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web oct 25 2019 das zunehmende verständnis molekularer mechanismen der duchenne muskeldystrophie die durch einen
defekt im dystrophingen bedingt ist und anderer muskeldystrophien führt dazu dass eine vielzahl neuer therapieoptionen
eingang in klinische studien und teilweise bereits in die klinische versorgung gefunden hat

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selbstverständnis und ihre haltungen gegenüber ihrer klientel das dritte kapitel zeigt ausgewählte aspekte des fachwissens
hinsichtlich der institutionen und lebensthemen im kontext des lebenslaufs auf

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web die muskeldystrophie des typs duchenne auch duchenne muskeldystrophie und oder dmd genannt ist die häufigste
muskuläre erbkrankheit im kindesalter sie tritt etwa in einer frequenz von 1 3600 bis 1 6000 auf 1 2 aufgrund des x
chromosomal rezessiven erbanges sind fast nur jungen betroffen