of dPPP(90)/HZ23 after evacuating 0.625%v NH₃ and exposed to N₂.

Table K1 Specific conductivity and sensitivity of 50:1dPPP and its composites when exposed to air, N_2 and NH_3 with various concentrations under chamber temperature (T_c) of 28 ± 1 °C, at 1 atm, t_i = the time that σ_{NH_3} reaches equilibrium, t_r = the time that σ reaches equilibrium after evacuate NH_3 , V = applied voltage, K = correction factor (2.19×10^{-5})

No.	Samples	Thickness (cm)	37	DATE 1	t _i (min)	t _r (min)		I ((A)			σ (\$	S/cm)		Response	Sensitivity
			(v)	[NH ₃] (% v.)			Air	N ₂ initial	NH ₃	N ₂ final	Air	N ₂ initial	NH ₃	N ₂ final		
1	50:1dPPP	0.04127	25	0.15625	106.68	228.54	(7.38± 0.12)E-7	(6.78± 0.04)E-7	(6.09± 0.02)E-7	(6.71± 0.03)E-7	(3.26± 0.05)E-2	(2.99± 0.02)E-2	(2.69± 0.01)E-2	(2.96± 0.01)E-2	(-3.04± 0.24)E-3	(-9.81± 0.71)E-2
2	50:1dPPP	0.02736	25	0.3125	65.84	145.04	(5.18± 0.07)E-7	(4.16± 0.03)E-7	(3.62± 0.06)E-7	(4.06± 0.02)E-7	(3.45± 0.05)E-2	(2.78± 0.02)E-2	(2.42± 0.04)E-2	(2.68± 0.02)E-2	(-3.61± 0.52)E-3	(-1.15± 0.07)E-1
3	50:1dPPP	0.02054	20	0.625	33.34	105.00	(9.86± 0.85)E-7	(5.39± 0.01)E-7	(4.6± 0.02)E-7	(4.71± 0.21)E-7	(1.09± 0.09)E-1	5.98± 0.01)E-2	(5.10± 0.03)E-2	(5.23± 0.02)E-2	(-8.85± 0.33)E-3	(-1.43± 0.02)E-1
4	50:1dPPP	0.02538	20	0.625	80.84	62.99	(4.29± 0.17)E-7	(4.17± 0.05)E-7	(3.62± 0.08)E-7	3.99± 0.04)E-7	(3.85± 0.15)E-2	(3.75± 0.05)E-2	(3.25± 0.07)E-2	(3.58± 0.03)E-2	(-4.98± 0.97)E-3	(-1.14± 0.15)E-1
5	50:1dPPP	0.02512	20	1.25	80.84	132.24	(1.17± 0.09)E-6	(6.28± 0.01)E-7	(5.43± 0.02)E-7	(5.99± 0.02)E-7	(1.06± 0.08)E-1	(5.70± 0.01)E-2	(4.93± 0.02)E-2	(5.43± 0.02)E-2	(-7.72± 0.22)E-3	(-1.33± 0.02)E-1
6	50:1dPPP	0.02252	20	1.25	107.5	68.1	(4.97± 0.10)E-7	(3.73± 0.03)E-7	(3.05± 0.03)E-7	(3.13± 0.04)E-7	(5.03± 0.10)E-2	(3.78± 0.03)E-2	(3.09± 0.03)E-2	(3.17± 0.04)E-2	(-6.87± 0.49)E-3	(-1.74± 0.09)E-1
7	50:1dPPP	0.02049	25	1.25	83.34	111.77	(3.50± 0.22)E-7	(4.73± 0.05)E-7	(3.96± 0.1)E-7	(4.59± 0.15)E-7	(3.11± 0.19)E-2	(4.21± 0.04)E-2	(3.52± 0.10)E-2	(4.08± 0.10)E-2	(-6.87± 1.24)E-3	(-1.40± 0.12)E-1
8	50:1dPPP	0.02296	20	2.5	91.67	150	(6.83± 0.29)E-7	(5.98± 0.03)E-7	(4.92± 0.01)E-7	(5.47± 0.01)E-7	(6.77± 0.29)E-2	(5.93± 0.03)E-2	(4.88± 0.01)E-2	(5.43± 0.01)E-2	(-1.05± 0.04)E-2	(-1.75± 0.07)E-1
9	50:1dPPP	0.02145	25	2.5	75.01	130.31	(4.93± 0.30)E-7	(4.53± 0.04)E-7	(3.69± 0.04)E-7	(4.14± 0.02)E-7	(4.19± 0.25)E-2	(3.85± 0.03)E-2	(3.14± 0.04)E-2	(3.52± 0.01)E-2	(-7.17± 0.55)E-3	(-1.76± 0.09)E-1
10	50:1dPPP	0.02256	20	5	75.84	86.67	(9.88± 0.75)E-7	(3.75± 0.04)E-7	(2.96± 0.08)E-7	(3.20± 0.02)E-7	(9.98± 0.76)E-2	(3.79± 0.04)E-2	(2.99± 0.09)E-2	(3.23± 0.02)E-2	(-8.04± 0.11)E-3	(-2.12± 0.12)E-1
11	50:1dPPP	0.0204	20	5	45	125	(9.71± 0.64)E-7	(5.73± 0.01)E-7	(4.71± 0.03)E-7	(5.15± 0.04)E-7	(1.08± 0.07)E-1	(6.40± 0.01)E-2	(5.26± 0.03)E-2	(5.75± 0.04)E-2	(-1.14± 0.04)E-2	(-1.77± 0.03)E-1
12	50:1dPPP	0.02097	25	10	32,51	96.49	(5.95± 0.08)E-7	(5.52± 0.05)E-7	(4.05± 0.09)E-7	(3.60± 0.01)E-7	(5.17± 0.07)E-2	(4.80± 0.04)E-2	(3.52± 0.07)E-2	(3.13± 0.01)E-2	(-1.28± 0.1)E-2	(-2.51± 0.10)E-1

re-ti-re	Samples	Thickness (cm)	v	DATE 1	(min)	t _r (min)		I(A)			σ (\$	S/cm)			Sensitivity
No.			(V)	[NH ₃] (% v.)			Air	N ₂ initial	NH ₃	N ₂ final	Air	N ₂ initial	NH ₃	N ₂ final	Response	
13	50:1dPPP(90)/NaZ23	0.02084	12	5	28.34	127.7	(1.05± 0.02)E-5	(3.09± 0.01)E-6	(2.07± 0.01)E-6	(3.02± 0.00)E-6	(1.92± 0.04)E0	(5.63± 0.01)E-1	(3.78± 0.07)E-1	(5.50± 0.00)E-1	(-1.85± 0.03)E-1	(-3.24± 0.03)E-1
14	50:1dPPP(90)/NaZ23	0.02977	12	5	53.34	148.04	(4.27± 0.18)E-6	(2.01± 0.03)E-6	(1.07± 0.01)E-6	(1.16± 0.00)E-6	(5.45± 0.23)E-1	(2.57± 0.04)E-1	(1.37± 0.00)E-1	(1.48± 0.00)E-1	(-1.20± 0.05)E-1	(-4.64± 0.21)E-1
15	50:1dPPP(80)/NaZ23	0.021	15	5	58.53	39.61	(5.74± 0.19)E-6	(2.32± 0.12)E-6	(4.79± 0.10)E-7	(2.92± 0.01)E-7	(8.30± 0.27)E-1	(3.35± 0.17)E-1	(6.93± 0.15)E-2	(4.23± 0.83)E-2	(-2.66± 0.20)E-1	(-7.91± 0.75)E-1
16	50:1dPPP(70)/NaZ23	0.02114	25	5	47.51	252.79	(1.41± 0.02)E-4	(3.71± 0.01)E-6	(4.34± 0.13)E-7	(6.19± 0.01)E-7	(1.21± 0.02)E+1	(3.20± 0.01)E-1	(3.74± 0.11)E-2	(5.34± 0.01)E-2	(-2.83± 0.02)E-1	(-8.80± 0.03)E-1
17	50:1dPPP(60)/NaZ23	0.02452	50	5	54.17	83.57	(9.53± 4.34)E-6	(2.73± 0.01)E-6	(7.54± 0.08)E-7	(1.00± 0.01)E-6	(3.54± 1.61)E-1	(1.01± 0.01)E-1	(2.80± 0.03)E-2	(4.35± 0.04)E-2	(-7.34± 0.06)E-2	(-7.21± 0.06)E-1
18	50:1dPPP(90)/NaZ23	0.01383	10	1.25	53.34	109.8	(1.70± 0.02)E-5	(9.19± 0.05)E-7	(6.41± 0.09)E-7	(6.92± 0.07)E-7	(5.59± 0.07)E0	(3.03± 0.02)E-1	(2.11± 0.03)E-1	(2,28± 0.02)E-1	(-9.16± 0.39)E-2	(-2.93± 0.06)E-1
20	50:1dPPP(80)/NaZ23	0.01969	30	1.25	53.34	137.24	(2.26± 0.32)E-5	(2.81± 0.03)E-6	(1.32± 0.01)E-6	(8.15± 0.02)E-7	(1.74± 0.25)E0	(2.17± 0.02)E-1	(1.02± 0.01)E-1	(6.29± 0.01)E-2	(-1.15± 0.03)E-1	(-5.26± 0.14)E-1
21	50:1dPPP(70)/NaZ23	0.01813	40	1.25	50.84	128.16	(2.40± 0.11)E-5	(3.42± 0.02)E-6	(6.80± 0.22)E-7	(9.19± 0.06)E-7	(1.51± 0.07)E0	(2.15± 0.01)E-1	(4.27± 0.14)E-2	(5.77± 0.04)E-2	(-1.72± -0.02)E-1	(-7.95± 0.09)E-1
22	50:1dPPP(60)/NaZ23	0.01929	50	1.25	50	193.48	(6.19± 0.47)E-6	(1.45± 0.01)E-6	(6.44± 0.15)E-7	(6.09± 0.01)E-7	(2.93± 0.45)E-1	(6.84± 0.02)E-2	(3.04± 0.01)E-2	(2.65± 0.00)E-2	(-3.8± 0.03)E-1	(-5.55± 0.04)E-1
23	50:1dPPP(90)/NaZ23	0.01504	8	0.625	50	36.67	(1.68± 0.09)E-6	(7.31± 0.02)E-7	(5.68± 0.02)E-7	(7.34± 0.02)E-7	(6.37± 0.32)E-1	(2.77± 0.01)E-1	(2.15± 0.01)E-1	(2.8± 0.06)E-1	(-6.20± 0.11)E-2	(-2.21± 0.02)E-1
24	50:1dPPP(80)/NaZ23	0.01778	30	0.625	53.34	56.27	(8,60± 0.05)E-6	(3.09± 0.02)E-6	(1.89± 0.02)E-6	(1.80± 0.01)E-6	(7.35± 0.04)E-1	(2.64± 0.01)E-1	(1.61± 0.02)E-1	(1.53± 0.01)E-1	(-1.03± 0.02)E-1	(-3.84± 0.05)E-1
25	50:1dPPP(70)/NaZ23	0.01639	50	0.625	55	164.04	(1.57± 0.10)E-5	(3.25± 0.03)E-6	(1.57± 0.01)E-6	(1.99± 0.01)E-6	(8.75± 0.56)E-2	(1.81± 0.02)E-1	(8.76± 0.04)E-1	(1.11± 0.07)E-1	(-9.33± 0.20)E-2	(-5.14± 0.12)E-1
26	50:1dPPP(60)/NaZ23	0.02963	70	0.625	55	273.94	(1.33± 0.04)E-5	(2.76± 0.54)E-6	(1.79± 0.04)E-6	(1.97± 0.02)E-6	(2.93± 0.13)E-1	(6.07± 1.18)E-1	(3.93± 0.11)E-2	(4.33± 0.06)E-2	(2.14± 1.12)E-2	(-3.22± 1.31)E-1
27	50:1dPPP(90)/KZ23	0.01449	10	0.625	58	105.39	(1.05± 0.03)E-5	(8.11± 0.02)E-7	(7.05± 0.02)E-7	(7.07± 0.01)E-7	(3.30± 0.10)E0	(2.55± 0.01)E-1	(2.22± 0.01)E-1	(2.22± 0.00)E-1	(-3.32± 0.12)E-2	(-1.27± 0.04)E-1
28	50:1dPPP(90)/NH4Z23	0.01675	10	0.625	51.46	111.67	(1.32± 0.00)E-6	(2.63± 0.04)E-7	(1.62± 0.04)E-7	(2.08± 0.10)E-7	(3.58± 0.01)E-1	(7.14± 0.11)E-3	(4.41± 0.10)E-3	(5.66± 0.03)E-2	(-2.73± 0.17)E-3	(-3.68± 0.19)E-1
29	50:1dPPP(90)/HZ23	0.01353	10	0.625	52	150	(5.07± 0.38)E-6	(9.87± 0.02)E-7	(5.83± 0.02)E-7	(5.77± 0.01)E-7	(1.71± 0.01)E0	(3.33± 0.01)E-1	(1.96± 0.01)E-1	(1.94± 0.00)E-1	(-1.36± 0.01)E-1	(-4.08± 0.03)E-1
30	50:1dPPP(90)/HZ23	0.02023	10	0.625	85.42	38.77	(1.30± 0.06)E-5	(7.64± 0.38)E-7	(5.15± 0.05)E-1	(4.80± 0.07)E-7	(2.92± 0.13)E0	(1.72± 0.09)E-1	(1.16± 0.01)E-1	(1.08± 0.02)E-1	(-5.61± 1.00)E-2	(-3.20± 0.61)E-1

Appendix L Ammonia Temperature Programmed Desorption

An ammonia temperature programmed desorption (NH₃-TPD) of zeolite was performed in a flow-through reactor with helium as a carrier. In order to remove water and undesired impurities, approximated 0.05 g of sample was pre-treated under He flow by heating from 30 °C to 500 °C at 10 °C/min. After holding temperature at 500 °C for 1 h and cooling to ambient temperature, the gas flow was switched to 1% NH₃/He. The NH₃ adsorption under room temperature was allowed for 1 h and the removal of weakly adsorbed NH₃ was then achieved by purging He gas into the system for 2 h. The NH₃ desorption was done by elevating temperature from 30 °C to 600 °C at ramp rate of 10 °C/min. In case of dPPP, NH₃ desorption was performed by elevating temperature from 30 °C to 300 °C. The desorbed NH₃ was detected by thermal conductivity detector (TCD).

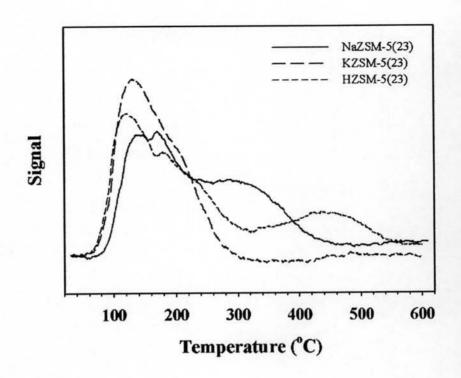


Figure L1 NH₃-TPD thermogram of ZSM-5(23) with various cation types.

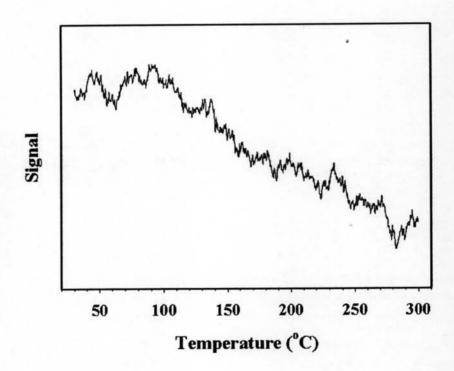


Figure L2 NH₃-TPD thermogram of 50:1 dPPP.

Appendix M Investigation of Interaction between dPPP, dPPP composite, NaZSM-5(23) and NH₃ by using FTIR technique

FTIR spectra of 50:1 dPPP, 50:1dPPP(70)/NaZ23 and NaZSM-5(23) were taken by using KBr pellet technique. The sample pellet was located on sample holder and put in the gas cell. The spectra of samples were collected, before, during and after NH₃ exposure, in order to study the interaction between these samples and NH₃.

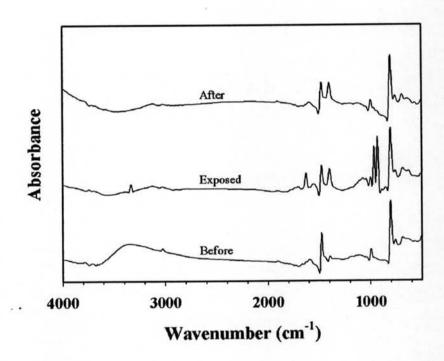


Figure M1 The FT-IR spectra of 50:1dPPP; before, during and after NH₃ exposure.

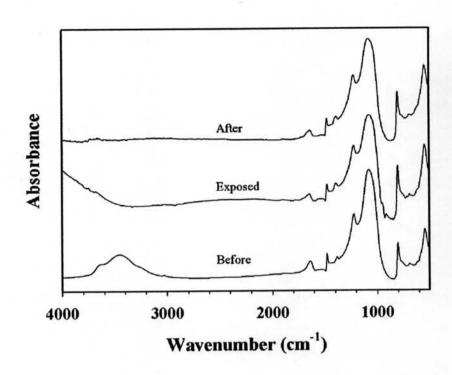


Figure M2 The FT-IR spectra of 50:1dPPP(70)/NaZ23; before, during and after NH₃ exposure.

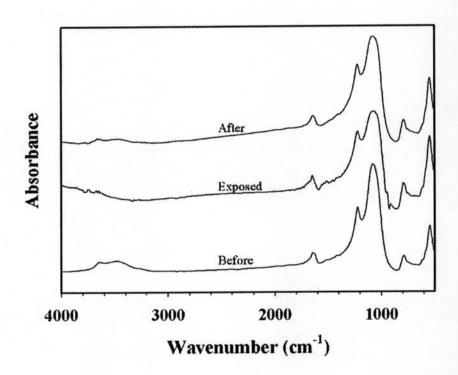


Figure M3 The FT-IR spectra of NaZSM-5(23); before, during and after NH₃ exposure.

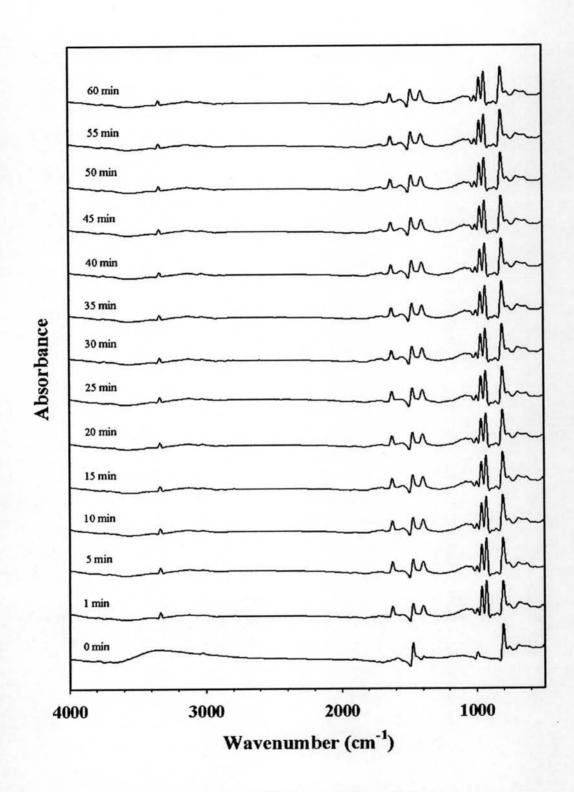


Figure M4 The FT-IR spectra of 50:1dPPP versus time of NH₃ exposure.

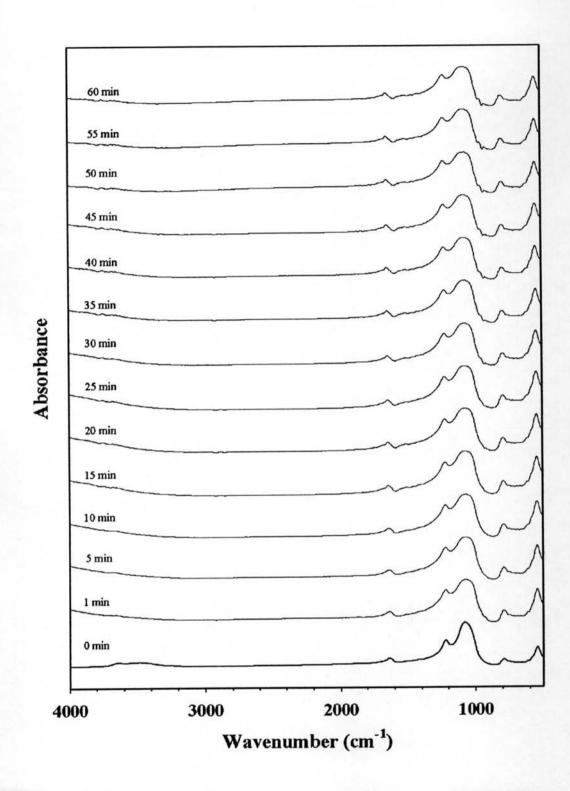


Figure M5 The FT-IR spectra of NaZSM-5(23) versus time of NH₃ exposure.

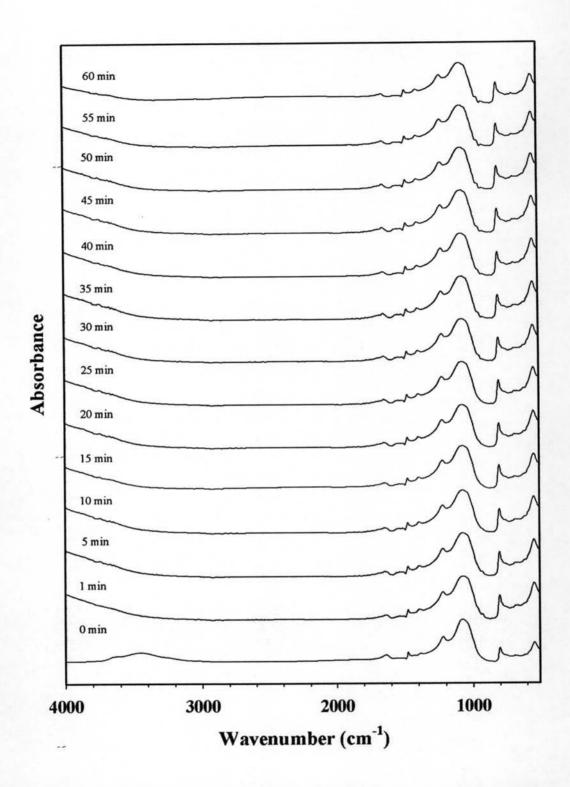


Figure M6 The FT-IR spectra of 50:1dPPP(70)/NaZ23 versus time of NH₃ exposure.

Table M1 Peak position in FT-IR spectra of 50:1dPPP before, during and after NH₃ exposure

Characteristic bands	Ref. (cm ⁻¹)	Wavenumber (cm ⁻¹)					
		before	exposed	after			
NH ₃ vibration	3336 ^{a,b}	-	3333.06	-			
C-H stretching of aromatic	3029°	3021.6	3021.6	3021.6			
NH ₃ vibration	1627.5 ^{a,b}	-	1624.47	-			
Quinoid structure	1600 ^d	1593	1554	1592			
C-C stretching of aromatic	1480 ^d	1475.8	1474.25	1473.96			
NH ₃ vibration	1397 ^b		1398.5	1399.3			
C-H in-plane vibration of para disubstituted phenyl rings	1000e	995.9	998.7	995			
NH ₃ vibration	968.1 ^{a,b}	-	964.3	/ -			
NH ₃ vibration	931.6 ^b	-	929.5	100			
C-H out of plane vibration of para disubstituted phenyl rings	817-805 ^f	803.7	803.5	803.2			
C-H out of plane vibration mono substituted phenyl rings	760 ^g	760.7	757.8	760.0			
C-H out of plane vibration of mono substituted phenyl rings	692 ^h	688.5	687.7	691.5			

Table M-2 Peak position in FT-IR spectra of NaZSM-5(23) before, during and after NH₃ exposure

Characteristic bands	Ref. (cm ⁻¹)	Wavenumber (cm ⁻¹)					
		before	exposed	after			
Silanol group	3638i	3645	-	3662			
3	N/A	1642	1645.6	1640.5			
Vibrations of T-O-T linkages	1232 ⁱ	1222	1222.8	1223.3			
Vibrations of T-O-T linkages	1062 ⁱ	1079	1063.2	1077.8			
NH ₃ vibration	968.1 ^{a,b}	-	951.8	-			
NH ₃ vibration	931.6 ^b	-	916.4	-			
Vibrations of T-O-T linkages	797 ⁱ	790	792.6	791			
Vibrations of T-O-T linkages	581 ¹	544	544.6	544			

Table M3 Peak position in FT-IR spectra of 50:1dPPP(70)/NaZ23 before, during and after NH₃ exposure

Characteristic bands	Ref. (cm ⁻¹)	Wavenumber (cm ⁻¹)					
U=		before	exposed	after			
Silanol group	3638 ⁱ	3639		3663			
5 1	N/A	1638	1645.1	1640.2			
C-C stretching of aromatic	1480 ^d	1479	1475.7	1476			
NH ₃ vibration	1397 ^b	1383	1392.2	1387.2			
Vibrations of T-O-T linkages	1232 ⁱ	1223	1223	1223			
Vibrations of T-O-T linkages	1062 ⁱ	1078	1079.7	1078			
NH ₃ vibration	968.1 ^{a,b}		950.9	-			
NH ₃ vibration	931.6 ^b	-	916.4				
C-H out of plane vibration of para disubstituted phenyl rings	817-805 ^f	803	803.3	803.4			
C-H out of plane vibration of mono substituted phenyl rings	692 ^h	692	690.2	688.1			
Vibrations of T-O-T linkages	581 ⁱ	543	545.1	543.9			

^aSu, et al., 2000, ^bYin et al., 1997, ^cDale, 1957 and Castiglioni et al., 1989,

^dPham et al., 1990, ^eSoubiran et al., 1998, ^fFroyer et al., 1985,

^gKvarnstrom et al. 1985, ^hRakovic et al., 1991, ⁱVenkatathri N., 2006

CURRICULUM VITAE

Name:

Ms. Pimchanok Phumman

Date of Birth:

August 6th, 1982

Nationality:

Thai

University Education:

2001-2005 Bachelor Degree of Science, Faculty of Science, King Mongkut's Institute of Technology of Ladkrabang. Bangkok, Thailand