

**Appendix A: Band 3 Mutations Resulting in Hereditary Spherocytosis**

(adapted from Walensky *et al*, 2003)

<b>Name</b>	<b>Affected Exon</b>	<b>Affected Codon</b>	<b>Designation</b>	<b>Genetic Mutation</b>	<b>Mutation Type</b>	<b>Consequence</b>
<b>Bohain</b>	4	81	241delT	GCTGGG→CGGGG	Frameshift	PTC
<b>Foggia</b>	4	55	163delC	ACCCAC→ACCAC	Frameshift	PTC
<b>Kagoshima</b>	4	56	167delA	AAG→AG	Frameshift	PTC
<b>Neapolis</b>	2 nucleotides after exon 2	-	15+2t→c	T→c	Splicing	EITHER exon 2 skipping (initiation codon abolished) OR Retention of intron 2 (PTC)
<b>Cape Town</b>	5	90	E90K	GAG→AAG	Missense	Replacement of glutamic acid 90 with lysine
<b>Fukayama I</b>	5	112-113	336-337del2	AGAGTC→AGTC	Frameshift	PTC
<b>Hodouin (Prague IV)</b>	5	81	W81X	TGG→TGA	Nonsense	Replacement of tryptophan 81 with PTC
<b>Napoli I</b>	5	100	298-299ins1	TCTCT→TCTTCT	Frameshift	PTC
<b>Fukuoka</b>	6	130	G130R	GGA→AGA	Missense	Replacement of glycine 130 with arginine

<b>Lyon (Osnabrück II)</b>	6	150	R150X	CGA→TGA	Nonsense	Replacement of arginine 150 with PTC
<b>Fukuyama II</b>	7	183	548-549insA	GAT→GAAT	Frameshift	PTC
<b>Wilson</b>	7	170-172	515delG	G <sub>(6)</sub> →G <sub>(5)</sub>	Frameshift	PTC
<b>Worcester</b>	7	170-172	515-516insG	G <sub>(6)</sub> →G <sub>(7)</sub>	Frameshift	PTC
<b>Boston</b>	9	285	A285D	GCT→GAT	Missense	Replacement of alanine 285 with aspartic acid
<b>Campinas</b>	<b>1nucleotide after exon 8</b>	204	694+1G>T	G→T	Splicing Frameshift	Eradication of splice site with the resultant skipping of exon 8 and PTC 13 amino acids downstream of the exon 7-exon 9 splice site as a result of a frameshift
<b>Princeton</b>	9	273-275	823-824insC	C <sub>(6)</sub> →C <sub>(7)</sub>	Frameshift	PTC
<b>Noirterre</b>	10	330	Q330X	CAG→TAG	Nonsense	Replacement of glutamine 330 with PTC

<b>Tuscaloosa</b>	10	327	P327R	CCC→CGC	Missense	Replacement of proline 327 with arginine
<b>Bruggen</b>	11	418-419	1257delC	CACCCG→CACCG	Frameshift	PTC
<b>Nachod (Hradec Kralove II)</b>	Exon 5 to exon 6 splice site	117-121	350-3c→a	Replacement of a cytosine with an adenine in the acceptor splice site in intron 5 (three nucleotides downstream of position 350)	Splicing	Deletion of amino acids 117 to 121 as result of modified splicing
<b>Yamagata</b>	12	455	G455R	GGG→AGG	Missense	Replacement of glycine 455 with arginine
<b>Benesov (Prague V)</b>	12	455	G455E	GGG→GAG	Missense	Replacement of glycine 455 with glutamic acid
<b>Bicetre II</b>	12	456	1366delG 1475	CTGGGGGCT→CT GGGGCT	Frameshift	PTC at codon 1527 or 1565

<b>Pribram (Prague VI)</b>	1 nucleotide after Intron 12	-	1431+1g→ a	g→a	Splicing	Intron 12 is retained after codon 477, resulting in seven abnormal amino acids and a premature stop codon at triplet 8 of intron 12
<b>Coimbra</b>	13	488	V488M	<b>GTG→ATG</b>	Missense	Replacement of valine 488 with methionine
<b>Bicetre I</b>	13	490	R490C	<b>CGC→TGC</b>	Missense	Replacement of arginine 490 with cysteine
<b>Pinhal (Bloemfontein)</b>	13	490	R490H	<b>CGC→CAC</b>	Missense	Replacement of arginine 490 with histidine
<b>Dresden</b>	13	518	R518C	<b>CGC→TGC</b>	Missense	Replacement of arginine 518 with cysteine
<b>Evry</b>	13	496	1486delT	<b>TTCTGG→TTCGG</b>	Frameshift	PTC at position 1733
<b>Milano</b>	13	499	1497- 1498ins69	Duplication of nucleotides 1432- 1500	Insertion	Duplication of amino acids 478 to 499 and insertion of a glutamine
<b>Smichov (Prague VII)</b>	15	616	1848delC	<b>ATC→AT</b>	Frameshift	PTC

<b>Trutnov</b>	15	628	Y628X	TAC→TAA	Nonsense	Replacement of a tyrosine with PTC
<b>Hobart</b>	16	646-647	1940delG	CGGGGC→CGGGC	Frameshift	PTC
<b>Osnabruck II</b>	16	663-664	DelM664	ATGATG→ATG	Deletion	Deletion of methionine 664
<b>Tochigi II</b>	17 -5 nucleotides	-	2058-5delA	delA	Splicing	PTC
<b>Most (Prague VIII)</b>	17	707	L707P	CTG→CCG	Missense	Replacement of leucine 707 with a proline
<b>Okinawa</b>	17	714	G714R	GGG→AAG	Missense	Replacement of glycine 714 with arginine
<b>Hradec Kralove (Tochigi I)</b>	17	760	R760W	CGG→TGG	Missense	Replacement of arginine 760 with tryptophan
<b>Kumamoto (Prague II)</b>	17	760	R760Q	CGG→CAG	Missense	Replacement of arginine 760 with glutamine
<b>Chur</b>	18	771	G771D	GGC→GAC	Missense	Replacement of glycine 771 with aspartic acid
<b>Napoli II</b>	18	783	I783N	ATC→AAC	Missense	Replacement of isoleucine 783 with asparagine

<b>Jablonec</b>	18	808	R808C	CGC→TGC	Missense	Replacement of arginine 808 with cysteine
<b>Nara</b>	18	808	R808H	CGC→CAC	Missense	Replacement of arginine 808 with histidine
<b>Prague I</b>	18	822	2464-2465ins10	TGTG→TG(10nt)TG	Frameshift	PTC
<b>Birmingham</b>	19	834	H834P	CAC→CCC	Missense	Replacement of histidine 834 with a proline
<b>Nagoya</b>	19	837	T837R	ACG→AAG	Missense	Replacement of threonine 837 with arginine
<b>Philadelphia</b>	19	837	T837M	ACG→ATG	Missense	Replacement of threonine 837 with methionine
<b>Tokyo</b>	19	837	T837A	ACG→GCG	Missense	Replacement of threonine 837 with alanine
<b>Prague III</b>	19	870	R870W	CGG→TGG	Missense	Replacement of arginine 870 with tryptophan
<b>Vesuvio</b>	20	894	2681delC	ACC→AC	Frameshift	Termination at codon 1026; elongated carboxy-terminal

PTC=premature termination codon