

MODERN CATALYSTS FOR ACETYLENE HYDROCHLORATION

Rakhmatov Doniyor Khudayorovich

Assistant of the Yangiyer Branch of The Tashkent Institute of Chemical Technology.

Sirdarya Region, Yangier, Tinchlik St. 1.

E-mail: doniyor87@mail.ru

Rakhimkulov Shokir Rizoevich

Assistant of the Yangiyer Branch of the Tashkent Institute of Chemical Technology.

Sirdarya Region, Yangier, Tinchlik St. 1.

E-mail: shokir0404@mail.ru

Abdullaev Bakhtishod Menglikul ugli

Master of the 2nd Course of the Karshi Engineering and Economic Institute.

Kashkadarya Region Karshi, St. Mustakillik 224

Norboev Farrukh Abdumalikovich

Master of the 2nd Course of the Karshi Engineering and Economic Institute.

Kashkadarya Region Karshi, St. Mustakillik 224

Rakhmatov Durdonakhon Khudoyor kizi

Head Office of the Yangiyer Branch of the Tashkent Institute of Chemical Technology.

Sirdarya Region, Yangiyer, Tinchlik St. 1.

E-mail: durdonakhon93@mail.ru

ABSTRACT

The paper presents the results of the analysis of patent literature on the development of mercury-free catalysts for the process of acetylene hydrochlorination. Catalysts based on gold chloride and base metal chlorides are considered, and a comparative analysis of these catalysts is carried out.

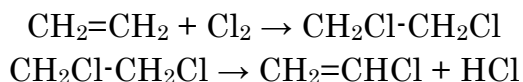
Keywords: vinyl chloride, acetylene hydrochlorination, gold chloride, acetylene, hydrochloride, catalyst, base metals.

INTRODUCTION

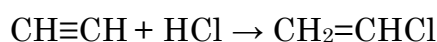
At the moment, in Russia, almost a third of vinyl chloride is obtained by hydrochlorination of acetylene [1]. There are two methods using acetylene as raw material:

1. Hydrochlorination of acetylene, where the raw material is pure acetylene and hydrogen chloride;
2. Combined method, where the raw material is pyrogas with an acetylene content of at least 8% and hydrogen chloride.

The latter method involves carrying out the process in several stages, the first of which is the direct chlorination of ethylene from pyrogas to 1,2-dichloroethane, which is then cracked to produce vinyl chloride and hydrogen chloride.



At the second stage, the formed hydrogen chloride enters the stage of acetylene hydrochlorination.



The hydrochlorination process uses a catalyst consisting of mercury(II) chloride on activated carbon. Mercury chlorides are toxic (hazard class 1), and they also form a mobile complex with hydrogen chloride, which is carried away from the catalyst surface.

Since 2013, the number of studies and patents on mercury-free acetylene hydrochlorination catalysts has increased dramatically. As a result of the analysis, it was found that the most active in relation to the hydrochlorination reaction are the chlorides of gold, iridium, platinum and palladium (Fig. 1). Since catalysts based on gold salts have shown the highest activity, most of the patents on non-mercury catalysts for acetylene hydrochlorination suggest using gold chlorides as the active substance [2].

With the new catalyst, the conversion of acetylene and the selectivity of the process were more than 99.9% [3], higher than that of the mercury catalyst. The disadvantage of this catalyst is the rapid deactivation due to the reduction of gold.

This problem is solved by introducing various additives: copper salts, salts of other noble metals, organic compounds such as phenanthroline, thiourea, etc. Additives also make it possible to reduce the gold content in the catalyst, thereby reducing its cost.

The addition of thiourea [4] to gold salts makes it possible to reduce the gold content in the catalyst to 0.25% wt., but the acetylene conversion and selectivity also become lower (90% and 95%, respectively).

Salts of ruthenium, palladium or platinum [5, 6] as additives to the catalyst exhibit a stabilizing effect - the service life of the catalyst increases from 200 h without loss of activity to 500 h. Using this catalyst, the acetylene conversion is 98%, and the selectivity is 99%. The disadvantage of these additives is the high price of precious metals.

The most promising are catalysts based on gold and copper chlorides [7, 8]. Copper chloride can catalyze the hydrochlorination reaction without gold chloride, which can reduce the amount of noble metal used and reduce the cost of the catalyst. Also, the addition of copper chloride makes it possible to increase the service life of the catalyst, acting as a stabilizer that prevents the reduction of Au_3^+ to Au_0 .

Since gold is an expensive noble metal, interest has increased in the catalytic ability of base metals in relation to the hydrochlorination of acetylene.

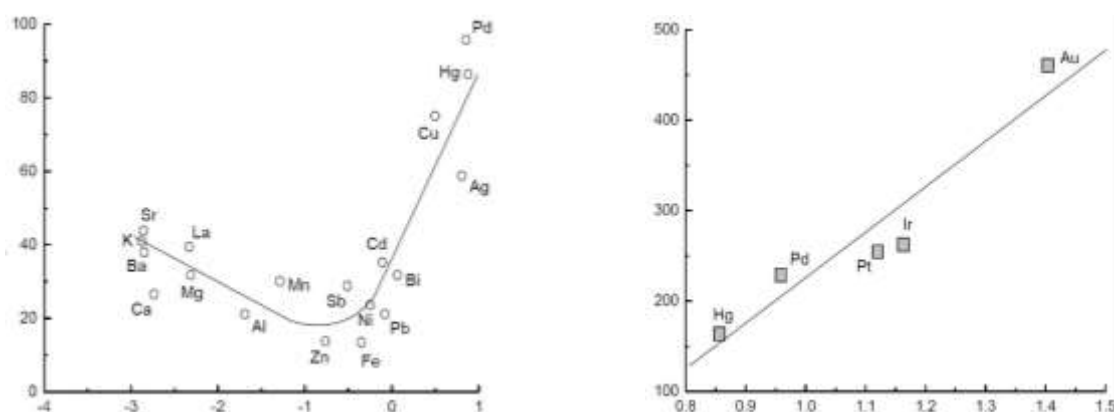


Fig. - Activities of various metal chlorides on AC for the process of acetylene hydrochlorination

Copper or barium is used as a catalyst component based on base metals [9, 10]. In this case, the conversion of acetylene is reduced to 30%.

Various additives increase the acetylene conversion to 80-90%, but the catalyst life is less than 100 hours without loss of activity. These additives include: phosphoric acid, nitrogen-containing activated carbon, fatty organic acids, complexes of rubidium chlorides with ammonia, etc.

At the moment, the studied catalysts without the addition of noble metals are less active than the catalysts containing gold.

The most promising is a catalyst containing gold and copper chlorides on activated carbon [11], where the total metal content is 3% wt., and the ratio Au:Cu = 1:5. This catalyst is more active, safer than mercury (hazard class 3, and for mercury - 1st) and has the ability to regenerate.

LITERATURE

1. Флид М. Р. Состояние и перспективы развития производства винилхлорида – мономера для получения ПВХ [Электронный ресурс]. – Режим доступа: http://www.creonconferences.com/upload/iblock/2f3/4.Flid_Sintez.pdf
2. 19. Davies C. Gold Catalysts for the Hydrochlorination of Acetylene: автореф. дис. ... канд. наук / C. Davies – 2012. – 160 с.
3. Hao XU, Guohua LUO Green Production of PVC from Laboratory to Industrialization: State-of-the-art Review of Heterogeneous Non-Mercury Catalysts for Acetylene Hydrochlorination / Hao XU, Guohua LUO – Текст: электронный // Journal of Industrial and Engineering Chemistry. – 2018. – № 10. – URL: <https://doi.org/10.1016/j.jiec.2018.05.009>
4. One-pot synthesis of nitrogen and sulfur co-doped activated carbon supported AuCl₃ as efficient catalysts for acetylene hydrochlorination // X.-X. Di, J. Zhao, Y. Yu, X.-L. Xu, S.-C. Gu, H.-H. He, T.-T. Zhang, X.-N. Li, Chin / Chemical Letters. – 2016. - № 27. – С. 1567-1571.
5. Пат. CN103623838A МПК C02C 21/06, B01J 27/10, B01J 27/14, B01J 27/16. Ru-Pt-Cu катализатор для гидрохлорирования ацетилена синтеза винилхлорида / заявитель и патентообладатель: Чжан Цзиньли, Шэн Вэй, Ли Вэй, Хан Ю, Дай Бин. - №201210307780A; заявл. 24.08.2012 ; опубл. 12.03.2014.
6. Пат. CN103894195A МПК B01J23 / 644; C07C17 / 08; C07C21 / 06. Катализатор Ru-Bi, используемый для гидрохлорирования ацетилена, а также способ получения и

- применения катализатора Ru-Bi / заявитель и патентообладатель: Guohua Luo, Kai Zhou, Wei Wang, Fei We. - №201410142951A ; заявл. 04.04.2014 ; опубл. 02.07.2014.
7. Пат. CN201810775463 МПК МПК B01J31 / 22; C07C17 / 08; C07C21 / 06. Катализатор и способ получения комплекса меди для реакции гидроксихлорирования ацетилена / заявитель и патентообладатель: Университет Шихэцзи. - №201810775463A ; заявл. 16.07.2018 ; опубл. 14.12.2018.
 8. A ligand coordination approach for high reaction stability of an Au–Cu bimetallic carbon-based catalyst in the acetylene hydrochlorination process // H. Xu, J. Si, K. Zhou, G. Luo / Catalytic Science and Technology. – 2016. - № 6. – С. 13571366.
 9. Hydrochlorination of acetylene using supported bimetallic Au-based catalysts / M. Conte, A.F. Carley, G. Attard, A.A. Herzing, C.J. Kiely, G.J. Hutchings // Journal of Catalysis. - 2008. - №257. – С. 190-198.
 10. A novel high-stability Au(III)/Schiff-based catalyst for acetylene hydrochlorination reaction // C. Huang, M. Zhu, L. Kang, B. Dai / Catalysis Communications. - 2014. - №54. - С. 61-65.
 11. A ligand coordination approach for high reaction stability of an Au–Cu bimetallic carbon-based catalyst in the acetylene hydrochlorination process // H. Xu, J. Si, K. Zhou, G. Luo / Catalytic Science and Technology. – 2016. - № 6. – С. 1357-1366.