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Supplementary Material

A Missense Mutation in *COL10A1* Gene in a Pakistani Consanguineous Family with Schmid Type Metaphyseal Chondrodysplasia

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Supplementary Table I. Clinical features and *COL10A1* mutation in 8 patients with Schmid type of Metaphyseal Dysplasia and in 2 control subjects.

Sr. No.	Position in pedigree	Pheno-type	Gender	Age (Years)	Height (cm)	Weight (Kg)	Mutation	Other abnormalities (If any)
1	IV-13	Dwarf	Female	40	129	41	c.2011A>G	Pain in pelvic region
2	IV-1	Dwarf	Female	40	132	34	c.2011A>G	Restlessness
3	V-17	Dwarf	Male	25	132	59	c.2011A>G	Not applicable
4	IV-6	Dwarf	Female	60	119	60	c.2011A>G	Pelvic and joint pain
5	V-12	Dwarf	Female	35	132	40	c.2011A>G	Asthma, miscarriage
6	V-10	Dwarf	Male	36	132	60	c.2011A>G	Not applicable
7	V-13	Normal	Male	65	157	65	c.2011A	Not applicable
8	V-8	Dwarf	Female	25	121	50	c.2011A>G	Not applicable
9	III-8	Normal	Female	60	142	41	c.2011A	Not applicable
10	IV-16	Dwarf	Male	25	134	46	c.2011A>G	Not applicable

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