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Short Communication

Scenario of Marfan Syndrome Non a Kashmir Family

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MARFAN Syndrome is a kind of connective tissue disorder which involves almost all structures of body like skin ligaments tendons joints bones b veesels valves eyes cvs DX of this syndrome complex is by clinical features blood test mr Ict scan echo etc.

M syndrome has got following manifestations

- OPHTHALMIC manifestations
- SKELTO MUSCULAR manifestations
- CVS manifestations
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Tall thin person tower skull macroglossia high arched palate positive metacarpal index pigeon shaped chest subluxation and dislocation of lens spherophakia ECTOPIA LENTES nystagmus squint keratoconus atrial septal defects p d a mitral valve prolapse syndrome.

Ophthalmic manifestations

Subluxation and dislocation of lens schizophasia ECTOPIA LENTES nystagmus squint cataract high myopia keratoconus retinal detachment thin cornea.

Skeletal muscular manifestations are

TaLl thin person towers skull macroglossia high arched palate arachnodactyly kyphosis SCOLIASIS SPAN of arm is greater than HEIGHT positive metacarpasal index pigeon shaped chest tendency for fractures.

CVS manifestations are

- AORTIC dilatation
- Mitral valve prolapse syndrome
- Severe mitral regurgitation
- Pda
- Atrial septal defects
- Case report
- I saw a family of MARFANS Syndrome in my office.

Father and elder son had typical features of MARFANS Syndrome in fact father had undergone cataract surgery both father and son had no clinical evidence of cvs daughter was normal.

Discussion

We do see cases of MARFANS may be they may not have all features like ophthalmic skelto muscular and c v s manifestations they may be kind of marfiod.

Conclusion

Once we see cases of m syndrome try to EXCLUDE all possible manifestations like OPHTGALMIC nscm and cvs manifestations with the possible presentation do the necessary treatment.

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