Table S1. Variants in the *CARD14* gene in patients with sporadic PRP

SNP rs	Variant			Minor allele frequency
	cDNA	Codon	Amino acid	
rs9895931	c.27C>C/T	TCC>TCT	p.S9S	T: 0.0%
rs114688446	c.599G>G/A	AGC>AAC	p.S200N	T: 0.8%
rs4889990	c.633G>A	GAG>GAA	p.E211E	A: 37.9%
rs28674001	c.676-6G>A*			A: 37.4%
rs142246283	c.683T>T/G	CTA>CGA	p.L228R	no data
rs61751629	c.1264G>A	GAG>AAG	p.E422K	A: 2.1%
rs11658460	c.1323C>T	GAC>GAT	p.D441D	T: 10.2%
rs34367357	c.1753G>A	GTC>ATC	p.V585I	A: 5.2%
rs2066964	c.1641G>C	AGG>AGC	p.R547S	C: 65.1%
rs117918077	c.2044C>C/T	CGG>TGG	p.R682W	T: 1.2%
rs11653893	c.2399-4A>G*			G: 40.4%
no data	c.2406C>C/A	AGC>AGA	p.S802R	no data
rs11652075	c.2458C>T	CGG>TGG	p.R820W	T: 39.9%
rs61757652	c.2481C>T	CCC>CCT	p.P827P	T: 6.2%
rs139789664	c.2495C>T,	CTC>CTT,	p.L832L	T: 0.5%

*Intronic sequences