

Girls with Turner syndrome have normal muscle force but decreased muscle power



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None of the authors have any conflict of interest.

Introduction

Turner syndrome (TS) is associated with **decreased bone mineral density** and **altered bone geometry**, which is assumed a risk factor leading to **increased fracture rate**. Although hypogonadism or SHOX gene haploinsufficiency are the probable causes, the exact mechanism remains unclear. Particularly, the **muscle function as an important determinant of bone strength** has yet **not been focused on** in TS.

We tested the hypothesis that there is muscle dysfunction in TS.

Secondary aim was to describe the **influence of pubertal stage, hormone therapy, fracture history and genotype**.

Table 1. Anthropometry characteristics of TS patients.

N=60	mean (SD)
Age (year)	13.7 (4.6)
Weight (kg)	43.2 (16.3)
Weight (Z-score)	-0.63 (1.2)***
Height (cm)	142.3 (17.3)
Height (Z-score)	-1.8 (0.93)***
BMI (kg/m ²)	20.3 (4.1)
BMI (Z-score)	0.44 (0.98)**

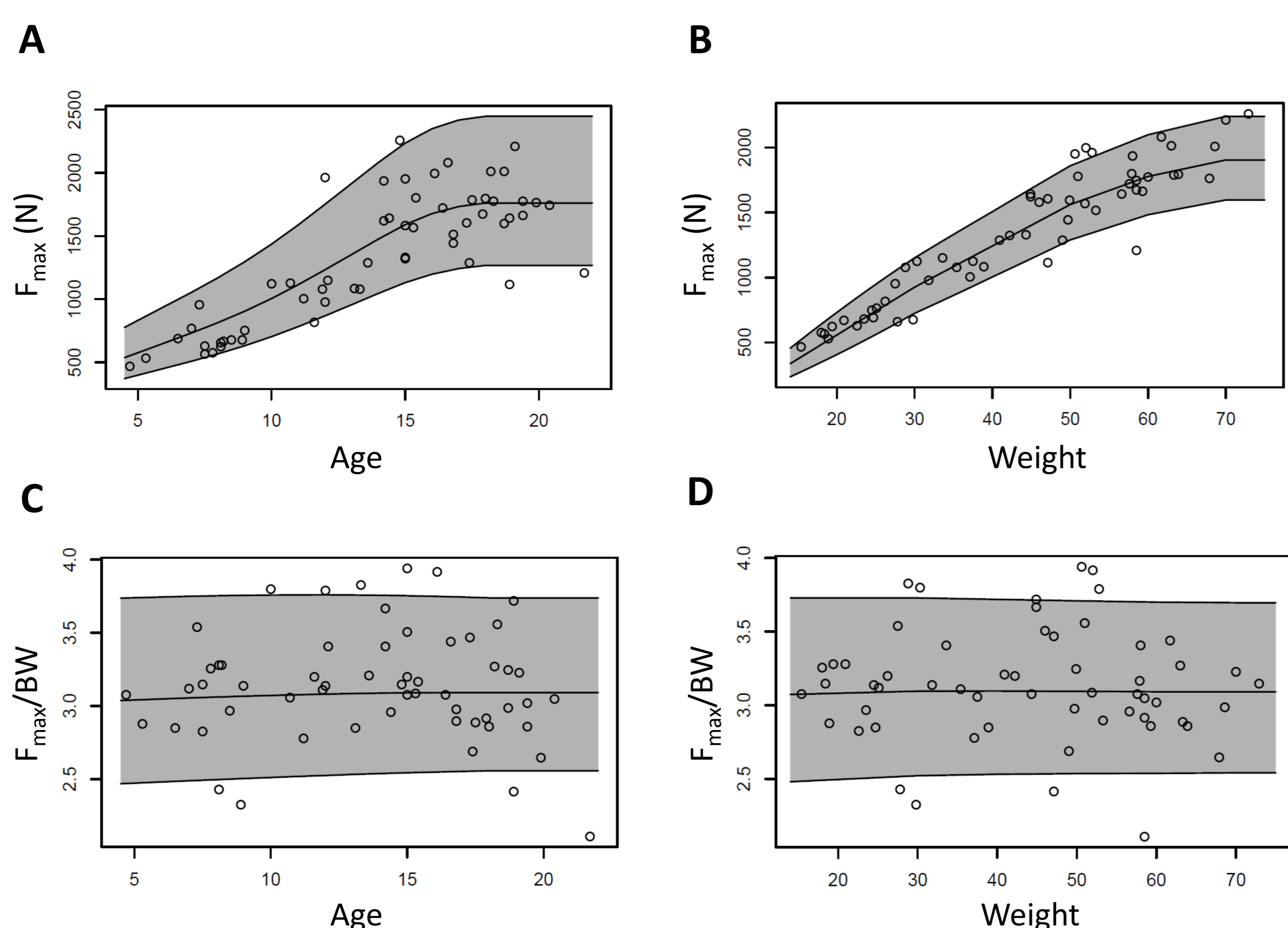
The Z-scores were tested for difference from 0 by one-sample T-test. *p<0.05 **p<0.01 ***p<0.001

Patients and Methods

All TS patients consenting to the study and having no other chronic disease were included (**60 patients, age 13.7±4.5 years**). Age- and weight-specific **z-scores** of muscle parameters were calculated **based on control group of 432 healthy girls**.

Leonardo Mechanograph® Ground Reaction Force Platform was used to assess **muscle force (F_{max})** by the multiple one-legged hopping test and **muscle power (P_{max})** by the single two-legged jump test. Muscle functions were related to body weight (F_{max}/BW) and body mass (P_{max}/mass), respectively.

Figure 1. F_{max} (A+B) and F_{max}/BW (C+D) of TS patients plotted into age- and weight-specific nomograms, respectively.

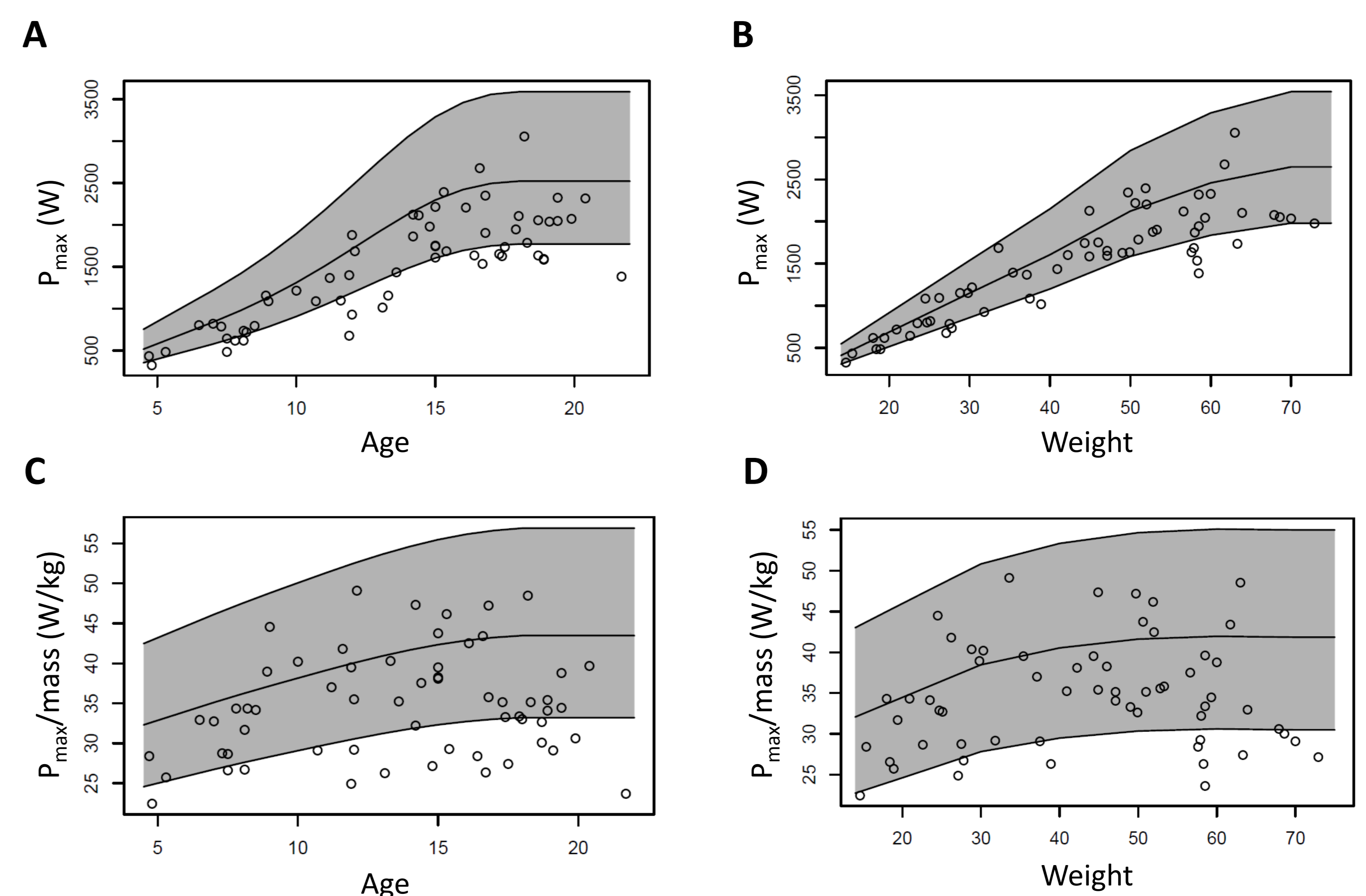


F_{max} was assessed through multiple one legged hopping. Lines are mean ± 2SD, based on 432 healthy girls.

Results

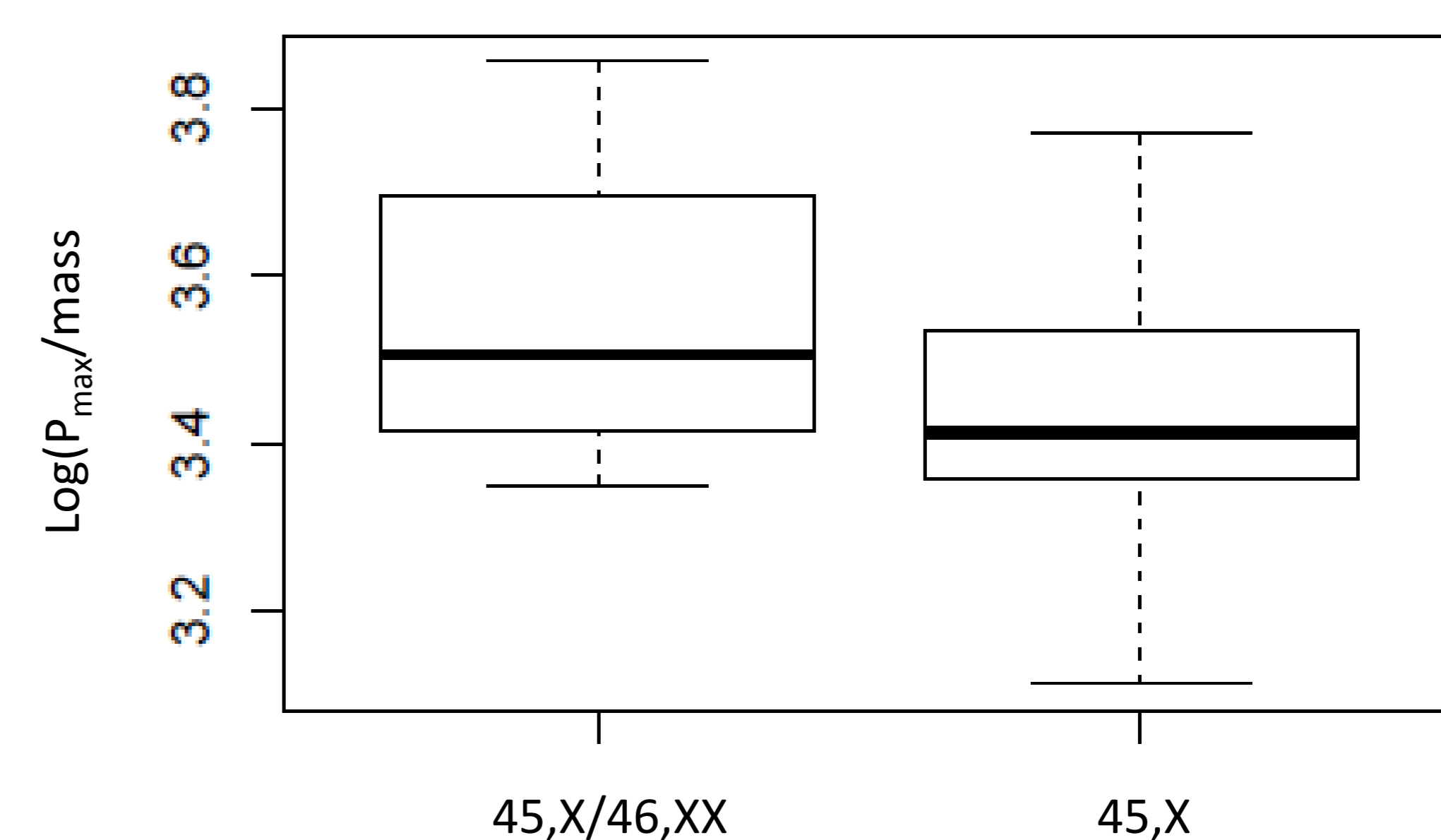
While F_{max} and F_{max}/BW were **normal** (mean weight-specific Z-scores 0.11±0.77, p=0.27, and 0.046±0.62, p=0.55; Fig. 1), P_{max} and P_{max}/mass were **decreased in TS** (Z-scores -0.93±1.5, p<0.001, and -0.45±0.58, p<0.001; Fig. 2), as compared to healthy controls. The **muscle functions were not significantly influenced by pubertal stage, hormone therapy, fracture history nor genotype** (linear regression, adjusted for age, weight and height, all p>0.05).

Figure 2. P_{max} (A+B) and P_{max}/mass (C+D) of TS patients plotted into age- and weight-specific nomograms, respectively.



P_{max} was assessed through single two legged jump. Lines are mean ± 2SD, based on 432 healthy girls.

Figure 3. The influence of karyotype on P_{max}/mass in TS girls.



The influence of karyotype (mosaic vs. X monosomy) was tested in linear regression model with adjustment for age, age², weight, weight² and height ($\beta = -0.108 \pm 0.085$, P = 0.22). Boxes represent median and interquartile range, lines are maximum and minimum.

Conclusions

F_{max} as a principal determinant of bone strength is normal in TS. The changes in bone quality and structure in TS are therefore not related to inadequate mechanical loading, but rather **represent a primary bone deficit**. Decreased P_{max} may represent a novel indicator of impaired muscle coordination in TS.