

附图 2 OsWRKY42 干扰植株中病程相关蛋白质的表达特征

以接种 3 天、6 天的叶片(病斑线±1cm)为材料,通过 SDS-PAGE 或者 Tricine-SDS-PAGE 分离总蛋白质,一抗为抗病程相关蛋白质抗体,分别为抗 OsPR1a、抗 OsPR1b 及抗 OsPR10a 多克隆抗体(Wu et al., 2011);抗 OsPR2、抗 OsPR5、抗 OsPR6、抗 OsPR15 及抗 OsPR16 多克隆抗体(Hou et al., 2012)和抗 OsHSP82 单克隆抗体(Li et al., 2011)。抗 OsGST 多克隆抗体(Bai et al., 2012)。其它抗体购自北京华大蛋白质研发中心有限公司。 TP309: 对照植株;4021:带有 Xa21 的转基因 TP309 植株;R01、R02 和 R03 是 OsWRKY42-RNAi 转基因株系。红框内为正文(图 6)展示的部分。

Appendix figure 2 Expression profiling of pathogenesis-related proteins in inoculated leaves

Equal amounts of total protein, isolated from inoculated leaves (±1cm within lesion line) of 3-day or 6-day, were resolved by SDS-PAGE or Tricine-SDS-PAGE. The antibodies were anti-*OsPR1a*, *OsPR1b* and *OsPR10a* polyclonal antibodies (Wu et al., 2011); anti-*OsPR2*, *OsPR6*, *OsPR15* and *OsPR16* polyclonal antibodies (Hou et al., 2012) and anti-*OsHSP82* monoclonal antibody (Li et al., 2011); anti-*OsGST* polyclonal antibody (Bai et al., 2012). Other antibodies used in this study were purchased from Beijing Protein innovation Co., Ltd. TP309: Wild type plant; 4021: transgenic TP309 plant carrying *Xa21*; R01, R02, and R03 are OsWRKY*42-RNAi* transgenic lines. The red frames indicate the portion demonstrated in figure 6.