Chapter from the book Miscellanea on Encephalopathies - A Second Look
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1. Introduction

Drug-induced encephalopathy is a disease entity often caused by impaired cerebral metabolism that is not attributed to structural brain lesions. However, some drug-induced encephalopathies can develop structural lesions and share other underlying pathophysiological mechanisms (table 2). Leading symptoms are acute or chronic disturbances of consciousness, brain function and personality changes with concomitant neurological symptoms such as asterixis, myoclonias, paresis or seizures (see table 3). Isoniazid-induced encephalopathy was one of the earliest descriptions of a drug-induced encephalopathy (Adams & White, 1965). Clinical symptoms depend on the type and severity of the drug-evoked encephalopathy. A well-described and frequently-reported drug-induced encephalopathy is valproic acid encephalopathy, first described in the late 1970s. This acute encephalopathy was characterized by altered behaviour, worsening seizure control and confusion. After a reduction in the valproate acid dose, the patient’s symptoms resolved completely (Chadwick et al., 1979). Encephalopathies have been reported after consumption of several types of drugs as depicted below (table 1).

2. Drugs

2.1 Analgesics and anaesthesia

Drug-induced encephalopathy was reported after morphine administered intrathecally and the use of propofol (Eran & Barak, 2009). Morphine has been described as having induced an encephalopathy characterized by a myoclonus, motor dysfunction, or vertigo (Goundrey, 1990; Kakinohana et al., 2003). Spinal anaesthesia with hyperbaric bupivacaine lead to an encephalopathy that developed a few days after the drug’s administration (Ho & Chan, 2007). A drug abuse-evoked encephalopathy was also reported after ketamine and gamma hydroxybutyrate (Virmani et al., 2010). Toxic encephalopathy has been described after intake of an acetaminophen overdose (Brusilow & Cooper, 2011).

2.2 Antibiotics

Drug-induced encephalopathy can occur after an intake of cefepime and metronidazole (Kim et al., 2011; Lin et al., 2011). The incidence of metronidazole-induced encephalopathy is unknown. Several studies addressed reversible brain changes caused by metronidazole...
induced-encephalopathy (Ahmed et al., 1995), and bilateral, symmetric brain abnormalities have been observed in patients (Ahmed et al., 1995; Kim et al., 2011). Ceftriaxone induced a reversible encephalopathy in an patient treated for a urinary tract infection (Rancon-Albuquerque et al., 2009). That encephalopathy was completely reversible. An early-onset encephalopathy a day and a half after linezolid therapy occurred in a male, thus clinicians must be aware of the potential of linezolid-induced encephalopathy, particularly in patients presenting risk factors (Fletcher et al., 2010). Clarithromycin has also induced an encephalopathy in adults characterized by symptoms appearing 1-10 days after drug intake and displaying clinical features ranging from delirium to non-convulsive status epilepticus (Bandettin di Poggio et al., 2011). Chinolones like ciprofloxacin and gemifloxacin are also reported to induce an encephalopathy (Rfidah et al., 1995; Barrett MJ 2009). Cephalosporin is reported to evoke an encephalopathy associated with a variety of electroencephalographic manifestations (Grill & Magati, 2008). Other cephalosporines such as cefuroxime, ceftazidime and cefazoline can result in an encephalopathy as well (Herishanu et al. 1988, Jackson et al., 1992; Ortiz et al., 1991). Cefoperazone is a cephalosporine that can cause a reversible encephalopathy characterized by triphasic waves in electroencephalography (Pro et al., 2011). Also penicillin-based antibiotics like penicillin itself, piperacillin and pivmecillinam caused an encephalopathy (Park-Matsumoto et al., 1996; Conway et al., 1968; Lokrantz et al., 2004).

2.3 Antiviral agents

Antiviral agents have seldom been reported to have induced an encephalopathy. As described in one case report, aciclovir can cause an encephalopathy. That patient had normal blood levels of aciclovir, and his renal function was normal (Delluc et al., 2004).

2.4 Antidepressants

Antidepressants can also result in an encephalopathy. The drug amitriptyline can cause an encephalopathy appearing as a neuroleptic malignant syndrome or a serotonin syndrome spectrum disorder (Miyaoka & Kamijama, 1995).

2.5 Anticonvulsants

The following anticonvulsants have been reported to induce a drug-induced encephalopathy: carbamazepine, gabapentin, levetiracetam, lamotrigine, phenytoine, primidone, topiramate, valproic acid and vigabatrine (Engel et al., 1971; Bauer & Elger, 1993; Hennessy & Miles, 1996; Garcia-Pastor et al., 2000; Sechi et al., 2004; Siniscalchi et al., 2004; Cheung et al., 2005; Horvath et al., 2005; Bauer, 2008). The most studied encephalopathy is valproic acid encephalopathy, which was first reported in epileptic and later psychiatric patients (Duarte et al., 1993; Settle, 1995). Valproic acid encephalopathy is often reversible after a week; prolonged time courses have been rare (Bauer & Elger, 1993). Antiepileptic drug-induced encephalopathies represent a seldom, but important side effect of antiepileptic drug therapy. There is an estimated 2% incidence of combined topiramate, valproate acid-induced hyperammonemnic encephalopathy (Cheung et al., 2005). An average age of 38.6 years has been reported in a long-term study of valproic acid-induced encephalopathy (Gerstner et al., 2007).
2.6 Antineoplastic drugs and chemotherapeutics

Capecitabine is an antineoplastic drug replacing 5-Fluouracil in clinical practice. This drug can result in an encephalopathy with seizures even if a conventional dosis is used. No correlation was found between the encephalopathy development and a dihydropyrimidinidene dehydrogenase mutation (Fantini et al., 2010). Carmofur, a 5-fluorouracil derivative that induced a subacute leukencephalopathy, revealed an unsteady gait and dementia (Kuzuhara et al., 1987). A rare complication associated with cisplatin therapy is an encephalopathy with or without seizures (Steeghs et al., 2003). In particular, high doses of ifosfamide can induce an encephalopathy. Ifofosfamide can result in myoclonus-encephalopathy syndrome (Savica et al., 2011). A cohort study revealed a prevalence of 10-40% of this drug-evoked encephalopathy. Female sex, low total albumin and haemoglobulin levels, as well as obesity appear to be risk factors associated with a ifosfamide-evoked encephalopathy (Sweiss et al., 2008). There are few reports of the CNS toxicity of paclitaxel. Seizures have been reported in two patients. Little or no blood brain barrier penetration were the result of confusion and word-finding difficulties, and the encephalopathy resolved itself (Perry & Warner, 1996). Vincristine is known to be an agent that may lead to consecutive sensory and motor dysfunction and eventually fatal myeloencephalopathy (Fawaz al, 1992). Cyclosporine encephalopathy has also been reported (Kwon et al., 2008). Methotrexate has rarely led to an acute encephalopathy, its incidence is 0.8% in leukaemia or lymphoma and 4.5% in osteosarcoma or malignant fibrous histiocytoma (Inaba et al., 2008).

2.7 Immunosuppressants

An encephalopathy occurred after tacrolimus administration and improved after the drug was discontinued. It clinically depicted a right-sided hemiplegia with responsible lesions on diffusion-tensor imaging and diffusion-tensor tractography of the white matter tract (Kim et al., 2011). Sorafenib was reported to induce an encephalopathy in a patient with hepatocellular carcinoma (Dogan et al., 2010). Furthermore, a posterior, reversible leukoencephalopathy syndrome was observed after an infusion of infliximab (Zamvar et al., 2009).

2.8 Immunomodulators

Intravenous immunoglobulins (IVIG) can induce an acute encephalopathy probably caused by a cytotoxic brain oedema (Wada et al., 2005).

2.9 Neuroleptics and lithium

Lithium can to lead to an encephalopathy characterized by seizures, choreiforme as well as parkinsonian movements with cerebellar signs. Three risk factors contribute to lithium toxicity: a nephrogenic diabetes insipidus, age over 50 years, and thyroid dysfunction (Smith et al., 2003). Haloperidol can evoke an encephalopathy characterised by an electroencephalography (EEG) with characteristics of toxic encephalopathy (Maxa et al., 1997). The combination of lithium-risperidone induces an reversible encephalopathy (Boora & Hyatt, 2008). Two patients presented a prolonged postictal encephalopathy with clozapine-induced seizures (Karper et al., 1992) which lasted 63-72 hours and caused electroencephalographic abnormalities.
<table>
<thead>
<tr>
<th>Drug</th>
<th>Dosage/d</th>
<th>Outcome after drug discontinuation</th>
<th>Reference</th>
</tr>
</thead>
<tbody>
<tr>
<td>aciclovir</td>
<td>standard</td>
<td>symptoms reversed after 72h</td>
<td>Delluc et al., 2004</td>
</tr>
<tr>
<td>capecitabine</td>
<td>2000mg/m²</td>
<td>improvement of symptoms</td>
<td>Fantini et al., 2010</td>
</tr>
<tr>
<td>carbamazepine</td>
<td>1200mg</td>
<td>symptoms and EEG normal after 2w</td>
<td>Horvath et al., 2005</td>
</tr>
<tr>
<td>ceftazidime</td>
<td>4g</td>
<td>symptoms resolved after a few days</td>
<td>Jackson et al., 1992</td>
</tr>
<tr>
<td>cefoperazone</td>
<td>2g</td>
<td>EEG and symptoms normal after 36h</td>
<td>Pro et al., 2011</td>
</tr>
<tr>
<td>dexamethasone</td>
<td>4mg p.o.</td>
<td>neuroimaging improvement after 4d</td>
<td>Irvin et al., 2007</td>
</tr>
<tr>
<td>Duodopa + entacapone</td>
<td>1200mg/d</td>
<td>100% mental status recovery after 48h</td>
<td>Manca et al., 2009</td>
</tr>
<tr>
<td>gabapentin</td>
<td>900mg</td>
<td>symptoms normalized after 4w</td>
<td>Sechi et al., 2004</td>
</tr>
<tr>
<td>gemifloxacin</td>
<td>320mg</td>
<td>full recovery 2d later</td>
<td>Barrett MJ, 2009</td>
</tr>
<tr>
<td>isoretinoin</td>
<td>80mg</td>
<td>full recovery 24h later</td>
<td>Wong et al., 2010</td>
</tr>
<tr>
<td>IVIG</td>
<td>1000mg</td>
<td>full recovery after 11d</td>
<td>Wada et al., 2005</td>
</tr>
<tr>
<td>lamotrigine</td>
<td>400mg</td>
<td>symptoms normalized after 4w</td>
<td>Sechi et al., 2004</td>
</tr>
<tr>
<td>levetiracetam</td>
<td>3000mg</td>
<td>symptoms normalized</td>
<td>Bauer, 2008</td>
</tr>
<tr>
<td>lithium</td>
<td>400mg</td>
<td>symptoms resolved after 1 w</td>
<td>Smith et al., 2003</td>
</tr>
<tr>
<td>metronidazole</td>
<td>45.5g</td>
<td>symptoms improved within 6.7d</td>
<td>Kim et al., 2011</td>
</tr>
<tr>
<td>morphine</td>
<td>0.5mg</td>
<td>recovery after 10d</td>
<td>Eran &amp; Barak, 2009</td>
</tr>
<tr>
<td>odansetron</td>
<td>4mg</td>
<td>full recovery of symptoms</td>
<td>Ritter et al., 2003</td>
</tr>
<tr>
<td>oxcarbazepine</td>
<td>1800mg</td>
<td>EEG and symptoms normalized after 20d</td>
<td>Siniscalchi et al., 2004</td>
</tr>
<tr>
<td>penicillin</td>
<td>i.v. 60 Mega Units</td>
<td>patient died</td>
<td>Conway et al., 1968</td>
</tr>
<tr>
<td>phenytoine</td>
<td>300mg</td>
<td>symptoms and EEG normalized</td>
<td>Engel et al., 1971</td>
</tr>
<tr>
<td>pivmecillinam</td>
<td>600mg</td>
<td>fast symptom recovery</td>
<td>Lokrantz et al., 2004</td>
</tr>
<tr>
<td>primidone</td>
<td>600mg</td>
<td>symptoms and EEG normalized after 2 w</td>
<td>Katano et al., 2002</td>
</tr>
<tr>
<td>propofol</td>
<td>150mg</td>
<td>recovery after 10d</td>
<td>Eran &amp; Barak, 2009</td>
</tr>
<tr>
<td>sorafenib</td>
<td>400mg/2x</td>
<td>all symptoms resolved after 5d</td>
<td>Dogan et al., 2010</td>
</tr>
<tr>
<td>tacrolimus</td>
<td>0.1mg/kg/d</td>
<td>imaging abnormalities normalized after 4m</td>
<td>Kim et al., 2011</td>
</tr>
<tr>
<td>topiramate</td>
<td>1400mg</td>
<td>EEG and symptoms normalized after 7d</td>
<td>Cheung et al., 2005</td>
</tr>
<tr>
<td>valproic acid</td>
<td>2400mg</td>
<td>improvement in EEG and symptoms</td>
<td>Bauer &amp; Elger, 1993</td>
</tr>
<tr>
<td>vigabatrine</td>
<td>3000mg</td>
<td>improvement in 2w</td>
<td>Garcia-Pastore et al., 2000</td>
</tr>
</tbody>
</table>

d = days, EEG = Electroencephalography, h = hours, IVIG intravenous immunoglobulines, m = month, w = weeks, p.o. = per os

Table 1. Drug-induced encephalopathy- clinics and outcome (not all drugs mentioned in the text are depicted in the table)
2.10 Other classes of drugs

There are several other classes of drugs that can result in a drug-induced encephalopathy. For instance, baclofen [a derivative of gamma-aminobutyric acid (GABA)] caused an encephalopathy with severe electroencephalographic abnormalities (Kumar et al., 2010). Another class of drugs such as duodopa (a combination of levodopa and carbidopa) has induced a reversible encephalopathy in Parkinson’s disease. An intermittent multifocal myoclonus was observed, and neurologic examination revealed a flaccid tetraparesis (Manca et al., 2009). An inadvertent injection of gadolinium (solutions of chelated organic gadolinium complexes) can result in grand mal seizures and mental changes due to an encephalopathy (Kapoor et al., 2010). This gadolinium encephalopathy probably occurred due to the inadvertent simultaneous entry of gadolinium and blood into the subarachnoid space. This case highlights the importance of using only a small amount of gadolinium. Agents like isoretinoin (medication in the therapy of Acne) may induce an encephalopathy with myoclonic jerks and confusion (Wong et al., 2010). Ondansetron (a 5-HT 3 receptor antagonist used mainly as an antiemetic drug) can produce a multifocal encephalopathy depicted by a transient pyramidal and extrapyramidal dysfunction with Babinski signs, oculogyric crisis, oromandibular and limb dystonia. The symptoms resolved after hours. Anaesthesiologists must take special care when administering ondansetron therapy because of this rare complication and the severe clinical manifestation reflecting transient structural brain damage that however results in a full resolution of neurological symptoms. Sulfasalazine is a drug used primarily as an anti-inflammatory agent in the treatment of inflammatory bowel disease and rheumatoid arthritis. It has caused an encephalopathy characterised by cerebrospinal fluid with a high protein level (Mut et al., 2008).

3. Pathophysiological mechanisms

The underlying causes of a drug-induced encephalopathy are not yet fully understood. Several mechanisms of drug-induced encephalopathy are discussed below (table 3).

3.1 Cytotoxic and neurotoxic effects

There are several pharmaceutical cytotoxic and neurotoxic side effects that can cause an encephalopathy. A rise in the glutamine and glutamate complex peak in MR spectroscopy suggests for example an excitotoxic injury in the neurons and astrocytes in an acute IVIG-induced encephalopathy (Wada et al., 2005), and it is one possible mechanism inducing neurotoxicity.

3.2 Electrolytic disturbances

There are electrolytic disturbances such as a hypo- or hypernatremia that can promote drug-induced encephalopathy. Hyponatremia may be a side effect of drugs such as oxcarbazepine or diuretics. Severe hyponatremia is commonly caused by the syndrome of inappropriate antidiuresis (SIADH), which can also be induced by drugs like cyclophosphamide, vincristine, vinblastine, thiotixene, thoridazine, haloperidol, monoamine oxidase inhibitors, tricyclic antidepressants, selective serotonin reuptake inhibitors and bromocripitine (Esposito et al., 2011). Hypopotassemia plays a role in the pathogenesis of convulsions and the high rate of mortality in theophylline encephalopathy (Suarez Ortega et al., 1995).
3.3 Hepatic enzyme interactions and hyperammonemia

Valproic acid can inhibit differential enzymes of the urea cycle, inducing a hyperammonemia (Sechi et al. 2004, Treem et al., 1994). Moreover, there is the potential of damage on an enzymatic level that can lead to hyperammonemia: (1) carbamylphosphat synthetase-, (2) ornithin-transcarbamylase-, (3) N-acetylglutamat-synthetase-, (4) argininosuccinat-synthetase- and (5) arginino-succinat-lyase deficiency. As the incidence of these enzyme defects is low, valproic-acid encephalopathy very seldom has a hereditary cause. A high level of ammonia can lead to hepatic necrosis in addition to encephalopathy. Hyperammonemia can be induced by the drugs depicted in table 2. Recent study data indicate that the hyperammonemia observed in patients under valproic-acid treatment is based on the direct inhibition of hepatic N-acetylglutamate synthase activity by valproyl-CoA (Aires et al., 2011). Hyperammonemia can induce an encephalopathy by inhibiting the glutamate uptake by astrocytes, thus provoking neuronal damage and cerebral oedema (Blindauer et al., 1998). Moreover, elevated extracellular glutamate reduces the size of the astrocytes, thereby inhibiting their function. The reduced synthesis of glutathione causes the neurons and glia cells to become more vulnerable to oxidative stress (Verotti et al., 2002). Finally, the over-production of glutamine leads to a swelling of the astrocytes followed by cerebral oedema and even higher cerebral pressure (Noremberg, 1996).

<table>
<thead>
<tr>
<th>Drug</th>
<th>Reference</th>
</tr>
</thead>
<tbody>
<tr>
<td>5-Flouracil</td>
<td>Advani &amp; Fakih et al., 2011</td>
</tr>
<tr>
<td>Acetazolamide</td>
<td>Kim et al., 2007</td>
</tr>
<tr>
<td>Carbamazepine</td>
<td>Adams et al., 2009</td>
</tr>
<tr>
<td>Haloperidol</td>
<td>Rubenstein et al., 1990</td>
</tr>
<tr>
<td>Lamotrigine</td>
<td>Fan et al., 2008</td>
</tr>
<tr>
<td>Primidone</td>
<td>Katano et al., 2002</td>
</tr>
<tr>
<td>Valproate acid</td>
<td>Aires et al., 2011</td>
</tr>
<tr>
<td>Zonisamide</td>
<td>Shaikh et al., 2009</td>
</tr>
</tbody>
</table>

Table 2. Drugs inducing hyperammonemia

An acute intermittent porphyria as one form of acute hepatic porphyria can present as a diffuse encephalopathy (Maramattom et al., 2005). Additionally, a mouse model has demonstrated that griseofulvin induces a hepatic porphyria characterized by psychiatric behavior sometimes observed in drug-induced encephalopathies (Satoh et al., 2008).

Drugs such as barbiturates, bernegride, chloramphenicol, chlordiazepoxide, chloroquine, chlorpropamide, danazol, diazepam, ergot preparations, estrogens, ethanol excess, griseofulvin, halothane, hydantoins, imipramine, ketamine, meprobamate, methyldopa, methyprylon, methsuximide, nikethamide, oral contraceptives, pentazocine, phensuximide, phenylbutazone, progestogens, pyrazinamide, pyrazolone derivatives, sulfonamides, theophylline derivatives, tolbutamide, troxidone and valproic acid (Bonkowsky et al., 1982) have been reported to exacerbate acute porphyrias. These drugs should thus be administered with caution in patients with an encephalopathy associated with porphyria. A further mechanism leading to an encephalopathy based on an increase in neuronal P450 CYP2E1 activity is induced by acetaminophen in an animal model (Posadas et al., 2010). Posadas et al. showed that acetaminophen can result in a concentration-dependent neuronal apoptosis on rat cortical neurons through a mitochondrial-mediated mechanism.
that includes cytochrome c release and caspase 3 activation (Posadas et al., 2010). Surprisingly, the neurotoxic action by acetoaminophen in rats is below those required to induce hepatotoxicity.

### 3.4 Effects on cerebral receptors

Effects on cerebral receptors play an important role as underlying pathomechanisms in drug-induced encephalopathy. The neurotoxicity in metronidazole encephalopathy is based on the RNA (Bradley et al., 1977) and DNA binding of intermediate metabolites of metronidazole (Wright & Tyler, 2003), modulating inhibitory GABA receptors in the cerebellar and vestibular systems (Evans et al., 2003).

Interaction with the GABA receptor plays a role in the intrinsic toxic effects of valproic acid encephalopathy (Miyazaki et al., 1988). Topiramate can induce a direct toxic effect on the central nervous system (CNS). Combined therapy with valproate acid produces this effect by reducing the metabolism of topiramate due to the interaction of valproic acid with the cytochrome-P450 effect. Gabapentin may cause a reversible encephalopathy clinically characterised by an asterixis. One candidate mechanism this encephalopathy is the agonistic interaction of gabapentin on cerebral GABA receptors in conjunction with increased inhibitory action (Fink et al., 2002). Cephalosporine-induced encephalopathy seems to involve GABA A receptor inhibition (Grill & Magati et al., 2008).

### 3.5 Metabolic effects

Severe diseases or malnutrition have a reduction in glucuron acid as a consequence. It is thus possible to inhibit the glucoronidation of valproate acid, resulting in a higher cumulative concentration of valproic acid, lamotrigine and oxcarbazepine in blood levels.

### 3.6 Vasogenic and cytotoxic brain oedema

Vasogenic and cytotoxic brain oedema as an underlying mechanism of a drug-induced encephalopathy is widespread. Metronidazole encephalopathy is probably caused by vasogenic and cytotoxic brain oedema. Most of the lesions in metronidazole encephalopathy correspond to areas of vasogenic oedema according to diffusion weighted imaging. Some lesions are located in the corpus callosum and correspond to cytotoxic oedema. Cytotoxic oedema is also a candidate mechanism in IVIG-induced encephalopathy. An intramyelinic oedema in the myelin sheath was observed in IVIG-induced encephalopathy (Wada et al., 2005). Many drug-induced encephalopathies share in common a posterior reversible leukoencephalopathy syndrome (PRES) possibly due to vasogenic oedema.

### 3.7 Posterior reversible leukoencephalopathy syndrome

The PRES has been described after the intake of immunosuppressants such as tacrolimus, cyclosporine or in association with acute hypertensive encephalopathy and eclampsia (Hinchey et al., 1996). It is characterised by capillary-leak syndrome in the brain caused by hypertension, liquid retention, immunosuppressants, and chemotherapeutics affecting the vascular endothelium. Clinical symptoms are headache, vomiting, confusion, seizures, cortical blindness and other visual symptoms. Neuroimaging reveals bilateral signal alterations in the posterior white matter suggesting oedema.
<table>
<thead>
<tr>
<th>Candidate Mechanism</th>
<th>Drugs</th>
<th>Reference</th>
</tr>
</thead>
<tbody>
<tr>
<td>cytotoxic brain edema</td>
<td>IVIG, metronidazole</td>
<td>Kim et al., 2011; Wada et al., 2005</td>
</tr>
<tr>
<td>effect on cerebral receptors</td>
<td>methotrexate</td>
<td>Sasazaki et al., 1992</td>
</tr>
<tr>
<td>electrolytic disturbance</td>
<td>theophylline</td>
<td>Suarez Ortega et al., 1995</td>
</tr>
<tr>
<td>hepatic enzyme interactions</td>
<td>valproic acid</td>
<td>Bauer &amp; Elger, 1993</td>
</tr>
<tr>
<td>hypoalbuminuria</td>
<td>ifosfamide</td>
<td>Sweiss et al., 2008</td>
</tr>
<tr>
<td>metabolic effects</td>
<td>lamotrigine, oxcarbazepine, valproic acid</td>
<td>Bauer &amp; Elger, 1993, Hennessy &amp; Miles, 1996</td>
</tr>
<tr>
<td>neurotoxic effect</td>
<td>IVIG</td>
<td>Wada et al., 2005</td>
</tr>
<tr>
<td>posterior reversible leukoencephalopathy syndrome</td>
<td>dexamethasone, tacrolimus,</td>
<td>Kim et al., 2011; Irvin et al., 2007; Zhang, 2011</td>
</tr>
<tr>
<td>vasogenic brain edema</td>
<td>metronidazole</td>
<td>Kim et al., 2011</td>
</tr>
</tbody>
</table>

Table 3. Drug-induced encephalopathy-pathophysiological mechanisms

4. Pathological studies

Mild gliosis of the white matter and ischemic lesions in the temporal area were observed in a patient’s postmortem analysis (Steeghs et al., 2003). Pathological-anatomic studies showed changes in the cerebellum and temporal lobe of predominantly the pyramidal and purkinje cells in rats after chronic administration of valproate acid (Sobaniek-Lotowska, 2003). Those studies reported damage to the hippocampal astrocytes and neocortex. All these abnormalities seemed to disappear three months after discontinuation of the drug.

5. Genetic susceptibility

A further factor contributing to the development of a drug-induced encephalopathy is genetic susceptibility. The individual’s genetic patrimony including ethnicity and gender influences the susceptibility to the risk of a drug-induced encephalopathy. Any genetic polymorphism may influence the metabolism, excretion or action of the drug depending on single or multiple genes or by changes in gene expression (Dodd et al., 2004). For instance, some mutations can promote development of an encephalopathy, i.e. a mutation in ETHE1, a mitochondrial matrix sulphur dioxygenase causing an ethylmalonic encephalopathy (Viscomi et al., 2010). In a patient with the rare missense variant methionine synthetase c.2756A>G (D919G), a methotrexate encephalopathy was observed probably due to a modified effect of methotrexate on homocysteine metabolism (Linnebank et al., 2007). A recent clinical study showed that the genetic polymorphism of the human thymidylate synthetase gene contributes to 5-fluorouracil-associated hyperammonemic encephalopathy. A GABA A receptor modification caused by knockout of the taurine transporter resulted in striatal disinhibition in mice. This animal study demonstrates that a genetic defect ending up in a lack of taurine partly explains the pathophysiology of a hepatic encephalopathy (Sergeeva et al., 2007). Mitochondrial dysfunction underlies different types of encephalopathy, for example, mitochondrial myopathy, encephalopathy, lactic acidosis, and stroke-like episodes (MELAS). As an example, the mutation of mitochondrial DNA (mtDNA) G13513A encoding the ND5 subunit of
respiratory chain complex 1 causes mitochondrial encephalopathy with lactic acidosis (Wang et al. 2008). Therefore, supplementation with the mitochondrial respiratory chain cofactor coenzyme Q10 has been demonstrated to advance recuperation following heroin-induced encephalopathy (Gacouin et al., 2003).

6. Epidemiology

The epidemiology of ifosfamide encephalopathy is well known. Ifosfamide encephalopathy occurred in 31.2% of patients with soft tissue and bone sarcomas (17/61) treated with ifosfamide, and in 13.6% of ifosfamide treatment courses. A history of cisplatin was identified as a potential risk factor for the development of an ifosfamide-induced encephalopathy (Tajino et al., 2010). Furthermore, a dose of ifosfamide at $>9g/m^2$ is a further risk factor of ifosfamide-induced encephalopathy (Tajino et al., 2010). In other previous studies, risk factors such as large tumors in the female pelvic cavity (Meanwell et al., 1986), poor performance status (Antman et al., 1989), hypoalbuminemia (Merimsky et al., 1992), high serum creatinine level (Antmann et al., 1990) and low iron bicarbonate level (Antmann et al., 1989) were identified as risk factors for ifosfamide-induced encephalopathy. It remains controversial as to whether there are risk factors of ifosfamide-induced encephalopathy, as another study showed no risk factors associated with this encephalopathy and that each patient has his own predisposition (Rieger et al. 2004). Alcoholism was identified as a risk factor in linezolid-induced encephalopathy (Fletcher et al., 2010). Renal failure and previous central nervous system disease may predispose to ceftriaxone- and cefepime-induced encephalopathy (Roncon-Albuquerque et al., 2009; Garces et al. 2008). Dialysis may be a risk factor in isoniazide-induced encephalopathy (Cheung et al. 1993). There are common risk factors affecting the neuronal health for different types of drug-induced encephalopathies such as environmental toxins, infectious diseases, traumatic events, brain tumors, brain ischemia, age (state of health, disease), nutritional deficiencies and intolerances, and even poverty (Virmani et al. 2010).

In conclusion, not everyone develops an encephalopathy after taking a certain drug, but those individuals who are at risk (see above) - although the extent and nature of the risk are often unknown - may be more apt to develop an encephalopathy.

Patients with metronidazole encephalopathy showed a mean age of 61 years (49-71 years) (Kim et al., 2007), whereas those with clarithromycin encephalopathy exhibited an average age of 51 years (19-87 years) (Bandetti di Poggio et al., 2011). These data are based on case series, and there are no long-term clinical studies addressing the epidemiology of drug-induced encephalopathies. In a cohort study of 19 patients, 8 patients (42%) developed an ifosfamide induced encephalopathy (Sweiss et al. 2008). The exact prevalence of a drug-induced encephalopathy is unknown, as case series with calculated epidemiologic data are rare. Furthermore, there are no studies larger in scale examining the specific age of a drug-induced encephalopathy. The average age for valproic-acid encephalopathy was 38.6 years in a long-term study (Gerstner et al. 2007).

Toxic encephalopathies are accompanied by high blood levels of the suspected drug, whereas drug-induced encephalopathies often reveal therapeutic blood levels of the drug. Thus we know of no dose-dependent effect of drugs that induce an encephalopathy. The symptoms can develop from within hours until a month after taking the drug.
7. Basic clinical features

The clinical spectrum of symptoms can result in slight disturbances of the mental state up to severely damaged consciousness (table 4). Transient acute encephalopathy has been observed in 3-15% of cancer patients after methotrexate therapy (Rubnitz et al. 1998). Chronic encephalopathy develops slowly, may progress, and can permanently impair neurological function. A drug-induced encephalopathy may reveal a varied spectrum of psychiatric symptoms, i.e. hallucinations (Sorafenib; Dogan et al., 2010), psychotic state (vigabatrine; Garcia-Pastore et al., 2000), depression (sodium valproate, Connacher et al., 1987) and neuropsychologic symptoms like reduced psychomotor speed and impaired working memory (levetiracetame, valproic acid; Bauer et al., 2008). The development of psychiatric symptoms may be acute, subacute or chronic.

<table>
<thead>
<tr>
<th>Clinics</th>
<th>Drugs</th>
<th>Reference</th>
</tr>
</thead>
<tbody>
<tr>
<td>abnormal sensation</td>
<td>metronidazole</td>
<td>Wada et al., 2005</td>
</tr>
<tr>
<td>aggravation of preexisting neurological deficits</td>
<td>carbamazepine, gabapentine, levetiracetame, lamotrigine, oxcarbazepine, primidone, topiramate</td>
<td>Bauer, 2008; Hennessy &amp; Miles, 1996, Horvath et al., 2005; Katano et al., 2002; Latour et al., 2004; Siniscalchi et al., 2004; Sechi et al., 2004</td>
</tr>
<tr>
<td>altered consciousness, concentration</td>
<td>capecitabine, gabapentine, IVIG, lithium, valproic acid</td>
<td>Bauer &amp; Elger 1993; Fantini et al., 2010; Katano et al., 2002; Sechi et al., 2004</td>
</tr>
<tr>
<td>anisocoria, diplopia</td>
<td>dexamethasone</td>
<td>Irvin et al., 2007</td>
</tr>
<tr>
<td>aphasia</td>
<td>carbamazepine, topiramate</td>
<td>Horvath et al., 2005, Latour et al., 2004</td>
</tr>
<tr>
<td>ataxia, apraxia</td>
<td>capecitabine, carbamazepine, phenytoine, topiramate, valproic acid</td>
<td>Fantini et al., 2010, Horvath et al., 2005</td>
</tr>
<tr>
<td>choreiform movements, athetosis</td>
<td>tiagabine, trimetazidine</td>
<td>Sivet et al., 2008; Tombini et al., 2006</td>
</tr>
<tr>
<td>delirium, coma</td>
<td>carmofur</td>
<td>Kuzuhara et al., 1987</td>
</tr>
<tr>
<td>dementia, memory loss</td>
<td>carbamazepine, carmofur, gabapentine, levetiracetame</td>
<td>Bauer, 2008; Horvath et al., 2005, Kuzuhara et al., 1987; Sechi et al., 2004</td>
</tr>
<tr>
<td>dysarthria</td>
<td>metronidazole, lithium, tacrolimus,</td>
<td>Smith et al., 2003; Wada et al., 2005,</td>
</tr>
<tr>
<td>gait disturbance</td>
<td>carmofur, sorafenib, trimetazidine,</td>
<td>Dogan et al., 2010; Kuzuhara et al., 1987; Sivet et al., 2008</td>
</tr>
<tr>
<td>headache</td>
<td>paclitaxel</td>
<td>Perry &amp; Warner, 1996</td>
</tr>
<tr>
<td>myoclonias</td>
<td>carbamazepine, levetiracetame, odansetron, vigabatrine, lithium</td>
<td>Bauer, 2008; Garcia-Pastor et al., 2000; Horvath et al., 2005, Ritter et al., 2003; Smith et al., 2003</td>
</tr>
</tbody>
</table>
8. Diagnostics

8.1 Electrophysiologic studies

EEG has often revealed signs of encephalopathy. The main characteristics are a diffuse, unusually mild to heavy general changes (Horvath et al., 2005, Tombini et al., 2006). Triphasic waves with intermittent frontal delta activity are sometimes observed (Gallmetzer et al., 2004; Rancon-Albuquerque et al., 2009). Focal and generalised slow waves, and generalised or focal epileptic spike-wave complexes have also been seen. Once the responsible drug is discontinued, the encephalopathy with general slowing and epileptic discharges resolve after days or weeks, sometimes after months (Bauer & Elger, 1993; Latour et al., 2004).

8.2 Laboratory investigations

The effective concentration of the drug in sera is often within the normal range in patients with valproic acid-induced encephalopathy (Bauer & Elger, 1993), whereas higher concentration of the drugs were noted in carbamazepine (Neumann et al. 1994) and lamotrigine encephalopathy (Hennesy & Miles, 1996). When clinical signs and symptoms of

Table 4. Drug-induced encephalopathy- Clinics

<table>
<thead>
<tr>
<th>Symptom</th>
<th>Drug</th>
<th>Reference</th>
</tr>
</thead>
<tbody>
<tr>
<td>oculogyric crisis, oromandibular and limb dystonia</td>
<td>odansetrone</td>
<td>Ritter et al., 2003</td>
</tr>
<tr>
<td>parkinsonism</td>
<td>carbamazepine</td>
<td>Horvath et al., 2005</td>
</tr>
<tr>
<td>psychiatric symptoms</td>
<td>valproic acid, sorafenib, levetiracetame, vigabatrine</td>
<td>Dogan et al.2010; Garcia-Pastor et al., 2000; Bauer, 2008</td>
</tr>
<tr>
<td>ptosis</td>
<td>dexamethasone</td>
<td>Irvin et al., 2007</td>
</tr>
<tr>
<td>seizures</td>
<td>cisplatin, cyclosporine, gadolinium levetiracetame, valproic acid, vigabatrine</td>
<td>Bauer, 2008; Bauer &amp; Elger, 1993, Dzudie et al., 2009; Garcia-Pastor et al., 2000; Kapoor et al., 2010; Steeghs et al., 2003;</td>
</tr>
<tr>
<td>sleep disturbance, hypersomnia, insomnia</td>
<td>capecitabine, carbamazepine, topiramate</td>
<td>Cheung et al., 2005, Fantini et al., 2010, Horvath et al., 2005, Cheung et al., 2005</td>
</tr>
<tr>
<td>stupor, agitated state</td>
<td>morphine</td>
<td>Eran &amp; Barak, 2009</td>
</tr>
<tr>
<td>tremor</td>
<td>tacrolimus</td>
<td>Kim et al., 2011</td>
</tr>
<tr>
<td>vertigo</td>
<td>sorafenib, valproic acid</td>
<td>Bauer &amp; Elger, 1993; Dogan et al., 2010</td>
</tr>
<tr>
<td>visual symptoms, nystagmus</td>
<td>metronidazole, phenytoine, primidone, trimetazidine, sorafenib, topiramate</td>
<td>Wada et al., 2005; Dogan et al., 2010; Engel, 1971; Katano et al., 2002; Latour et al., 2004; Sivet et al., 2008;</td>
</tr>
<tr>
<td>vomiting, nausea</td>
<td>valproic acid</td>
<td>Bauer &amp; Elger, 1993</td>
</tr>
</tbody>
</table>
a drug-induced encephalopathy are present, relevant clinical routine tests for natremia, ammonemia or glycemia should always be performed to identify the reason for the encephalopathy and develop a treatment strategy. A hypoglycemic encephalopathy can be detected by measuring the blood glucose level, thereby differentiating it from drug-induced encephalopathies. The clinical spectrum of hypoglycemic encephalopathy ranges from simple neurological deficits and mental changes to severe coma and death (Lo et al., 2006). A specific lesion pattern is frequently detected in hypoglycemic encephalopathy, often affecting the cerebral cortex, basal ganglia, hippocampus, splenium and bilateral internal capsula (Aoki et al., 2004; Chan et al., 2003; Terakawa et al., 2007; Cho et al., 2006). This selective vulnerability in hypoglycemic encephalopathy may be associated with the extent to which the metabolism necessary to conserve the function of brain structures and neuronal integrity has been compromised (Lee et al., 2011).

8.3 Imaging patterns

Cerebral atrophy has been observed in valproic-acid encephalopathy, especially in chronic encephalopathy as some authors have described in cranial computertomography (CT) and magnetic resonance imaging (MRI) (Baganz et al., 1994; Papazian et al., 1995). Lacunar lesions were found in gabapentin-induced encephalopathy (Sechi et al., 2004). Metronidazole-induced encephalopathy induced bilateral symmetric T2-hyperintense lesions in the cerebellar dentate nucleus, midbrain, dorsal pons, medulla, and splenium of the corpus callosum. Except for the corpus callosum, all lesions were irreversible. The lesions are often bilateral and symmetric. High signal intensity in T2-weighted images appeared, but the signal alterations did not demonstrate contrast enhancement and were reversible after drug discontinuation. Dexamethasone encephalopathy in MRI resulted in diffuse cortical and subcortical white matter lesions in symmetric bilateral distribution involving predominantly occipital areas, the cerebellum and focal areas of bilateral thalami not evident in the T1-weighted images characteristic of PRES. In cyclosporine-induced encephalopathy, the lesions show vasogenic oedema apparently in diffusion coefficient maps (Bartynski et al., 2007). Some imaging pattern are shown in table 5.

<table>
<thead>
<tr>
<th>Anatomic lesion pattern</th>
<th>Drugs</th>
<th>Reference</th>
</tr>
</thead>
<tbody>
<tr>
<td>bilateral temporal periventricular white-matter lesions</td>
<td>sulfasalazine</td>
<td>Mut et al., 2008</td>
</tr>
<tr>
<td>centrum semi-oval atrophy</td>
<td>tacrolimus</td>
<td>Kim et al., 2011</td>
</tr>
<tr>
<td>cerebellum</td>
<td>metronidazole</td>
<td>Kim et al., 2011</td>
</tr>
<tr>
<td>corpus callosum</td>
<td>metronidazole</td>
<td>Kim et al., 2011</td>
</tr>
<tr>
<td>cortical and subcortical parietal, occipital and frontal white matter lesions</td>
<td>dexamethasone, IVIG</td>
<td>Irvin et al., 2007; Wada et al., 2005</td>
</tr>
<tr>
<td>deep white matter lesions</td>
<td>capecitabine, paclitaxel</td>
<td>Fantini et al., 2010; Perry &amp; Warner, 1996</td>
</tr>
<tr>
<td>midbrain, pons, medulla lesions</td>
<td>metronidazole</td>
<td>Kim et al., 2011</td>
</tr>
</tbody>
</table>

Table 5. Neuroimaging of drug-induced encephalopathies
9. Differential diagnosis

The most important differential diagnosis of metronidazole-induced encephalopathy is Wernicke encephalopathy. In the early stages of the disease, the two entities may be confounded because they can produce similar clinical features. In metronidazole-induced encephalopathy (unlike Wernicke encephalopathy), lesions of cerebellar dentate nuclei are supported by pathological studies (Troncoso et al., 1981). Further differential diagnoses are acute infectious encephalitis and demyelinating disease including Marchiafava-Bignami disease. Other types of encephalopathies must be differentiated from drug-induced encephalopathies such as hepatic, heavy metal, uremic, septic, and mitochondrial encephalopathy.

10. Therapeutic options

Therapy consists in the immediate discontinuation of the suspected drug when first signs of encephalopathy appear. A complete reversal of symptoms should take place soon after drug discontinuation. The atrophy in valproate-induced encephalopathy can also be reversed in individual cases. The administration of L-carnitine makes therapeutic sense in cases of carnitine deficiency (Kelley, 1994). Intravenous carnitine was recently shown to be useful in the treating hyperammonemic encephalopathy (Bøhmer & Hoymark, 2010). Moreover, the treatment of a valproate-induced encephalopathy via haemodialysis has succeeded (Tsai & Chen, 2008). Short-term hemodialysis often helps to reverse the symptoms in cefepime encephalopathy (Lin et al., 2011). Drug-induced encephalopathy often has a good prognosis. Methylben blue is an therapeutic option for an ifosfamide-induced encephalopathy (Patel, 2006). A drug-induced encephalopathy can sometimes be prevented by adjusting the dosage and monitoring serum concentrations of the suspected drug. Normally, the blood level of the suspected drug is within the therapeutic range, so that treating an overdose would make no sense. Only in particular situations are certain measures to treat overdoses necessary, i.e. gastric lavage, activated charcoal, hemodialysis, hyperhydration or forced diuresis.

11. Conclusions

Several drugs can induce drug-induced encephalopathies. They seldom occur in clinical practice, but are important pharmacological side effects. Drug-induced encephalopathies are a key differential diagnosis when a disturbance of the consciousness is initially unclear. The effective levels of the drug in the blood often fall within the reference and not in the toxic range. Effective therapy consists in immediately discontinuing the drug.

12. References


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The book project “Miscellanea on Encephalopathies—a second look” aims to cover some of the important aspects regarding metabolic, hypoxic, neoplasm- and drug-related encephalopathies, by transmitting valuable information filtered through the real life clinical and research experience of the authors.

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