Revealing new phosphoarginine binding proteins using chemoproteomic methods

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Arginine phosphorylation, an underexplored post-translational modification, is crucial for many cellular functions, including protein degradation.¹ While protein arginine kinase² and pArg-specific phosphatase³ have recently been reported, phosphoarginine (pArg)-binding proteins remain elusive. Finding new pArg-binding proteins will be important in understanding the functions of protein arginine phosphorylation. In this study, we developed chemoproteomic methods to capture and isolate pArg-binding proteins from proteomes. Our analysis identified several candidate proteins in *B. subtilis* and *M. smegmatis*, which are highly important in cellular stress responses. These proteins are potentially linking Arg phosphorylation and metabolism.

References

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