

SURGICAL PATHOLOGY REPORT

No. 07-154 Hosp.: 大阪市立総合医療センター DATE: August 12/2007

NAME: [REDACTED] AGE: 6y SEX: female

CLINICAL DIAGNOSIS: ^{筋萎縮}筋ジストロフィー

MICROSCOPIC FINDINGS (SPECIMEN:):

Serial frozen sections were stained with hematoxylin & eosin (H&E), modified Gomori trichrome (mGT) and a battery of histochemical methods.

On H&E, there is a marked variation in fiber size, measuring from a few from 80 micrometer. Necrotic and regenerating fibers are scattered. Moderate endomysial fibrosis is seen. There is no cellular infiltration. A few small angular fibers are observed. Muscle fibers with internal nuclei comprise around 1%.

On mGT, there are no nemaline bodies, and rimmed vacuoles, or ragged-red fibers. Peripheral nerve bundles are well myelinated.

On oxidative enzyme stains (NADH-TR and SDH), intermyofibrillar network is well organized except in necrotic and regenerating fibers. No strongly SDH-reactive blood vessel (SSV) is highlighted.

On ATPase, type 1, 2A, 2B, and 2C comprise 43%, 13%, 16%, and 28%, respectively. There is no fiber type grouping. Type 2 fiber atrophy is seen.

Other stains, including PAS, acid phosphatase, alkaline phosphatase, AChE, NSE, AMP, FJK, Oil red O, Cyt.o.c.o. and MAG, show no additional abnormalities.

PATHOLOGICAL DIAGNOSIS:

Myopathic change, with:

- 1) variation in fiber size, marked
- 2) necrotic and regenerating process, active
- 3) endomysial fibrosis, marked
- 4) type 2C fibers, 28%
- 5) type 2 fiber atrophy

PATHOLOGICAL DIAGNOSIS

The above findings are suggestive of muscular dystrophy. Based upon clinical information, FSHD should be ruled out in addition to LGMD.

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