Canadian Journal of Bioethics Revue canadienne de bioéthique



Review of: Kaposy C. (2018) Choosing Down Syndrome

Meghan Chevalier

Volume 2, Number 2, 2019

URI: https://id.erudit.org/iderudit/1058145ar DOI: https://doi.org/10.7202/1058145ar

See table of contents

Publisher(s)

Programmes de bioéthique, École de santé publique de l'Université de Montréal

ISSN

2561-4665 (digital)

Explore this journal

Cite this review

Chevalier, M. (2019). Review of [Review of: Kaposy C. (2018) Choosing Down Syndrome]. *Canadian Journal of Bioethics / Revue canadienne de bioéthique*, 2(2), 21–22. https://doi.org/10.7202/1058145ar Article abstract

With the advent of Non-Invasive Prenatal Testing, Chris Kaposy believes that more people should choose to parent children with Down Syndrome. Kaposy advocates for the Social Disability Model and recommends a normative pragmatic approach as standard. He makes use of both quantitative and qualitative evidence to support his position.

All Rights Reserved © Meghan Chevalier, 2019



érudit

This document is protected by copyright law. Use of the services of Érudit (including reproduction) is subject to its terms and conditions, which can be viewed online.

https://apropos.erudit.org/en/users/policy-on-use/

This article is disseminated and preserved by Érudit.

Érudit is a non-profit inter-university consortium of the Université de Montréal, Université Laval, and the Université du Québec à Montréal. Its mission is to promote and disseminate research.

https://www.erudit.org/en/

COMPTE RENDU / REVIEW

Review of: Kaposy C. (2018) *Choosing Down Syndrome*

Meghan Chevalier¹

Résumé

Avec l'avènement du test prénatal non invasif, Chris Kaposy croit que plus de gens devraient choisir d'élever des enfants atteints du syndrome de Down. Kaposy plaide en faveur du modèle de handicap social et recommande une approche normative pragmatique comme norme. Il utilise des données quantitatives et qualitatives pour étayer sa position.

Mots-clés

(c) (i)

trisomie 21, test prénatal, avortement sélectif

Abstract

With the advent of Non-Invasive Prenatal Testing, Chris Kaposy believes that more people should choose to parent children with Down Syndrome. Kaposy advocates for the Social Disability Model and recommends a normative pragmatic approach as standard. He makes use of both quantitative and qualitative evidence to support his position.

Keywords

Down syndrome, prenatal testing, selective termination

Choosing Down Syndrome [1] by Chris Kaposy situates itself within a modern era of prenatal testing. The book is motivated by the advent and proliferation of Non-Invasive Prenatal Testing (NIPT), which seems to be occurring without sufficient reflection. Kaposy questions the potential repercussions of widespread use of NIPT in the larger framework of prenatal testing for Down syndrome and selective termination. Choosing Down Syndrome puts forth thorough and convincing arguments for choosing to parent children with Down syndrome, addressing previously published sources on the topic. Kaposy clearly outlines the structure of each chapter in the book's introduction, and what Kaposy intends to argue is clear.

Choosing Down Syndrome begins by establishing itself within a context of ongoing debates regarding the ethics of prenatal genetic testing for Down syndrome. NIPT offers a low-risk, highly accurate screen for Down syndrome. Ultimately, Kaposy suggests that more people should choose to parent a child with Down syndrome following a prenatal diagnosis, rather than terminate their pregnancy. To support this claim, Kaposy begins with an analysis of several autobiographical accounts written by parents of children with Down syndrome. He subsequently presents empirical research about families of children with Down syndrome. He subsequently presents empirical research about families of children with Down syndrome, while discrediting suggestions that the happiness that these families experience is due to adaptive preferences, i.e., the preferences that people develop based on a limited amount of potentially oppressive and less than ideal options. Kaposy then discusses the morality of bringing into the world a child with a disability, as well as the morality of selective termination. The penultimate chapter considers how the identity of a person with Down syndrome is perceived by society, and Kaposy recommends a normative pragmatic approach as standard. This means that we should employ the Social Disability Model (the perspective that disability results from the failure of society to accommodate individual needs, rather than from the individual themselves) while recognizing their disability as a medical condition when it is in their best interest, such as in therapeutic or educational contexts. The concluding chapter of the book examines the capitalist influence on selective termination. Kaposy argues that biases against people with Down syndrome result from their difficulty in achieving the type of monetary success that is valued in a capitalist society.

The primary strength of Kaposy's arguments comes from his decision to use both qualitative and quantitative supporting evidence. The qualitative analysis of autobiographical accounts written by parents of children with Down syndrome and quantitative empirical findings are mutually supportive. Kaposy includes autobiographical accounts (Chapter 3) before addressing empirical research statistics (Chapter 4), emphasizing the importance of these autobiographical accounts and avoiding this chapter being read through solely a statistical lens. The points raised in Chapter 3 are later supported by statistics, but the order in which they are presented allows the autobiographical accounts to be read independently as legitimate and valuable evidence.

After discussing and evaluating autobiographical accounts and empirical statistics, Kaposy considers questions of morality with respect to continuing pregnancies and selective termination. He clarifies and questions potential motives for selective termination. Kaposy writes with a nuanced understanding of context. For example, while some could view legislation that has been passed to promote accessibility and inclusivity as indicators that society is progressing away from biased attitudes, Kaposy discusses the presence of informal bias. Further, when addressing previously published literature, he examines and critiques their definitions of important concepts, such as disability, well-being and quality of life. Kaposy effectively challenges assumptions about people with disabilities, simplistic arguments, and arguments based on intuition in other works. In a well-rounded approach, *Choosing Down Syndrome* calls into question both individual and structural issues that contribute to the bias toward people with Down syndrome.

Throughout *Choosing Down Syndrome*, Kaposy writes with transparency. He acknowledges a perceived conflict of interest, the fact that he parents a son with Down syndrome, and handles potential criticism well, demonstrating that this perception is a by-product of the bias he is arguing exists. Further, he admits the faults of the Social Disability Model to which he often alludes. From my perspective, these critiques do not weaken Kaposy's arguments, but rather demonstrate that he has thoroughly considered the validity of his arguments and understands the social and academic context in which he is writing.



En collaboration avec / In collaboration with

While Kaposy acknowledges disabilities other than Down syndrome and explains that they are beyond the scope of this book, a broader context of prenatal testing and disability would lead to a better understanding of populations with disabilities, as well as the Down syndrome advantage, that appears when families of children with Down syndrome function better than families of children with other disabilities. Though passages are occasionally wordy, Choosing Down Syndrome is written in accessible language. A shorter, practical and therefore more accessible resource for prospective parents would be a welcome companion to Choosing Down Syndrome.

Conflicts of Interest

None to declare

Conflit d'intérêts Aucun à déclarer

Édition/Editors: Aliya Affdal & Patrick Gogognon

Affiliations

¹ Schulich School of Medicine, Western University, London, Ontario, Canada

Correspondance / Correspondence: Meghan Chevalier, meghanchevalier01@gmail.com

Reçu/Received: 22 Oct 2018 Publié/Published: 12 Mar 2019

Les éditeurs suivront les recommandations et les procédures décrites The editors follow the recommendations and procedures outlined in dans le Code of Conduct and Best Practice Guidelines for Journal Editors de COPE. Plus précisément, ils travaillent pour s'assurer des plus hautes normes éthiques de la publication, y compris l'identification et la gestion des conflits d'intérêts (pour les éditeurs et pour les auteurs), la juste évaluation des manuscrits et la publication of manuscripts, and the publication of manuscripts that meet the de manuscrits qui répondent aux normes d'excellence de la revue.

the COPE Code of Conduct and Best Practice Guidelines for Journal Editors. Specifically, the editors will work to ensure the highest ethical standards of publication, including: the identification and management of conflicts of interest (for editors and for authors), the fair evaluation journal's standards of excellence.

References

1. Kaposy C. Choosing Down Syndrome: Ethics and New Prenatal Testing Technologies. The MIT Press; 2018