

纳米MoS₂ 催化剂的合成及其在加氢脱硫反应中的应用

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摘要 通过羰基钼和升华硫反应制备了晶片层数为3~5, 比表面积为 71 m²/g 的纳米 MoS₂ 催化剂, 并考察了其催化十二硫醇或二苯并噻吩的加氢脱硫活性。结果表明, 在 3.0 MPa 初始氢气压力下, 该催化剂在 200 °C 和 280 °C 就可使十二硫醇和二苯并噻吩转化接近 100%。

关键词: 二硫化钼 纳米催化剂 加氢脱硫 十二硫醇 二苯并噻吩

Abstract: The MoS₂ nanocatalyst of 3~5 layers and surface area as high as 71 m²/g was synthesized via a chemical reaction between molybdenum carbonyl and chalcogen. The hydrodesulfurization (HDS) of n-dodecanethiol and dibenzothiophene (DBT) was studied over the MoS₂ nanocatalyst. Compared with the commercial MoS₂ with lower surface area of 3 m²/g, the MoS₂ nanocatalyst showed much higher activity. 100% conversion in HDS at temperatures of 200 °C for dodecanethiol and 280 °C for DBT was achieved under p(H₂) = 3.0 MPa.

Keywords: molybdenum disulfide, nanocatalyst, hydrodesulfurization, n-dodecanethiol, dibenzothiophene

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- [1] Babich I V, Moulijn J A. Fuel, 2003, 82: 607
- [2] Breysse M, Djega-Mariadassou G, Pessaire S, Geantet C, Vrinat M, Perot G, Lemaire M. Catal Today, 2003, 84: 129
- [3] Zhao D B, Wu M, Kou Y, Min E Z. Catal Today, 2002, 74: 157
- [4] Cho A, Koh J H, Lee S I, Moon S H. Catal Today, 2010, 149: 47
- [5] Zhao D B, Wu M, Kou Y, Min E Z. Catal Today, 2002, 74: 157
- [6] Yan N, Xiao Ch X, Kou Y. Coord Chem Rev, 2010, 254: 1179
- [7] Yan N, Xiao Ch X, Kou Y. Coord Chem Rev, 2010, 254: 1179
- [8] Daage M, Chianelli R R. J Catal, 1994, 149: 414
- [9] Daage M, Chianelli R R. J Catal, 1994, 149: 414
- [10] van Veen J A R, Colijn H A, Hendriks P A J M, van Welsenes A J. Fuel Proc Technol, 1993, 35: 137
- [11] van Veen J A R, Colijn H A, Hendriks P A J M, van Welsenes A J. Fuel Proc Technol, 1993, 35: 137

- [12] Devers E, Afanasiev P, Jouguet B, Vrinat M. Catal Lett, 2002, 82: 13
- [13] Devers E, Afanasiev P, Jouguet B, Vrinat M. Catal Lett, 2002, 82: 13
- [14] Tye C T, Smith K J. Catal Today, 2006, 116: 461
- [15] Tye C T, Smith K J. Catal Today, 2006, 116: 461
- [16] Duphil D, Bastide S, Rouchaud J C, Pastol J L, Legendrel B, Levy-Clement C. Nanotechnology, 2004, 15: 828
- [17] Duphil D, Bastide S, Rouchaud J C, Pastol J L, Legendrel B, Levy-Clement C. Nanotechnology, 2004, 15: 828
- [18] Ma X L, Sakanishi K, Mochida I. Ind Eng Chem Res, 1994, 33: 218
- [19] Ma X L, Sakanishi K, Mochida I. Ind Eng Chem Res, 1994, 33: 218
- [20] Singhal G H, Espino R L, Sobel J E, Huff G A Jr. J Catal, 1981, 67: 457
- [21] Singhal G H, Espino R L, Sobel J E, Huff G A Jr. J Catal, 1981, 67: 457
- [22] Vanrysselberghe V, Froment G F. Ind Eng Chem Res, 1996, 35: 3311
- [23] Vanrysselberghe V, Froment G F. Ind Eng Chem Res, 1996, 35: 3311
- [24] Dinter N, Rusanen M, Raybaud P, Kasztelan S, da Silva P, Toulhoat H. J Catal, 2009, 267: 67
- [25] Dinter N, Rusanen M, Raybaud P, Kasztelan S, da Silva P, Toulhoat H. J Catal, 2009, 267: 67
- [26] Lee J J, Kim H, Koh J H, Jo A, Moon S H. Appl Catal B, 2005, 58: 89
- [27] Lee J J, Kim H, Koh J H, Jo A, Moon S H. Appl Catal B, 2005, 58: 89
- [28] Lee J J, Kim H, Moon S H. Appl Catal B, 2003, 41: 171
- [29] Lee J J, Kim H, Moon S H. Appl Catal B, 2003, 41: 171
- [30] Liu D, Liu L H, Li G C, Liu Ch G. J Nat Gas Chem, 2010, 19: 530
- [31] Liu D, Liu L H, Li G C, Liu Ch G. J Nat Gas Chem, 2010, 19: 530
- [32] Lan L, Ge Sh H, Liu K H, Hou Y D, Bao X J. J Nat Gas Chem, 2011, 20: 117
- [33] Lan L, Ge Sh H, Liu K H, Hou Y D, Bao X J. J Nat Gas Chem, 2011, 20: 117
- [1] 杨祝红, 李力成, 王艳芳, 刘金龙, 冯新, 陆小华.磷化镍/介孔 TiO_2 催化剂的制备及其催化加氢脱硫性能[J].催化学报, 2012,33(3): 508-517
- [2] 亓雪, 石秋杰, 谌伟庆, 张荣斌.Mo 对非晶态合金 Ni-B/薄水铝石催化剂上噻吩加氢脱硫性能的影响[J].催化学报, 2012,33(3): 543-549
- [3] 李巧灵 1, 张元华 1, 陈世萍 1, 方维平 1,2, 杨意泉 1,2.反映工艺条件对管式反应器催化反应影响的转化率方程[J].催化学报, 2011,32(3): 446-450
- [4] 齐和日玛 1,3, 李会峰 2, 袁蕙 2, 张韫宏 1, 徐广通 2. Al_2O_3 性质对加氢脱硫催化剂 Co-Mo/ Al_2O_3 活性相形成的影响[J].催化学报, 2011,32(2): 240-249
- [5] 石国军, 赵鹤, 黄玉安, 沈俭一.介孔碳担载的 Co-Mo 和 Ni-Mo 加氢脱硫催化剂[J].催化学报, 2010,26(8): 961-964
- [6] 陈闪山;朱银华;李伟;刘维佳;李力成;杨祝红;刘畅;姚文俊;陆小华;冯新.含 $TiO_2(B)$ 介孔氧化钛材料的制备、特性和应用[J].催化学报, 2010,31(6): 605-614
- [7] 宋华;于洪坤;武显春;郭云涛. $TiO_2-Al_2O_3$ 载体的制备及Ni2P/ $TiO_2-Al_2O_3$ 催化剂上的同时加氢脱硫和加氢脱氮反应[J].催化学报, 2010,31(4): 447-453
- [8] 郭亚男;曾鹏晖;季生福;魏妮;刘辉;李成岳.Mo助剂含量对Mo-Ni2P/SBA-15/堇青石整体式催化剂加氢脱硫性能的影响[J].催化学报, 2010,31(3): 329-334
- [9] 葛晖;李学宽;王国富;秦张峰;吕占军;王建国.硫代硫酸铵对 CoMo 和 NiMoP 催化剂的预硫化[J].催化学报, 2010,31(1): 18-20
- [10] 葛晖;李学宽;樊卫斌;秦张峰;吕占军;王建国.氮气热处理对硫代硫酸铵预硫化的Mo/ Al_2O_3 催化剂的影响[J].催化学报, 2009,30(2): 111-118
- [11] 安莹;陆亮;李才猛;程时富;高国华.磷钼杂多酸离子液体催化氧化脱硫[J].催化学报, 2009,30(12): 1222-1226
- [12] 范闽光;李斌;张飞跃;方金龙;李望良;邢建民;刘自力 .镧改性的CuHY分子筛中铜离子的分布及其对吸附脱硫性能的影响[J].催化学报, 2009,30(10): 1041-1048
- [13] 张健;白秀梅;李翔;王安杰;马学虎 .溶胶-凝胶法制备MoO₃-CeO₂-SiO₂氧化脱硫催化剂[J].催化学报, 2009,30(10): 1017-1021
- [14] 马崇乐;金鑫;丁玲;张秋民;邱介山;梁长海.高比表面积二硫化钼的制备及其对喹啉选择加氢反应的催化性能[J].催化学报, 2009,30(1): 78-82
- [15] 葛晖;李学宽;王建国;秦张峰;吕占军;杨英.硫代硫酸铵预硫化的MoO₃/ Al_2O_3 催化剂的活化和加氢脱硫活性[J].催化学报, 2008,29(9): 921-927