

# Conversion of Isobutane in Presence of Carbon Dioxide over Molybdenum Oxide Catalysts Obtained from Heteropolymolybdate Precursors

Ahmed AOUISSI \*, Daif ALDHAYAN, Saad ALKAHTANI

Chemistry Department, College of Science, King Saud University, Riyadh, Kingdom of Saudi Arabia

Ahmed AOUISSI \*, Daif ALDHAYAN, Saad ALKAHTANI

Chemistry Department, College of Science, King Saud University, Riyadh, Kingdom of Saudi Arabia

- 摘要
- 参考文献
- 相关文章

Download: PDF (333KB) [HTML](#) (1KB) Export: BibTeX or EndNote (RIS) Supporting Info

**摘要** Molybdenum based oxide catalysts Mo-H, Mo-Fe, Mo-Ce, and Mo-Sn were prepared by calcining  $H_3PMo_{12}O_{40}$ ,  $Fe_{1.5}PMo_{12}O_{40}$ ,  $Ce_{1.5}PMo_{12}O_{40}$ , and  $Sn_{1.5}PMo_{12}O_{40}$  heteropolyanion precursors at  $700^\circ C$ , respectively. The prepared oxides have been characterized and tested for the dehydrogenation of isobutane (IB) to isobutene in the presence of  $CO_2$ . The effects of temperature, time on stream, and  $CO_2/IB$  ratio were investigated. It was found that  $\alpha$ - and  $\beta$ - $MoO_3$  phases were present in all catalysts. Catalytic tests showed that increasing the reaction temperature increased both the conversion and isobutene selectivity, whereas increasing the  $CO_2/IB$  molar ratio increased the conversion but decreased the selectivity for isobutene. Iron was found to be an effective additive element for the enhancement of catalytic activity compared with Ce and Sn.

**关键词:** [isobutane](#) [isobutene](#) [heteropoly compound](#) [mixed oxide](#) [carbon dioxide](#)

**Abstract:** Molybdenum based oxide catalysts Mo-H, Mo-Fe, Mo-Ce, and Mo-Sn were prepared by calcining  $H_3PMo_{12}O_{40}$ ,  $Fe_{1.5}PMo_{12}O_{40}$ ,  $Ce_{1.5}PMo_{12}O_{40}$ , and  $Sn_{1.5}PMo_{12}O_{40}$  heteropolyanion precursors at  $700^\circ C$ , respectively. The prepared oxides have been characterized and tested for the dehydrogenation of isobutane (IB) to isobutene in the presence of  $CO_2$ . The effects of temperature, time on stream, and  $CO_2/IB$  ratio were investigated. It was found that  $\alpha$ - and  $\beta$ - $MoO_3$  phases were present in all catalysts. Catalytic tests showed that increasing the reaction temperature increased both the conversion and isobutene selectivity, whereas increasing the  $CO_2/IB$  molar ratio increased the conversion but decreased the selectivity for isobutene. Iron was found to be an effective additive element for the enhancement of catalytic activity compared with Ce and Sn.

**Keywords:** [isobutane](#), [isobutene](#), [heteropoly compound](#), [mixed oxide](#), [carbon dioxide](#)

收稿日期: 2012-03-20; 出版日期: 2012-07-19

引用本文:

Ahmed AOUISSI, Daif ALDHAYAN, Saad ALKAHTANI .Conversion of Isobutane in Presence of Carbon Dioxide over Molybdenum Oxide Catalysts Obtained from Heteropolymolybdate Precursors[J] 催化学报, 2012,V33(9): 1474-1479

Ahmed AOUISSI, Daif ALDHAYAN, Saad ALKAHTANI .Conversion of Isobutane in Presence of Carbon Dioxide over Molybdenum Oxide Catalysts Obtained from Heteropolymolybdate Precursors[J] Chinese Journal of Catalysis, 2012,V33(9): 1474-1479

链接本文:

[http://www.chxb.cn/CN/10.1016/S1872-2067\(11\)60412-9](http://www.chxb.cn/CN/10.1016/S1872-2067(11)60412-9) 或 <http://www.chxb.cn/CN/Y2012/V33/I9/1474>

- [1] Matsuda T, Koike I, Kubo N, Kikuchi E. Appl Catal A, 1993, 96: 3
- [2] Al-zahrani S M, Elbashir N O, Abasaeed A E, Abdulwahed M. J Mol Catal A, 2004, 218: 179
- [3] Li L, Yan Z F. Progr Chem, 2005, 17: 651
- [4] Dias C R, Zavoianu R, Portela M F. Catal Commun, 2002, 3: 85
- [5] Bi Y L, Zhen K J, Valenzuela R X, Jia M J, Corberán V C. Catal Today, 2000, 61: 369
- [6] Ogonowski J, Skrzńska E. Catal Lett, 2006, 111: 79
- [7] Sun A, Qin Z, Chen S, Wang J. J Mol Catal A, 2004, 210: 189
- [8] Krylov O V, Mamedov A K, Mirzabekov S R. Catal Today, 1995, 24: 371
- [9] Sun A, Qin Z, Wang J. Appl Catal A, 2002, 234: 179
- [10] Cai Y, Chou L, Li S, Zhang B, Zhao J. Catal Lett, 2003, 86: 191

## Service

- ▶ 把本文推荐给朋友
- ▶ 加入我的书架
- ▶ 加入引用管理器
- ▶ Email Alert
- ▶ RSS

## 作者相关文章

- ▶ Ahmed AOUISSI
- ▶ Daif ALDHAYAN
- ▶ Saad ALKAHTANI

- [11] Wang Y, Ohtsuka Y. *Appl Catal A*, 2001, 219: 183
- [12] Xu B, Zeng B, Hua W, Yue Y, Gao Z. *J Catal*, 2006, 239: 470
- [13] Aouissi A, Apblett A W, AL-Othman Z A, Al-Amro A. *Transi-tion Met Chem*, 2010, 35: 927
- [14] Rocchiccioli-Deltcheff C, Fournier M. *J Chem Soc, Faraday Trans*, 1991, 87: 3913
- [15] Andersson G, Magnéli A. *Acta Chem Scand*, 1950, 4: 793
- [16] Mc Carron III E M. *J Chem Soc, Chem Commun*, 1986: 336
- [17] Huang S, Liu S, Zhu Q, Zhu X, Xin W, Liu H, Feng Z, Li C, Xie S, Wang Q, Xu L. *Appl Catal A*, 2007, 323: 94
- [18] Massarotti V, Flor G, Marini A. *J Appl Cryst*, 1981, 14: 64
- [19] Carriazo J G, Molina R, Moreno S. *Appl Catal A*, 2008, 334: 168
- [20] Rashad M M. *Mater Sci Eng B*, 2006, 127: 123
- [21] Biernacki L, Pokrzywnicki S J. *J Therm Anal Calorim*, 1999, 55: 227
- [22] Ogonowski J, Skrzyńska E. *Catal Commun*, 2009, 11: 132
- [23] Ding J F, Qin Z F, Li X K, Wang G F, Wang J G. *J Mol Catal A*, 2010, 315: 221
- [24] Shimada H, Akazawa T, Ikenaga N, Suzuki T. *Appl Catal A*, 1998, 168: 243
- [25] Sun A L, Qin Z F, Chen S W, Wang J G. *J Mol Catal A*, 2004, 210: 189
- [26] Chen S, Qin Z, Xu X, Wang J G. *Appl Catal A*, 2006, 302: 185
- [27] Saito M, Kimura H, Mimura N, Wu J, Murata K. *Appl Catal A*, 2003, 239: 71
- [28] Marques S P D, Pinheiro A L, Braga T P, Valentini A, Filho J M, Oliveira A C. *J Mol Catal A*, 2011, 348: 1
- [1] Kimfung LI, David MARTIN, Junwang TANG. Conversion of Solar Energy to Fuels by Inorganic Heterogeneous Systems[J]. *催化学报*, 2011,32(6): 879-890
- [2] K. KOCI, K. ZATLOUKALOVA, L. OBALOVA, S. KREJCIKOVA, Z. LACNY, L. CAPEK, A. HOSPODKOVA, O. SOLCOVA. Wavelength Effect on Photocatalytic Reduction of CO<sub>2</sub> by Ag/TiO<sub>2</sub> Catalyst[J]. *催化学报*, 2011,32(5): 812-815