

# Adsorption of HCN from Pyrolysis of Tobacco Leaves with Na<sub>2</sub>CO<sub>3</sub>, NaHCO<sub>3</sub>, and Proline

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## Abstract

Hydrogen cyanide (HCN) was produced by flameless combustion of dried tobacco leaves for investigating the adsorption on  $Na_2CO_3$ ,  $NaHCO_3$  and proline (HNC<sub>4</sub>H<sub>7</sub>COOH). The concentration of HCN could be produced steadily at 0.73 mg/dm<sup>3</sup>. For the given 245 cm<sup>3</sup> of smoke with the flow rate of 17.5 cm<sup>3</sup>/s, the efficiency of adsorption of HCN was improved to 80% by the addition of the optimal amounts of  $Na_2CO_3$ ,  $NaHCO_3$  and proline (HNC<sub>4</sub>H<sub>7</sub>COOH), which were 10, 20, and 30 mg respectively. In the circumstances, the breakthrough concentration was reduced to 0.15 - 0.20 mg/dm<sup>3</sup>.

## **Keywords**

Adsorption, HCN, Tobacco Leaves, Sodium Carbonate, Sodium Bicarbonate, Proline

## **1. Introduction**

The increase in population requires more energy and food. To reduce the costs, industrial factories look for changing waste materials; such as wood, bark, bagasse, rice husk and agricultural straw, to energy instead of conventional fossil fuels. Meanwhile, cultivators might burn the remaining crop after cultivation for replantation. Various toxic gases can be produced by combustion of this biomass [1]-[13]. As this biomass has nitrogen as a constituent, besides carbon and hydrogen, nitrogen compounds, such as HNCO, HCN,  $NO_x$ ,  $N_2O$ , and  $NH_3$ , can be generated by combustion [1] [2] [3].

Hydrogen cyanide, HCN, is generally produced by the combustion of various synthetic materials containing nitrogen constituent, such as polyamide (nylon), 1,3,5-triazine-2,4,6-triamine (melamine) and polyacrylonitrile. It can also be produced by the pyrolysis of biomass containing amino acids, heterocyclic compounds of nitrogen or dicarboxylic acids [8] [9] [10] [11] [12]. Even though

the amount of hydrogen cyanide, HCN, from biomass pyrolysis is small, it is harmful to public health. It is so toxic asphyxiant that it inhibits the metabolism process. Moreover, cyanide ions might block oxidative respiration and might also cause cancer [14] [15] [16].

Historically, various substances, such as carbon, activated carbon, zeolite, catalysts, nanotube and architectural materials, were studied for the adsorption of hydrogen cyanide [17]-[33]. Oliver *et al.* [30] revealed that copper containing synthetic activated carbons produced from porous sulfonated styrene/divinylbenzene resins could adsorb hydrogen cyanide gas effectively without the formation of  $(CN)_2$ . Rajakovic *et al.* [31] researched that the carbon materials impregnated with metal organic compounds could adsorb hydrogen cyanide, especially copper (II)-tartrate. Furthermore, the impregnated organic compounds had more efficiency than inorganic compounds. While R. R. Kotdawala *et al.* [32] exposed that the polar compounds, containing carboxyl, hydroxyl or carbonyl groups, had high efficiency for adsorbing hydrogen cyanide, depending on their induced dipole and charge-dipole interactions. Peter Branton *et al.* [33] reported that the efficiency of the adsorption of hydrogen cyanide depended on physisorption or chemisorption.

According to the previous researches on the pyrolysis of tobacco leaves, hydrogen cyanide, HCN, could be generated directly by the pyrolysis of dried tobacco leaves, which was used for this study. The direct adsorption of hydrogen cyanide on sodium carbonate (Na<sub>2</sub>CO<sub>3</sub>), sodium hydrogen carbonate (NaHCO<sub>3</sub>) and proline (pyrrolidine-2-carboxylic acid, HNC<sub>4</sub>H<sub>7</sub>COOH) compounds was investigated without producing porous adsorbents. In addition to their chemical structures, the influence of both amounts of compounds and pyrolysis gas on the adsorption of hydrogen cyanide could be investigated more conveniently than porous adsorbents.

## 2. Material and Methods

#### 2.1. Material

Dried tobacco leaves were used as raw materials for producing hydrogen cyanide (HCN). They were cut into fine strips. A gram of strips of dried leaves was packed in a paper cylinder with a diameter of 8 mm. and length of 84 mm. All packs of dried leaves were kept in a container at 22°C with 60% relative humidity in order to control moisture contents of dried leaves.

The glass fiber filter pads, with 2 mm thick and 44 mm diameter, were dipped into the solutions of sodium carbonate ( $Na_2CO_3$ ), sodium hydrogen carbonate ( $NaHCO_3$ ) or proline (pyrrolidine-2-carboxylic acid,  $HNC_4H_7COOH$ ) for adsorption experiment. The concentration of these solutions could be varied for variation of amounts of compounds on the pads for investigating the influence on the adsorption of HCN. The pads were dried at 100°C for 3 hours. They were put in the adsorption experimental set, consisting of hydrogen cyanide generation with flameless combustion HCN, adsorption on an investigated compound,

collecting the unadsorbed hydrogen cyanide in the 0.625 M sodium hydroxide solution, and a piston vacuum pump, respectively, as shown in **Figure 1**.

## 2.2. The Adsorption Experimental

Each pack of dried leaves was combusted flamelessly, while the combustion gas was pumped by the piston vacuum pump through the pad containing a selected adsorbing compound with the flow rate of 17.5 cm<sup>3</sup>/s. The unadsorbed hydrogen cyanide was collected in the 40 cm<sup>3</sup> of 0.625 M NaOH solution for analyzing the breakthrough amount of hydrogen cyanide. Each condition of adsorption was repeated at least 4 times.

#### 2.3. Quantifications

The NaOH solution, which collected the unadsorbed amount of hydrogen cyanide, was analyzed by the continuous flow analyzer, Skalar, with the detection limits of  $CN^-$  of 0.02 mg/dm<sup>3</sup>. In addition, the amount of hydrogen cyanide, which was adsorbed on the pad was analyzed in the same manner as the unadsorbed hydrogen cyanide above, by dissolving the adsorbing pad in 40 cm<sup>3</sup> of 0.625 M NaOH solution. The adsorption efficiency was evaluated by Equation (1).

Adsorption Efficiency =  $\frac{\text{Amount HCN adsorbed} \times 100}{(\text{Amount HCN adsorbed} + \text{Amount HCN unadsorbed})} (1)$ 

#### 3. Results and Discussion

#### 3.1. The Adsorption of HCN on Na<sub>2</sub>CO<sub>3</sub>, NaHCO<sub>3</sub>, and Proline

Hydrogen cyanide was generated by flameless combustion at the end of a cylindrical pack of dried strips of tobacco leaves. Meanwhile, the combustion gas was pumped steadily through the other end of the pack via the adsorption zone at the flow rate of 17.5 cm<sup>3</sup>/s by the piston vacuum pump. The unadsorbed hydrogen cyanide was collected cumulatively in the 0.625 M NaOH solution after passing the adsorption zone as shown in Figure 1. With consecutive flameless combustion of up to 6 packs of dried leaves, the total amount of hydrogen cyanide which was, collected in the NaOH solution, increased proportionally to the cumulative amount of smoke passing the adsorption zone, as shown in Figure 2. This demonstrated that hydrogen cyanide was produced steadily at the average concentration about 0.73 mg/dm<sup>3</sup> by the apparatus for adsorption investigation. In addition, the analysed results of the amount of HCN adsorbed on the pad revealed that the cumulative amount adsorbed on the pad was slightly more than a half of the amount of HCN flowing through the adsorption zone for the whole range of investigation. This has demonstrated the unsaturated adsorption for the total amount of HCN investigated.

With only the filter pad in the adsorption zone, the cumulative amount of hydrogen cyanide adsorbed on the pad increased proportional to the cumulative volume of smoke flowing through the adsorption zone in the same manner as



**Figure 1.** The experimental set for adsorption of hydrogen cyanide from combustion of dried stripes of tobacco leaves.



**Figure 2.** The relation of cumulative total and adsorbed amount of hydrogen cyanide generated by flameless combustion with the cumulative smoke volume flowing through the adsorption zone with flow rate of 17.5 cm<sup>3</sup>/s.

the total amount of HCN produced as shown in **Figure 2**. The concentration of HCN was reduced at least 70% by adsorption on the filter pad for the first 245 cm<sup>3</sup> of smoke, which was produced by the first pack of dried leaves, hence the breakthrough concentration of HCN dropped below 0.20 mg/dm<sup>3</sup>, as shown in **Figure 3**. For additional amount of smoke produced by the second to the sixth pack of dried leaves, the concentration of HCN was reduced slightly more than 50% by further adsorption on the filter pad. Consequently, the breakthrough concentration of HCN increased to just below 0.35 mg/dm<sup>3</sup>. In the circumstances, the amount of HCN adsorbed on the pad was increased from 0.24 mg/g of pad for the first pack of dried leaves.



**Figure 3.** The breakthrough concentrations of hydrogen cyanide of the adsorption systems containing 10 mg  $Na_2CO_3$ ,  $NaHCO_3$  and proline (HNC<sub>4</sub>H<sub>7</sub>COOH), respectively, relative to the filter pad only, with flow rate of 17.5 cm<sup>3</sup>/s.

With 10 mg of sodium carbonate (Na<sub>2</sub>CO<sub>3</sub>) on the filter pad, the breakthrough concentration of HCN dropped below 0.10 mg/dm<sup>3</sup> for the first 200 cm<sup>3</sup> of smoke, as shown in **Figure 3**. This corresponded to at least 85% of HCN in the smoke was adsorbed on the pad containing 10 mg sodium carbonate. The breakthrough concentration of HCN increased gradually to just above 0.25 mg/dm<sup>3</sup> with further adsorption of HCN in smoke beyond 200 cm<sup>3</sup> up to 1500 cm<sup>3</sup>. The breakthrough concentration approached to the values of the filter pad without sodium carbonate. The results indicated that most sodium carbonate on the pad almost reached the saturated adsorption. With the similar amount of sodium bicarbonate (NaHCO<sub>3</sub>) or proline (HNC<sub>4</sub>H<sub>7</sub>COOH) on the filter pad instead of sodium carbonate, the breakthrough concentrations of HCN were similar to that of sodium carbonate. The similar breakthrough concentration of HCN might relate to the presence of common carbonyl group on these compounds [32].

#### 3.2. The Effect of Amounts of Na<sub>2</sub>CO<sub>3</sub>, NaHCO<sub>3</sub>, and Proline

As the breakthrough concentration of HCN increased drastically within the range of the first 245 cm<sup>3</sup> of smoke, of which could be produced by each pack of dried tobacco leaves with the flow rate of 17.5 cm<sup>3</sup>/s, the influence of the amount of these compounds on the adsorption of HCN was investigated. For sodium carbonate (Na<sub>2</sub>CO<sub>3</sub>), the adsorption efficiency was almost constant at 80% by increasing the amount more than 10 mg, as shown in **Figure 4**. In the circumstances, the breakthrough concentration of HCN was reduced to just below 0.15 mg/dm<sup>3</sup>, as shown in **Figure 5**. By reducing the amount of sodium carbonate on the filter pad, the efficiency dropped drastically to 70%, 60% and just above 50% respectively. The maximum adsorption efficiency for the smoke produced by



**Figure 4.** The total adsorption efficiency of hydrogen cyanide on  $Na_2CO_3$ ,  $NaHCO_3$  and proline (HNC<sub>4</sub>H<sub>7</sub>COOH), for 245 cm<sup>3</sup> of smoke with flow rate of 17.5 cm<sup>3</sup>/s.



**Figure 5.** The breakthrough concentration of hydrogen cyanide of the adsorption on  $Na_2CO_3$ ,  $NaHCO_3$  and proline (HNC<sub>4</sub>H<sub>7</sub>COOH), for 245 cm<sup>3</sup> of smoke with flow rate o17.5 cm<sup>3</sup>/s.

each pack of dried tobacco leaves was achieved by 10 mg sodium carbonate. In other words, the optimum amount of sodium carbonate for adsorption of HCN from 245  $\text{cm}^3$  of smoke was 10 mg.

By using 10 mg of sodium bicarbonate (NaHCO<sub>3</sub>) on the filter pad, the adsorption efficiency was about 70%, as shown in Figure 4. With the double amount of sodium bicarbonate (NaHCO<sub>3</sub>) on the filter pad, the adsorption efficiency was improved from 70% to almost 80%. In the circumstances, the breakthrough concentration of HCN dropped in the same manner as that of sodium carbonate but was above 0.15 mg/dm<sup>3</sup>, as shown in **Figure 5**. By reducing the amount of sodium bicarbonate to a half of the base condition, the adsorption efficiency was just above 50%, while the breakthrough concentration became just below 0.3 mg/dm<sup>3</sup>. The atomic polarization of H atom on sodium bicarbonate was much lower than that of Na atom on sodium carbonate [34] [35], therefore, the adsorption efficiency on sodium bicarbonate was less than that of sodium carbonate for the same amount on the filter pad. The maximum adsorption efficiency could be achieved by adding the amount of sodium bicarbonate as double amount of the optimum amount of sodium carbonate.

With 10 mg of proline on the filter pad, the adsorption efficiency was just 60% under the given conditions. This was influenced by the presence of organic functional group ( $HNC_4H_7$ -) on proline molecule, of which the polarizability was much less than those of Na and H atoms sodium carbonate and sodium bicarbonate, respectively. It could be improved quite proportionally to the increasing amount of proline. The maximum adsorption efficiency at 80% could be achieved by adding 30 - 40 mg proline on the filter pad, as shown in **Figure 4**. In the circumstances, the breakthrough decreased gradually from just above 0.2 mg/dm<sup>3</sup> to just above 0.15 mg/dm<sup>3</sup>, as shown in **Figure 5**. By reducing the amount proline on the pad, the adsorption efficiency dropped drastically to just above 50% in the same manner as both sodium carbonate and sodium bicarbonate. Therefore, the optimum amount of proline for adsorption of HCN from 245 cm<sup>3</sup> smoke was 30 mg.

## 4. Conclusion

Hydrogen cyanide could be produced steadily by flameless combustion of dried stripped tobacco leaves with concentration of 0.73 mg/dm<sup>3</sup>. With the flow rate of 17.5 cm<sup>3</sup>/s, a half of hydrogen cyanide produced could be adsorbed on the filter pad, with specific adsorption of 0.24 mg HCN/ g pad, under which the break-through was reduced to 0.3 - 0.4 mg/dm<sup>3</sup> at the room temperature. With the addition of Na<sub>2</sub>CO<sub>3</sub>, NaHCO<sub>3</sub>, or proline (HNC<sub>4</sub>H<sub>7</sub>COOH) on the pad, the maximum efficiency for adsorption of HCN in the smoke of 245 cm<sup>3</sup> could be improved to about 80%. In the circumstances, the optimum amounts of Na<sub>2</sub>CO<sub>3</sub>, NaHCO<sub>3</sub>, and proline were 10, 20, and 30 mg, respectively. While the break-through concentration of HCN was reduced to the range of 0.15 - 0.20 mg/dm<sup>3</sup>.

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## **Conflicts of Interest**

The authors declare no conflicts of interest regarding the publication of this paper.

#### References

- Lindsay, A.E., Greenbaumb, A.R. and O'Hare, D. (2004) Analytical Techniques for Cyanide in Blood and Published Blood Cyanide Concentrations from Healthy Subjects and Fire Victims. *Analytica Chimica Acta*, 511, 185-195.
- Siegler, D.S. (1976) Plants of the Northeastern United States that Produce Cyanogenic Compounds. *Economic Botany*, 30, 395-407. https://doi.org/10.1007/BF02904662
- Karlsson, H.L. (2004) Ammonia, Nitrous Oxide and Hydrogen Cyanide Emissions from Five Passenger Vehicles. *Science of the Total Environment*, **334-335**, 125-132. https://doi.org/10.1016/j.scitotenv.2004.04.061
- [4] Yi, S.-C., Hajaligol, M.R. and HoonJeong, S. (2005) The Prediction of the Effects of Tobacco Type on Smoke Composition from the Pyrolysis Modeling of Tobacco Shreds. *Journal of Analytical and Applied Pyrolysis*, 74, 181-192.
- [5] Barber, T.R., Lutes, C.C., Doorn, M.R.J., Fuchsman, P.C., Timmenga, H.J. and Crouch, R.L. (2003) Aquatic Ecological Risks Due to Cyanide Releases from Biomass Burning, *Chemosphere*, 50, 343-348.
- [6] Bekkeringa, J., Broekhuis, A.A. and van Gemert, W.J.T. (2010) Optimisation of a Green Gas Supply Chain—A Review. *Bioresource Technology*, **101**, 450-456. https://doi.org/10.1016/j.biortech.2009.08.106
- [7] Yuan, S., Zhou, H., Li, J., Chen, X. and Wang, F. (2011) HCN and NH<sub>3</sub> (NOx Precursors) Released under Rapid Pyrolysis of Biomass/Coal Blends. *Journal of Analytical and Applied Pyrolysis*, **92**, 463-469. <u>https://doi.org/10.1016/j.jaap.2011.08.010</u>
- [8] Johnson, W.R. and Kang, J.C. (1971) Mechanisms of Hydrogen Cyanide Formation from the Pyrolysis of Amino Acids and Related Compounds. *The Journal of Organic Chemistry*, **36**, 189-192.
- [9] Tso, T.C., Chaplin, J.F., Adams, J.D. and Hoffmann, D. (1982) Simple Correlation and Multiple Regression among Leaf and Smoke Characteristics of Burley Tobaccos. *Beiträgezur Tabakforschung International*, 11, 141-150.
- [10] Ren, Q., Zhao, C., Chen, X., Duan, L., Li, Y. and Ma, C. (2011) NO<sub>x</sub> and N<sub>2</sub>O Precursors (NH<sub>3</sub> and HCN) from Biomass Pyrolysis: Co-Pyrolysis of Amino Acids and Cellulose, Hemicellulose and Lignin. *Proceedings of the Combustion Institute*, **33**, 1715-1722. <u>https://doi.org/10.1016/j.proci.2010.06.033</u>
- [11] Hansson, K.-M., Samuelsson, J., Tullin, C. and Åmand, L.-E. (2004) Formation of HNCO, HCN, and NH<sub>3</sub> from the Pyrolysis of Bark and Nitrogen Containing Model Compounds. *Combustion and Flame*, **137**, 265-277. https://doi.org/10.1016/j.combustflame.2004.01.005
- [12] Rickert, W.S. and Stockwell, P.B. (1979) Automated Determination of Hydrogen Cyanide Acrolein and Total Aldehydes in the Gas Phase of Tobacco Smoke Labstat Incorporated, 262 Manitou Drive, Kitchener, Ontario, Canada, N2C 1L3. *The Journal of Automatic Chemistry*, **3**, 152-154.
- [13] Zhang, Z.-W., Xu, Y.-B., Wang, C.-H., Che, K.-B., Tong, H.-W. and Liu, S.-M. (2011) Direct Determination of Hydrogen Cyanide in Cigarette Mainstream Smoke by Ion Chromatography with Pulsed Amperometric Detection. *Journal of Chromatography A*, **1218**, 1016-1019. <u>https://doi.org/10.1016/j.chroma.2010.12.100</u>

- [14] (2013) The Emergency Response Safety and Health Database. http://www.cdc.gov/niosh/ershdb/EmergencyResponseCard\_29750038.html
- [15] Bhattacharya, R. and Flora, S.J.S. (2009) Cyanide Toxicity and Its Treatment: Section II Agents That Can Be Used as Weapons of Mass Destruction. In: *Handbook of Toxicology of Chemical Warfare Agents*, Elsevier Inc., Amsterdam, 255-270.
- [16] World Health Organisation (WHO) (2004) Hydrogen Cyanide and Cyanides: Human Health Aspects. Concise International Chemical Assessment Document, 61.
- [17] Zhao, H., et al. (2006) Catalytic Oxidation of HCN over a 0.5 % Pt/Al<sub>2</sub>O<sub>3</sub> Catalyst. *Applied Catalysis B: Environmental*, 65, 282-290. <u>https://doi.org/10.1016/j.apcatb.2006.02.009</u>
- [18] KrÖcher, O. and Elsener, M. (2009) Hydrolysis and Oxidation of Gaseous HCN over Heterogeneous Catalysts. *Applied Catalysis B: Environmental*, **92**, 75-89. <u>https://doi.org/10.1016/j.apcatb.2009.07.021</u>
- [19] Seredych, M., van der Merwe, M. and Bandosz, T.J. (2009) Effects of Surface Chemistry on the Reactive Adsorption of Hydrogen Cyanide on Activated Carbons. *Carbon*, 47, 2456-2465.
- [20] Hou, H., Zhu, Y., Tang, G. and Hu, Q. (2012) Lamellar *y*-AlOOH Architectures: Synthesis and Application for the Removal of HCN. *Materials Characterization*, 68, 33-41.
- [21] Gupta, N., Balomajumder, C. and Agarwal, V.K. (2010) Enzymatic Mechanism and Biochemistry for Cyanide Degradation: A Review. *Journal of Hazardous Materials*, 176, 1-13.
- [22] Zhao, Q., Tian, S., Yan, L., Zhang, Q. and Ning, P. (2015) Novel HCN Sorbents Based on Layered Double Hydroxides: Sorption Mechanism and Performance. *Journal of Hazardous Materials*, 285, 250-258. https://doi.org/10.1016/j.jhazmat.2014.11.045
- [23] Deveci, H., Yazıcı, E.Y., Alp, I. and Uslu, T. (2006) Removal of Cyanide from Aqueous Solutions by Plain and Metal-Impregnated Granular Activated Carbons. *International Journal of Mineral Processing*, **79**, 198-208. <u>https://doi.org/10.1016/j.minpro.2006.03.002</u>
- [24] Samanta, P.N. and Das, K.K. (2013) Adsorption Sensitivity of Zigzag GeC Nanotube towards N<sub>2</sub>, CO, SO<sub>2</sub>, HCN, H<sub>3</sub>, and H<sub>2</sub>CO Molecules. *Chemical Physics Letters*, 577, 107-113. <u>https://doi.org/10.1016/j.cplett.2013.05.055</u>
- [25] Shi, L.B., Wang, Y.P. and Dong, H.K. (2015) First Principle Study of Structural, Electronic, Vibrational and Magnetic Properties of HCN Adsorbed Graphene Doped with Cr, Mn and Fe. *Applied Surface Science*, **329**, 330-336.
- [26] Colin-Garcia, M., Ortega-Gutierrez, F., Ramos-Bernal, S. and Negron-Mendoza, A.
  (2010) Heterogeneous Radiolysis of HCN Adsorbed on a Solid Surface. *Nuclear Instruments and Methods in Physics Research*, 619, 83-85. https://doi.org/10.1016/j.nima.2009.10.074
- [27] Smith, J.W.H., Westreich, P., Abdellatif, H., Filbee-Dexter, P., Smith, A.J., Wood, T.E., Croll, L.M., Reynolds, J.H. and Dahn, J.R. (2010) The Investigation of Copper-Based Impregnated Activated Carbons Prepared from Water-Soluble Materials for Broad Spectrum Respirator Applications. *Journal of Hazardous Materials*, 180, 419-428. <u>https://doi.org/10.1016/j.jhazmat.2010.04.047</u>
- [28] Ahmadi Peyghan, A., Hadipour, N. and Bagheri, Z. (2013) Effects of Al-Doping and Double Antisite Defect on the Adsorption of HCN on a BC<sub>2</sub>N Nanotube: DFT Studies. *The Journal Physical Chemistry C*, **117**, 2427-2432.

- [29] Soltani, A., Baei, M.T., Ghasemi, A.S., Tazikeh Lemeski, E. and Komail, H.A. (2014) Adsorption of Cyanogen Chloride over Al- and Ga-Doped BN Nanotubes. *Superlattices and Microstructures*, **75**, 564-575. https://doi.org/10.1016/j.jhazmat.2010.04.047
- [30] Oliver, T.M., Jugoslav, K., Aleksandar, P. and Nikola, D. (2005) Synthetic Activated Carbons for the Removal of Hydrogen Cyanide from Air. *Chemical Engineering* and Processing, 44, 1181-1187. <u>https://doi.org/10.1016/j.cep.2005.03.003</u>
- [31] Rajakovic, L.J.V., Ilic, M.R., Jovanic, P.B. and Radosevic, P.B. (1995) Stoichiometric Analysis of Chemisorption of Hydrogen-Cyanide onto Activated Carbon Cloth. *Carbon*, 33, 1433-1441.
- [32] Kotdawala, R.R., Kazantzis, N. and Thompson, R.W. (2008) Molecular Simulation Studies of Adsorption of Hydrogen Cyanide and Methyl Ethyl Ketone on Zeolite NaX and Activated Carbon. *Journal of Hazardous Materials*, **159**, 169-176. https://doi.org/10.1016/j.jhazmat.2008.01.045
- [33] Branton, P., Lu, A.-H. and Schuth, F. (2009) The Effect of Carbon Pore Structure on the Adsorption of Cigarette Smoke Vapor Phase Compounds. *Carbon*, 47, 1005-1011. <u>https://doi.org/10.1016/j.carbon.2008.12.003</u>
- [34] Mason, E.A. and McDaniel, E.W. (1988) Transport Properties of Ions in Gases. John Wily and Sons, Hoboken.
- [35] Schwerdtfeger, P. (2006) Atomic Static Dipole Polarizabilities. In: Maroulis, G., Ed., Computational Aspects of Electric Polarizability Calculations. Atoms, Molecules and Clusters, IOS Press, Amsterdam, 1-32. https://doi.org/10.1142/9781860948862\_0001

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