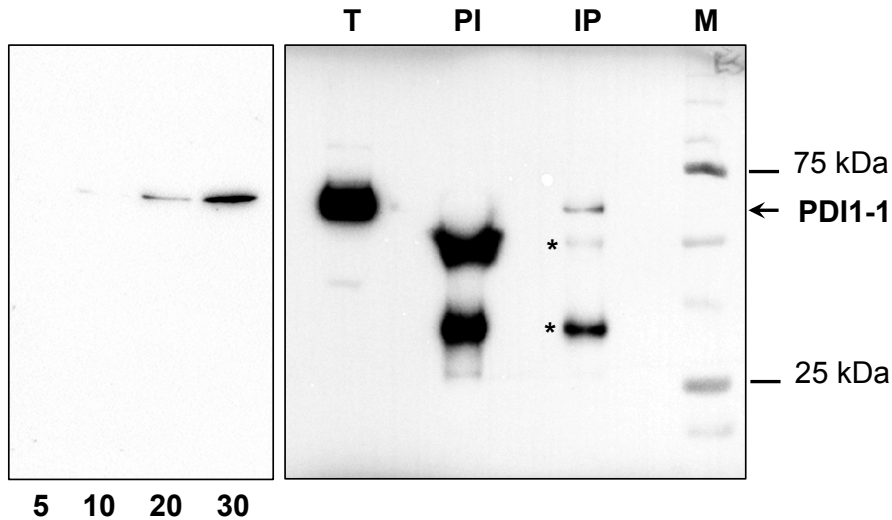
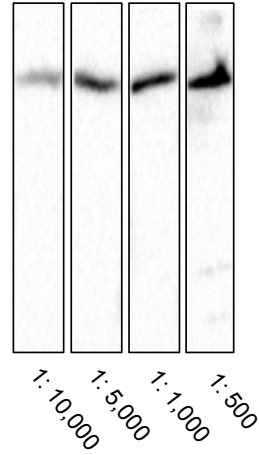


Barley PDI1-1 protein sequence

MAISKVWISLLLLALAVVLSAPAARAEEAAAAEE
 AAAPAVLTLHADNFDDAIGQHPFILVEFYAPW
 CGHCKSLAPEYEKAAQLLSKHDPAILVAKVDAN
 DEKNKPLAGKYEVQGFPTLKI FRNGGKSIQEYK
 GPRAEAGIVEYLLKQVGPASKEIKAPEDATYLE
 DGKIHIVGVFTEFSGPEFTNFLEVAEKLSYYD
 FGHTVHANHLPRGDAVERPVVRLFKPFDELVV
 DSKDFDVSALEKFDASSTPKVVI FDKNPDNHP
 YLLKFFQSNAPKAMFLNFSTGPFESFKSAYYG
 AVEEFSGKDVKFLIGDISSQGAFAQYFGLKVDQ
 APLILIQDGSKKFLKEHVEAGQIVAWLKDYFD
 GKLTPEFKSEPI PEANNEPVKVVVADNVHDVVF
 KSGKNVLIIEFYAPWCGHCKKLAPILDEAAATLQ
 SEEDVVIKMDATENDVPGEFDVQGYPTLYFVT
 PSGKKVSYEGGRTADEIVDYIRKNKETAGQAAA
 ATEKAAEPAATEPLKDEL



S2 Fig. Description and validation of the PDI1-1 antiserum. The carboxi-terminus peptide shown in red was used to generate polyclonal antiserum against PDI1-1 in rabbit (GenScript, USA Inc.). The antiserum was purified against the peptide as recommended by the provider. Different dilutions of the purified antiserum were used in western blot (right upper corner) with total protein extracts from barley seeds (30 µg) and a single band was detected at the expected molecular weight for PDI1-1 (55 kDa). Different amounts (µg) of total protein were tested for the 1:5000 dilution (lower left corner). Immunoprecipitation (IP) with the antiserum showed the same PDI1-1 band, detected in total extracts (T), whereas the pre-immune serum (PI) did not detect the band. High and low molecular weight IgGs were detected in both immunoprecipitate reactions (*). In addition, the band corresponding to PDI1-1 in the IP was identified by mass spectrometry.