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Bereiche

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Funktion

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Publikationen (58)

Wendebourg M, Weigel M, Weidensteiner C, Sander L, Kesenheimer E, Naumann N, Haas T, Madoerin P, Braun N, Neuwirth C, Weber M, Jahn K, Kappos L, Granziera C, Schweikert K, Sinnreich M, Bieri O, Schlaeger R. Cervical and thoracic spinal cord gray matter atrophy is associated with disability in patients with amyotrophic lateral sclerosis. *Eur J Neurol* 2024:e16268.

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Günther R, Wurster C, Brakemeier S, Osmanovic A, Schreiber-Katz O, Petri S, Uzelac Z, Hiebeler M, Thiele S, Walter M, Weiler M, Kessler T, Freigang M, Lapp H, Cordts I, Lingor P, Deschauer M, Hahn A, Martakis K, Steinbach R, Ilse B, Roediger A, Bellut J, Nentwich J, Zeller D, Muhandes M, Baum T, Koch J, Schrank B, Fischer S, Hermann A, Kamm C, Naegel S, Mensch A, Weber M, Neuwirth C, Lehmann H, Wunderlich G, Stadler C, Tomforde M, George A, Gross M, Pechmann A, Kirschner J, Türk M, Schimmel M, Bernert G, Martin P, Rauscher C, Meyer Zu Hörste G, Baum P, Löscher W, Flotats-Bastardas M, Köhler C, Probst-Schendzielorz K, Goldbach S, Schara-Schmidt U, Mueller-Felber W, Lochmüller H, von Velsen O, SMARtCARE study group, Kleinschnitz C, Ludolph A, Hagenacker T. Long-term efficacy and safety of nusinersen in adults with 5q spinal muscular atrophy: a prospective European multinational observational study. *Lancet Reg Health Eur* 2024; 39:100862.

Megat S, Mora N, Sanogo J, Roman O, Catanese A, Alami N, Freischmidt A, Mingaj X, De Calbiac H, Muratet F, Dirrig-Grosch S, Dieterle S, Van Bakel N, Müller K, Sieverding K, Weishaupt J, Andersen P, Weber M, Neuwirth C, Margelisch M, Sommacal A, van Eijk K, Veldink J, PROJECT MINE ALS SEQUENCING CONSORTIUM, Lautrette G, Couratier P, Camuzat A, Le Ber I, Grassano M, Chio A, Boeckers T, Ludolph A, Roselli F, Yilmazer-Hanke D, Millecamps S, Kabashi E, Storkebaum E, Sellier C, Dupuis L. Author Correction: Integrative genetic analysis illuminates ALS heritability and identifies risk genes. *Nat Commun* 2023; 14:8026.

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Projekte (18)

A Phase 3, Randomized, Double-blind, Placebo-controlled, Parallel, Multicenter Study to Evaluate the Safety and Efficacy of ALXN1720 in Adults with Generalized Myasthenia Gravis

Klinische Forschung - 01.02.2024 - 04.04.2027

Laufend

Sicherheit und Wirksamkeit von ION363 bei Patienten mit genetischer ALS mit Fused in Sarcoma Mutation (FUS-ALS), Phase 1-3

Klinische Forschung - 03.04.2023 - 03.04.2028

Laufend

TEAR und EARLY ALS

Grundlagenforschung - 01.09.2021 - 31.03.2024

Automatisch geschlossen

Study to Assess the Safety, Tolerability, Pharmacokinetics, and Effect on Disease Progression of BIIB078 Administered to Previously Treated Adults C9ORF72-Associated Amyotrophic Lateral Sclerosis (ALS)

Klinische Forschung - 28.04.2020 - 28.07.2023

Automatisch geschlossen

ALXN1210-ALS-308: An Efficacy and Safety Study of Ravulizumab in ALS Patients

Klinische Forschung - 30.03.2020 - 17.10.2021

Abgebrochen

SMARtCare: Longitudinal data assessment from patients with spinal muscular atrophy

Klinische Forschung - 09.12.2019 - 31.12.2030

Laufend

Arimoclomol in Amyotrophic Lateral Sclerosis - Open Label Extension Trial

Klinische Forschung - 19.09.2019 - 02.07.2021

Abgebrochen

A Phase 1 Multiple-Ascending-Dose Study to Assess the Safety, Tolerability, and Pharmacokinetics of BIIB078 Administered Intrathecally to Adults with C9ORF72-Associated Amyotrophic Lateral Sclerosis

Klinische Forschung - 10.09.2018 - 17.11.2021

Abgeschlossen

Arimoclomol in Amyotrophic Lateral Sclerosis

Klinische Forschung - 31.07.2018 - 18.12.2020

Abgebrochen

Methodologische Studie neuartiger elektrophysiologischer, physikalischer und bildgebender Messgrößen zur Beurteilung der Krankheitsprogression bei amyotropher Lateralsklerose

Klinische Forschung - 01.05.2016 - 01.06.2018

Automatisch geschlossen

Methodology Study of Novel Outcome Measures to Assess Progression of ALS

Klinische Forschung - 02.04.2016 - 14.08.2019

Abgeschlossen

Survival, Trigger and Risk, Epigenetic, eNviromental and Genetic Targets for motor neuron Health (STRENGTH)

Klinische Forschung - 01.01.2015 - 31.12.2017

Abgeschlossen

Randomized placebo-controlled crossover trial with quinine sulfate for the treatment of cramps in amyotrophic lateral sclerosis (ALS)

Klinische Forschung - 01.11.2013 - 26.03.2020

Automatisch geschlossen

Swiss Registry for Neuromuscular Disorders

Klinische Forschung - 01.01.2012 - 30.12.2030

Laufend

A novel neurophysiological measurement (MUNIX) in neuromuscular disorders

Klinische Forschung - 26.03.2011 - 26.03.2020

Automatisch geschlossen

Validierung der Awaji-Kriterien

Klinische Forschung - 20.03.2011 - 20.03.2014

Abgeschlossen

PEG: Perkutane Endoskopische Gastrostomie-Register für Patienten mit amyotropher Lateralsklerose (ALS) in Deutschland und Schweiz. Multizentrische Beobachtungsstudie

Klinische Forschung - 15.12.2010 - 14.12.2014

Abgeschlossen

Autopsy study in ALS

Klinische Forschung - 01.01.1999 - 01.01.2003

Automatisch geschlossen

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