CHAPTER 21.

DELIRIUM

Introduction

Delirium is from Latin and literally means the individual is not at the top of his/her form and travelling at a lower level than normal [de – (off, away from) + lira (a ridge between ploughed furrows)].

“Delirium is a common clinical syndrome characterized by inattention and acute cognitive dysfunction” (Fong et al, 2009a). Inattention means poor ability to concentrate.

Delirium can be an outcome of a general medical conditions, head injury and drug intoxication or withdrawal. It may be the result of the dysfunction of various bodily organs such as kidneys and liver, but it may also be the result of primary pathological processes in the brain.

Delirium is not fully understood. There are problems with terminology; delirium synonyms have included ‘acute confusional state’, ‘organic brain syndrome’, and even, ‘reversible dementia’.

Delirium is a common life-threatening disorder (Inouye, 2006) – 80% of mechanically ventilated patients experience delirium (Field & Wall, 2013).

It is a distressing (to patients, family and staff) and financially costly. Unfortunately, it often goes unrecognized and is poorly managed.

Delirium is seen more commonly in medical and surgical wards than in psychiatric wards. It complicates the hospital stays of 20% of the people over the age of 65 years, and is found in up to 87% of older patients in intensive care wards (Pisani et al, 2003). For reasons which are not always clear, the one year mortality rate following delirium may be as high as 40% (Morgan & Dorevitch, 2001).

[Delirious mania is a unique condition is so far as the only insult to the brain is a psychiatric disorder. The condition can be overlooked because it is difficult to communicate with highly disturbed manic people. However, up to 15% of people with mania may be delirious – which carries some mortal risk. ECT may be indicated. Benzodiazepines are generally considered contra-indicated in delirium, however, in delirious mania, intravenous lorazepam can have dramatic, beneficial effects (Jacobowski et al, 2013).]

DSM-5 criteria Delirium

A. Disturbance of attention (reduced ability to focus, sustain, or shift attention).
B. Develops over a short time (hours or a few days) – a change from baseline attention and awareness, fluctuates in severity in the course of a day.
C. An additional disturbance in cognition (such as memory deficit, disorientation, language disturbance).
Confusion Assessment Method (CAM)

CAM (Inouye et al, 1990) is a remarkable instrument – it is a brief structured assessment - with a sensitivity of 94%, a specificity of 89%, and moderate-to-high inter-rater reliability. It is simple and widely used by nursing staff.

Four questions to be answered with: Yes/No?
The diagnosis of delirium by CAM requires the presence of features 1 and 2 and either 3 or 4.

1. *The history of acute onset and fluctuating course*
   Obtained from family member or nurse as is shown by positive response to the questions:
   Is there evidence of acute change in mental status from the patient’s baseline?
   Does the (abnormal) behaviour fluctuate during the day, that is, does it tend to come and go or increase or decrease in severity?

2. *Inattention*
   This feature is shown by a positive response to the following question:
   Does the patient have difficulty focusing attention such as are they easily distracted or do they have difficulty keeping track of what is being said?

3. *Disorganised thinking*
   This feature is shown by a positive response to the following questions:
   In the patient’s thinking disorganised or incoherent?
   I the conversation rambling or incoherent, unclear with an illogical flow of ideas or unpredictable switching from one subject to another?

4. *Altered level of consciousness*
   This feature is shown by any answer other than ‘alert’ to the following question:
   Overall, how would you rate the patient’s level of consciousness? (alert [normal], vigilant [hyper alert], lethargic [drowsy, easily aroused], stupor [difficult to arouse], or coma [unrousable])

Sub-types

Three clinical subtypes of delirium, based on arousal and psychomotor behaviour are described (Trezepacz et al, 1999)
1. Hyperactive (hyperaroused, hyperalert, or agitated)
2. Hypoactive (hypooroused, hypoalert, or lethargic)
3. Mixed (alternating features of hyperactive and hypoactive types)
While the “classic” presentation of delirium is considered to be the wildly agitated patient, the hyperactive type represents only about 25% of cases. Over half all delirious patients have the hypoactive “quite” type. These people attract less attention and may pass undiagnosed, which is unfortunate, as this (hypoactive) type has the poorer prognosis.

Another “classic” feature is widely believed to be “sundowning”, by which is meant, the mental status deteriorates in the evening. Recent work, however, demonstrated that more symptoms were demonstrated in the morning (47%) than in the afternoon, evening and night (37%).

**Subsyndromal delirium (SSD)**

Subsyndromal delirium (SSD), at the moment, is a research rather than a clinical diagnosis. It has been variously described, and is said to include the presence of one or more core symptoms of CAM delirium, but not meeting the full criteria for delirium. It introduces the notion of early signs of delirium. Not surprisingly, studies suggest a better prognosis for people with 1 rather than 2 core symptoms of delirium (Cole et al, 2011).

**Other delirium scales**

There is a large number of other delirium scales. The Delirium Rating Scale (DRS) was a widely used 10 item scale (Trzepacz et al, 1988). This has recently been updated and expanded into a 16 item scale: Delirium Rating Scale-Revised-98 (Trzepacz et al, 2001).
Testing attention

A commonly used method of testing attention is to ask the patient to perform the serial 7’s test. Rudolph & Marcantonio (2003) make the point that this test requires more calculation skill than attention. Accordingly, they recommend the following:

- Days of the week backwards
- Months of the year backwards
- Digit span (forwards and backwards)
- Spell “world” backwards
- Trailmaking test A

Predisposing and precipitating factors

Predisposing and precipitating factors have been identified. Placement under these headings is somewhat arbitrary, and there is overlap. The large number of factors sets the scene for the next section which points out that multiple factors are involved in most cases.

Predisposing factors

- Advanced age
- Dementia
- Functional impairment in activities of daily living
- Medical comorbidity
- History of alcohol abuse
- Male gender
- Sensory impairment (blindness, deafness)

Precipitating factors

- Acute myocardial events
- Acute pulmonary events
- Bed rest
- Fluid and electrolyte disturbance (including dehydration)
- Drug withdrawal (sedatives, alcohol)
- Infection (especially respiratory, urinary)
- Medications (wide range, esp. psychoactive, anticholinergics and opioids)
- Uncontrolled pain
- Urinary retention, faecal impaction
- Indwelling devices (urinary catheters)
- Severe anaemia
- Use of restraints
- Intracranial events (stroke, bleeding, infection)
Pathophysiology

Delirium arises from different aetiologies, and frequently, in a particular patient, more than one factor is operating (Maldonado, 2013). Thus, a single pathophysiology cannot be identified at this point (may not exist).

Probable mechanisms include:

1. Leaky blood-brain barrier. Recent evidence suggests the blood-brain barrier becomes leaky as the brain ages, allowing exposure to drugs and toxins (Zeevi et al, 2010).

2. Cholinergic deficiency. This is one of the best documented mechanisms. It is seen in overdose of anticholinergic drugs, such as atropine. It may also be seen with the use of drugs not primarily classified as anticholinergics, but with clear cholinergic action: antihistamines, some opioids and antidepressants. However, significant anticholinergic activity has been found in the serum of patients who are not taking drugs with anticholinergic properties - this suggests an endogenous anticholinergic activity may predispose certain patients to delirium.

3. Imbalance of neurotransmitter production. Serotonin is a major CNS neurotransmitter. Production depends on transport of tryptophan across the blood-brain barrier. Tryptophan competes with the amino acid phenylalanine for transport across the blood-brain barrier. Disturbance of the tryptophan:phenalanine ratio may increase or decrease the level of serotonin resulting in delirium. Disturbance of the tryptophan:phenalanine ratio has been observed in post traumatic states and other medical and surgical conditions.

4. Inflammation. Trauma and infection leads to increased production of proinflammatory cytokines, which may produce delirium. Peripherally secreted cytokines can cause responses from microglia, causing inflammation of the brain. Cytokines affect the synthesis and release of a wide range of neurotransmitters and also have neurotoxic (Eikelenboom et al, 2002; Cavallazzi et al, 2013).

5. Elevated cortisol. Acute stress has been hypothesized as a cause of delirium. This is consistent with the notion that elevated cortisol seen in PTSD results in hippocampal shrinkage. The role of cortisol in delirium is under investigation (Maclullich et al, 2008).

6. Neuronal injury caused by a variety of metabolic or ischaemic insults.

7. Other neurotransmitter abnormalities associated with delirium include elevated dopamine function (haloperidol is effective in controlling symptoms). Possibly, also NA and GABA.
Differential diagnosis

The main disorders to consider include dementia (Chapter 20), depression, anxiety and other psychotic disorders.

Hypoactive delirium may look like severe depression, with lack of movement and interest in the surroundings. (This carries the risk of adding an antidepressant medication which may compound the problem (Rathier & Baker, 2011)). Depression is usually preceded by a history of mood disorder, and the thought content may be helpful. Hyperactive delirium is rarely taken to be agitated depression, however, it may be difficult to exclude a severe anxiety disorder. Hallucinations and delusions associated with delirium may suggest a “functional” psychosis, but the picture is clarified by looking for clouding of consciousness (concentration), cognitive difficulties (memory and orientation difficulties) and a fluctuating course.

Delirium and dementia

Where delirium is termed acute brain failure/disorder, dementia is termed chronic brain failure/disorder.

The traditional view is that delirium and dementia are separate disorders. However, evidence suggests they may represent points along a continuum of cognitive decline.

These conditions are interrelated. Dementia is a risk factor for delirium; over half the patients who develop delirium have an underlying dementia. And, acute delirium may leave dementia in its wake. Recent studies indicate that delirium, once considered a brief disorder, may persist for months or even years (McCusker et al, 2003). The line between persistent delirium and reversible dementia is blurred.

Both conditions are associated with decreased cerebral metabolism, cholinergic deficiency and inflammation (Eikelenboom & Hoogendijk, 1999). Imaging studies demonstrate both conditions feature regions of hypoperfusion (Yokota et al, 2003). It is remembered from Chapter 20 that in dementia with Lewy bodies, fluctuating cognition and hallucinations are core features. Thus, similar mechanisms may be involved.

An episode of delirium can dramatically worsen the trajectory of an underlying dementia (Inouye, 2006; Fong et al, 2009b).

Prevention

In efforts to prevent delirium, the following points are recommended:

- Routine cognitive testing on admission and during hospitalization
- Ensure the continued use of glasses and hearing aids as appropriate
- Ensure adequate intake of fluids and nutrition by providing assistance as necessary
- Early identification and treatment of dehydration
- Early mobilization
• Avoid physical restraints (Fick, 2011).
• Involving family members or one-to-one nursing to calm and reorientate.
  (Freter & Rockwood, 2004)
• Cease or minimize use of potentially problematic medications -
  o Minimize benzodiazepine use; dexmedetomidine, an alpha-adrenergic
    receptor agonist, appears to be a suitable sedative alternative (Riker &
    Fraser, 2011).
  o Adequate pain relief - inadequately treated pain increases the
    likelihood of delirium. With respect to older person post hip surgery
    management, opioid use is not associated with delirium in patients
    with or without dementia (Sieber et al, 2011).
• Prophylactic perioperatively antipsychotics (haloperidol, risperidone,
  olanzapine) use has been successful, but is not yet routinely recommended

Multifactorial aetiology and management

Until recently, when faced with a patient with delirium, we looked for the (single)
cause. There is now evidence that delirium most commonly has multifactorial
aetiology. A recent study revealed 16% of a sample had a single aetiological factor,
27% had two, and 90% had up to four aetiological factors (Camus et al, 2000).
Nevertheless, the aetiology in up to 75% of cases remains unknown (Stiefel et al,

The most commonly observed aetiologies of delirium are infection, drug intoxication
and withdrawal, brain injury, low brain perfusion rate, and metabolic disturbances.

Investigations are guided by a comprehensive assessment of the patient, advice on
baseline functioning from people who know the patient, and a careful review of
prescribed medications (with particular attention to recent additions and changes).

Basic laboratory testing includes complete blood count, electrolytes and renal
function tests, oxygen saturation, ECG, urinalysis and chest X-ray. Somewhat
unexpectedly, intracranial factors are rare and should be considered only when all
other factors have been excluded, or if there are focal neurological signs.

Curative pharmacological agents such as antibiotics should be applied as indicated.

A multifactorial non-pharmacological approach is indicated in prevention and
management and includes attention to fluid and electrolyte balance, nutrition and
bladder and bowel function.

The presence of family members at the bed-side is most reassuring. One-to-one
nursing is recommended if possible. Orientation is assisted by having a clock and
calendar nearby, and a window with a view. Disturbance of the sleep-wake cycle is
assisted by discouraging day-time naps and providing a quiet, but softly lit room.
Patients should be encouraged to use glasses and hearing aids when appropriate.
Reassurance will reduce anxiety and assist orientation. The importance of “sitters” has
received some attention (Carr, 2013).
Anxiolytic medication (particularly benzodiazepine) is best avoided, because of the real risk of worsening matters.

Symptom controlling pharmacological agents may be necessary with combative and disturbed behaviour. Drugs with a high anticholinergic effect are avoided. Haloperidol (in small does) remains the most studied treatment of psychotic symptoms in the elderly (Breitbart et al, 1996). The atypical antipsychotics have little advantage over haloperidol and may have additional side-effects such as prolongation of the QT interval. Quetiapine, however, has recently been described as being effective and safe for the treatment of delirium in both general medicine and intensive care units (Hawkins et al, 2013).

A recent review (Bathula & Gonzales, 2013) states “There is a lack of evidence supporting pharmacologic treatment for ICU delirium”. That may be so, but in the real world patients must be managed, and the above paragraph touches on current practice.

Ramelton is a novel selective melatonin receptor agonist. A short case series suggest ramelton has a place in treating delirium (Furuya et al, 2012). Such improvement could be connected with the correction of circadian rhythm disturbance.

Dexmedetomidine is a highly selective 2 adrenergic receptor agonist which provides pain relief and sedation, and has been used in the treatment of delirium. However, there is a caution where there is evidence of heart disease (Tan and Ho, 2013).

Cholinesterase inhibitors apparently have little to offer as a treatment or preventative (Sampson et al, 2007).

References

Sampson E, Raven P, Ndhlouvü P et al. A randomized, double-blind, placebo-controlled trial of denepezil hydrochloride for reducing the incidence of postoperative