Clinical and Genetic Features of Women's Cohort with Turner Syndrome from Lviv Region (West Ukraine) for 1997-2017

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Abstract

Turner syndrome (TS) is a chromosomal disorder caused by complete or partial X chromosome monosomy that manifests various clinical features depending on the karyotype and on the genetic background of affected girls. Aim of our study was to assess the clinical and the cytogenetic characteristics in patient with Turner syndrome from Lviv region (West Ukraine) for 1997-2017.

Methods: 135 female patients aged 1 month-49 years from Lviv region were clinical and cytogenetically

Mosaicism Isochromosome X	with	-	-	-	-	1	0.7	11	8.1	-	-	-	-	12	8.9
45,X/47,XXX mosaicism		1	0.7	-	-	1	0.7	1	0.7	2	1.5	1	0.7	6	4.4
Mosaicism with X ring		-	-	-	-	-	-	3	2.2	2	1.5	-	-	5	3.7
Other		-	-	-	-	1	0.7	4	3.0	2	1.5	-	-	7	5.2
All mosaicism forms		4	3.0	1	0.7	9	6.7	22	16.3	13	9.6	2	1.5	51	37.8
All		17	12.6	3	2.2	20	14.8	56	41.5	27	20	12	8.9	135	100

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3 E C BD 3; I 1 C1 6 D G:1 G B F1 3 C CD = C 31 B F1 C3 E 1 B B C B1 D B I 1 CDB D CD 1 G B 1 D DC G D ; 1 B D C 1 = C

3 D D 3 3 1 B 1 3 D B CD 3 1 2 = G D E B B C B = 1 B 1 B 1 D 6 D C C CD = = D G = G D = C = C = B ; D 1 F 1 2 B = 1 D 6 C; D1 C CD = = C = C 1 D 6 1 D B B 1 C 1

2 DG 1 2

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References

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 2 DG C- B 3 6 1 DE B C ; 1 B D 1 31 B 1 BD 3

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- F BCI 1 1 1 B E 31 D1 D 31

 D 31C 3DC 61ED = E D B C B BC E 1D DC

 G D E B B C B = D1 1DB
- B1 F D BC G1 ; BC ; C PC C1

 31 B13D3 E C6BD 31B 6 BC1 G = GD

 EB B C B = B 3 C6B = D 3 1D DB1D 1

 EB B E EB 3B
- B1 F D = 31 3 B 1 = D12 3 6 1 DE B C
 EB B C B = EB 3 B

- C; 1 F3 ; 2 1 1 1 F1 1 D G

 CDE 6 E B B C B = ; B1 3 DB 3 B C E B3

 2 CDB1 3 DC D E 1 D 6 D E B 1 3 D 6 B 1 DB 3
- E 1 1 D1 31 = 1 6 CDLD 1

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- C = AE1 B E3DF CCE C = G D EB B
- E D 2 E D 13 6 2 F B BE 2 D 1

 BD D 1 B 1 3 E B B B 2 CD D 1 3 1
- 1 BD = D E C BB C 3 3 E2 BD 1 6 BD D 1 1 D D 1 1

 D B1 DE B B F G = D
- B 1 G = 1 1 DD C E B6 B

 B = C = B 1 B1 D C 6 E; 3 D C 3 E DE B 6 B = E = 1
- 1 2 B D 1 2 1 D 3 AE 6 B E = 1 3 B = C = C