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Forskningsinteresser

Sjældne Sygdomme, dysmorfologi, genomisk medicin, genetisk diagnostik af sjældne sygdomme og syendormer med og uden mentale handicap, Klinisk Genetik, Genetisk rådgivning.

Ansættelse

Klinisk Institut

SDU

1. sep. 2024 → 31. jul. 2025

Klinisk professor

KI, OUH, Forskningsenhed for Human Genetik (Odense)

SDU

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Cheflæge

Odense Universitetshospital

Odense, Danmark

1. jul. 2022 → present

Ledende overlæge

Odense Universitetshospital

Odense, Danmark

1. mar. 2010 → 30. jun. 2022

Publikationer

Late diagnosis of partial 3 β -hydroxysteroid dehydrogenase type 2 deficiency - characterization of a new genetic variant

Øzdemir, C. M., Nielsen, M. M., Liimatta, J., Voegel, C. D., Elzenaty, R. N., Wasehuus, V. S., Lind-Holst, M., Ornstrup, M. J., Gram, S. B., Ousager, L. B., Flück, C. E. & Gravholt, C. H., 1. jul. 2024, I: *Endocrinology, Diabetes & Metabolism Case Reports*. 2024, 3, 8 s., e230090.

Klinisk genetik

Ousager, L. B., Kibæk Nielsen, I. & Sunde, L., 30. maj 2024, *Medicinsk Kompendium*. Hauge, E. M., Ainsworth, M. A. & Poulsen, S. D. (red.). 20. udg. Munksgaard, Bind 2. s. 1947-1957

Comprehensive Noninvasive Fetal Screening by Deep Trio-Exome Sequencing

Miceikaitė, I., Hao, Q., Brasch-Andersen, C., Fagerberg, C. R., Topping, P. M., Kristiansen, B. S., Ousager, L. B., Sperling, L., Ibsen, M. H., Löser, K. & Larsen, M. J., 1. maj 2024, I: *Obstetrical and Gynecological Survey*. 79, 5, s. 261-263

Identification of a founder variant AAGAB c.370C>T, p.Arg124Ter in patients with punctate palmoplantar keratoderma in Southern Denmark

Gram, S. B., Jørgensen, A. S. F., Bygum, A., Brusgaard, K. & Ousager, L. B., maj 2024, I: *Clinical Genetics*. 105, 5, s. 561-566

Plantar keratoderma and curly hair as a diagnostic clue of cardiomyopathy risk

Gram, S. B., Brusgaard, K., Bygum, A., Christensen, A. H. & Ousager, L. B., maj 2024, I: *Journal of Dermatology*. 51, 5, s. e143-e144

Risk of Cancer and Mortality in Peutz-Jeghers Syndrome and Juvenile Polyposis Syndrome—A Nationwide Cohort Study With Matched Controls

Jelsig, A. M., Wullum, L., Kuhlmann, T. P., Ousager, L. B., Burisch, J. & Karstensen, J. G., dec. 2023, I: *Gastroenterology*. 165, 6, s. 1565-1567.e2

Comprehensive Noninvasive Fetal Screening by Deep Trio-Exome Sequencing

Miceikaitė, I., Hao, Q., Brasch-Andersen, C., Fagerberg, C. R., Topping, P. M., Kristiansen, B. S., Ousager, L. B., Sperling, L., Ibsen, M. H., Löser, K. & Larsen, M. J., 23. nov. 2023, I: *The New England Journal of Medicine*. 389, 21, s. 2017-2019

Structural and non-coding variants increase the diagnostic yield of clinical whole genome sequencing for rare diseases

Pagnamenta, A. T., Camps, C., Giacomuzzi, E., Taylor, J. M., Hashim, M., Calpena, E., Kaisaki, P. J., Hashimoto, A., Yu, J., Sanders, E., Schwessinger, R., Hughes, J. R., Lunter, G., Dreau, H., Ferla, M., Lange, L., Kesim, Y., Ragoussis, V., Vavoulis, D. V. & Allroggen, H. & 77 flere, Ansorge, O., Babbs, C., Banka, S., Baños-Piñero, B., Beeson, D., Ben-Ami, T., Bennett, D. L., Bento, C., Blair, E., Brasch-Andersen, C., Bull, K. R., Cario, H., Cilliers, D., Conti, V., Davies, E. G., Dhalla, F., Dacal, B. D., Dong, Y., Dunford, J. E., Guerrini, R., Harris, A. L., Hartley, J., Hollander, G., Javaid, K., Kane, M., Kelly, D., Kelly, D., Knight, S. J. L., Kreins, A. Y., Kvikstad, E. M., Langman, C. B., Lester, T., Lines, K. E., Lord, S. R., Lu, X., Mansour, S., Manzur, A., Maroofian, R., Marsden, B., Mason, J., McGowan, S. J., Mei, D., Milcochova, H., Murakami, Y., Németh, A. H., Okoli, S., Ormondroyd, E., Ousager, L. B., Palace, J., Patel, S. Y., Pentony, M. M., Pugh, C., Rad, A., Ramesh, A., Riva, S. G., Roberts, I., Roy, N., Salminen, O., Schilling, K. D., Scott, C., Sen, A., Smith, C., Stevenson, M., Thakker, R. V., Twigg, S. R. F., Uhlig, H. H., van Wijk, R., Vona, B., Wall, S., Wang, J., Watkins, H., Zak, J., Schuh, A. H., Kini, U., Wilkie, A. O. M., Popitsch, N. & Taylor, J. C., 9. nov. 2023, I: *Genome Medicine*. 15, 25 s., 94.

Cancer risk and mortality in patients with solitary juvenile polyps-A nationwide cohort study with matched controls

Jelsig, A. M., Wullum, L., Kuhlmann, T. P., Ousager, L. B., Burisch, J. & Karstensen, J. G., okt. 2023, I: *United European Gastroenterology Journal*. 11, 8, s. 745-749

Cancer risk and mortality in patients with solitary Peutz-Jeghers polyps

Jelsig, A. M., Wullum, L., Ousager, L. B., Burisch, J., Kuhlmann, T. P. & Karstensen, J. G., 22. sep. 2023, I: *Gastroenterology Report*. 11, goad056.

Is punctate palmoplantar keratoderma type 1 associated with malignancy? A systematic review of the literature

Gram, S. B., Bjerrelund, J., Jelsig, A. M., Bygum, A., Leboeuf-Yde, C. & Ousager, L. B., 13. sep. 2023, I: *Orphanet Journal of Rare Diseases*. 18, 11 s., 290.

Comprehensive prenatal diagnostics: Exome versus genome sequencing

Miceikaite, I., Fagerberg, C., Brasch-Andersen, C., Topping, P. M., Kristiansen, B. S., Hao, Q., Sperling, L., Ibsen, M. H., Löser, K., Bendsen, E. A., Ousager, L. B. & Larsen, M. J., aug. 2023, I: *Prenatal Diagnosis*. 43, 9, s. 1132-1141

Survival, surveillance, and genetics in patients with Peutz-Jeghers syndrome: A nationwide study

Jelsig, A. M., van Overeem Hansen, T., Gede, L. B., Qvist, N., Christensen, L.-L., Lautrup, C. K., Frederiksen, J. H., Sunde, L., Ousager, L. B., Ljungmann, K., Bertelsen, B. & Karstensen, J. G., jul. 2023, I: *Clinical Genetics*. 104, 1, s. 81-89

En sjælden differentialdiagnose til vorter: Ugens billede

Gram, S. B., Brusgaard, K., Ousager, L. B. & Bygum, A., 27. feb. 2023, (E-pub ahead of print) I: *Ugeskrift for Læger*. V71166.

Further clinical and molecular characterization of an XLID syndrome associated with BRWD3 variants, a gene implicated in the leukemia-related JAK-STAT pathway

Delanne, J., Lecat, M., Blackburn, P. R., Klee, E. W., Stumpel, C. T. R. M., Stegmann, S., Stevens, S. J. C., Nava, C., Heron, D., Keren, B., Mahida, S., Naidu, S., Babovic-Vuksanovic, D., Herkert, J. C., Topping, P. M., Kibæk, M., De Bie, I., Pfundt, R., Hendriks, Y. M. C. & Ousager, L. B. & 15 flere, Bend, R., Warren, H., Skinner, S. A., Lyons, M. J., Pöe, C., Chevarin, M., Jouan, T., Garde, A., Thomas, Q., Kuentz, P., Tisserant, E., Duffourd, Y., Philippe, C., Faivre, L. & Thauvin-Robinet, C., jan. 2023, I: *European Journal of Medical Genetics*. 66, 1, 7 s., 104670.

A novel heterozygote LORICRIN variant in a father and daughter with palmoplantar keratoderma

Fredberg, S., Gram, S. B., Lei, U., Bygum, A., Brusgaard, K. & Ousager, L. B., 2023, I: *European Journal of Human Genetics*. 31, Suppl. 1

Genetic testing of 76 families with palmoplantar keratoderma reveals a monogenic cause in more than 78 %

Gram, S. B., Brusgaard, K., Lei, U., Sommerlund, M., Vinding, G. R., Fast, S., Overgaard Bach, R., Bygum, A. & Ousager, L. B., 2023, I: *European Journal of Human Genetics*. 31, Suppl. 1

Identification of a Novel PLCD1 Variant in a Danish Family with Hereditary Leukonychia

Lieberoth, S., Kumar, S., Brusgaard, K., Ousager, L. B., Betz, R. C. & Bygum, A., 2023, I: *Skinmed*. 21, 1, s. 44-46

Loss-of-function variants in SRRM2 cause a neurodevelopmental disorder

Cuinat, S., Nizon, M., Isidor, B., Stegmann, A., van Jaarsveld, R. H., van Gassen, K. L., van der Smagt, J. J., Volker-Touw, C. M. L., Holwerda, S. J. B., Terhal, P. A., Schuhmann, S., Vasileiou, G., Khalifa, M., Nugud, A. A., Yasaei, H., Ousager, L. B., Andersen, C. B., Deb, W., Besnard, T. & Simon, M. E. H. & 31 flere, Amsterdam, K.H.-V., Verbeek, N. E., Matalon, D., Dykzeul, N., White, S., Spiteri, E., Devriendt, K., Boogaerts, A., Willemsen, M., Brunner, H. G., Sinnema, M., De Vries, B. B. A., Gerkes, E. H., Pfundt, R., Izumi, K., Krantz, I. D., Xu, Z. L., Murrell, J. R., Valenzuela, I., Cusco, I., Rovira-Moreno, E., Yang, Y., Bizaoui, V., Patat, O., Faivre, L., Tran-Mau-Them, F., Vitobello, A., Denommé-Pichon, A.-S., Philippe, C., Bezieau, S. & Cogné, B., aug. 2022, I: *Genetics in medicine : official journal of the American College of Medical Genetics*. 24, 8, s. 1774-1780

von Hippel-Lindau disease: Updated guideline for diagnosis and surveillance

Louise M Binderup, M., Smerdel, M., Borgwadt, L., Beck Nielsen, S. S., Madsen, M. G., Møller, H. U., Kiilgaard, J. F., Friis-Hansen, L., Harbud, V., Cortnum, S., Owen, H., Gimsing, S., Friis Juhl, H. A., Munthe, S., Geilswijk, M., Rasmussen, Å. K., Møldrup, U., Graumann, O., Donskov, F. & Grønbæk, H. & 11 flere, Stausbøl-Grøn, B., Schaffalitzky de Muckadell, O., Knigge, U., Dam, G., Wadt, K. A., Bøgeskov, L., Bagi, P., Lund, L., Stochholm, K., Ousager, L. B. & Sunde, L., aug. 2022, I: *European Journal of Medical Genetics*. 65, 8, 12 s., 104538.

Familial Multiple Lipomatosis - analyses of genetic etiology by whole genome sequencing and delineation of the clinical phenotype

Bjerrelund, J., Larsen, M. J., Schroder, H. D., Frost, M., Kassem, M., Ousager, L. B. & Frederiksen, A. L., 1. apr. 2022, I: *European Journal of Human Genetics*. 30, Suppl. 1, s. 205 1 s., P06.014.C.

Klinisk genetik: Genetisk udredning og rådgivning

Ousager, L. B. & Kibæk Nielsen, I., 2022, *Medicinsk genetik*. Østergaard, E. & Sunde, L. (red.). 3. udg. FADL's Forlag, s. 431-452 1

Klinisk genetik: Etik og jura

Ousager, L. B., Gerdes, A.-M. A. & Kristensen, K., 2022, *Medicinsk genetik*. Østergaard, E. & Sunde, L. (red.). 3. udg. FADL's Forlag, s. 477-500 4

Sygdommes genetik: Syndromer, medfødte misdannelser og mental retardering

Kibæk Nielsen, I. & Ousager, L. B., 2022, *Medicinsk genetik*. Østergaard, E. & Sunde, L. (red.). 3. udg. FADL's Forlag, s. 393-422 16

Truncating SRCAP variants outside the Floating-Harbor syndrome locus cause a distinct neurodevelopmental disorder with a specific DNA methylation signature

Rots, D., Chater-Diehl, E., Dingemans, A. J. M., Goodman, S. J., Siu, M. T., Cytrynbaum, C., Choufani, S., Hoang, N., Walker, S., Awamleh, Z., Charkow, J., Meyn, S., Pfundt, R., Rinne, T., Gardeitchik, T., de Vries, B. B. A., Deden, A. C., Leenders, E., Kwint, M. & Stumpel, C. T. R. M. & 67 flere, Stevens, S. J. C., Vermeulen, J. R., van Harssel, J. V. T., Bosch, D. G. M., van Gassen, K. L. I., van Binsbergen, E., de Geus, C. M., Brackel, H., Hempel, M., Lessel, D., Denecke, J., Slavotinek, A., Strober, J., Crunk, A., Folk, L., Wentzensen, I. M., Yang, H., Zou, F., Millan, F., Person, R., Xie, Y., Liu, S., Ousager, L. B., Larsen, M., Schultz-Rogers, L., Morava, E., Klee, E. W., Berry, I. R., Campbell, J., Lindstrom, K., Pruniski, B., Radley, J. A., Phornphutkul, C., Schmidt, B., Wilson, W. G., Öunap, K., Reinson, K., Pajusalu, S., van Haeringen, A., Ruivenkamp, C., Cuperus, R., Santos-Simarro, F., Palomares-Bralo, M., Pacio-Míguez, M., Ritter, A., Bhoj, E., Tønne, E., Tveten, K., Cappuccio, G., Brunetti-Pierri, N., Rowe, L., Bunn, J., Saenz, M., Platzer, K., Mertens, M., Caluseriu, O., Nowaczyk, M. J. M., Cohn, R. D., Kannu, P., Alkhunaizi, E., Chitayat, D., Scherer, S. W., Brunner, H. G., Vissers, L. E. L. M., Kleefstra, T., Koolen, D. A. & Weksberg, R., 3. jun. 2021, I: *American Journal of Human Genetics*. 108, 6, s. 1053-1068

Hereditary leiomyomatosis and renal cell carcinoma: a case series and literature review

Chayed, Z., Kristensen, L. K., Ousager, L. B., Rønlund, K. & Bygum, A., 18. jan. 2021, I: Orphanet Journal of Rare Diseases. 16, 9 s., 34.

A novel homozygous SLURP1 variant in a boy with Mal de Meleda born to non-consanguineous parents

Gram, S. B., Blechingberg, J., Brusgaard, K., Bygum, A. & Ousager, L. B., 1. dec. 2020, I: European Journal of Human Genetics. 28, Suppl. 1, s. 840 1 s., E-P04.47.

Location, location, location: protein truncating variants in different loci of SRCAP cause three distinct neurodevelopmental disorders, associated with distinctive DNA methylation signatures

Rots, D., Chater-Diehl, E., Dingemans, A. J. M., Siu, M., Cytrynbaum, C., Hoang, N., Walker, S., Scherer, S., Pfundt, R., Rinne, T., Gardeitchik, T., de Vries, B. B. A., Stumpel, C. T. R. M., Stevens, S. J. C., van Harsseel, J., Bosch, D. G. M., van Gassen, K. L. I., van Binsbergen, E., de Geus, C. M. & Hempel, M. & 30 flere, Lessel, D., Denecke, J., Slavotinek, A., Strober, J., Ousager, L. B., Larsen, M., Schultz-Rogers, L., Morava, E., Klee, E. W., Berry, I. R., Campbell, J., Lindstrom, K., Neumeyer, A. M., Radley, J. A., Phornphutkul, C., Wilson, W. G., Schmidt, B., Meyn, S., Ounap, K., Reinson, K., Pajusalu, S., Ruivenkamp, C., van Haeringen, A., Cuperus, R., Vissers, L. E. L. M., Brunner, H. G., Kleefstra, T., Koolen, D. A., Weksberg, R. & Inc, G., 1. dec. 2020, I: European Journal of Human Genetics. 28, Suppl. 1, s. 34-35

Chromosomal translocation disrupting the SMAD4 gene resulting in the combined phenotype of Juvenile polyposis syndrome and Hereditary Hemorrhagic Telangiectasia

Aagaard, K. S., Brusgaard, K., Miceikaite, I., Larsen, M. J., Kjeldsen, A. D., Lester, E. B., Ousager, L. B. & Tørring, P. M., nov. 2020, I: Molecular Genetics and Genomic Medicine. 8, 11, 8 s., e1498.

Low frequency of parental mosaicism in de novo COL4A5 mutations in X-linked Alport syndrome

Helle, O. M. B., Pedersen, T. H., Ousager, L. B., Thomassen, M. & Hertz, J. M., 1. okt. 2020, I: Molecular Genetics & Genomic Medicine. 8, 10, 7 s., e1452.

Lessons learned from 40 novel PIGA patients and a review of the literature

Bayat, A., Knaus, A., Pendziwiat, M., Afenjar, A., Stefan Barakat, T., Bosch, F., Callewaert, B., Calvas, P., Ceulemans, B., Chassaing, N., Depienne, C., Endziniene, M., Ferreira, C. R., Moura de Souza, C. F., Freihuber, C., Ganesan, S., Gataullina, S., Guerrini, R., Guerrot, A.-M. & Hansen, L. & 34 flere, Jezela-Stanek, A., Karsenty, C., Kievit, A., Kooy, F. R., Korff, C. M., Kragh Hansen, J., Larsen, M., Layet, V., Lesca, G., McBride, K. L., Meuwissen, M., Mignot, C., Montomoli, M., Moore, H., Naudion, S., Nava, C., Nougues, M.-C., Parrini, E., Pastore, M., Schelhaas, J. H., Skinner, S., Szczałuba, K., Thomas, A., Thomassen, M., Tranebjaerg, L., van Slegtenhorst, M., Wolfe, L. A., Lal, D., Gardella, E., Bomme Ousager, L., Brünger, T., Helbig, I., Krawitz, P. & Møller, R. S., jun. 2020, I: Epilepsia. 61, 6, s. 1142-1155

c.1227_1228dupGG (p.Glu410Glyfs), a frequent variant in Tunisian patients with MUTYH associated polyposis

Kdissa, A., Brusgaard, K., Ksaa, M., Golli, L., Hallara, O., Ousager, L. B., Manoubi, W., Seghaier, R. B., Adala, L., Halleb, Y., Saad, A., Hmila, F. & Gribaa, M., jan. 2020, I: Cancer Genetics. 240, s. 45-53

Estimating the effect size of the 15Q11.2 BP1-BP2 deletion and its contribution to neurodevelopmental symptoms: recommendations for practice

Jønch, A. E., Douard, E., Moreau, C., Van Dijck, A., Passeggeri, M., Kooy, F., Puechberty, J., Campbell, C., Sanlaville, D., Lefroy, H., Richetin, S., Pain, A., Geneviève, D., Kini, U., Le Caignec, C., Lespinasse, J., Skytte, A. B., Isidor, B., Zweier, C. & Caberg, J. H. & 8 flere, Delrue, M. A., Møller, R. S., Bojesen, A., Hjalgrim, H., Brasch-Andersen, C., Lemyre, E., Ousager, L. B. & Jacquemont, S., okt. 2019, I: Journal of Medical Genetics. 56, 10, s. 701-710

Characterizing and quantifying the effect of the recurrent copy number variants between BP1-BP2 at chromosome 15q11.2

Jønch, A. E., Douard, E., Van Dijck, A., Kooy, F. R., Puechberty, J., Campbell, C., Salanville, D., Lefroy, H., Geneviève, D., Kini, U., Le Caignec, C., Lespinasse, J., Skytte, A., Isidor, B., Zweier, C., Caberg, J., Brasch-Andersen, C., Lemyre, E., Ousager, L. B. & Jacquemont, S., 1. jul. 2019, I: European Journal of Human Genetics. 26, Suppl 1, s. 363-364 2 s., P09.007C.

Danish Cytogenetic Central Register

Al-Zehhawi, L., Hansen, J., Kjaergaard, S., Ousager, L., Fagerberg, C., Sunde, L. & Rasmussen, M., 1. jul. 2019, I: European Journal of Human Genetics. 26, Suppl 1, s. 792 1 s., P19.13A.

THE TRUE CONTRIBUTION OF THE 15Q11.2 BP1-BP2 DELETION TO NEURODEVELOPMENTAL SYMPTOMS

Jonch, A., Douard, E., Moreau, C., Van Dijck, A., Kooy, F., Sanlaville, D., Brasch-Andersen, C., Lemyre, E., Ousager, L. B. & Jacquemont, S., 2019, I: *European Neuropsychopharmacology*. 29, 4 Supplement, s. 1150-1151 2 s., F75.

De Novo Variants in the F-Box Protein FBXO11 in 20 Individuals with a Variable Neurodevelopmental Disorder

University of Washington Center for Mendelian Genomics, 2. aug. 2018, I: *American Journal of Human Genetics*. 103, 2, s. 305-316

Autosomal dominant stapes fixation, syndactyly, and symphalangism in a family with NOG mutation: Long term follow-up on surgical treatment

Westergaard-Nielsen, M., Amstrup, T., Wanscher, J. H., Brusgaard, K. & Ousager, L. B., maj 2018, I: *International Journal of Pediatric Otorhinolaryngology*. 108, s. 208-212

Pigmentary mosaicism: A review of original literature and recommendations for future handling

Kromann, A. B., Ousager, L. B., Ali, I. K. M., Aydemir, N. & Bygum, A., 5. mar. 2018, I: *Orphanet Journal of Rare Diseases*. 13, 10 s., 39.

Chromosomal translocation as a cause of JP/HHT

Brusgaard, K., Tørring, P. M., Kjeldsen, A. D., Timshel, S., Hammer, T. & Ousager, L. B., 1. feb. 2018, I: *Angiogenesis*. 21, 1, s. 146 1 s.

Mosaicism in a patient with HHT

Brusgaard, K., Tørring, P. M., Kjeldsen, A. D. & Ousager, L. B., 1. feb. 2018, I: *Angiogenesis*. 21, 1, s. 146-147

ENG mutational mosaicism in a family with hereditary hemorrhagic telangiectasia

Tørring, P. M., Kjeldsen, A. D., Ousager, L. B. & Brusgaard, K., jan. 2018, I: *Molecular Genetics & Genomic Medicine*. 6, 1, s. 121-125

Chromosomal contacts connect loci associated with autism, BMI and head circumference phenotypes

Ousager, L. B. (Medlem af forfattergruppering) & 2p15 Consortium, jun. 2017, I: *Molecular Psychiatry*. 22, 6, s. 836-849

Familial cerebral abscesses caused by hereditary hemorrhagic telangiectasia

Tørring, P. M., Lauridsen, M. F., I Dali, C., Andersen, P. E., Ousager, L. B., Brusgaard, K. & Kjeldsen, A. D., jun. 2017, I: *Clinical Case Reports*. 5, 6, s. 805-808

Genetic screening of the FLCN gene identify six novel variants and a Danish founder mutation

Rossing, M., Albrechtsen, A., Skytte, A.-B., Jensen, U. B., Ousager, L. B., Gerdes, A.-M., Nielsen, F. C. & Hansen, T. V., feb. 2017, I: *Journal of Human Genetics*. 62, 2, s. 151-157

Danish Cytogenetic Central Register

Al-Zehhawi, L. I. K., Hansen, J., Kjærgaard, S., Ousager, L. B., Fagerberg, C., Sunde, L. & Rasmussen, M., 2017.

De Novo Variants in GRIA4 Lead to Intellectual Disability with or without Seizures and Gait Abnormalities

Martin, S., Chamberlin, A., Shinde, D. N., Hempel, M., Strom, T. M., Schreiber, A., Johannsen, J., Ousager, L. B., Larsen, M. J., Hansen, L. K., Fatemi, A., Cohen, J. S., Lemke, J., Sørensen, K. P., Helbig, K. L., Lessel, D. & Abou Jamra, R., 2017, I: *American Journal of Human Genetics*. 101, 6, s. 1013-1020

Novel ELN mutation in a family with supraaortic stenosis and intracranial aneurysm

Jelsing, A. M., Urban, Z., Huchtagowder, V., Nissen, H. & Ousager, L. B., 2017, I: *European Journal of Medical Genetics*. 60, 2, s. 110-113

Germline mutations in BMP9 are not identified in a series of Danish and French patients with hereditary hemorrhagic telangiectasia

Tørring, P. M., Dupuis-Girod, S., Giraud, S., Brusgaard, K., Ousager, L. B. & Kjeldsen, A. D., 1. dec. 2016, I: *Gene Reports*. 5, s. 30-33

Juvenile Polyps in Denmark From 1995 to 2014

Jelsig, A. M., Ousager, L. B., Brusgaard, K. & Qvist, N., aug. 2016, I: *Diseases of the Colon and Rectum*. 59, 8, s. 751-757

JP-HHT phenotype in Danish patients with SMAD4 mutations

Jelsig, A. M., Tørring, P. M., Kjeldsen, A. D., Qvist, N., Bojesen, A., Jensen, U. B., Andersen, M. K., Gerdes, A. M., Brusgaard, K. & Ousager, L. B., jul. 2016, I: *Clinical Genetics*. 90, 1, s. 55-62

Acromelic frontonasal dysostosis and ZSWIM6 mutation: phenotypic spectrum and mosaicism

Twigg, S. R. F., Ousager, L. B., Miller, K. A., Zhou, Y., Elalaoui, S. C., Sefiani, A., Bak, G. S., Hove, H., Kjærsgaard Hansen, L., Fagerberg, C. R., Tajir, M. & Wilkie, A. O. M., 2016, I: *Clinical Genetics*. 90, 3, s. 270-275

Bone structure in two adult subjects with impaired minor spliceosome function resulting from RNU4ATAC mutations causing microcephalic osteodysplastic primordial dwarfism type 1 (MOPD1)

Krøigård, A. B., Frost, M., Larsen, M. J., Ousager, L. B. & Frederiksen, A. L., 2016, I: *Bone*. 92, s. 145-149

Disease pattern in Danish patients with Peutz-Jeghers syndrome

Jelsig, A. M., Qvist, N., Sunde, L., Brusgaard, K., Hansen, T., Wikman, F. P., Nielsen, C. B., Nielsen, I. K., Gerdes, A. M., Bojesen, A. & Ousager, L. B., 2016, I: *International Journal of Colorectal Disease*. 31, 5, s. 997-1004

Germline variants in Hamartomatous Polyposis Syndrome-associated genes from patients with one or few hamartomatous polyps

Jelsig, A. M., Brusgaard, K., Hansen, T. P., Qvist, N., Larsen, M., Bojesen, A., Nielsen, C. B. & Ousager, L. B., 2016, I: *Scandinavian Journal of Gastroenterology*. 51, 9, s. 1118-1125

Two novel mutations in RNU4ATAC in two siblings with an atypical mild phenotype of microcephalic osteodysplastic primordial dwarfism type 1

Krøigård, A. B., Jackson, A. P., Bicknell, L. S., Baple, E., Brusgaard, K., Hansen, L. K. & Ousager, L. B., 2016, I: *Clinical Dysmorphology*. 25, 2, s. 68-72

X-exome sequencing of 405 unresolved families identifies seven novel intellectual disability genes

Hu, H., Haas, S. A., Chelly, J., Van Esch, H., Raynaud, M., de Brouwer, A. P. M., Weinert, S., Froyen, G., Frints, S. G. M., Laumonnier, F., Zemojtel, T., Love, M. I., Richard, H., Emde, A.-K., Bienek, M., Jensen, C., Hambrock, M., Fischer, U., Langnick, C. & Feldkamp, M. & 64 flere, Wissink-Lindhout, W., Lebrun, N., Castelnaud, L., Rucci, J., Montjean, R., Dorseuil, O., Billuart, P., Stuhlmann, T., Shaw, M., Corbett, M. A., Gardner, A., Willis-Owen, S., Tan, C., Friend, K. L., Belet, S., van Roozendaal, K. E. P., Jimenez-Pocquet, M., Moizard, M.-P., Ronce, N., Sun, R., O'Keeffe, S., Chenna, R., van Bömmel, A., Göke, J., Hackett, A., Field, M., Christie, L., Boyle, J., Haan, E., Nelson, J., Turner, G., Baynam, G., Gillissen-Kaesbach, G., Müller, U., Steinberger, D., Budny, B., Badura-Stronka, M., Latos-Bieleńska, A., Ousager, L. B., Wieacker, P., Rodríguez Criado, G., Bondeson, M.-L., Annerén, G., Dufke, A., Cohen, M., Van Maldergem, L., Vincent-Delorme, C., Echenne, B., Simon-Bouy, B., Kleefstra, T., Willemsen, M., Fryns, J.-P., Devriendt, K., Ullmann, R., Vingron, M., Wrogemann, K., Wienker, T. F., Tzschach, A., van Bokhoven, H., Gecz, J., Jentsch, T. J., Chen, W., Ropers, H.-H. & Kalscheuer, V. M., 2016, I: *Molecular Psychiatry*. 21, 1, s. 133-148 16 s.

Research participants in NGS studies want to know about incidental findings

Jelsig, A. M., Qvist, N., Brusgaard, K. & Ousager, L. B., okt. 2015, I: *European Journal of Human Genetics*. 23, 10, s. 1423-1426

Noonans syndrom kan diagnosticere klinisk og molekylærgenetisk

Krab Henningsen, M., Jelsig, A. M., Andersen, H., Brusgaard, K., Ousager, L. B. & Hertz, J. M., 3. aug. 2015, I: *Ugeskrift for Læger*. 177, 32, s. V12140755

Noonans syndrom kan diagnosticeres klinisk og molekylærgenetisk

mk, H., Jelsig, A. M., Andersen, H., Brusgaard, K., Ousager, L. B. & Hertz, J. M., 3. aug. 2015, I: *Ugeskrift for Læger*. 177, 24, s. 2-7 V12140755.

Patienter med basalcellenævussyndrom bør tilbydes tidlig interdisciplinær opfølgning og behandling

Bay, C., Ousager, L. B. & Jelsig, A. M., 13. jul. 2015, I: Ugeskrift for Læger. 178, 4, s. 2-6 V12140701.

Global gene expression profiling of telangiectasial tissue from patients with hereditary hemorrhagic telangiectasia

Tørring, P. M., Larsen, M. J., Kjeldsen, A. D., Ousager, L. B., Tan, Q. & Brusgaard, K., 1. maj 2015, I: Microvascular Research. 99, s. 118-126

De novo heterozygous mutations in SMC3 cause a range of Cornelia de Lange syndrome-overlapping phenotypes

Gil-Rodríguez, M. C., Deardorff, M. A., Ansari, M., Tan, C. A., Parenti, I., Baquero-Montoya, C., Ousager, L. B., Puisac, B., Hernández-Marcos, M., Teresa-Rodrigo, M. E., Marcos-Alcalde, I., Wesselink, J.-J., Lusa-Bernal, S., Bijlsma, E. K., Braunholz, D., Bueno-Martinez, I., Clark, D., Cooper, N. S., Curry, C. J. & Fisher, R. & 42 flere, Fryer, A., Ganesh, J., Gervasini, C., Gillessen-Kaesbach, G., Guo, Y., Hakonarson, H., Hopkin, R. J., Kaur, M., Keating, B. J., Kibæk, M., Kinning, E., Kleefstra, T., Kline, A. D., Kuchinskaya, E., Larizza, L., Li, Y. R., Liu, X., Mariani, M., Picker, J. D., Pié, Á., Pozojevic, J., Queralt, E., Richer, J., Roeder, E., Sinha, A., Scott, R. H., So, J., Wusik, K. A., Wilson, L., Zhang, J., Gómez-Puertas, P., Casale, C. H., Ström, L., Selicorni, A., Ramos, F. J., Jackson, L. G., Krantz, I. D., Das, S., Hennekam, R. C. M., Kaiser, F. J., FitzPatrick, D. R. & Pié, J., apr. 2015, I: Human Mutation. 36, 4, s. 454-462 9 s.

Arvelig palmopantar keratodermi

Kamaleswaran, S., Ousager, L. B., Overgaard Bach, R. & Bygum, A., 2015, I: Ugeskrift for Læger. 177, 5, s. 457-461 V05130280.

Germline mutations in BMP9 are not identified in Danish patients with hereditary haemorrhagic telangiectasia

Tørring, P. M., Brusgaard, K., Ousager, L. B. & Kjeldsen, A. D., 2015, I: Angiogenesis. 18, 4, s. 537 1 s., OR31 .

Long non-coding rna expression profiles in hereditary haemorrhagic telangiectasia

Tørring, P. M., Larsen, M. J., Kjeldsen, A. D., Ousager, L. B., Tan, Q. & Brusgaard, K., 2015, I: Angiogenesis. 18, 4, s. 569-569 1 s., P76.

Phenotype and genotype in 103 patients with tricho-rhino-phalangeal syndrome

Maas, S. M., Shaw, A. C., Bikker, H., Lüdecke, H.-J., van der Tuin, K., Badura-Stronka, M., Belligni, E., Biamino, E., Bonati, M. T., Carvalho, D. R., Cobben, J., de Man, S. A., Den Hollander, N. S., Di Donato, N., Garavelli, L., Grønberg, S., Herkert, J. C., Hoogeboom, A. J. M., Jamsheer, A. & Latos-Bielenska, A. & 27 flere, Maat-Kievit, A., Magnani, C., Marcelis, C., Mathijssen, I. B., Nielsen, M., Otten, E., Ousager, L. B., Pilch, J., Plomp, A., Poke, G., Poluha, A., Posmyk, R., Rieubland, C., Silengo, M., Simon, M., Steichen, E., Stumpel, C., Szakszon, K., Polonkai, E., van den Ende, J., van der Steen, A., van Essen, T., van Haeringen, A., van Hagen, J. M., Verheij, J. B. G. M., Mannens, M. M. & Hennekam, R. C., 2015, I: European Journal of Medical Genetics. 58, 5, s. 279-292

Tidlig epileptisk encefalopati forårsaget af CDKL-5-mutation

Hjalmsgrim, H., Hansen, L. K., Ousager, L. B. & Møller, R. S., 15. dec. 2014, I: Ugeskrift for Læger. 176, 25A, s. 46-47

Juvenil polypose-syndrom er en sjælden årsag til kræft i gastrointestinkanalen

Jelsig, A. M., Tørring, P. M., Qvist, N., Bernstein, I. & Ousager, L. B., 27. okt. 2014, I: Ugeskrift for Læger. 176, 22, s. 2077-2080 V06130391.

Juvenil polypose-syndrom og arvelig hæmorrhagisk telangiectasi hos en patient med SMAD4-mutation

Jelsig, A. M., Tørring, P. M., Wikman, F., Mortensen, M. B., Qvist, N. & Ousager, L. B., 27. okt. 2014, I: Ugeskrift for Læger. 176, 44, s. 2081-2082

National mutation study among Danish patients with hereditary haemorrhagic telangiectasia

Tørring, P. M., Brusgaard, K., Ousager, L. B., Andersen, P. E. & Kjeldsen, A. D., aug. 2014, I: Clinical Genetics. 86, 2, s. 123-133 11 s.

Hamartomatous polyposis syndromes: a review

Jelsig, A. M., Qvist, N., Brusgaard, K., Nielsen, C. B., Hansen, T. P. & Ousager, L. B., 15. jul. 2014, I: Orphanet Journal of Rare Diseases. 9, July, 10 s., 101.

On the formation of 7-ketocholesterol from 7-dehydrocholesterol in patients with CTX and SLO

Björkhem, I., Diczfalussy, U., Lövgren-Sandblom, A., Starck, L., Jonsson, M., Tallman, K., Schirmer, H., Ousager, L. B., Crick, P. J., Wang, Y., Griffiths, W. J. & Guengerich, F. P., 25. apr. 2014, I: *Journal of Lipid Research*. 55, 6, s. 1165-1172

Mutations in Danish patients with long QT syndrome and the identification of a large founder family with p.F29L in KCNH2

Christiansen, M., Hedley, P. L., Theilade, J., Stoevring, B., Leren, T. P., Eschen, O., Sørensen, K. M., Tybjærg-Hansen, A., Ousager, L. B., Pedersen, L. N., Frikke-Schmidt, R., Aidt, F. H., Hansen, M. G., Hansen, J., Bloch Thomsen, P. E., Toft, E., Henriksen, F. L., Bundgaard, H., Jensen, H. K. & Kanfers, J. K., 7. mar. 2014, I: *BMC Medical Genetics*. 15, s. 31

Long non-coding RNA expression profiles in hereditary haemorrhagic telangiectasia

Tørring, P. M., Larsen, M. J., Kjeldsen, A. D., Ousager, L. B., Tan, Q. & Brusgaard, K., 6. mar. 2014, I: *PLOS ONE*. 9, 3, e90272.

Is Colorectal Neoplasia Part of the Birt-Hogg-Dub Syndrome?

Boman, P. S., Ousager, L. B., Friis-Hansen, L., Hansen, T. V. O., Broesby-Olsen, S. & Gerdes, A.-M., 2014, I: *Journal of Gastroenterology and Hepatology Research*. 3, 4, s. 1039-40

To know or not to know: Research participants want to know about incidental findings in WES-studies

Jelsig, A. M., Qvist, N., Brusgaard, K. & Ousager, L. B., 2014.

Histiocytic disorders of the gastrointestinal tract

Detlefsen, S., Fagerberg, C. R., Ousager, L. B., Lindebjerg, J., Marcussen, N., Nathan, T. & Sørensen, F. B., maj 2013, I: *Human Pathology*. 44, 5, s. 683-696

Heart defects and other features of the 22q11 distal deletion syndrome

Fagerberg, C. R., Graakjaer, J., Heintz, U. D., Ousager, L. B., Dreyer, I., Kirchhoff, E. M., Rasmussen, A. A., Lautrup, C. K., Birkebaek, N. & Sorensen, K., feb. 2013, I: *European Journal of Medical Genetics*. 56, 2, s. 98-107

Juvenil polypose er en sjælden årsag til gastrointestinal cancer

Jelsig, A. M., Tørring, P. M., Qvist, N., Bernstein, I. T. & Ousager, L. B., 2013, I: *Ugeskrift for Læger*. 175, s. 2-5 V06130391.

Von Hippel-Lindau disease (vHL): National clinical guideline for diagnosis and surveillance in Denmark

Binderup, M. L. M., Bisgaard, S. M. L., Harbud, V., Møller, H. U., Gimsing, S., Friis-Hansen, L. J., Hansen, T. V. O., Bagi, P., Knigge, U. P., Kosteljanetz, M., Bøgeskov, L., Thomsen, C., Gerdes, A.-M., Ousager, L. B., Sunde, L. & Ousager, L. B., 2013, I: *Danish Medical Journal*. 60, 12, B4763.

Xeroderma Pigmentosum-Trichothiodystrophy overlap patient with novel XPD/ERCC2 mutation

Kralund, H. H., Ousager, L., Jaspers, N. G., Raams, A., Pedersen, E. B., Gade, E. & Bygum, A., 2013, I: *Rare diseases (Austin, Tex.)*. 1, e24932.

Allelic Dropout in the ENG Gene, Affecting the Results of Genetic Testing in Hereditary Hemorrhagic Telangiectasia

Tørring, P. M., Kjeldsen, A. D., Ousager, L. B., Brasch-Andersen, C. & Brusgaard, K., 1. dec. 2012, I: *Genetic Testing and Molecular Biomarkers*. 16, 12, s. 1419-1423 5 s.

Calcaneonavicular Coalition in Patients with Gorlin Syndrome

Kristiansen, B. S., Jelsig, A. M., Gerdes, A.-M. & Ousager, L. B., 23. jun. 2012.

Distinct mutations in STXPB2 are associated with variable clinical presentations in patients with familial hemophagocytic lymphohistiocytosis type 5 (FHL5)

Pagel, J., Beutel, K., Lehmeberg, K., Koch, F., Maul-Pavicic, A., Rohlf, A.-K., Al-Jefri, A., Beier, R., Ousager, L. B., Ehlert, K., Gross-Wieltsch, U., Jorch, N., Kremens, B., Pekrun, A., Sparber-Sauer, M., Mejstrikova, E., Wawer, A., Ehl, S., zur Stadt, U. & Janka, G., 2012, I: *Blood*. 119, 25, s. 6016-24 9 s.

Duplication of 7q36.3 encompassing the Sonic Hedgehog (SHH) gene is associated with congenital muscular hypertrophy
Kristensen, L. K., Kjaergaard, S., Kirchhoff, M., Kock, K., Brasch Andersen, C., Kibæk, M. & Ousager, L. B., 2012, I: European Journal of Medical Genetics. 55, 10, s. 557-60 4 s.

Genetisk udredning ved infantile spasmer

Kjærsgaard Hansen, L., Ousager, L. B., Møller, R. S., Uldall, P. V. & Hjalgrim, H., 2012, I: Ugeskrift for Læger. 174, 17, s. 1152-1155

Identification of a novel S249C FGFR3 mutation in a keratinocytic epidermal nevus syndrome

Ousager, L. B., Bygum, A. & Hafner, C., 2012, I: British Journal of Dermatology. 167, 1, s. 202-4 3 s.

Mutations in ENG, ALK1 and SMAD4 among Danish patients with HHT

Mathiesen Tørring, P., Kjeldsen, A. D., Ousager, L. B. & Brusgaard, K., 2012.

Mutation spectrum in the large GTPase dynamin 2, and genotype-phenotype correlation in autosomal dominant centronuclear myopathy

Böhm, J., Biancalana, V., Dechene, E. T., Bitoun, M., Pierson, C. R., Schaefer, E., Karasoy, H., Dempsey, M. A., Klein, F., Dondaine, N., Kretz, C., Haumesser, N., Poirson, C., Toussaint, A., Greenleaf, R. S., Barger, M. A., Mahoney, L. J., Kang, P. B., Zanoteli, E. & Vissing, J. & 48 flere, Witting, N., Echaniz-Laguna, A., Wallgren-Pettersson, C., Dowling, J., Merlini, L., Oldfors, A., Ousager, L. B., Melki, J., Krause, A., Jern, C., Oliveira, A. S. B., Petit, F., Jacquette, A., Chaussonot, A., Mowat, D., Leheup, B., Cristofano, M., Poza Aldea, J. J., Michel, F., Furby, A., Llona, J. E. B., Van Coster, R., Bertini, E., Urtizberea, J. A., Drouin-Garraud, V., Bérout, C., Prudhon, B., Bedford, M., Mathews, K., Erby, L. A. H., Smith, S. A., Roggenbuck, J., Crowe, C. A., Brennan Spitala, A., Johal, S. C., Amato, A. A., Demmer, L. A., Jonas, J., Darras, B. T., Bird, T. D., Laurino, M., Welt, S. I., Trotter, C., Guicheney, P., Das, S., Mandel, J.-L., Beggs, A. H. & Laporte, J., 2012, I: Human Mutation. 33, 6, s. 949-59 11 s.

Mutation update for the PORCN gene

Lombardi, M. P., Bulk, S., Celli, J., Lampe, A., Gabbett, M. T., Ousager, L. B., van der Smagt, J. J., Soller, M., Stattin, E.-L., Mannens, M. A. M. M., Smigiel, R. & Hennekam, R. C., 2011, I: Human Mutation. 32, 7, s. 723-8 6 s.

Selenoproteinrelateret muskeldystrofi

Hansen, L. K., Schrøder, H. & Ousager, L. B., 2011, I: Ugeskrift for Læger. 173, 48, s. 3116-3117 2 s.

Diagnostik af Dravet syndrom

Hansen, L. K., Rasmussen, N. H. & Ousager, L. B., 22. feb. 2010, I: Ugeskrift for Læger. 172, 8, s. 626-7 1 s.

Dravets syndrom

Hansen, L. K., Rasmussen, N. H. & Ousager, L. B., 22. feb. 2010, I: Ugeskrift for Læger. 172, 8, s. 622-5 3 s.

CNTNAP2 and NRXN1 are mutated in autosomal-recessive Pitt-Hopkins-like mental retardation and determine the level of a common synaptic protein in Drosophila

Zweier, C., de Jong, E. K., Zweier, M., Orrico, A., Ousager, L. B., Collins, A. L., Bijlsma, E. K., Oortveld, M. A. W., Ekici, A. B., Reis, A., Schenck, A. & Rauch, A., 1. nov. 2009, I: American Journal of Human Genetics. 85, 5, s. 655-66 11 s.

Two siblings with microcephaly, growth retardation, cataract, hearing loss, and unusual appearance

Hansen, L. K., Bygum, A. & Ousager, L. B., 1. jul. 2009, I: Clinical Dysmorphology. 18, 3, s. 181-3 2 s.

Skin manifestations in a case of trisomy 16 mosaicism

Ousager, L. B., Brandrup, F., Andersen, C. B. & Erlandsson, A., 2006, I: British Journal of Dermatology. 154, 1, s. 172-6

Assessments of clonal composition of colorectal adenomas by fish analysis of chromosomes 1, 7, 13 and 20

Ousager, L. B., Lothe, R. A., Bardi, G., Fenger, C., Kronborg, O. & Heim, S., 2001, I: International Journal of Cancer. 92, s. 816-23

Cytogenetic analysis of Colorectal Polyps

Ousager, L. B., 1999, Odense: Syddansk Universitet. Det Sundhedsvidenskabelige Fakultet.

Cytogenetic analysis of colorectal adenomas: karyotypic comparisons of synchronous tumors

Bomme, L., Bardi, G., Pandis, N., Fenger, C., Kronborg, O. & Heim, S., 1. okt. 1998, I: Cancer Genetics and Cytogenetics. 106, 1, s. 66-71 5 s.

Allelic imbalance and cytogenetic deletion of 1p in colorectal adenomas: a target region identified between DIS199 and DIS234

Bomme, L., Heim, S., Bardi, G., Fenger, C., Kronborg, O., Brøgger, A. & Lothe, R. A., 1. mar. 1998, I: Genes, Chromosomes & Cancer. 21, 3, s. 185-94 9 s.

Fluorescence in situ hybridization of old G-banded and mounted chromosome preparations

Gerdes, A. M., Pandis, N., Bomme, L., Dietrich, C. U., Teixeira, M. R., Bardi, G. & Heim, S., 1. okt. 1997, I: Cancer Genetics and Cytogenetics. 98, 1, s. 9-15 6 s.

Cytogenetic findings in metastases from colorectal cancer

Bardi, G., Parada, L. A., Bomme, L., Pandis, N., Johansson, B., Willén, R., Fenger, C., Kronborg, O., Mitelman, F. & Heim, S., 7. aug. 1997, I: International Journal of Cancer. 72, 4, s. 604-7 3 s.

Cytogenetic comparisons of synchronous carcinomas and polyps in patients with colorectal cancer

Bardi, G., Parada, L. A., Bomme, L., Pandis, N., Willén, R., Johansson, B., Jeppsson, B., Beroukas, K., Heim, S. & Mitelman, F., 1. jan. 1997, I: British Journal of Cancer. 76, 6, s. 765-9 4 s.

Chromosome abnormalities in colorectal adenomas: two cytogenetic subgroups characterized by deletion of 1p and numerical aberrations

Bomme, L., Bardi, G., Pandis, N., Fenger, C., Kronborg, O. & Heim, S., 1. nov. 1996, I: Human Pathology. 27, 11, s. 1192-7 5 s.

Clonal karyotypic abnormalities in colorectal adenomas: clues to the early genetic events in the adenoma-carcinoma sequence

Bomme, L., Bardi, G., Pandis, N., Fenger, C., Kronborg, O. & Heim, S., 1. jul. 1994, I: Genes, Chromosomes & Cancer. 10, 3, s. 190-6 6 s.

Deletion of 1p36 as a primary chromosomal aberration in intestinal tumorigenesis

Bardi, G., Pandis, N., Fenger, C., Kronborg, O., Bomme, L. & Heim, S., 15. apr. 1993, I: Cancer Research. 53, 8, s. 1895-8 3 s.

Presse/medie

Erfaren overlæge bliver professor i genetik

Ousager, L. B.

02/09/2020

1 Mediebidrag

Fagfolk frygter slingrekurs på vejen mod personlig medicin

Ousager, L. B.

04/08/2023

1 Mediebidrag

Lillian Bomme Ousager er udnævnt til professor

Ousager, L. B.

07/09/2020

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Ny professor i genetik på SDU og OUH

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PRM / Ny professor i genetik på SDU og OUH

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TV-2 Fyn artikel 25. januar 2020. 'Nyt center skal skabe tryghed hos patienter med sjældne sygdomme'

Ousager, L. B.
25/01/2020
1 element af Mediedækning

TV-2 Fyn Nyheder 25 januar 2020. Nyt center skal skabe tryghed hos patienter med sjældne sygdomme.

Ousager, L. B.
25/01/2020
1 element af Mediedækning