<scp>A</scp>shkenazi <scp>J</scp>ewish population s
<scp>T</scp>ayâ€“<scp>S</scp>achs disease: The Inter experience

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#	Article	IF	Citations
1	Preconception risk assessment for thalassaemia, sickle cell disease, cystic fibrosis and Tay-Sachs disease. The Cochrane Library, 2015, , CD010849.	1.5	12
2	Tay-Sachs disease: current perspectives from Australia. The Application of Clinical Genetics, 2015, 8, 19.	1.4	33
3	Points to Consider: Ethical, Legal, and Psychosocial Implications of Genetic Testing in Children and Adolescents. American Journal of Human Genetics, 2015, 97, 6-21.	2.6	453
4	Public Health and Rare Diseases: Oxymoron No More. Preventing Chronic Disease, 2016, 13, E05.	1.7	68
5	Approach to diagnosis ofÂmetabolic diseases. Translational Science of Rare Diseases, 2016, 1, 3-22.	1.6	8
6	Perspective: The rare must become common. Nature, 2016, 537, S151-S151.	13.7	3
7	Preconception and prenatal genetic counselling. Best Practice and Research in Clinical Obstetrics and Gynaecology, 2017, 42, 2-10.	1.4	6
8	Roads to Health in Developing Countries: Understanding the Intersection of Culture and Healing. Current Therapeutic Research, 2017, 86, 13-18.	0.5	29
9	Tay-Sachs Disease. , 2017, , 2725-2732.		0
10	New era in genetics of early-onset muscle disease: Breakthroughs and challenges. Seminars in Cell and Developmental Biology, 2017, 64, 160-170.	2.3	24
11	Preconception risk assessment for thalassaemia, sickle cell disease, cystic fibrosis and Tay-Sachs disease. The Cochrane Library, 2018, 3, CD010849.	1.5	14
12	Emptying the stores: lysosomal diseases and therapeutic strategies. Nature Reviews Drug Discovery, 2018, 17, 133-150.	21.5	191
13	Tay-Sachs disease. Revista Facultad De Medicina, 2019, 67, 323-329.	0.0	1
14	Consanguinity and genetic diseases among the Bedouin population in the Negev. Journal of Community Genetics, 2020, 11, 13-19.	0.5	19
15	Equitable Expanded Carrier Screening Needs Indigenous Clinical and Population Genomic Data. American Journal of Human Genetics, 2020, 107, 175-182.	2.6	24
16	Survey on patientsâ€™ organisationsâ€™ knowledge and position paper on screening for inherited neuromuscular diseases in Europe. Orphanet Journal of Rare Diseases, 2021, 16, 75.	1.2	3
17	The Core Outcome DEvelopment for Carrier Screening (CODECS) study: protocol for development of a core outcome set. Trials, 2021, 22, 480.	0.7	7
18	Global epidemiology of mucopolysaccharidosis type III (Sanfilippo syndrome): an updated systematic review and meta-analysis. Journal of Pediatric Endocrinology and Metabolism, 2021, 34, 1225-1235.	0.4	12

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19	Tay-Sachs Disease; Report of 6 Iranian Patients and Review of Literature. Sarem Journal of Reproductive Medicine, 2017, 2, 35-38.	0.0	1
20	Preconception risk assessment for thalassaemia, sickle cell disease, cystic fibrosis and Tay-Sachs disease. The Cochrane Library, 2021, 2021, CD010849.	1.5	4
21	Tay-Sachs Disease. , 2016, , 1-8.		1
23	Community Genetics screening in a pandemic: solutions for pre-test education, informed consent, and specimen collection. European Journal of Human Genetics, 0, , .	1.4	0