



**Socialstyrelsen**

**Litteraturliste til temaet  
Børn og unge som  
pårørende**

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**Viden til gavn**

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# Litteraturliste til temaet Børn og unge som pårørende

## Baggrund for søgning

År for søgning: august 2021

Tidsafgrænsning: 2016-2021

Geografisk afgrænsning: Nordisk og internationalt

## Den samlede litteraturliste

Aldridge, J. (2018). Where are we now? Twenty-five years of research, policy and practise on young carers. *Critical Social Policy*, Vol. 38(1): 155-165.

Amilon, A. et al. (2021). *Mennesker med handicap: Hverdagsliv og levevilkår 2020*. Kbh.: VIVE.

Berardini, Y. et al. (2021). Unfolding What Self-Compassion Means in Young Carers' Lives. *Child and Adolescent Social Work Journal*, Vol. 38: 533-545.

Björgvinsdóttir, K. & Halldórsdóttir, S. (2014). Silent, invisible and unacknowledged: Experiences of young caregivers of single parents diagnosed with multiple sclerosis. *Scandinavian Journal of Caring Sciences*, Vol. 28(1): 38-48.

Bogosian, A. et al. (2011). How do adolescents adjust to their parent's multiple sclerosis?: An interview study. *British Journal of Health Psychology*, Vol. 16(2): 430-444.

Chen, C. Y.-C. & Panebianco, A. (2020). Physical and psychological conditions of parental chronic illness, parentification and adolescent psychological adjustment. *Psychology & Health*, Vol. 35(9): 1075-1094.

Cunningham, L. C et al. (2017). A qualitative evaluation of an innovative resilience-building camp for young carers. *Child and Family Social Work*, Vol. 22(2): 700-710.

Elliott, L. et al. (2020). A systematic review of somatic symptoms in children with a chronically ill family member. *Psychosomatic Medicine*, Vol. 82(4): 366-376.

Eriksson, M. (2018). *Användning av BRA- Barns rätt som anhöriga*. Eskilstuna: Mälardalens Högskola.

Finch, J. & Mason, J. (1993). *Negotiating Family Responsibilities*. London: Routledge.

Heins, Agneta Berghamre (2019). *Barn som anhöriga: Alla barn ska få möjlighet till hälsa och personlig utveckling*. Kalmar: Nationellt kompetenscentrum anhöriga.

Hendricks, B. A. et al. (2021). How far have we come? An updated scoping review of young carers in the US. *Child & Adolescent Social Work Journal*, Vol. 38(5): 491-504.

Hendricks, B. A. et al. (2021). Parentification among young carers: A concept analysis. *Child & Adolescent Social Work Journal*, Vol. 38(5): 519-531.

Iezzoni, L. I. et al. (2016). Adults' Recollection and Perceptions of Childhood Caregiving to a Parent with Significant Physical Disability. *Disability and Health Journal*, Vol. 9(2): 208-217.

Ireland, M. J. & Pakenham, K. I. (2010). Youth adjustment to parental illness or disability: The role of illness characteristics, caregiving, and attachment. *Psychology, Health & Medicine*, Vol. 15(6): 632-645.

- Jørgensen, S. E. et al. (2019). Trivsel og hverdagsliv blandt børn og unge som pårørende og efterladte. Kbh.: Statens Institut for Folkesundhed, SDU.
- Jørgensen, S. E. et al. (2021). Identifying and Characterizing Children of Parents with a Serious Illness Using Survey and Register Data. *Clinical Epidemiology*, Vol. 13: 253-263.
- Jørgensen, S. E. et al. (2021). Why Do Some Adolescents Manage Despite Parental Illness? Identifying Promotive Factors. *Journal of Adolescent Health*, Vol. 69 (2): 335-341.
- Kjoelaas, S. et al. (2020). The Ripple Effect: A Qualitative Overview of Challenges When Growing Up in Families Affected by Huntington's Disease. *Journal of Huntington's Disease*, Vol. 9(2): 129 -141.
- Knight, T. (2018). Social Identity in Hearing Youth who have Deaf Parents. *International Journal of Business and Social Science*, Vol. 9(9): 1-12.
- Knutsson, S. et al. (2017). Children as relatives to a sick parent: Healthcare professionals' approaches. *Nordic Journal of Nursing Research*, Vol. 37(2): 61-69.
- Kvalificeringsgruppe (2021). *Møde i kvalificeringsgruppen. 01/11/2021 og 04/11/2021*. Onlinemøde. Odense: Socialstyrelsen.
- Leu, A. et al. (2018). Young carers and young adult carers in Switzerland: Caring roles, ways into care and the meaning of communication. *Health & Social Care in the Community*, Vol. 26(6): 925-934.
- Manskow, U. S. et al. (2015). Factors Affecting Caregiver Burden 1 Year After Severe Traumatic Brain Injury: A Prospective Nationwide Multicenter Study. *The Journal of Head Trauma Rehabilitation*, Vol. 30(6): 411-423.
- Mason, J. (1996). Gender, Care and Sensibility in Family and Kin Relations, I: Holland, J. & Adkins, L., Sex, Sensibility and the Gendered Body (s. 15-36). London: Palgrave MacMillan.
- Masterson-Algar, P. & Williams, S. (2020). "Thrown into the deep end": Mapping the experience of young people living in a family affected by a neurological condition. *Qualitative Health Research*, Vol. 30(5): 717-729.
- Mattsson, Å.L. (2018). *Guide för BRA samtal*. Stockholm: Stiftelsen Allmänna Barnhuset.
- Mauseth, T. & Hjälmhult, E. (2016). Adolescents' experiences on coping with parental multiple sclerosis: A grounded theory study. *Journal of Clinical Nursing*, Vol. 25(5-6): 856-865.
- McGibbon, M. et al. (2019). Young Carers in Northern Ireland: Perceptions of and Responses to Illness and Disability within the Family. *British Journal of Social Work*, Vol. 49(5): 1162-1179.
- Merry, S. et al. (2004). A randomized placebo-controlled trial of a school-based depression prevention program. *Journal of the American Academy of Child & Adolescent Psychiatry*, Vol. 43 (5): 538-547.
- Moberg, J.Y. et al. (2017). Striving for balance between caring and restraint: Young adults' experiences with parental multiple sclerosis. *Journal of Clinical Nursing*, Vol. 26(9-10): 1363-1374.
- Morley, D. & Jenkinson, C. (2012). The Importance of Recognising Depression in Adolescents Affected by Parental Illness. *Psychology*, Vol. 3(9A): 756-757.
- Norge Helse- og omsorgsdepartementet (2020). Vi – de pårørende: Regjeringens pårørendestrategi og handlingsplan. Oslo: Departementenes sikkerhets- og serviceorganisasjon.

- Pakenham K. I. & Cox, S. (2012). The nature of caregiving in children of a parent with multiple sclerosis from multiple sources and the associations between caregiving activities and youth adjustment over time. *Psychology & Health*, Vol. 27(3): 324-346.
- Pakenham, K. I. & Cox, S. (2014). Comparisons between youth of a parent with MS and a control group on adjustment, caregiving, attachment and family functioning. *Psychology & Health*, Vol. 29(1): 1-15.
- Pakenham, K. I. & Cox, S. (2015). The effects of parental illness and other ill family members on youth caregiving experiences. *Psychology & Health*, Vol. 30(7): 857-878.
- Pakenham, K. I. & Cox, S. (2018). Effects of Benefit Finding, Social Support and Caregiving on Youth Adjustment in a Parental Illness Context. *Journal of Child and Family Studies*, Vol. 27(8): 2491-2506.
- Razaz, N. et al. (2014). Children and adolescents adjustment to parental multiple sclerosis: A systematic review. *BMC Neurology*, Vol. 14: ArtID: 107.
- Rivet-Duval, E. et al. (2011). Preventing adolescent depression in Mauritius: A universal school-based program. *Child and Adolescent Mental Health*, Vol. 16(2): 86-91.
- Shochet, I. M. & Ham, D. (2004). Universal School-based Approaches to Preventing Adolescent Depression: Past Findings and Future Directions of the Resourceful Adolescent Program. *International Journal of Mental Health Promotion*, Vol. 6(3): 17-25.
- Shochet, I. M. et al. (2001). The efficacy of a universal school-based program to prevent adolescent depression. *Journal of Clinical Child Psychology*, Vol. 30 (3): 303-315.
- Sieh, D. S. et al. (2010). Problem Behavior in Children of Chronically Ill Parents: A Meta-Analysis. *Clinical Child and Family Psychology Review*, Vol. 13(4): 384-397.
- Sieh, D. S. et al. (2012). Risk factors for problem behavior in adolescents of parents with a chronic medical condition. *European Child & Adolescent Psychiatry*, Vol. 21(8): 459-471.
- Socialstyrelsen (2020). *Stärkt stöd till barn som anhöriga: Slutrapport från regeringsuppdrag 2017-2020*. Stockholm: Socialstyrelsen.
- Socialstyrelsen (2021). *Interne drøftelser ifm. udarbejdelse af temaet Børn og unge som pårørende. Interne drøftelser – ikke publiceret*. Handicap. Odense: Socialstyrelsen.
- Stamatopoulos, V. (2015). One million and counting: The hidden army of young carers in Canada. *Journal of Youth Studies*, Vol. 18(6): 809-822.
- Stiftelsen Allmänna Barnhuset (2021). *Rapport till FN:s kommitté för barnets rättigheter*. Stockholm: Stiftelsen Allmänna Barnhuset.
- Wepf, H. et al. (2021). Pathways to Mental Well-Being in Young Carers: The Role of Benefit Finding, Coping, Helplessness, and Caring Tasks. *Journal of Youth and Adolescence*, Vol. 50(9): 1911-1924.
- Wind, G. & Jørgensen, G. K. (2020). "It has been fun. Super-duper fantastic": Findings from a Danish respite programme to support young carers. *Health and Social Care in the Community*, Vol. 28(1): 100-109.



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