

## The Chinese Barley Malt Filterability-Related Proteins Explored by the Proteomic Strategy

The filterability defect of domestic barley malt is a long term disturbing problem in current Chinese brewing industry. In contrast, beer manufacture using the Canadian and Australian barley malts with superior filterability seldom met this problem. The direct factor of filterability was historically considered to be certain incomplete hydrolyzed macromolecules, such as  $\beta$ -glucan, arabinoxylan (AX) and prolamine, which was attributed to complex reasons. To systematically characterize filterability-related proteins in barley malt, the Canadian Metcalfe barley malt with superior filterability, and the Chinese Dan'er barley malt encountering filterability problem, were chosen for comparative proteomic using 2D-DIGE. The most different proteins, β-amylase and AXAH-I have been verified to be positively related with filterability, while POD (peroxidase) has a negative effect on filterability.

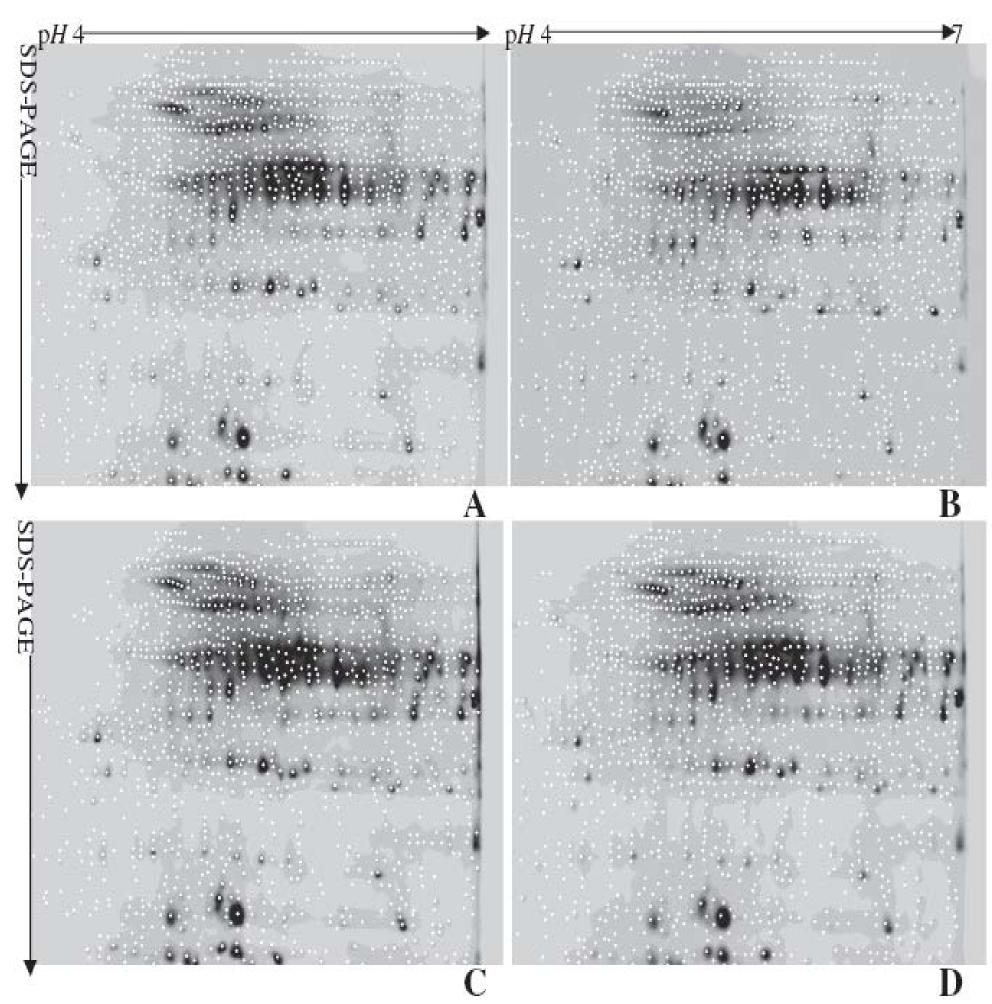


Figure 1. Representative DIGE maps of low-salt soluble proteins (labeled with Cy3) from malts in a pI range of 4~7 and a molecular mass range of 10~60 kDa. A, Metcalfe malt I; B, Metcalfe malt II; C, Dan'er malt I; D, Dan'er malt II.

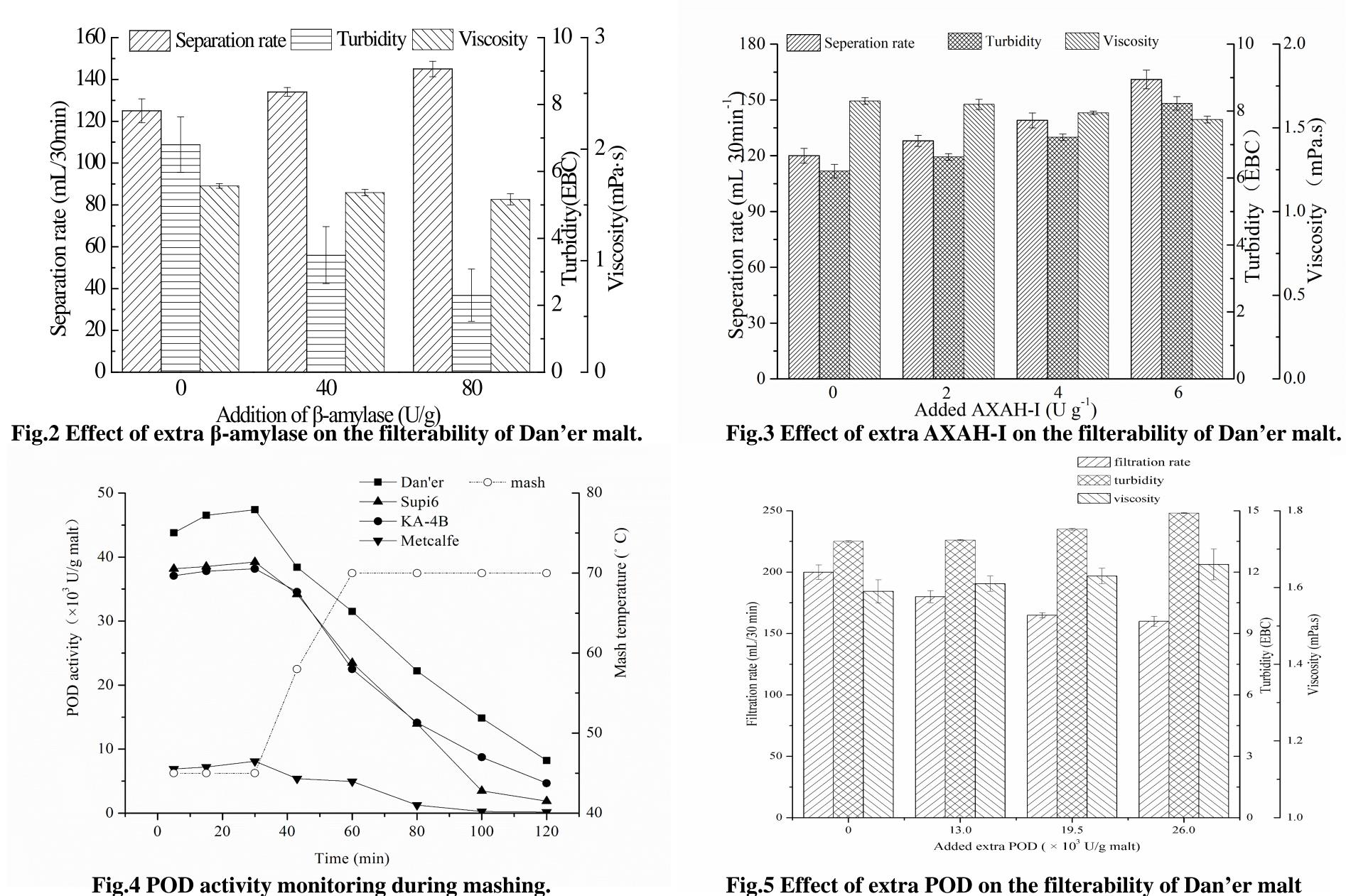
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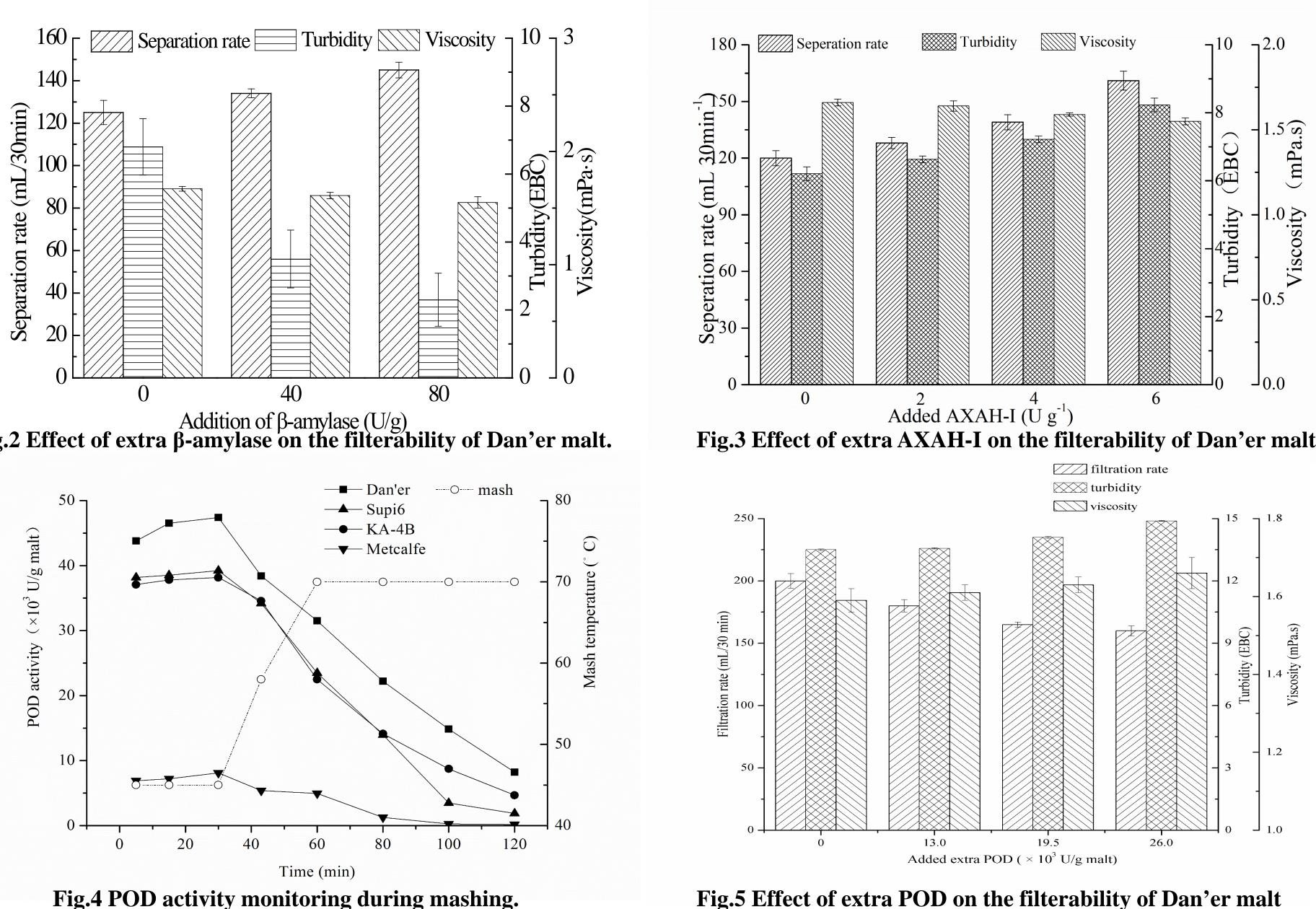
X. M. Li, J. Y. Sun, T. Zhou, Z. Jin, F. Gao, G. L. Cai, and J. Lu

National Engineering Laboratory for Cereal Fermentation Technology, School of Biotechnology, Jiangnan University, Wuxi, Jiangsu, China

Table 1 Major quantitative differential proteins between	
Dan'er and Metcalfe malts	

Protein name Spot Av. ratio profiles (T1, T2)/(C1, C2)no. 94 5.55 β-amylase AXAX-I 1.88 68 **C1** malts 451 LTP1 -2.51 C1 C2 •<u>T1</u> <u>T2</u> • malts -4.28 115 Peroxidase BP1 •C1 **C2** malts 262 Serpin Z7 -7.68 **C2 C1** <sup>™</sup> .0.4 **T1 T2** malts 353 Serpin Z4 -5.36 **T1** malts 438 -2.20 α-amylase/trypsin **₽**C1 inhibitor CMb 





## **2015 ASBC Annual Meeting** June 14–17, 2015 La Quinta Resort and Club La Quinta, CA