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Valerie J. Page and E. Wesley Ely
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Delirium in Critical Care

Second Edition

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Delirium in critical care

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We dedicate this book to our fathers,
both of whom suffered delirium during critical illness.

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[More information](#)

CONTENTS

<i>Foreword to the first edition</i>	<i>page ix</i>
David K. Menon	
<i>Foreword to the second edition</i>	xi
Alasdair M. J. MacLulich	
<i>Delirium, a patient testimony</i>	xiv
1. What is delirium in critical care?	1
2. How common is delirium in critical care?	14
3. What does delirium look like in critical care?	27
4. Delirium in critical care: how does it happen?	44
5. Delirium: what causes it? Risk factors	68
6. Delirium in critical care: why is it important?	94
7. Delirium in critical care: monitoring tools	109
8. How to prevent delirium?	140
9. Treatment of delirium in critical care	157
10. Mental capacity and restraints	186
11. End-of-life care	197
12. What is the future?	208
<i>Selected references</i>	214
<i>Index</i>	216

Cambridge University Press

978-1-107-43365-6 - Delirium in Critical Care: Second Edition

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Frontmatter

[More information](#)

Cambridge University Press
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Frontmatter
[More information](#)

FOREWORD TO THE FIRST EDITION

The identification of delirium as an important entity in acutely and critically ill patients has been one of major advances in intensive care over the last decade. There is increasing recognition that the condition has an important impact on morbidity, health economics and patient outcome, not just in critical care, but also in the perioperative period, during acute medical illness, and at the end of life. However, there has also been a realization that the condition is under diagnosed, and that its prevention and treatment are frequently neglected.

Given this context, this book is a welcome resource for clinicians who are involved in treating patients who are at risk of delirium or require treatment for the condition. The authors are practising clinicians with complementary backgrounds in critical care. Professor Wesley Ely is perhaps the best recognized expert in this field worldwide, whose publications have put delirium on the critical care agenda. Dr Valerie Page runs a busy general intensive care unit and brings her experience of everyday critical care to the problem, along with knowledge of the background literature. While this book does provide some information on the clinical science and neurobiology underpinning the condition, this is not its main attraction, and there is a refreshing candor about the

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Valerie J. Page and E. Wesley Ely
Frontmatter
[More information](#)

x Foreword to the first edition

substantial and large lacunae in knowledge about delirium. Its great strength lies in its practicality, and in the robust clinical sense that it displays in guiding the practising ICU doctor or nurse through the process of detecting, classifying, quantifying, preventing and treating delirium. The resource that it provides should make individual clinicians and ICU teams more aware of the condition, and in doing this, could help improve patient outcomes.

David K. Menon
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Valerie J. Page and E. Wesley Ely
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[More information](#)

FOREWORD TO THE SECOND EDITION

As authors Valerie Page and Wes Ely remind us in their introduction of Chapter 1, delirium is an ‘an acute episode of brain failure’. As with acute failures of other organs, delirium is ‘common . . . dangerous . . . even life-threatening’. Yet delirium has lagged behind the other major organ failure syndromes in its status in routine clinical practice. This book is an important resource in addressing this gap through providing a highly user-friendly distillate of the best in clinical practice and research.

Valerie Page and Wes Ely are practising clinicians and researchers. Alongside her clinical work as an intensivist in a busy general hospital in Watford, Dr Page has contributed several important trials to the field. She also founded and edits *Annals of Delirium*, the newsletter of the European Delirium Association, and is an active member of the Association’s board. Professor Ely and his team at Vanderbilt University have produced the world’s largest body of globally influential studies on delirium care in the ICU. Professor Ely is also an exceptionally energetic advocate for the well-being of patients with delirium through his multiple international lectures and site visits, and his support of the work of the American Delirium Society.

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Valerie J. Page and E. Wesley Ely
Frontmatter
[More information](#)

xii Foreword to the second edition

This book is comprehensive and all key areas are covered. Two highlights to mention here are detection, and the appropriate use of drugs. Though it is axiomatic that optimal care cannot be provided without a diagnosis, gross underdetection of delirium is a persisting challenge in clinical practice. The authors tackle both the *why* and the *how* of detection. Evidence confirming the high prevalence of ICU delirium is summarized, and major findings from the new BRAIN-ICU study on outcomes of ICU delirium from Professor Ely's team are described. This study shows that increased duration of delirium predicts worse cognitive outcomes 3 and 12 months after the episode of delirium, with the implication that detection and then good care has the potential to make a difference in the long term. Another compelling *why* is patient distress, and this is richly illustrated through many case histories. The authors also provide readers with valuable and pragmatic information on the *how*. They discuss the new diagnostic criteria for delirium (DSM-5, published in May 2013), providing clear guidance on interpretation of these new criteria. They also offer the reader valuable detail on the use of the two major clinical rating scales used to detect and monitor delirium in the ICU, the Confusion Assessment Method for the ICU (CAM-ICU) and the Intensive Care Delirium Screening Checklist (ICDSC). The book also gives substantial coverage of the current state of the art in the appropriate use of drugs in delirium treatment alongside the differential risks of sedatives and other drugs in causing delirium. New findings showing relative benefits of dexmedetomidine over benzodiazepines, the likely lack of

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978-1-107-43365-6 - Delirium in Critical Care: Second Edition
Valerie J. Page and E. Wesley Ely
Frontmatter
[More information](#)

Foreword to the second edition **xiii**

efficacy of haloperidol, and others are discussed. Other notable new content in the second edition relates to new methods of prediction of delirium risk, the role of the ICU environment in modifying delirium risk and the patient experience, and novel evidence on practical and effective methods of delirium prevention in the ICU.

In summary, this second edition builds upon the great success of the first. The content is grounded in both the recent scientific evidence as well as hands-on clinical knowledge. Busy clinicians in the ICU will find no better concise and yet wide-ranging guide to this challenging area of medicine.

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Frontmatter
[More information](#)

DELIRIUM, A PATIENT TESTIMONY

As birthdays go, this one was absolute rubbish. It was 8 o'clock on a May evening in 2007, and where I should have been enjoying an evening out with my husband and friends, here I was sitting in A & E with a broken nose, the result of the most mundane of domestic accidents – falling over some washing while I was completely sober.

Two weeks later I was summoned for day surgery to sort the nose out. My conversation with a porter about the next day's FA Cup final, while making my way down to theatre, is the last memory I have before being plunged into the most terrifying experience of my life.

The next occasion when I had any perception of time was 12 days later, when I found myself being stared at by two middle-aged men in dark suits and bright ties. One was busily explaining to me that I was in the Intensive Care Unit and that I was quite safe.

However, I knew better. I knew they were lying. For me, the reason I was in a bed, on a ventilator, hardly able to move, was that I had been drugged and kidnapped. It had all started in Portugal; at least I thought it was Portugal, where I'd been abducted. At some point I'd managed to escape but was re-captured and taken to a hospital, a few miles from my home.

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978-1-107-43365-6 - Delirium in Critical Care: Second Edition
Valerie J. Page and E. Wesley Ely
Frontmatter
[More information](#)

Delirium, a patient testimony **xv**

I knew that I must have done something wrong, to be held with no hope of escape, but I had no idea what it was. I'd tried on several occasions to pull the tube out of my mouth, but had always been instantly plunged back into darkness.

It never crossed my mind that there might have been a medical reason for my predicament, and I had no knowledge that severe aspiration pneumonia following my routine surgery had landed me in ICU and was putting my family through hell.

What I did know with certainty though was that I would die. One particular doctor would visit me every time I started to wake. He always wore the same clothes and would always speak slowly and deliberately. 'You've been dying to know what that sign on the wall says, haven't you?', he asked one day. He was right; I had wondered what it said but the problem was I could see two of everything and objects and people were frequently blurred so I hadn't been able to read it. 'I'll tell you what it says' he continued. 'It says "moron". We put it there so every time you open your eyes it's the first thing you see; so that for every minute of every day you know exactly what you are.

Unfortunately I'm not allowed to turn your ventilator off, but I want you out of here, and you will leave soon, in a body bag. You're not going to live, but just remember every time you open your eyes you will see exactly what we think of you – moron'.

From then on, that sign was the only thing I could see that wasn't blurred.

The rest of my stay in ICU was filled with more incidents of despair, humiliation and terror. I saw a patient stabbed to death by his wife, and two people committing suicide. I witnessed arguments, in my mind all caused by me, and the

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978-1-107-43365-6 - Delirium in Critical Care: Second Edition
Valerie J. Page and E. Wesley Ely
Frontmatter
[More information](#)

xvi Delirium, a patient testimony

pain I felt as my lungs started to recover was all part of a plan to give me pain-inducing drugs – in fact I had seen doctors laughing about it.

The day after I was extubated I found myself in the High Dependency Unit, where the sheer terror of the execution attempts began. Initially I thought I was in the morgue as I was lying flat and was extremely cold. There was a plain clothes policeman watching me because I'd witnessed a murder on ICU. Someone spoke to me 'I can control your mind' they said, and then proceeded to demonstrate they had power over me by causing me pain and by interrupting my supply of oxygen at will.

The following morning a tall and distinguished looking man sat down and explained to me that I had Pershing's disease. This was a rare congenital heart condition that can lie dormant for many years. Once a sufferer shows symptoms, however, their life expectancy is less than a year. In my mind I might as well just die where I was, and the doctor encouraged me to do exactly that. Pershing's disease of course, does not exist, but just like everything else that happened to me, the hallucination I had where it was explained was so convincing that I was still trying to find evidence of its existence weeks later.

I was put into a side room in the High Dependency Unit, allegedly for infection control, but I knew it was for my execution. I'd heard the nurses talk about CTO, which was a Compulsory Termination Order, and one had been issued for me. As the blinds were pulled down over the room's windows and door, smoke appeared through every vent. A voice told me it was cyanide and I would die more quickly if I relaxed and

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978-1-107-43365-6 - Delirium in Critical Care: Second Edition
Valerie J. Page and E. Wesley Ely
Frontmatter
[More information](#)

Delirium, a patient testimony **xvii**

inhaled it. I watched it creep closer, paralysed with terror, and all I could think about was that I would never see my family again.

Having somehow survived, the execution attempts continued. They included suffocation, poisoning, drug overdoses and being forced to hold my breath until four lights went out on my monitor. By the time I moved down to a respiratory ward I had given up trying to convince my family that the hospital staff were trying to kill me. I still had no idea of what was wrong with me, and when my husband explained, I was sure that all my problems had been induced and had not simply happened. It was then that I decided to keep quiet about my views as no one believed me or was prepared to help me, so I planned my escape alone. By this time I could take some of my medications orally, so this gave me some element of control, as I would wait until the nurse left the room, then would throw the pills containing poison into the medical waste. I ate nothing at mealtimes, but instead stole pieces of cutlery that would help me prise my window open. I was utterly oblivious to the fact that I was four floors up. The hallucinations had stopped by now but were replaced by paranoia and deviousness. The day I planned to leave via the window was the day I was discharged. That might seem lucky, but I spent the next few months wishing that I had died that day.

My physical recovery once I was home moved forward very quickly, but inside I was in meltdown. I couldn't tell anyone how I really felt – my family and friends had been through so much already. My delirium and its consequences caused me to believe that I was still being poisoned, even after leaving hospital, so I stopped taking my medication.

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978-1-107-43365-6 - Delirium in Critical Care: Second Edition
Valerie J. Page and E. Wesley Ely
Frontmatter
[More information](#)

xviii Delirium, a patient testimony

What do I wish had been different for me? Well, when I received a copy of my medical notes following a suggestion that it would help me to make some sense of what had happened, I read through the hundreds of pages that comprised a file 6 inches thick. Only once did I find a relevant reference. It was one about me being severely paranoid, and that note was made by a physiotherapist. Although I have no doubts whatsoever that the care I received was of the highest order, I still feel today that my delirium was seen as an acceptable side effect of my illness and treatment. Months later, during my ICU follow-up appointment, they were not at all surprised that I had suffered prolonged and extreme delirium; in fact they appeared to know I had. To them it was 'normal'. To me it was anything but.

Two years have now passed since my experience of delirium. In that time I've been able to make sense of at least some of what happened to me. My mother reminded me that my belief I'd been kidnapped could well have been my brain confusing my situation with something that had been constantly in the news. Unfortunately, I'd shared my birthday, the day of my accident, with the day that Madeleine McCann was taken from her family's holiday apartment in Portugal, and the tragedy had been at the top of every news bulletin leading up to the day of my surgery.

I know too that my almost total lack of memory for the routine events you would expect to experience on a hospital ward, and which would have reassured me about where I was and what was happening, stopped me from challenging my warped and terrifying perception of the world.

Whatever the future holds though, I'm never doing the washing on my birthday again.