### **Supplementary Materials**

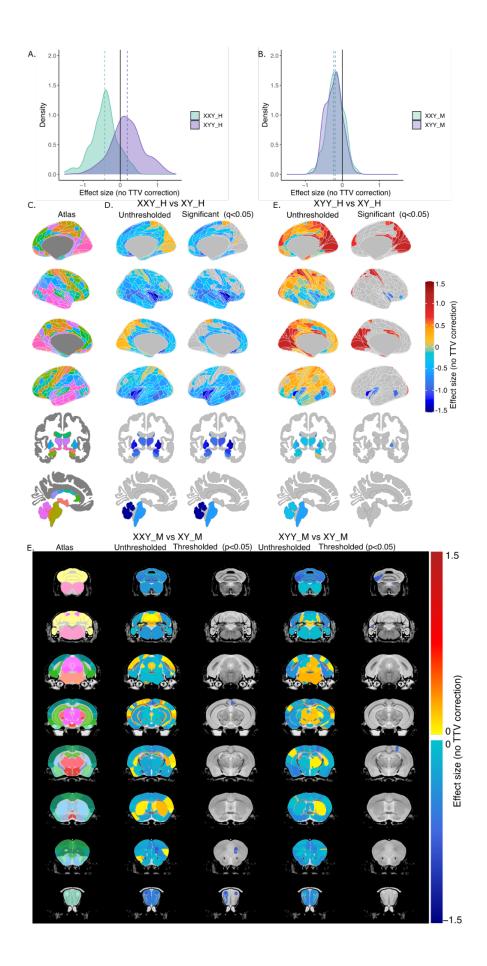
### 1. Results

## 1.1 Regional brain volume changes due to added X- or Y- chromosome in humans and mice without total tissue volume (TTV) correction

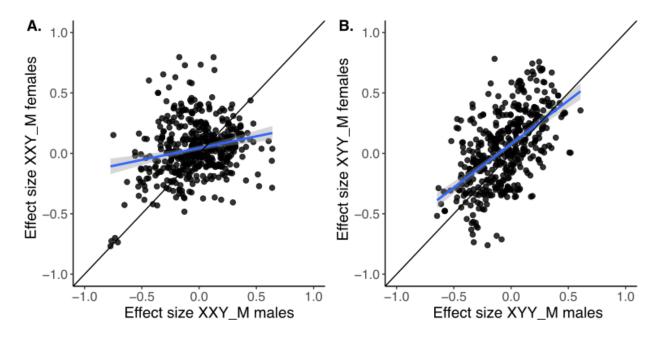
In humans, without TTV correction, the effect of an added X-chromosome on standardized effects sizes across all brain regions were largely negative, (range of  $\beta$ =-1.448 to 0.464), while those for an added Y-chromosome were more evenly distributed, with both positive and negative effects (range of  $\beta$ =-1.239 to 1.564) (**Supplementary Figure 1A**). Indeed, mean standardized effect sizes for XXY\_H ( $\beta$ =-0.422) were significantly smaller than those for the XYY\_H group ( $\beta$ =0.184), which were close to zero (comparison of effect size distributions, t=-20.838, df=714.58, p< 2.2e-16). The range of standardized effect sizes was also significantly different (Levene's test, F(1, 754) =18.145, p=2.306e-05). In contrast, in mice, the range of standardized effect sizes across brain regions was similar for the effect of an added X- ( $\beta$ =-0.848 to 0.385) and added Y-chromosome ( $\beta$ =-0.792 to 0.313) (**Supplementary Figure 1B**). In contrast to the TTV corrected effects, the range and mean standardized effect sizes for XXY\_M ( $\beta$ =-0.186) and XYY\_M ( $\beta$ =-0.228) were shifted towards more negative, and significantly different from zero (t=2.799, df=904.82, p=0.005), but not significantly different from each other (Levene's test: F(1,906) =6e<sup>-4</sup>, p=0.981).

Without TTV correction, large decreases in regional brain volume were observed in XXY\_H driven by global volume reductions identified in **2.1**. These include widespread reductions in insula, orbitofrontal, fusiform face, frontal, piriform, visual, temporal cortex, cingulate, somatosensory association cortices, and in the cerebellum cortex, putamen, pallidum, amygdala, brainstem, thalamus, hippocampus, and caudate. Volume increases were observed in the somatosensory cortex (5m), visual cortex, paracentral lobule, and posterior cingulate cortex (POS2). Interestingly, the spatial patterning and directionality of XYY\_H effects on regional brain anatomy were similar to results with TTV correction applied (**2.2**). Volume of the insula, fusiform face complex, inferior frontal cortex, and right pallidum was decreased, while volume of the visual, parieto-occipital, posterior cingulate, retrosplenial cortex, and somatosensory association cortex was increased. The effects of added X- or Y-chromosomes were not as spatially similar without TTV correction due to a global volume reduction in XXY\_H.

In mice, non-significant (unthresholded) brain wide reductions were observed in both XXY\_M and XYY\_M, with increased volume in the striatum, auditory and visual cortices, and brainstem of XXY\_M, and increased mid- and hind-brain volume in the XYY\_M. At an uncorrected (t=2.042, p<0.05) threshold, we observed volume decreases in the left main olfactory bulb granular layer ( $\beta$ =-0.662) and accessory olfactory bulb glomerular layers ( $\beta$ =-0.790), as well as the right cingulate cortex (area 29c) ( $\beta$ =-0.780), ventral orbital area ( $\beta$ =-0.871), and right primary somatosensory cortex ( $\beta$ =-0.643) for XXY\_M. In XYY\_M, volume decreases were observed in the left piriform-amygdalar area ( $\beta$ =-0.600), right primary somatosensory area (also decreased in XXY\_M;  $\beta$ =-0.726), right cuneate nucleus ( $\beta$ =-0.675), and left paramedian lobule of the cerebellum ( $\beta$ =-0.851). Results for XXY\_M and XYY\_M without TTV were similar to observations made with TTV correction (**Supplementary Figure 1F**).



Supplementary Figure 1. Effect of added X- or Y- chromosomes on regional brain volume in humans and mice, without total tissue volume (TTV) correction. Distribution of standardized effect size for the effect of an euploidy are displayed for humans ( $\mathbf{A}$ ) and for mice ( $\mathbf{B}$ ).  $\mathbf{C}$ . Representative views for the cortical Glasser atlas for lateral and medial slices of the left and right hemispheres, and subcortical FreeSurfer atlas with both axial and sagittal views. Unthresholded (left) and significant (q<0.05; right) standardized effect size for the effect of added X ( $\mathbf{D}$ ) and added Y ( $\mathbf{E}$ ) in the human brain.  $\mathbf{F}$ . Mouse brain coronal slices with the representative atlas on the left, followed by unthresholded, then thresholded (p<0.05) standardized effect size for the effect of added X, followed by unthresholded, then thresholded (p<0.05) effects of added Y.

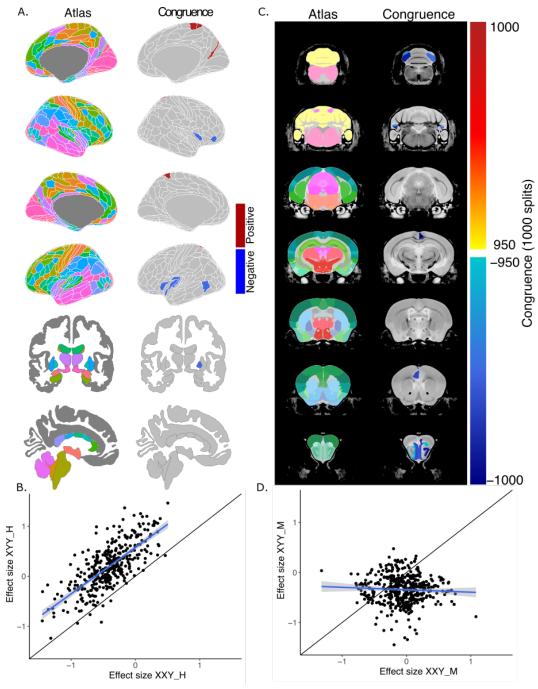


Supplementary Figure 2. Similarity of SCT effects on neuroanatomy across gonadal background. Correlation of standardized effect sizes for the effect of an added ( $\bf A$ ) X chromosome (XXY vs XY) (r=0.22) and added ( $\bf B$ ) Y chromosome (XXY vs XY) (r=0.56) for mice with testicular background or ovarian background.

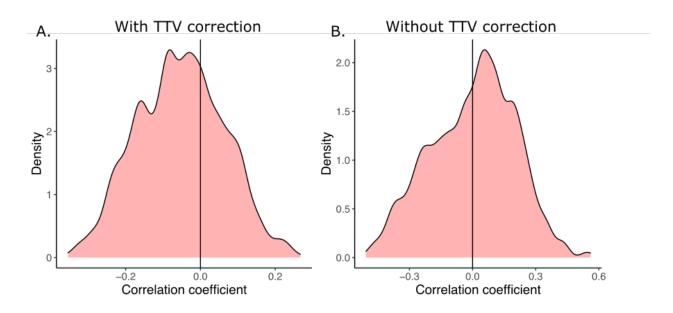
# 1.2 Spatial similarity of added X- or Y-chromosome effects without TTV correction in humans but not mice

A similar pattern of altered brain anatomy was observed in both human SCA groups, although to a lesser degree than in the TTV corrected analysis (2.3). A few subregions within the parietal and visual cortices (n=6 regions) were significantly increased in both XXY\_H and XYY\_H, while the pallidum, insula, temporal, and inferior frontal (inferior frontal gyrus and fusiform face complex) were both decreased due to SCA (n=11 regions) (Supplementary Figure 3A). As with the TTV corrected analysis, the overall similarity of effects was still high (cross-ROI r=0.73) in humans (Supplementary Figure 3B). In the mouse, the bootstrap analysis revealed no brain regions consistently increased (>95% of the time) in both SCA groups (Supplementary Figure 3C). However, several regions were consistently and congruently decreased across both murine

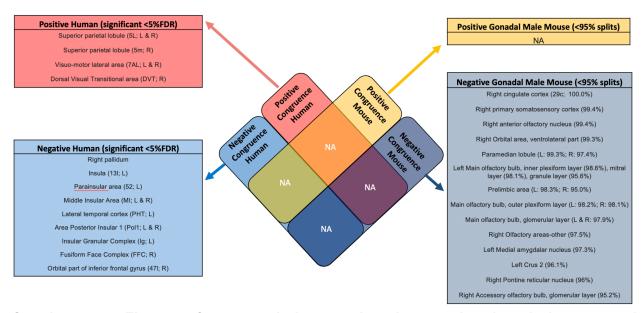
SCT groups, including the right cingulate cortex (29c), right primary somatosensory cortex, ventrolateral orbital areas, paramedian lobule of the cerebellum, and several olfactory bulb regions, which were also consistently and congruently decreased in the TTV corrected analyses. However, there were congruent decreases in the prelimbic area, left medial amygdalar nucleus, left crus 2, and right pontine reticular nucleus of both XXY and XYY mice, which were not observed when correcting for TTV effects. The distribution of cross-ROI correlations was centered close to zero (cross-ROI mean r= 0.024) (**Supplementary Figure 4B**). As with the TTV-corrected analysis, and representative example (split #34) is plotted to (**Supplementary Figure 3D**) illustrate the lack of congruence (r=0.071). Furthermore, there were no regions in humans and mice that showed similar or opposite directionality in effects (**Supplementary Figure 5**).



Supplementary Figure 3. Spatial convergence between added X- or Y- chromosomes on the human and mouse brain without total tissue volume (TTV) correction. A. Human brain regions whose volume was either increased (red) or decreased (blue) in both aneuploidy groups. B. Correlation between standardized effect size for the XXY\_H and XYY\_H group across all brain regions. C. Mouse brain regions whose volume was consistently increased (red colours) or decreased (blue colours) in 95% (950/1000 splits) of the XY\_M control splits in both aneuploidy groups based on bootstrap analysis. D. Correlation between standardized effect size for a representative split (34) for XXY\_M and XYY\_M groups across all brain regions. All plots use the standardized effect sizes for the effect of aneuploidy with total brain volume correction.



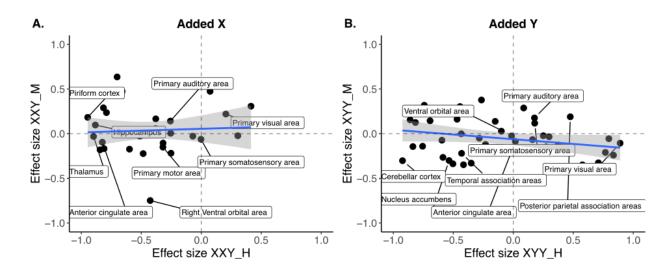
Supplementary Figure 4. Bootstrap analysis shows low spatial correlation for effects of an added X- or Y-chromosome in the mouse brain. The coefficient for the correlation between cross-ROI standardized effect size (β) for XXY\_M (relative to XY\_M sample A) and XYY\_M (relative to XY\_M sample B) both with TTV correction (A) and without (B).



Supplementary Figure 5. Convergently increased or decreased regions in humans and mice without total tissue volume (TTV) correction. Venn diagram of overlapping ROIs highlighting the regions congruently (for both XXY and XYY) increased or decreased in humans and mice. Brain regions that were statistically significantly impacted by both XXY and XYY in humans alone (<5%FDR) are listed in the red (increased volume) and light blue (decreased volume) boxes on the left. Brain regions that showed significantly convergent impacts of XXY and XYY (based on bootstrap resampling) in mice alone are listed in the yellow (increased volume) and dark blue (decreased volume) boxes. There were no regions showing convergence (increased or decreased), or opposite (increased in one species and decreased in another, or vice versa) highlighted by the "NA" in the intersecting cells.

#### 1.3 Human-mouse comparison

As explained in **2.4**, the standardized effect sizes for adding an X- or Y- chromosome were compared in the regions identified as human-mouse homologs without correction for TTV. There was low similarity between the effects on the human and mouse brain based on low correlation of standardized effect sizes across regions (added X: r=-0.08; added Y: r=-0.27; **Supplementary Figure 6**). For both humans and mice, an added X-chromosome decreased volume of the right amygdala, bilateral anterior cingulate area, bilateral cerebellum, left nucleus accumbens, right piriform cortex, bilateral primary motor, somatosensory, and retrosplenial cortices, left thalamus, and bilateral ventral orbital area. The left posterior parietal association area, and bilateral primary visual areas were both increased in volume. An added Y-chromosome decreased volume of the left angular insula cortex, bilateral amygdala, bilateral cerebellar cortex, left globus pallidus, left hippocampus, left nucleus accumbens, bilateral piriform cortex, and left temporal association areas, and increased volume in the right hippocampus, right perirhinal cortex, left posterior parietal association area, left primary somatosensory cortex, and left thalamus (**Supplementary Figure 6 & Supplementary Table 1**).



Supplementary Figure 6. Correlation of effects of added X- or Y- chromosomes in human and mouse homologous brain regions when total tissue volume (TTV) is not controlled for. A. Standardized effect size correlation for the effect of added X-chromosome (cross-ROI r=-0.07) or B. added Y-chromosome (cross-ROI r=-0.27). Points are labeled only if they have the same directionality of standardized effect size in humans and mice (i.e., both positive or both negative). For simplicity, labels are provided only for left hemisphere regions given that effects were similar across hemispheres.

Supplementary Table 1. Mapping of homologous human-mouse brain regions (standardized effect sizes with TTV correction). Blue shade highlights cells with negative standardized effect sizes, while peach highlights positive ones. Human = \_Y; mouse = \_M. Bold and \* denotes regions that are statistically significantly different relative to controls.

Label	Glasser/Freesurfer# names	Mouse atlas	Hemisphere	Human (XXY_H)	Mouse (XXY_M)	Human (XYY_H)	Mouse (XYY_M)
Agranular insula	AVI, AAIC, MI	Agranular insular area	L	-1.412 *	0.216	-0.516	-0.141
			R	-1.503 *	0.299	-0.542	0.125
Amygdala	Amygdala#	Cortical subplate	L	-1.023 *	0.140	-0.168	-0.330
			R	-1.147 *	-0.012	-0.396	-0.073
Anterior cingulate area	A24pr, a24, p24pr, p24, 24dd, 24dv, p32pr, d32, a32pr, p32, s32	Anterior cingulate area	L	-0.825 *	-0.096	0.302	-0.087
			R	-0.810 *	-0.168	0.459	-0.067
Caudoputamen	Caudate#, Putamen#	Caudoputamen	L	-1.185 *	0.241	-0.514	0.144
			R	-1.369 *	0.375	-0.504	0.273
Cerebellar cortex	Cerebellar cortex#	Cerebellar cortex	L	-1.353 *	-0.079	-0.666	-0.302
			R	-1.444 *	-0.223	-0.307	-0.337
Entorhinal cortex	EC	Entorhinal area	L	-0.255	0.005	0.780 *	-0.349
			R	-0.318 *	-0.105	0.690	-0.371
Globus pallidus	Globus Pallidus#	Pallidum	L	-1.288 *	0.000	-0.640	-0.144
			R	-1.270 *	0.116	-0.735 *	0.155
Hippocampus	Hippocampus#	Hippocampal region	L	-0.883 *	0.097	-0.252	-0.001
			R	-0.790 *	0.236	0.052	0.138
Nucleus accumbens	Nucleus accumbens#	Striatum ventral region	L	-0.512 *	-0.028	-0.376	-0.301
			R	-0.519 *	0.043	-0.037	0.023
Perirhinal area	PeEc, TF, PHA2, PHA3	Perirhinal area	L	-0.701 *	0.635	0.449	-0.024
			R	-0.660 *	0.474	0.270	0.288
Piriform cortex	Pir	Piriform cortex	L	-0.947 *	0.183	-0.247	-0.348

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			R	-0.844 *	-0.180	-0.461	-0.266
Posterior parietal association areas  Primary auditory	5m, 5mv, 5L A1	Posterior parietal association areas  Primary auditory	L	0.414 *	0.308	0.565 *	0.189
			R	0.304 *	-0.024	0.863 *	-0.210
			L	-0.257	0.141	0.342	0.176
area Primary motor	4	area Primary motor	R	-0.331 *	0.041	-0.187	0.377
			L	-0.320 *	-0.151	0.115	-0.026
Primary somatosensory area	1, 2, 3a, 3b V1	Primary somatosensory area  Primary visual	R	-0.598 *	-0.173	0.491	-0.035
			L	-0.002	-0.063	0.292	0.117
			R	-0.070	-0.028	0.897 *	-0.056
			L	0.205	0.220	1.046 *	-0.106
area Retrosplenial area	RSC	area Retrosplenial area	R	0.073	0.474	0.995 *	-0.241
			L	-0.455 *	-0.003	0.503	-0.148
Subiculum	PreS	Subiculum	R	-0.252	-0.219	0.880 *	-0.326
			L	-0.379 *	0.056	0.480	-0.128
Temporal	FFC, PIT, TE1a, TE1p, TE2a, TF,	Temporal	R	-0.379 *	0.166	-0.218	-0.051
			L	-1.086 *	0.246	-0.163	-0.220
association areas	STV, STSvp, STSva	association areas	R	-0.815 *	0.289	-0.177	0.304
Thalamus	Thalamus#	Thalamus	L	-0.898 *	-0.031	-0.275	0.162
inalamus	maiamon	maiamas	R	-1.045 *	0.023	-0.244	0.259
Ventral orbital area	10r, 10v	Ventral orbital area	L	-0.484 *	-0.223	0.114	0.028
			R	-0.426 *	-0.749	0.581	-0.416
Brain stem (midline)	Brainstem#	Midbrain, Hindbrain	М	-0.620 *	0.040	-0.585	0.318
Total tissue volume (midline)	BrainSegVolNotVent#			-0.944 *	0.000	0.423	1.44282 E-16