

Anette Drøhse Kjeldsen
Department of Clinical Research
KI, OUH, Research unit of Oto Rhino Laryngology (Odense)
Email: anette.kjeldsen@rsyd.dk, jakjeldsen@dadlnet.dk

Publications

Skin and mucosal telangiectatic lesions in hereditary hemorrhagic telangiectasia patients

Hyldahl, S. J., El-Jaji, M. Q., Schuster, A. & Kjeldsen, A. D., 6. Jul 2022, (E-pub ahead of print) In: International Journal of Dermatology.

Fastklemt batteri i næsen hos en fireårig pige

Juul Ladegaard, P. B., Bräuner Skansing, D. & Kjeldsen, A. D., 16. May 2022, (E-pub ahead of print) In: Ugeskrift for Læger. V02220078.

Kutane kapillære malformationer med cerebral involvering

Brix, A. T. H., Tørring, P. M., Kjeldsen, A. D., Bygum, A., Schuster, A. & Nielsen, T. H., 9. May 2022, (E-pub ahead of print) In: Ugeskrift for Læger. V03210220.

A retrospective cohort study of patients with eosinophilia referred to a tertiary centre

Hougaard, M., Thomsen, G. N., Kristensen, T. K., Lindegaard, H. M., Davidsen, J. R., Hartmeyer, G. N., Kjeldsen, A. D., Martin-Iguacel, R., Maiborg, M., Assing, K., Andersen, C. L., Broesby-Olsen, S., Møller, M. B., Vestergaard, H. & Bjerrum, O. W., Apr 2022, In: Danish Medical Journal. 69, 4, A07210558.

Clinical evaluation of antibiotic regimens in patients with surgically verified parapharyngeal abscess: a prospective observational study

Klug, T. E., Andersen, C., Hahn, P., Danstrup, C. S., Petersen, N. K., Mikkelsen, S., Døssing, H., Christensen, A-L., Rusan, M., Kjeldsen, A. & Greve, T., Apr 2022, In: European Archives of Oto-Rhino-Laryngology. 279, 4, p. 2057-2067

A retrospective cohort study on European Reference Network for Rare Vascular Diseases 5 outcome measures for Hereditary Haemorrhagic Telangiectasia in Denmark

Hvelplund, T., Lange, B., Bird, S. D., Korsholm, M. & Kjeldsen, A. D., 6. Jan 2022, In: Orphanet Journal of Rare Diseases. 17, 8.

The European Rare Disease Network for HHT Frameworks for management of hereditary haemorrhagic telangiectasia in general and speciality care

Shovlin, C. L., Buscarini, E., Sabbà, C., Mager, H. J., Kjeldsen, A. D., Pagella, F., Sure, U., Ugolini, S., Torring, P. M., Suppressa, P., Rennie, C., Post, M. C., Patel, M. C., Nielsen, T. H., Manfredi, G., Lenato, G. M., Lefroy, D., Kariholu, U., Jones, B., Fiolla, A. D. & 8 others, Eker, O. F., Dupuis, O., Droege, F., Coote, N., Boccardi, E., Alsafi, A., Alicante, S. & Dupuis-Girod, S., Jan 2022, In: European Journal of Medical Genetics. 65, 1, 13 p., 104370.

Kronisk sinusitis maksillaris med en odontogen baggrund

Ghaws, S., Bakshaie Philipsen, B. & Kjeldsen, A. D., Nov 2021, In: Tandlaegebladet. 125, 11, p. 1076-1080

Microbiology of parapharyngeal abscesses in adults: in search of the significant pathogens

Klug, T. E., Greve, T., Andersen, C., Hahn, P., Danstrup, C., Petersen, N. K., Ninn-Pedersen, M., Mikkelsen, S., Pauli, S., Fuglsang, S., Døssing, H., Christensen, A. L., Rusan, M. & Kjeldsen, A., Jul 2021, In: European Journal of Clinical Microbiology and Infectious Diseases. 40, 7, p. 1461-1470

Successful treatment of massive haemoptysis in a young woman with anastomosis of right internal mammary artery to right superior pulmonary vein fistula

Bechsgaard, T., Midtgaard, A., Jakobsen, E. & Kjeldsen, A. D., 25. May 2021, In: BMJ Case Reports. 14, 5, e240739.

Screening for cerebrale arteriovenøsemalformationer ved mb. Osler

Kjær Simonsen, S., Nielsen, T. H., Dalby, R. B., Dam Fiolla, A., Lange, B., Tørring, P. M., Diaz, A. & Kjeldsen, A. D., 17. May 2021, In: Ugeskrift for Læger. 183, 19, V12200905.

Hereditary hemorrhagic telangiectasia

Berg, J. N. & Kjeldsen, A. D., 2021, *Cassidy and Allanson's Management of Genetic Syndromes*. Carey, J. C., Battaglia, A., Viskochil, D. & Cassidy, S. B. (eds.). 4. ed. Wiley, p. 475-486

Second International Guidelines for the Diagnosis and Management of Hereditary Hemorrhagic Telangiectasia

Faughnan, M. E., Mager, J. J., Hetts, S. W., Palda, V. A., Lang-Robertson, K., Buscarini, E., Deslandres, E., Kasthuri, R. S., Lausman, A., Poetker, D., Ratjen, F., Chesnutt, M. S., Clancy, M., Whitehead, K. J., Al-Samkari, H., Chakinala, M., Conrad, M., Cortes, D., Crocione, C., Darling, J. & 35 others, de Gussem, E., Derksen, C., Dupuis-Girod, S., Foy, P., Geisthoff, U., Gossage, J. R., Hammill, A., Heimdal, K., Henderson, K., Iyer, V. N., Kjeldsen, A. D., Komiyama, M., Korenblatt, K., McDonald, J., McMahon, J., McWilliams, J., Meek, M. E., Mei-Zahav, M., Olitsky, S., Palmer, S., Pantalone, R., Piccirillo, J. F., Plahn, B., Porteous, M. E. M., Post, M. C., Radovanovic, I., Rochon, P. J., Rodriguez-Lopez, J., Sabba, C., Serra, M., Shovlin, C., Sprecher, D., White, A. J., Winship, I. & Zarrabeitia, R., 15. Dec 2020, In: *Annals of Internal Medicine*. 173, 12, p. 989-1001

EUFOREA treatment algorithm for allergic rhinitis

Hellings, P. W., Scadding, G., Bachert, C., Bjermer, L., Canonica, G. W., Cardell, L. O., Carney, A. S., Constantinidis, J., Deneyer, L., Diamant, Z., Durham, S., Gevaert, P., Harvey, R., Hopkins, C., Kjeldsen, A., Klimek, L., Lund, V. J., Price, D., Rimmer, J., Ryan, D. & 11 others, Roberts, G., Sahlstrand-Johnson, P., Salmi, S., Samji, M., Scadding, G., Smith, P., Steinsvik, A., Wagenmann, M., Seys, S., Wahn, U. & Fokkens, W. J., 1. Dec 2020, In: *Rhinology*. 58, 6, p. 618-622

High output cardiac failure in 3 patients with hereditary hemorrhagic telangiectasia and hepatic vascular malformations, evaluation of treatment

Olsen, L. B., Kjeldsen, A. D., Poulsen, M. K., Kjeldsen, J. & Fialla, A. D., Dec 2020, In: *Orphanet Journal of Rare Diseases*. 15, 7 p., 334.

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, In: *Molecular Genetics and Genomic Medicine*. 8, 11, 8 p., e1498.

Real-life assessment of chronic rhinosinusitis patients using mobile technology: The mySinusitisCoach project by EUFOREA

Seys, S. F., De Bont, S., Fokkens, W. J., Bachert, C., Alobid, I., Bernal-Sprekelsen, M., Bjermer, L., Callebaut, I., Cardell, L. O., Carrie, S., Castelnovo, P., Cathcart, R., Constantinidis, J., Cools, L., Cornet, M., Clement, G., Cox, T., Delsupehe, L., Correia-de-Sousa, J., Deneyer, L. & 41 others, De Vos, G., Diamant, Z., Doulaptsi, M., Gane, S., Gevaert, P., Hopkins, C., Hox, V., Hummel, T., Hosemann, W., Jacobs, R., Jorissen, M., Kjeldsen, A., Landis, B. N., Lemmens, W., Leunig, A., Lund, V., Mariën, G., Mullol, J., Onerci, M., Palkonen, S., Proano, I., Prokopakis, E., Ryan, D., Riechelmann, H., Sahlstrand-Johnson, P., Salmi-Toppila, S., Segboer, C., Speleman, K., Steinsvik, A., Surda, P., Tomazic, P. V., Vanderveken, O., Van Gerven, L., Van Zele, T., Verfaillie, J., Verhaeghe, B., Vierstraete, K., Vlamincx, S., Wagenmann, M., Pugin, B. & Hellings, P. W., Nov 2020, In: *Allergy: European Journal of Allergy and Clinical Immunology*. 75, 11, p. 2867-2878

Granulomatous inflammation in lymph nodes of the head and neck—a retrospective analysis of causes in a population with very low incidence of tuberculosis

Flyger, T. F., Larsen, S. R. & Kjeldsen, A. D., 1. Aug 2020, In: *Immunologic Research*. 68, 4, p. 198-203

European Reference Network for Rare Vascular Diseases (VASCERN) position statement on cerebral screening in adults and children with hereditary haemorrhagic telangiectasia (HHT)

Eker, O. F., Boccardi, E., Sure, U., Patel, M. C., Alicante, S., Alsafi, A., Coote, N., Droege, F., Dupuis, O., Fialla, A. D., Jones, B., Kariholu, U., Kjeldsen, A. D., Lefroy, D., Lenato, G. M., Mager, H. J., Manfredi, G., Nielsen, T. H., Pagella, F., Post, M. C. & 8 others, Rennie, C., Sabbà, C., Suppressa, P., Toerring, P. M., Ugolini, S., Buscarini, E., Dupuis-Girod, S. & Shovlin, C. L., 29. Jun 2020, In: *Orphanet Journal of Rare Diseases*. 15, 10 p., 165.

Allergic respiratory disease care in the COVID-19 era: A EUFOREA statement

Scadding, G. K., Hellings, P. W., Bachert, C., Bjermer, L., Diamant, Z., Gevaert, P., Kjeldsen, A., Kleine-Tebbe, J., Klimek, L., Muraro, A., Roberts, G., Steinsvik, A., Wagenmann, M. & Wahn, U., May 2020, In: *World Allergy Organization Journal*. 13, 5, 4 p., 100124.

Patient-recorded benefit from nasal closure in a Danish cohort of patients with hereditary haemorrhagic telangiectasia
Andersen, J. H. & Kjeldsen, A. D., Mar 2020, In: *European Archives of Oto-Rhino-Laryngology*. 277, 3, p. 791-800

European Position Paper on Rhinosinusitis and Nasal Polyps 2020

Fokkens, W. J., Lund, V. J., Hopkins, C., Hellings, P. W., Kern, R., Reitsma, S., Toppila-Salmi, S., Bernal-Sprekelsen, M., Mullol, J., Alobid, I., Terezinha Anselmo-Lima, W., Bachert, C., Baroody, F., von Buchwald, C., Cervin, A., Cohen, N., Constantinidis, J., De Gabor, L., Desrosiers, M., Diamant, Z. & 108 others, Douglas, R. G., Gevaert, P. H., Hafner, A., Harvey, R. J., Joos, G. F., Kalogjera, L., Knill, A., Kocks, J. H., Landis, B. N., Limpens, J., Lebeer, S., Lourenco, O., Matricardi, P. M., Meco, C., O Mahony, L., Philpott, C. M., Ryan, D., Schlosser, R., Senior, B., Smith, T. L., Teeling, T., Tomazic, P. V., Wang, D., Wang, D., Zhang, L., Agius, A. M., Ahlstrom-Emanuelsson, C., Alabri, R., Albu, S., Alhabash, S., Aleksic, A., Aloulah, M., Al-Qudah, M., Alsaleh, S., Baban, M. A., Baudoin, T., Balvers, T., Battaglia, T., Bedoya, J. D., Beule, A., Bofares, K. M., Braverman, I., Brozek-Madry, E., Richard, B., Callejas, C., Carrie, S., Caulley, L., Chussi, D., de Corso, E., Coste, A., Devyani, L., El Hadi, U., Elfarouk, A., Eloy, P. H., Farrokhi, S., Felisati, G., Ferrari, M. D., Fishchuk, R., Grayson, W., Goncalves, P. M., Grdnic, B., Grgic, V., Hamizan, A. W., Heinichen, J. V., Husain, S., Ping, T. I., Ivaska, J., Jakimovska, F., Jovancevic, L., Kakande, E., Kamel, R., Karpischenko, S., Kariyawasam, H. H., Kjeldsen, A., Klimek, L., Kim, S. W., Letort, J. J., Lopatin, A., Mahdjoubi, A., Netkovski, J., Nyenbue Tshipukane, D., Obando-Valverde, A., Okano, M., Onerci, M., Ong, Y. K., Orlandi, R., Ouenoughy, K., Ozkan, M., Peric, A., Plzak, J., Prokopakis, E., Prepageran, N., Psaltis, A., Pugin, B., Raftopoulos, M., Rombaux, P., Sahtout, S., Sarafoleanu, C. C., Searyoh, K., Rhee, C. S., Shi, J., Shkoukani, M., Shukuryan, A. K., Sicak, M., Smyth, D., Snidvongs, K., Soklic Kosak, T. & Stjarne, P., 20. Feb 2020, In: *Rhinology*. 58, Suppl S29, p. 1-464

Safety of direct oral anticoagulants in patients with hereditary hemorrhagic telangiectasia

Shovlin, C. L., Millar, C. M., Droege, F., Kjeldsen, A., Manfredi, G., Suppressa, P., Ugolini, S., Coote, N., Fiella, A. D., Geithoff, U., Lenato, G. M., Mager, H. J., Pagella, F., Post, M. C., Sabbà, C., Sure, U., Topping, P. M., Dupuis-Girod, S., Buscarini, E. & VASCERN HHT, 28. Aug 2019, In: *Orphanet Journal of Rare Diseases*. 14, 8 p., 210.

Does severe bleeding in HHT patients respond to intravenous bevacizumab? Review of the literature and case series

Rosenberg, T., Fiella, A. D., Kjeldsen, J. & Kjeldsen, A. D., 1. Aug 2019, In: *Rhinology*. 57, 4, p. 242-251

Long-Term Single-Center Retrospective Follow-Up After Embolization of Pulmonary Arteriovenous Malformations Treated Over a 20-year Period: Frequency of Re-canalization with Various Embolization Materials and Clinical Outcome

Andersen, P. E., Duvnjak, S., Gerke, O. & Kjeldsen, A. D., Aug 2019, In: *Cardiovascular and Interventional Radiology*. 42, 8, p. 1102-1109

Sjældnen årsag til hævet lymfeknude på halsen hos ellers raske børn: Månedens case

Agger-Nielsen, H. E., Rosenberg, T. & Kjeldsen, A. D., Aug 2019, In: *Maanedsskrift for Almen Praksis*. 2019, 8, p. 535-542

Evaluation of day-care tonsil surgery in young children

Bruaset, I. A. M., Dahlstrøm, M., Möller, S. & Kjeldsen, A. D., 1. Mar 2019, In: *Danish Medical Journal*. 66, 3, 7 p., A5536.

Safety of thalidomide and bevacizumab in patients with hereditary hemorrhagic telangiectasia

Buscarini, E., Botella, L. M., Geithoff, U., Kjeldsen, A. D., Mager, H. J., Pagella, F., Suppressa, P., Zarrabeitia, R., Dupuis-Girod, S., Shovlin, C. L. & on behalf of VASCERN-HHT, 4. Feb 2019, In: *Orphanet Journal of Rare Diseases*. 14, 14 p., 28.

Prevention of serious infections in hereditary hemorrhagic telangiectasia: roles for prophylactic antibiotics, the pulmonary capillaries-but not vaccination

Shovlin, C., Bamford, K., Sabbà, C., Mager, H.-J., Kjeldsen, A., Droege, F., Buscarini, E., Dupuis-Girod, S. & VASCERN HHT, Feb 2019, In: *Haematologica*. 104, 2, p. e85-e86

Fordøjelseskanalens sygdomme

Schaffalitzky, O. B., Bytzer, P., Havelund, T., Kjeldsen, A. D., Kjeldsen, J. & Rumessen, J., 2019, *Medicinsk Kompendium*. de Muckadell, O. B. S., Hastrup Svendsen, J. & Vilstrup, H. (eds.). 19. ed. Munksgaard, Vol. 1. p. 939-1087

Glomuvenøse malformationer

Brix, A. T. H., Tørring, P. M., Kamaleswaran, S., Kjeldsen, A. D., Kjærskov, M. W. & Bygum, A., 2019, In: Ugeskrift for Læger. 181, 22, V10180740.

Comorbidity among HHT patients and their controls in a 20years follow-up period

Aagaard, K. S., Kjeldsen, A. D., Tørring, P. M. & Green, A., 14. Dec 2018, In: Orphanet Journal of Rare Diseases. 13, 9 p., 223.

Pulmonary arteriovenous malformations: a radiological and clinical investigation of 136 patients with long-term follow-up

Andersen, P. E., Tørring, P. M., Duvnjak, S., Gerke, O., Nissen, H. & Kjeldsen, A. D., Nov 2018, In: Clinical Radiology. 73, 11, p. 951-957

European Reference Network for Rare Vascular Diseases (VASCERN) Outcome Measures for Hereditary Haemorrhagic Telangiectasia (HHT)

Shovlin, C. L., Buscarini, E., Kjeldsen, A. D., Mager, H. J., Sabba, C., Droege, F., Geisthoff, U., Ugolini, S. & Dupuis-Girod, S., 15. Aug 2018, In: Orphanet Journal of Rare Diseases. 13, 1, 5 p., 136.

Middle ear disease in Danish toddlers attending nursery day-care – Applicability of OM-6, disease specific quality of life and predictors for middle ear symptoms

Indius, J. H., Alqaderi, S. K., Kjeldsen, A. D. & Heidemann, C. H., Jul 2018, In: International Journal of Pediatric Otorhinolaryngology. 110, p. 130-134

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, In: Angiogenesis. 21, 1, p. 146 1 p.

Hereditary haemorrhagic telangiectasia (HHT) is more than a bleeding nose, results from the Danish HHT database

Kjeldsen, A., Tørring, P. M. & Andersen, P. E., 1. Feb 2018, In: Angiogenesis. 21, 1, p. 141 1 p.

Mosaicism in a patient with HHT

Brusgaard, K., Tørring, P. M., Kjeldsen, A. D. & Ousager, L. B., 1. Feb 2018, In: Angiogenesis. 21, 1, p. 146-147

VASCERN HHT priority evaluations 2016-2017

Shovlin, C. L., Botella, L., Geisthoff, U. W., Kjeldsen, A., Mager, H. J., Pagella, F., Sabba, C., Buscarini, E. & Girod, S. D., 1. Feb 2018, In: Angiogenesis. 21, 1, p. 123 1 p.

Vascern HHT survey 2: drug registry-part 1

Buscarini, E., Botella, L. M., Geisthoff, U. W., Kjeldsen, A., Mager, H. J., Pagella, F., Suppressa, P., Girod, D. S., Shovlin, C. L. & Grp, VASCERN-HHT. W., 1. Feb 2018, In: Angiogenesis. 21, 1, p. 163 1 p.

The validity of nasal endoscopy in patients with chronic rhinosinusitis: An inter-rater agreement study

Larsen, K. L., Lange, B., Darling, P., Jørgensen, G. & Kjeldsen, A. D., Feb 2018, In: Clinical Otolaryngology. 43, 1, p. 144-150

ENG mutational mosaicism in a family with hereditary hemorrhagic telangiectasia

Tørring, P. M., Kjeldsen, A. D., Ousager, L. B. & Brusgaard, K., Jan 2018, In: Molecular Genetics & Genomic Medicine. 6, 1, p. 121-125

Odontogenic sinusitis among patients surgically treated for maxillary sinus disease

Philipsen, B. B., Ghawsi, S. & Kjeldsen, A. D., 2018, In: Rhinology Online. 1, p. 60-66

Sinogenic intracranial complications: is adalimumab a culprit?

Kofoed, M. S., Fisker, N., Christensen, A. E. & Kjeldsen, A. D., 2018, In: BMJ Case Reports. 2018, 6 p.

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, In: Clinical Case Reports. 5, 6, p. 805–808

EUFOREA Rhinology Research Forum 2016: report of the brainstorming sessions on needs and priorities in rhinitis and rhinosinusitis

Hellings, P. W., Akdis, C. A., Bachert, C., Bousquet, J., Pugin, B., Adriaensen, G., Advani, R., Agache, I., Anjo, C., Anmolsingh, R., Annoni, E., Bieber, T., Bizaki, A., Braverman, I., Callebaut, I., Castillo Vizueté, J. A., Chalermwatanachai, T., Chmielewski, R., Cingi, C., Cools, L. & 58 others, Coppije, C., Cornet, M. E., De Boeck, I., De Corso, E., De Greve, G., Doulaptsi, M., Edmiston, R., Erskine, S., Gevaert, E., Gevaert, P., Golebski, K., Hopkins, C., Hox, V., Jaeggi, C., Joos, G., Khwaja, T. S., Kjeldsen, A. D., Klimek, L., Koennecke, M., Kortekaas Krohn, I., Krysko, O., Kumar, B. N., Langdon, C., Lange, B., Lekakis, G., Levie, P., Lourijesen, E., Lund, V. J., Martens, K., Mösges, R., Mullol, J., Nyembue, T. D., Palkonen, S., Philpott, C., Aguilar-Pimentel, J. A., Poirrier, A., Pratas, A. C., Prokopakis, E., Pujols, L., Rombaux, P., Schmidt-Weber, C. B., Segboer, C., Spacova, I., Staikuniene, J., Steelant, B., Steinsvik, E. A., Teufelberger, A., Van Gerven, L. P. A., Van Gool, K., Verbrugge, R., Verhaeghe, B., Virkkula, P., Vlamincx, S., Vries-Uss, E., Wagenmann, M., Zuberbier, T., Seys, S. F. & Fokkens, W. J., 2017, In: Rhinology. 55, 3, p. 202-210

Prevalence of hereditary hemorrhagic telangiectasia in patients operated for cerebral abscess: a retrospective cohort analysis

Larsen, L., Rostgaard Marker, C., Kjeldsen, A. D. & Rom Poulsen, F., 2017, In: European Journal of Clinical Microbiology & Infectious Diseases. 36, 10, p. 1975–1980

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, In: Gene Reports. 5, p. 30-33

20-year follow-up study of Danish HHT patients-survival and causes of death

Kjeldsen, A., Aagaard, K. S., Tørring, P. M., Möller, S. & Green, A., 22. Nov 2016, In: Orphanet Journal of Rare Diseases. 11, p. 1-8 157.

JP-HHT phenotype in Danish patients with SMAD4 mutations

Jelsing, 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, In: Clinical Genetics. 90, 1, p. 55-62

The Sinonasal Outcome Test 22 score in persons without chronic rhinosinusitis

Lange, B., Thilsing, T., Baelum, J. & Kjeldsen, A. D., Apr 2016, In: Clinical Otolaryngology. 41, 2, p. 127-130

Efficacy of ESS in chronic rhinosinusitis with and without nasal polyposis: a Danish cohort study

Lind, H., Joergensen, G., Lange, B., Svendstrup, F. & Kjeldsen, A. D., 2016, In: European Archives of Oto-Rhino-Laryngology. 273, 4, p. 911-919

Laryngeal telangiectatic lesions in a patient diagnosed with Hereditary Hemorrhagic Telangiectasia

Kjeldsen, A. D., Printz, T., Slot Mehlum, C. & Grøntved, Å. M., 2016. 1 p.

Quality-of-Life Differences among Diagnostic Subgroups of Children Receiving Ventilating Tubes for Otitis Media

Heidemann, C. H., Lauridsen, H. H., Kjeldsen, A. D., Faber, C. E., Johansen, E. C. J. & Godballe, C., 1. Oct 2015, In: Otolaryngology - Head and Neck Surgery. 153, 4, p. 636-643

Do patients with chronic rhinosinusitis benefit from consultation with an ENT-doctor?

Lange, B., Thilsing, T., Baelum, J., Pedersen, O. F., Holst, R. & Kjeldsen, A. D., Jul 2015, In: Acta Oto-Laryngologica. 135, 7, p. 706-712

Treatment of Laryngeal Telangiectatic Lesions in a Patient Diagnosed with Hereditary Haemorrhagic Telangiectasia
Kjeldsen, A. D., Printz, T., Slot Mehlum, C. & Grøntved, Å. M., 11. Jun 2015, In: Journal of Otolaryngology-ENT Research. 2, 6, 00044.

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. May 2015, In: Microvascular Research. 99, p. 118-126

Prevalence of chronic rhinosinusitis in a population of patients with gastroesophageal reflux disease
Bohnhorst, I., Jawad, S., Lange, B., Kjeldsen, J., Hansen, J. M. & Kjeldsen, A. D., 1. May 2015, In: American Journal of Rhinology & Allergy. 29, 3, p. e70-e74 5 p.

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, In: Angiogenesis. 18, 4, p. 537 1 p., 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, In: Angiogenesis. 18, 4, p. 569-569 1 p., P76.

Surfactant Proteins A, B, C and D in the Human Nasal Airway: Associated with Mucosal Glands and Ciliated Epithelium but Absent in Fluid-Phase Secretions and Mucus
Gaunbaek, M. Q., Kjeldsen, A. D., Svane-Knudsen, V., Henriksen, M. L. & Hansen, S., 10. Dec 2014, In: O R L. 76, 5, p. 288-301

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, In: Clinical Genetics. 86, 2, p. 123-133 11 p.

Otitis Media and Caregiver Quality of Life: Psychometric Properties of the Modified Danish Version of the Caregiver Impact Questionnaire
Heidemann, C. H., Godballe, C., Kjeldsen, A. D., Johansen, E. C. J., Faber, C. E. & Lauridsen, H. H., 27. Jul 2014, In: Otolaryngology - Head and Neck Surgery. 151, 1, p. 142-149

Propranolol revolutionerer behandlingen af infantile subglottiske hæmangiomer
Lind, H., Kjeldsen, A. D., Schomerus, E., Schytte, S., Charabi, B., Pedersen, H. B. & Godballe, C., 2. Jun 2014, In: Ugeskrift for Laeger. 176, 11A, V66227.

Caregiver Quality of Life and Daily Functioning in Relation to Ventilating Tube Treatment
Heidemann, C. H., Lauridsen, H. H., Kjeldsen, A. D., Faber, C. E., Johansen, E. C. J. & Godballe, C., 14. Apr 2014, In: Otolaryngology - Head and Neck Surgery. 151, 2, p. 341-347

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, In: PLOS ONE. 9, 3, e90272.

Morbus Osler diagnosticeret hos 16-årig i forbindelse med en trafikulykke
Sivapalan, P., Demény, A. K., Almind, M. & Kjeldsen, A. D., 17. Feb 2014, In: Ugeskrift for Laeger. 176, 8A, p. 2-3

Acoustic rhinometry in persons recruited from the general population and diagnosed with chronic rhinosinusitis according to EPOS
Lange, B., Thilsing, T., Baelum, J., Pedersen, O. F., Holst, R. & Kjeldsen, A., 2014, In: European Archives of Oto-Rhino-Laryngology. 271, 405, p. 1961-1966

Quality of life and associated factors in persons with Chronic Rhinosinusitis in the general population: A prospective questionnaire and clinical cross-sectional study

Lange, B., Holst, R., Thilsing, T., Baelum, J. & Kjeldsen, A., Dec 2013, In: *Clinical Otolaryngology*. 38, 6, p. 474-480

The Otitis Media-6 questionnaire: psychometric properties with emphasis on factor structure and interpretability

Heidemann, C. H., Godballe, C., Kjeldsen, A. D., Johansen, E. C., Faber, C. E. & Lauridsen, H. H., 20. Nov 2013, In: *Health and Quality of Life Outcomes*. 11, 1, 201.

Cerebral abscesses among Danish patients with hereditary haemorrhagic telangiectasia

Kjeldsen, A. D., Tørring, P. M., Nissen, H. & Andersen, P. E., 2013, In: *Acta Neurologica Scandinavica*. 129, 3, p. 192-197 6 p.

Diagnosing chronic rhinosinusitis: comparing questionnaire-based and clinical-based diagnosis

Lange, B., Thilsing, T., Baelum, J., Holst, R. & Kjeldsen, A. D., 2013, In: *Rhinology*. 51, 2, p. 128-136

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, In: *Genetic Testing and Molecular Biomarkers*. 16, 12, p. 1419-1423 5 p.

Septale hæmatomer og abscesser, en follow-up undersøgelse ved Øreafdelingen på Odense Universitetshospital

Gade, S. & Kjeldsen, A. D., May 2012.

Chronic Rhinosinitis and Occupational risk factors among 20-75 year old Danes: A GA2LEN based study

Thilsing, T., Lange, B., Kjeldsen, A. D., Al-kalemji, A. & Bælum, J., 20. Mar 2012.

Chronic rhinosinitis and occupational risk factors among 20- to 75-year-old Danes-A GA(2) LEN-based study

Thilsing, T., Rasmussen, J., Lange, B., Kjeldsen, A. D., Al-Kalemji, A. & Baelum, J., 2012, In: *American Journal of Industrial Medicine*. 55, 11, p. 1037-43

Complement defects in patients with chronic rhinosinitis

Gaunsbaek, M. Q., Lange, B., Kjeldsen, A. D., Svane-Knudsen, V., Skjoedt, K., Henriksen, M. L., Nielsen, C., Palarasah, Y. & Hansen, S., 2012, In: *P L o S One*. 7, 11, p. e47383

Correlation between symptom-based and clinical-based Chronic Rhinosinitis.

Lange, B., Thilsing, T., Bælum, J., Holst, R. & Kjeldsen, A. D., 2012, In: *Rhinology*.

Embolization of pulmonary AVMs of feeding arteries less than 3 mm: reports of two cases and an 8-year follow-up without embolization

Andersen, P. E. & Kjeldsen, A. D., 2012, In: *Acta Radiologica Short Reports*. 1, 2, 5 p., 10.

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

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

Efficacy of laser treatment in HHT patients

Jørgensen, G., Lange, B., Wanscher, J. & Kjeldsen, A. D., 18. Aug 2011.

Arvelige sygdomme: Mb. Osler hereditær hæmorrhagisk teleangiæktasi (HHT)

Kjeldsen, A. D., 2011, *Lærebog i Øre-Næse-Halssygdomme og Hoved-Halskirurgi*. Ovesen, T. & von Buchwald, C. (eds.). 1 ed. Munksgaard, p. 142-143 2 p.

Efficiency of laser treatment in patients with hereditary hemorrhagic telangiectasia

Jørgensen, G., Lange, B., Wanscher, J. H. & Kjeldsen, A. D., 2011, In: *European Archives of Oto-Rhino-Laryngology*. 268, p. 1765-70 6 p.

Infektiøse/Inflammatoriske tilstande: Pyro- og mukoceler

Kjeldsen, A. D., 2011, *Lærebog i Øre-Næse-Halssygdomme og Hoved-Halskirurgi*. Ovesen, T. & von Buchwald, C. (eds.). 1 ed. Munksgaard , p. 173-174 2 p.

Kongenitte sygdomme: Deformitet af ydre næse og septum

Kjeldsen, A. D., 2011, *Lærebog i Øre-Næse-Halssygdomme og Hoved-Halskirurgi*. Ovesen, T. & von Buchwald, C. (eds.). 1 ed. Munksgaard , p. 139-140

Kongenitte sygdomme: Cyster og fistler

Kjeldsen, A. D., 2011, *Lærebog i Øre-Næse-Halssygdomme og Hoved-Halskirurgi*. Ovesen, T. (ed.). 1 ed. Munksgaard , p. 140-141 2 p.

Kongenitte sygdomme: Choanal atresi

Kjeldsen, A. D., 2011, *Lærebog i Øre-Næse-Halssygdomme og Hoved-Halskirurgi*. Ovesen, T. & von Buchwald, C. (eds.). Munksgaard , p. 141-142 2 p.

Lærebog i Øre-næse-halssygdomme og hoved- og halskirurgi

Godballe, C., Bilde, A., Kjeldsen, A. D., Petersen, C. G., Tauris, J. H., Kirkegaard, J., Lambertsen, K., Larsen, K., Nielsen, L. H., Klokke, M., Rasmussen, N., Hilberg, O., Homøe, P., Bundgaard, T., Nielsen, V. E., Grøntved, Å. M., Klug, T. E., Pedersen, U., Berg, S., Svensson, P. & 2 others, Ovesen, T. (ed.) & von Buchwald, C. (ed.), 2011, Munksgaard . 352 p.

Sygdomme med blandet genese: Epistaxis

Kjeldsen, A. D., 2011, *Lærebog i Øre-Næse-Halssygdomme og Hoved-Halskirurgi*. Ovesen, T. & von Buchwald, C. (eds.). Munksgaard , p. 180-183 4 p.

Udredning og behandling af morbus Osler

Kjeldsen, A. D., Andersen, P. E. & Tørring, P. M., 2011, In: *Ugeskrift for Læger*. 173, 7, p. 490-5 6 p.

Interventional treatment of pulmonary arteriovenous malformations

Andersen, P. E. & Kjeldsen, A. D., 28. Sep 2010, In: *World Journal of Radiology*. 2, 9, p. 339-44 6 p.

Livstruende abscedering af halsglandel tuberculose

Fleischer, J. G. & Kjeldsen, A. D., 30. Nov 2009, In: *Ugeskrift for læger*. 171, 49, p. 3620-1 1 p.

Ny metode til tonsillektomi

Kjeldsen, A. & Godballe, C., 9. Feb 2009, In: *Ugeskrift for læger*. 171, 7, p. 537; author reply 537

Homozygosity for a novel mutation in ENG discovered in a patient with Hereditary Hemorrhagic Teleangiectasia (HHT)

Kjeldsen, A. D., 2009, In: *Haematology Meeting Reports*. 4, p. 15-16 1 p.

Post-tonsillectomy hemorrhage: assessment of risk factors with special attention to introduction of coblation technique

Heidemann, C., Wallén, M., Aakesson, M., Skov, P., Kjeldsen, A. & Godballe, C., 25. Oct 2008, In: *European Archives of Oto-Rhino-Laryngology*. 266, 7, p. 1011-1015

Long-Term Follow-up After Embolization of Pulmonary Arteriovenous Malformations with Detachable Silicone Balloons

Andersen, P. E. & Kjeldsen, A. D., 2008, In: *Cardiovascular and Interventional Radiology*. 31, 3, p. 569-574 6 p.

Reposition af næsefrakturer i lokal anæstesi versus generel anæstesi: En retrospektiv patienttilfredshedsundersøgelse

Koch, K. U., Gano, L. & Kjeldsen, A. D., 11. Jun 2007, In: *Ugeskrift for læger*. 169, 24, p. 2322-5 4 p.

Behandling af hereditær hæmragisk telangiectasi. Dansk Rhinologisk Selskab.

Kjeldsen, A. D., 19. Mar 2007, In: *Ugeskrift for læger*. 169, 12, p. 1132

Long-Term Follow-up After Embolization of Pulmonary Arteriovenous Malformations with Detachable Silicone Balloons
Andersen, P. E. & Kjeldsen, A. D., 2007, In: Cardiovascular and Interventional Radiology. 31, 3, p. 569-74

Occlusion of Pulmonary Arteriovenous Malformations by Use of Vascular Plug
Andersen, P. E. & Kjeldsen, A. D., 2007, In: Acta Radiologica. 48, 5, p. 496-499 4 p.

Clinical and radiological long-term follow-up after embolization of pulmonary arteriovenous malformations
Andersen, P. E. & Kjeldsen, A. D., 1. Oct 2005, In: Cardiovascular and Interventional Radiology. 29, 1, p. 70-74

Brain abscess and hypoxia in a previously healthy young female.
Gaini, S., Kjeldsen, A. D. & Andersen, P. J., 2005, In: Scandinavian Journal of Infectious Diseases. 37, p. 301-303

Clinical symptoms according to genotype among patients with hereditary haemorrhagic telangiectasia.
Kjeldsen, A. D., Kjeldsen, A. D., Moeller, T. R., Brunsgaard, K., Vase, P. & Andersen, P. E., 2005, In: Journal of Internal Medicine. 258, p. 349-355

Clinical symptoms according to genotype among patients with hereditary haemorrhagic telangiectasia.
Kjeldsen, A. D., Møller, T. R., Brunsgaard, K., Vase, P. & Andersen, P. E., 2005, In: Journal of Internal Medicine. 258, p. 349-355

Clinical symptoms according to genotype amongst patients with hereditary haemorrhagic telangiectasia
Kjeldsen, A. D., Møller, T. R., Brusgaard, K., Vase, P. & Andersen, P. E., 2005, In: Journal of Internal Medicine. 258, 4, p. 349-355

Clinical symptoms according to genotype HHT1 or HHT2 in patients with Hereditary Haemorrhagic Telangiectasia (HHT).
Kjeldsen, A. D., Møller, T. R., Brusgaard, K., Vase, P. & Andersen, P. E., 2005, In: Journal of Internal Medicine. 258, 4, p. 349-355

Selective embolization in the treatment of intractable epistaxis
Andersen, P. J., Kjeldsen, A. D. & Nepper-Rasmussen, J., 2005, In: Acta Oto-Laryngologica. 125, 3, p. 293-297

Selective embolization in treatment of intractable epistaxis.
Andersen, P. J., Kjeldsen, A. D. & Nepper-Rasmussen, J., 2005, In: Acta Oto-Laryngologica. 125, p. 293-297

Månedens billede
Kjeldsen, A. D. & Marker, P., 23. Aug 2004, In: Ugeskrift for læger. 166, 35, p. 3010

Mutations in endoglin and in activin receptor-like kinase 1 among Danish patients with hereditary haemorrhagic telangiectasia
Brusgaard, K., Kjeldsen, A. D., Poulsen, L., Moss, H., Vase, P., Rasmussen, K., Kruse, T. A. & Hørder, M., Jan 2004, In: Clinical Genetics. 66, 6, p. 556-61 6 p.

Behandling af invert papillom med endonasal endoskopisk sinuskirurgi
Andersen, P. J., Kjeldsen, A. D. & Pedersen, A. T., 9. Sep 2002, In: Ugeskrift for læger. 164, 37, p. 4283-7 4 p.

Mutations in the ALK-1 gene and the phenotype of hereditary hemorrhagic telangiectasia in two large Danish families
Kjeldsen, A. D., Brusgaard, K., Poulsen, L., Kruse, T., Rasmussen, K., Green, A. & Vase, P., 2001, In: American Journal of Medical Genetics. Part B: Neuropsychiatric Genetics. 98, 4, p. 298-302 5 p.

Perkutan transluminal embolisation af pulmonale arteriovenøse malformationer
Andersen, P. E., Kjeldsen, A. D., Oxhøj, H. & Vase, P., 2001, In: Ugeskrift for læger. 163, 7, p. 925-8 4 p.

Perkutan transluminal embolisering af pulmonale arteriovenøse malformationer

Andersen, P. E., Kjeldsen, A. D., Oxhøj, H. & Vase, P., 2001, In: Ugeskrift for læger. 163, p. 925

Pulmonale arteriovenøse malformationer: Moderne behandlingsprincipper

Andersen, P. E. & Kjeldsen, A. D., 2001, In: Ugeskrift for læger. 163, p. 4398-4401

Pulmonale arteriovenøse malformationer, moderne behandlingsprincipper

Andersen, P. E. & Kjeldsen, A. D., 2001, In: Ugeskrift for læger. 163, 33, p. 4398-4401

Tympanoplasty with ionomeric cement

Kjeldsen, A. D. & Grøntved, A. M., 1. Jan 2000, In: Acta Oto-Laryngologica. Supplement. 543, p. 130-1 2 p.

Concerning the original article of C. Weik and L. Greiner, 'The liver in hereditary hemorrhagic telangiectasia (Weber-Rendu's disease)'

Kjeldsen, A. D. & Kjeldsen, J., 2000, In: Scandinavian Journal of Gastroenterology. 35, 7, p. 784

Diagnostic criteria for hereditary hemorrhagic telangiectasia (Rendu-Osler-Weber syndrome)

Shovlin, C. L., Guttmacher, A. E., Buscarini, E., Faughnan, M. E., Hyland, R. H., Westermann, C. J., Kjeldsen, A. D. & Plauchu, H., 2000, In: American Journal of Medical Genetics. Part B: Neuropsychiatric Genetics. 91, 1, p. 66-7 2 p.

Gastrointestinal bleeding in patients with hereditary hemorrhagic telangiectasia

Kjeldsen, A. D. & Kjeldsen, J., 2000, In: American Journal of Gastroenterology. 95, 2, p. 415-8 4 p.

Hereditaer haemoragisk telangiectasi. Et populations-baseret studium af praevalens og mortalitet blandt danske HHT-patienter

Kjeldsen, A. D., Vase, P. & Green, A., 2000, In: Ugeskrift for Laeger. 162, 25, p. 3597-601 5 p.

Månedens billede

Kjeldsen, A. D., Andersen, P. E. & Oxhøj, H., 2000, In: Ugeskrift for Laeger. 162, 25, p. 3618

Prevalence of pulmonary arteriovenous malformations (PAVMs) and occurrence of neurological symptoms in patients with hereditary haemorrhagic telangiectasia (HHT)

Kjeldsen, A. D., Oxhøj, H., Andersen, P. E., Green, A. & Vase, P., 2000, In: Journal of Internal Medicine. 248, 3, p. 255-62 8 p.

Screening for pulmonary arteriovenous malformations: contrast echocardiography versus pulse oximetry

Oxhøj, H., Kjeldsen, A. D. & Nielsen, G., 2000, In: Scandinavian Cardiovascular Journal. 34, 3, p. 281-285

Hereditary haemorrhagic telangiectasia: a population-based study of prevalence and mortality in Danish patients

Kjeldsen, A. D., Vase, P. & Green, A., 1999, In: Journal of Internal Medicine. 245, 1, p. 31-39 9 p.

Percutaneous transluminal treatment of pulmonary arteriovenous malformations

Andersen, P. E., Kjeldsen, A. D., Oxhøj, H. & Vase, P., 1999, In: J. Intervent. Radiol.. 14, p. 164-70

Pulmonary Arteriovenous Malformations: Screening Procedures and Pulmonary Angiography in Patients With Hereditary Hemorrhagic Telangiectasia

Kjeldsen, A. D., Oxhøj, H., Andersen, P. E., Elle, B., Jacobsen, J. P. & Vase, P., 1999, In: Chest. 116, 2, p. 432-439 8 p.

Embolotherapy for pulmonary arteriovenous malformations in patients with hereditary hemorrhagic telangiectasia (Rendu-Osler-Weber syndrome)

Andersen, P. E., Kjeldsen, A. D., Oxhøj, H., Vase, P. & White, R. I., Nov 1998, In: Acta Radiologica. 39, 6, p. 723-6 4 p.

Embolotherapy for pulmonary arteriovenous malformations in patients with hereditary hemorrhagic telangiectasia (Rendu-Osler-Weber syndrome)

Andersen, P. E., Kjeldsen, A. D., Oxhøj, H., Vase, P. & White, R. I., 1998, In: Acta Radiologica. 39, p. 723-6

Perkutan transluminal embolisering af pulmonale arteriovenøse misdannelser

Kjeldsen, A. D., Andersen, P. E., Oxhøj, H. & Vase, P., 1998, In: Ugeskrift for Læger. 160, 10, p. 1465-9 5 p.

Hereditary hemorrhagic telangiectasia

Kjeldsen, A. D., Vase, P. & Oxhøj, H., 1996, In: The New England Journal of Medicine. 334, 5, p. 331-2 2 p.

Group specific component in serum and otosclerosis: no association

Kjeldsen, A. D., Vase, P., Thymann, M., Green, A. & Morling, N., 1994, In: Acta Oto-Laryngologica. 114, 3, p. 303-4 2 p.

Mephenytoin and sparteine oxidation: genetic polymorphisms in Denmark

Kjeldsen, A. D., Bathum, L., Brøsen, K. & Gram, L. F., 1989, In: British Journal of Clinical Pharmacology. 27, 5, p. 620-5 6 p.

Press clippings

Det Sundhedsvidenskabelige Fakultet

Anette Drøhse Kjeldsen

13/01/2015

1 item of Media coverage

Det Sundhedsvidenskabelige Fakultet

Anette Drøhse Kjeldsen

13/01/2015

1 item of Media coverage

Det Sundhedsvidenskabelige Fakultet

Anette Drøhse Kjeldsen

13/01/2015

1 item of Media coverage

Det Sundhedsvidenskabelige Fakultet

Anette Drøhse Kjeldsen

13/01/2015

1 item of Media coverage

Måling af lægers kompetencer

Anette Drøhse Kjeldsen

29/04/2019

1 Media contribution

Activities

Allergisk rhinit

Anette Drøhse Kjeldsen (Lecturer)

14. Oct 2015

University of Southern Denmark

Anette Drøhse Kjeldsen (Visiting lecturer)

28. Sep 2015 → 29. Sep 2015

Excel kursus

Anette Drøhse Kjeldsen (Participant)
8. Sep 2015

11th International Scientific HHT Meeting

Anette Drøhse Kjeldsen (Participant)
9. Jun 2015 → 15. Jun 2015

Årsmøde i Dansk Rhinologisk Selskab

Anette Drøhse Kjeldsen (Participant)
26. Mar 2015

St. Vincent's Medical Center

Anette Drøhse Kjeldsen (Visiting researcher)
26. Nov 2012 → 30. Nov 2012

Udredning og behandling af patienter med Mb Osler, resultater fra Database og præsentation af pakkeforløb HHT centeret OUH

Anette Drøhse Kjeldsen (Lecturer)
25. May 2012

XXXI Congress of the Nordic Association of Otolaryngology

Anette Drøhse Kjeldsen (Participant)
18. Aug 2011 → 20. Aug 2011

XXXI Congress of the Nordic Association of Otolaryngology

Anette Drøhse Kjeldsen (Participant)
17. Aug 2011 → 20. Aug 2011

Årsmøde i Dansk Rhinologisk Selskab

Anette Drøhse Kjeldsen (Participant)
26. May 2011 → 27. May 2011

9th HHT Scientific Conference

Anette Drøhse Kjeldsen (Participant)
20. May 2011 → 25. May 2011

Dansk rhinologisk selskab

Anette Drøhse Kjeldsen (Visiting researcher)
27. Apr 2011 → 29. Apr 2011

Årsmøde i Dansk Selskab for Otolaryngologi - Hoved & Halskirurgi

Anette Drøhse Kjeldsen (Participant)
14. Apr 2011 → 15. Apr 2011

8th HHT scientific conference

Anette Drøhse Kjeldsen (Participant)
28. May 2009 → ...

Young scientist award - 8th HHT Scientific Conference (External organisation)

Anette Drøhse Kjeldsen (Member)
28. May 2009

Evaluering af kronisk rhinosinuit med Sinonasal Outcome Test 22

Anette Drøhse Kjeldsen (Speaker)

15. May 2009

Rhinology (Journal)

Anette Drøhse Kjeldsen (Editor)

1. Jan 2009

Euro HHT (External organisation)

Anette Drøhse Kjeldsen (Member)

22. Oct 2008 → ...

Week of the Face

Anette Drøhse Kjeldsen (Lecturer)

25. Aug 2008

Europæisk Rhinology Society Congress

Anette Drøhse Kjeldsen (Participant)

15. Jun 2008 → 19. Jun 2008

Dansk Rhinologisk Selskabs årsmøde

Anette Drøhse Kjeldsen (Participant)

8. May 2008 → 9. May 2008

Dansk rhinologisk selskab (External organisation)

Anette Drøhse Kjeldsen (Member)

1. May 2008 → ...

DSOHH årsmøde

Anette Drøhse Kjeldsen (Participant)

25. Apr 2008 → 26. Apr 2008

EuroHHT møde

Anette Drøhse Kjeldsen (Participant)

23. Apr 2008 → 25. Apr 2008

Medical advisory board international HHT organization (External organisation)

Anette Drøhse Kjeldsen (Member)

1. Jan 2007 → ...