Overview of literature and information on ‘khat-related’ mortality: a call for recognition of the issue and further research

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Abstract (word count = 261)

During the past 20 years or so, more has become known about the properties of khat, its pharmacology, physiological and psychological effects on humans. However, at the same time its reputation of social and recreational use in traditional contexts has hindered the dissemination of knowledge about its detrimental effects in terms of mortality. This paper focuses on this particular deficit and adds to the knowledge-base by reviewing the scant literature that does exist on mortality associated with the trade and use of khat. We sought all peer-reviewed papers relating to deaths associated with khat. From an initial list of 111, we identified 15 items meeting our selection criteria. Examination of these revealed 61 further relevant items. These were supplemented with published reports, newspaper and other media reports. A conceptual framework was then developed for classifying mortality associated with each stage of the plant’s journey from its cultivation, transportation, consumption, to its effects on the human body. The model is demonstrated with concrete examples drawn from the above sources. These highlight a number of issues for which more substantive statistical data are needed, including population-based studies of the physiological and psychological determinants of khat-related fatalities. Khat-consuming communities, and health professionals charged with their care should be more aware of the physiological and psychological effects of khat, together with the risks for morbidity and mortality associated with its use. There is also a need for information to be collected at international and national levels on other causes of death associated with khat cultivation, transportation, and trade. Both these dimensions need to be understood.

Key words: khat/qat/Catha edulis mortality/fatalities/death over-view/review

[word count for main text, exc acknowledgements, tables & refs = 9658]
Introduction

Botany and chemistry

Khat is a herbal product consisting of the leaves and shoots of the shrub Catha edulis Forsk, a member (genera) of the evergreen celastracae (moonseed or spindle-tree) family or tribe (Simmons et al, 2008). Under natural conditions, khat grows in and on the margins of dry evergreen forest and mist forest. It grows naturally at elevations of 1,500-2,000m but is found at altitudes of 1,200–2,500m. It is best cultivated at high elevations, with high rainfall in acidic, well-drained clay soils; but can survive long periods of drought. Khat's natural range extends throughout East Africa from Ethiopia, Eritrea and Somalia, through to South Africa; it is also found in Rwanda, Zaire, Malawi and Zimbabwe. Catha edulis is also found sporadically in Afghanistan, Israel, Saudi Arabia, Syria and Turkistan (Brooke, 1960; Balint and Balint, 1994). In Arabia, it is found principally in the Yemen. Khat grows in clumps, in beds or rows; often mixed in or interspersed with other crops, such as coffee and conifers, when cultivated. The primary areas for cultivation remain Eastern Africa – mainly Ethiopia, Kenya and to a lesser extent the Comoros, Madagascar and Tanzania - and the Arabian Peninsula, especially Yemen.

Fresh leaves of khat contain the alkaloids of the phenylpropylamine type of which the two psychoactive constituents are the stimulants cathinone (S-(−)-α-aminopropiophenone) and cathine (S,S-(+)-norpseudoephedrine). The psychoactive substances in khat act on two main neurochemical pathways – dopamine and noradrenalin. It has been suggested that cathinone, like amphetamine, releases serotonin in the CNS. Both these substances induce the release of dopamine from CNS dopamine terminals thereby increasing the activity of dopaminergic pathways (Kalix and Braenden, 1985). Cathinone is a
more powerful stimulant than cathine and is generally regarded as the most important element. However, cathinone is unstable in the presence of oxygen, oxidising at room temperature, and decomposes within a few days of harvesting or if dried (Griffiths et al, 1997). The stored product loses activity rapidly, becoming physiologically inactive after about 36 hours. For maximum potency, khat must be picked in the morning and chewed that afternoon (Baron, 1999).

Pharmacology

The psychotropic effects of khat are caused by the amphetamine-like compounds. Although amphetamine and cathinone act on different parts of the brain, they share common effects, including an interaction with the dopaminergic pathways (Pehek et al, 1990). Central stimulation by khat is manifested by euphoria, increased alertness, garrulousness, hyperactivity, excitement, aggressiveness, anxiety, elevated blood pressure, and manic behaviour. The simulation lasts for about 3 hours (Kalix, 1996). A depressive phase, including insomnia, malaise, and a lack of concentration, almost always follows (Al-Motarreb et al, 2002a; Hassan et al, 2002). True psychotic reactions occur with much less frequency than with amphetamines. This is most likely because of the self-limiting dose of khat, which does not permit blood vessels of the active compounds to rise high enough for toxic psychosis to occur. However, paranoid delusions, usually persecutory, have been seen (Jager and Sireling, 1994).

Several case studies have described Catha edulis-induced psychotic conditions (Giannini and Castellani, 1982; Critchlow and Seifert, 1987; McLaren, 1987; Yousef et al, 1995). Psychotic behaviour induced by khat use is sufficiently common in the khat-growing areas of eastern Ethiopia to have acquired a designated term – jezba (Kalix, 1988). In many instances, such psychoses disappear within a few days if khat consumption is stopped and/or anti-psychotic treatment initiated. However, recidivism typically results in a return of symptoms. However, the relationship between use of khat and psychiatric problems has been little explored by population-based studies.
It has been postulated that khat use can exacerbate psychotic symptoms in individuals with pre-existing conditions, and precipitate psychiatric disorders in vulnerable subjects (Yousef et al, 1995). Some commentators suggest that it is not clear whether khat use may cause a psychotic disorder in an otherwise healthy individual or trigger the onset of schizophrenia in an individual with high vulnerability to the condition (Poole and Brabbins, 1996; Phillips and Johnson, 2001). There have been suggestions that khat chewing exacerbates symptoms in patients with pre-existing psychiatric conditions; possibly caused by the sympathomimetic actions of cathinone on the CNS (Hassan et al, 2002). Odenwald et al (2005) propose that it is not khat consumption per se but specific patterns of its use are associated with the development of psychotic symptoms: early onset in life, excessive chewing (more than 2 bundles per day), and use as self-medication for war trauma-related symptoms. Other studies have found morbidity increased when daily consumption exceeds two bundles (Dhadphale and Omolo, 1988) or are dose-related (Critchlow and Seifert, 1987; Alem and Shibre, 1997). A recent review of the literature on the relationship between khat use and mental illness in general found that only clinical case studies revealed a causal connection between excessive khat use and psychosis. Such studies are unsuitable for providing evidence as to causality. A few case-control studies found an interaction between khat use and mental disorder but the findings were inconclusive. To date there have been no large-scale studies examining the relationship between dose-effect of khat and mental illness (Warfa et al, 2007).

There is some debate as to whether or not Catha edulis can actually cause dependence. Physical dependence to khat does not occur. The available literature suggests that it is likely that khat use has the potential to develop into dependence (Halbach, 1972). However it seems this is much less likely than dependence on stimulants such as amphetamine and more like the type of dependence seen with caffeine. The vast majority of people who chew khat do not use in a dependent fashion and there is no evidence of more widespread drug misuse amongst khat users. Development of tolerance to the effects of cathinone is more rapid than to that of amphetamine, and there is cross-tolerance between cathinone
and amphetamine (Schuster and Johanson 1979; Kalix, 1992). When a chronic khat chewer stops he feels hot, especially in the lower extremities, lethargic and with a great desire to chew khat in the first two days after cessation. Other documented withdrawal symptoms include nightmares and slight trembling, which appear several days after ceasing to chew (Al-Habori, 2005).

Context of khat use

It is important to ensure, so far as possible, that phenomena related to khat consumption are considered in the context of individuals and their environment(s). Other factors may impact on usage patterns and thus risks of both mental and physical health issues. A number of factors may be involved in the different patterns of use in the new countries of settlement of groups like Somalian refugees, especially younger generations. Economic, political, and social upheaval in such countries of origin has led to a large influx of traumatised individuals, often with little or no education. Young refugees enter an alien educational system, frequently exiting without qualifications that will afford them good employment opportunities. This leads to frustration and a lack of sense of direction. These factors make them vulnerable to the use of alcohol, tobacco and other psychoactive substances. Further, life in a new environment with different cultural and social contexts can be stressful and reinforce the need for the use of such substances (Cragg Ross Dawson, 2003).

Some research points to higher use amongst Somalis who are refugees or immigrants than in their native country. Bashford et al. (2003:23) suggest that cultural dislocation within the Somali refugee community may play a role in initiation into drug use, and “lead individuals to seek khat as a refuge” i.e. as a method of self-medication to mediate stress (although also prolonging and adding to existing stress). Alcohol and drugs play a role in ‘killing time’ for asylum seekers, and helping to forget past trauma (Dupont et al., 2005). Many start using before coming to the UK, but a significant proportion after their arrival – perhaps to help compensate for the sense of isolation, uncertainty about the future, unemployment, etc. (Griffiths.
et al., 1997; Tewolde et al., 2001; Gatiso and Jembere, 2001; Al-Osaimi et al., 2001; Mela and McBride, 2002; Patel et al., 2005; Kassim and Croucher, 2006).

In the past the use of this plant was geographically limited to these areas because it loses its efficacy after a few days. However, with developments in communications and transportation, especially by air, and the migration of khat-using communities to other continents, its use has spread over the last two decades. Large shipments of khat now arrive several times a week at major airports in western Europe, including the British Isles, as well to North America and Australia.

Scientific knowledge about khat

During the past 20 years or so, more has become known about the properties of khat, its pharmacology, physiological and psychological effects on humans. However, at the same time its reputation of social and recreational use in traditional contexts has hindered the dissemination of knowledge about its detrimental effects in terms of mortality. This aspect of the potential harms of khat has received little attention in the past and no systematic treatment or discussion. This paper focuses on this particular deficit and adds to the knowledge-base by reviewing the scant literature that does exist on mortality associated with the trade and use of khat. With khat being increasingly brought under domestic regulation in many countries, and its legal status being considered in others, it is important that such aspects are not overlooked so that balanced assessments can be taken based on evolving knowledge. It is hoped that this information will be noted by policy-makers, khat-using communities, and health professionals who deal with khat users. Deaths are associated with the trade in khat as well as from its consumption. Both these dimensions need to be understood.

This approach could also be applied to other substances, but is used here to generate awareness of the range of potential contexts in which khat cultivation, transportation, trade and consumption can lead to
fatalities. This paper is not just about pharmacology, toxicology, etc. but is about the wider phenomenon of deaths linked with a particular substance – khat.

The specific aims of this paper are to: (a) create a framework for discussion around the topic of khat-related death by generating a broad-based definition; (b) suggest dimensions of phenomena/aspects to consider without a priori theoretical paradigms/mindsets; (c) elucidate such dimensions by means of theoretical and actual examples.

Apart from our own paper looking at a series of 12 khat-related fatalities in the UK (Corkery et al, 2010), there are no known published reviews of, or statistics on, khat-related mortality howsoever defined. This is an important gap in the knowledge-base. This paper is an overview of what has been reported.

Methodology

We undertook a literature search of all relevant databases as well as the Internet to identify relevant reports and information on ‘khat-related’ mortality. The databases searched included: Medline, PubMed, psycINFO, Public Library of science (PLoS), Embase, INGENTA, African Journals on Line, Web of Science Citation Index, Allied & Complementary Medicine Database (AMED), the Cumulative Index to Nursing and Allied Health Literature (CINAHL®), Science & Social Science Citation Index, Cochrane, Toxnet and Google Scholar™ (2009) search engine. As khat is known by several names, these were also included as text search terms: cat, Catha edulis, chat, herbal ecstasy, jaad, khat, miraa, m(a)irungi, qaad, qat, tschat. The terms for the principal psychoactive substances were also employed: cathine, cathinone, and (nor)pseudoephedrine. The search terms in relation to mortality were: chok*, death, fatal*, intoxication, lethal*, mortality, overdose, poison*, suffocat*, toxic* - where * denotes wildcards. All of these sets of terms were systematically used in combination to conduct the searches.
These terms were used in English and also translated into other major European languages (e.g. Dutch, French, German, Italian, and Spanish); documents that were written in English or in other major European languages were examined, as well as abstracts written in those languages. Where necessary, on-line Internet translation engines (e.g. Babel Fish and Babylon) were employed to translate some of the search terms. Since it is possible that some reports and abstracts may not be available in non-Roman scripts, especially those relating to regions in Arabia, the search terms were converted to Arabic using babylon® before conducting the research; the results were then translated back into English. This method of on-line translation appears to have identified satisfactorily many relevant sources. The resultant titles and abstracts were then verified for inclusion. The time period covered was up to April 2011.

The database searches led to 111 hits in total. Papers in peer-reviewed journals, conference abstracts/posters, and studies from governments and non-governmental organisations that contained information on a link between khat and death were all eligible for inclusion in this review: specific incidents of deaths or fatalities; proposition of explanations for factors leading to death as a result of khat trade or consumption. Items were excluded because they: related to ‘cell death’; concerned original laboratory experiments; matched an irrelevant text word search; items identified did not relate to the khat plant but other topics such as ‘khat’ the ancient Egyptian concept of death, or were otherwise unconnected with the study parameters. Other reasons for ignoring items were where suggestions were vague, unsubstantiated, or misinformed. After exclusion, a final figure of 29 relevant items was obtained: of these, 14 were excluded because of duplication. Due to the paucity of relevant peer-reviewed items so identified, backward and forward citation tracking and cross-reference checking were also conducted. Non-peer-reviewed resources were also included in the Internet searches because exploratory work revealed that data-driven literature on our search topic was very limited. However, it has to be recognised that these were ‘snapshots’ at specific points in time, the items may be ephemeral and no longer
accessible. They may also have produced hits of grey literature that were not covered by the databases searched. This additional strategy led to a further 61 relevant articles being identified. So as not to be too restrictive in our research, it was also necessary to draw on media reports identified by Internet searches to illustrate various types of deaths associated with khat trade and consumption since these are not included in the above databases. Some of these references were provided by personal contacts.

There are risks in using anecdotal sources of information, as they are not always subject to scientific rigour in terms of reporting and interpretation. This has the potential to weaken the robustness of the information presented. However, such data provide a means of filling the lacunae in knowledge until such time as more robust information is generated. For a more detailed discussion of the use of such data see Corkery et al. (2010).

There is a range of deaths that can be directly and indirectly associated with khat, some of which can overlap. Different approaches could be employed to describe the links between khat and death, including a purely medical one concentrating on physiological, pharmacological and psychiatric issues. Since the aim of this paper is to broaden the theoretical conceptualisation of what might constitute a ‘death related to khat’, it is important to think outside the restricting confines of existing paradigms, especially those based on the ‘medical model’.

In order to assist an understanding of the different dimensions of this phenomenon it is necessary to derive themes or ‘motifs’ emerging from the results (Glaser and Strauss, 1967). This study looked at phenomena to describe types of deaths and their potential associations with khat, without being based on any a priori hypotheses or preconceptions, in order to see what emerged. Such a phenomenological approach is robust in demonstrating the presence of factors and their effects, but can only be tentative in suggesting their extent in respect of the sample population from which cases are drawn.
Findings

Following the journey of khat from its start to finish, from cultivation to consumption, provides a useful vehicle to identify such themes and to derive a taxonomy. Table 1 summarises the themes and categories that emerged from this process. The taxonomy is based on context(s) in which death occurred or was caused, e.g. its aetiology and manner of occurrence, based on the observation of reported incidences/events and post hoc abstraction from observed phenomena.

< Insert Table 1 about here >

Trade-related

Cultivation/production

There is concern that people who do not wash khat before consuming it will ingest pesticides (Date et al., 2004). Such pesticides contribute to cancers of the digestive system and kidney failure among khat chewers (IRIN, 2007). Deaths have been reported that were caused by poisoning following the ingestion of pesticides/fertilisers used in the cultivation of khat (see, for example, Almasamri, 2003; Al-thawra, 2002; Yemen Times, 2002).

Leaves are normally chewed fresh without any treatments (such as washing, soaking in hot water, or thermal treatments) to reduce the concentration of such chemicals, both in producing and consuming countries. For instance, an analysis of khat grown in different parts of Ethiopia in 2009 found excessive levels of pesticide residues in some producing areas (Daba et al, 2011). Diazinon levels in Jimma were found to be 462.6µg/kg (range 173.9 – 751.4µg/kg) compared to the level of 10µg/kg set for vegetables.
and fruit by the European Union (EU). The average total amount of DDT and metabolites was found to be 755.57 ±394.0 and 709.0 ±457.9 µg/kg respectively for Hararge and Butujera. Maximum levels found were between 240 and 1200 times the EU maximum recommended levels for foodstuffs (vegetables 10µg/kg, cereals 50µg/kg). The use of organochlorine pesticides has been attributed, in part, as contributing to the incidence of oral and oesophageal cancers amongst khat chewers (Al-Habori, 2005; Daba et al, 2011).

Some patients have been admitted to hospital in Yemen after chewing poisoned khat and within 3 days they have experienced jaundice, coma and sometimes death (Moghram, 2006). The sudden death of some previously healthy young people in Yemen while chewing khat sprayed with insecticide (such as dimethoate) may be explained by the accumulation of acetylcholine from the poison in the heart and brain involving bradycardia and cardiac arrest (Alhifi, 2006). No such cases have been reported in the UK. Clearly, farmers and their workers applying pesticides to khat and other inter-spaced crops may also be at risk. One example cited is that of a self-inflicted poisoning where a farmer ingested the pesticide to show others that it was harmless (Al-Hadrani and Thabet, 2000). Parasitic infections, such as fascioliasis, can also be transmitted through the consumption of contaminated khat (Doherty et al., 1995; Cats et al., 2000).

At least one violent death associated with disputes between khat-pickers over wages has been reported (Kamau, 2009). As is the case with other substances, khat can lead to the mind becoming overactive and exaggerating problems, with attendant misunderstanding or misinterpretation of the motives and actions of others. Chewers can become anxious, excited and display an aggressive attitude. Such effects on perception and mood during the intoxicated phase can last about three hours (Kalix, 1996). After this stage, a depressive phase almost always follows (Al-Motarreb et al., 2002a; Hassan et al; 2002). Users come to realise in the cold light of day that what they thought or believed during the khat session was incorrect and not thought through, and they often regret what they said or did.
Drug control

There is a body of evidence that supports the suggestion that drug law enforcement activities can contribute to drug market violence and increased homicide rates (Werb et al, 2011). For example, the imposition by Islamic militia in Somalia of bans on importation of and trading in khat has led to deaths. Opposition to the banning (by the Union of Islamic Courts) of importation of khat by air into Mogadishu in November 2006 by khat traders resulted in shots being fired to disperse protestors. At least one person was killed during the incident (Radio Somaliland, 2006; BBC, 2006). Similarly in November 2008 when a local militia took up arms to protest against a ban on khat by radical Shehab fighters imposing Sharia law in Eldher town (350 km north of Mogadishu) the ensuing clash led to the two lives being lost by both sides (Soomaalidamaanta, 2008). Breach of controls designating where khat trading should take place resulted in a trader being shot dead by a Hizbul-Islam militiaman in Afgoye. The trader had refused to sell the product in a designated market on the town’s outskirts (isltimes, 2010).

Distribution/marketing/wholesale/retail

Deaths related to the trade in khat have occasionally been reported by the media. For example, in the UK, several members of the Chohan family were murdered by gangsters attempting to take over an air-freight company which imported khat into the country; the gangsters sought to use the company as a front for trafficking illicit drugs within the European Union (Thompson, 2003). In the USA, a female vendor/dealer was killed and her money and plants stolen during a robbery (Adams, 2004; Collins, 2005; Lougawie, 2005; Padilla and McKinney, 2004; Zack, 2005a, 2005b). A similar fate befell a Somali dealer in England in 2002, when he was attacked and stabbed several times and left to bleed to death by
members of a rival clan who then stole his khat and sold it. However, it was unclear as to whether this was a fight over ‘turf’ or a robbery that went wrong (McBride, 2003).

Deaths have occurred much nearer the source countries. In the Somali region of Ethiopia a vehicle transporting khat was attacked by members of a different clan who intended to steal the cargo; in retaliation, the khat trader’s clan killed two relatives of the thieves (Devereux, 2006). Drug cartels in Kenya now control much of the transportation of khat from Kenya to Somalia; new entrepreneurs trying to establish themselves are permanently prevented from being in competition (Thatiah, 2008). Even consumers of khat face danger of death for failing to share it (AP Worldstream, 2005; Capitalfm, 2009; Sky News, 2005), or to agree a price (Devereux, 2006).

Deaths have also occurred in khat markets as a result of violence and accidents involving hand grenades (al-bab.com, 1998; Yemen Observer, 2003, 2006; US Dept of State, 2007). In another incident, a suicide bomber had been chewing khat immediately before detonating explosives that killed innocent by-standers (Al-Alaya’a and Al-Qiri, 2009). In other instances in markets, disputes over sales have led to children selling khat being killed (US Dept of State, 2007).

Transportation

Fatalities have occurred whilst khat has been transported by air. For example, in May 2004 a plane flying from Nairobi, Kenya, to Mogadishu, Somalia, carrying 55 bags of khat weighing 1.5 tons collided in mid-air with another plane. It crashed in an open field in Kenya killing the two occupants (Aviation Safety Network, 2004; Daily Nation, 2004). In August 2008, another aircraft en route with 5.4 tons of khat from Kenya to an airport 50km south-west of Mogadishu struck a telecommunications antenna during bad weather on its approach. It crashed killing all three crew members (Aviation Safety Network, 2008; Daily Nation, 2008). More recently, a plane carrying khat from Kenya to Mogadishu crash-landed shortly after
take-off from Wilson Airport in Kenya, having developed mechanical problems, killing the two co-pilots (Rajab, 2009, *Daily Nation*, 2009).

The need to transport khat to markets as rapidly as possible, so that the produce arrives on time and fresh, leads to those transporting them in vehicles to drive at dangerous speeds, on poor roads, in over-loaded, and poorly-maintained vehicles - often with fatal results (Ameen and Naji, 2001; *Yemen Observer*, 2003; Carrier, 2005). In many countries where khat is grown and chewed, there is a lack of reliable official road traffic accident data; furthermore, there is poor appreciation of the extent and significance of road accident fatalities. For Yemen, for example, it was possible to derive a causal model for such events taking khat into consideration (Ameen and Naji, 2001). An examination of available data for the period 1978-95 showed that khat consumption increases the risk of road traffic fatalities. The authors concluded that this was due to two factors: the anxiety and tensions that are experienced by khat chewers several hours after consumption; and, drivers consuming khat are often busy drinking water, smoking, or preparing and cleaning khat leaves whilst actually engaged in driving. These factors lead to decreased attention to driving and increased likelihood of accidents, especially on rural surfaced roads where drivers tend to drive faster.

Qualitative research from Ethiopia points to the chewing of khat as a major contributor to road traffic accidents, and hence potential fatalities. It increases driver confidence and vehicle speed, and simultaneously makes those drivers irritable and impairing concentration, with a misplaced sense of alertness (Eckersley et al., 2010). Research on African/Middle Eastern drivers stopped by German police on suspicion of driving under the influence of drugs found driving ability severely reduced by khat consumption in some cases, whereas in others it had no noticeable effect (Toennes and Kauert, 2004). However, because of concerns about the possibility of impairment, Ethiopian law implicitly makes it an offence to drive under the influence of khat (Eckersley et al., 2010). There is still no clear evidence from
population-based studies of a causal relationship between khat chewing and fatalities arising from road traffic accidents; however, the potential for khat to impact on driving and flying does exist.

**Law enforcement agencies**

In other instances, drivers have been killed in accidents trying to escape from law enforcement officers (Al-Jubelli, 2009). Smugglers also risk death when encountering guards at borders with countries which prohibit the possession, use and sale of khat (Yemeni News Agency, 2009b). It has been reported that in at least one country, capital punishment had been imposed in the past on a woman who was in possession of, and had used, khat (Fisk, 1995; Amnesty International, 2000).

**Consumption-related**

This section draws on a case-series of UK deaths associated with the use of khat, summarised in Table 2. Details of 13 out of the 15 cases have been previously described (Corkery et al., 2010).

< Insert Table 2 about here >

**Mechanical**

Through communication with various experts we have been informed of several mechanisms of deaths arising from khat use that are familiar to physicians and the public in Yemen, but no research has been undertaken or cases documented in the literature. The mechanisms described are: (a) a very strong irritation from substances in khat that causes a laryngeal spasm; and (b) whilst chewing, a piece of khat gets stuck in the trachea and cannot be dislodged by repeated coughing. Both types of incident can lead to choking and asphyxia. There have been isolated reports from Yemen of deaths resulting from individuals choking on khat. An 11-year old boy choked to death while playing soccer with a wad of khat
in his mouth (Al-Mugahed, 2008). A male patient got khat (smuggled into the hospital by his wife) stuck in his trachea and choked to death (Yemeni News Agency, 2009a).

**Toxicity**

We have been able to identify only one previously reported death from medical complications directly related to the use of khat in the literature or from surveys of contacts in the European Union, the eastern Mediterranean and Arabian Gulf regions. The solitary death was a case study reported over 60 years ago by Heisch (1945) who described the death from poisoning by *Catha edulis* of an elderly Arab trader in Wajir, Northern Kenya, after consuming an excessive amount of miraa stalks. The man was admitted to hospital and exhibited a number of symptoms including dilation of the pupil, cold sweats, convulsions, and hyperaesthesia. He fell into a coma (clinical observation) and died after 4 days. A post-mortem showed that the stomach was full of khat. Death could not be attributed to any cause other than khat consumption. Substantial advances in toxicological and medical knowledge have been made since then, but attribution can still be difficult in some cases. However, several deaths in the UK due to medical complications following the consumption of khat have been reported to or identified by the National Programme on Substance Abuse Deaths (np-SAD). These cases, including relevant discussion of mechanism of death for these as well as psychopathological factors, are described in more detail in a complementary paper (Corkery et al., 2010). Three of these were previously briefly described (Ghodse et al., 2006, 2007). No deaths relating to khat toxicity as overdose have been reported, but cases due to its toxic effects are now known.

It is thought that khat produces acute myocardial infarction in some patients and triggers such events in others, leading to fatal heart attacks. Ten out of twelve male Djiboutis treated for haemorrhagic stroke between September 2005 and March 2007 were daily consumers of khat, nine of whom also smoked (Benois et al., 2009). A clinical study of 120 Yemeni patients with myocardial infarction showed that khat
chewers were three times more likely to have experienced such an event than non-chewing controls (Alkadi et al., 2002). A multi-centre study of 8176 patients presenting to hospital with acute coronary syndrome in 6 Middle Eastern countries found that 11.4% were khat chewers (mostly of Yemeni origin and male). At admission, usually more than 12 hours after onset of symptoms compared to non-khat chewers, khat consumers had higher heart rate, Killip class and Global Registry of Acute Coronary Events risk scores. Khat-chewing was an independent risk factor for an in-hospital death (Odds Ratio 1.9, CI 95% 1.3-5.9, p < 0.001) and stroke (OR 2.7, CI 95% 1.3-5.9, P<0.01) (Ali et al, 2010). The incidence of myocardial infarction increases after chewing khat, demonstrating that it is a risk factor for this medical condition.

One mechanism suggested for this is the increased release of catecholaminens triggered by the cathinone content of khat, leading to hypertension and acute myocardial infarction (Al-Motarreb and Broadley, 2003; Al-Motarreb et al. 1997, 2002a and b, 2005; Croles et al., 2009; Health Canada, 2007). Cathinone, and to a lesser extent cathine, appears to act on the beta-1 adrenoceptor causing increased systolic blood pressure and pulse rate. However, these effects can be prevented with the use of beta-blockers such as atenolol (Hassan et al., 2005). Khat use is associated with acute coronary vasospasm and myocardial infarction (Al-Motarreb et al., 2002a, 2002b). Coronary vasospasm, induced by the cathinone in khat, causing vasoconstriction may occlude coronary arteries sufficiently to precipitate myocardial infarction.

Increased thrombogenicity could also be a key factor for myocardial infarction (de Ridder et al., 2007). Differences in diurnal patterns of acute myocardial infarction have been observed between non-khat users and khat users, with peak presentation amongst the second group shifting to the late afternoon and evening, coinciding with khat chewing sessions – 2pm to midnight (Al-Motarreb et al., 2002b; Al-Habori, 2005). It has been suggested that the cathinone content of khat is an independent dose-related risk factor for the development of acute myocardial infarction; heavy chewers having a 39-fold increased risk (Al-
Motarreb et al., 2005). The risk of acute myocardial infarction is also increased when the khat ‘session’ lasts more than 4 hours. Daily khat chewers are significantly more at risk, as are those whose use is recent and chronic. One study found that in those suffering from a myocardial infarction, khat use was associated with earlier age at presentation (Al-Motarreb et al., 2002b).

The np-SAD (Corkery et al, 2010) has been notified of the death of a young adult Somali male khat-user which was attributed by the pathologist to high norephedrine levels from khat inducing left ventricular failure with development of pulmonary oedema, which was regarded as mode of death. Another young Somali male who had consumed khat and alcohol, and had assaulted another male, collapsed and had difficulty breathing. Cardio-respiratory resuscitation was started but on the way to hospital by ambulance he suffered a cardiovascular event and was pronounced dead. There was no underlying disease which could be attributed as having contributed to death. The pathologist could not exclude a contribution of khat to the cardiovascular event with either an infarction or electrical instability (arrhythmia) causing death. The positive toxicological findings for cathine, cathinone, and norephedrine meant that it was not possible to attribute death to Sudden Adult Death Syndrome. We are also aware of another, non-notified, death involving khat-induced myocardial infarction.

Khat can also produce acute myocardial infarctions by direct coronary vasoconstriction (Al-Motarreb and Broadley, 2003). Rat studies conducted by Admassie and Engidawork (2011) suggest that frequent and chronic khat consumption may lead to transient but repetitive coronary vasoconstriction causing a reduction in blood flow and myocardial damage. Myocardial oxygen demand is increased by increased heart rate, force of contractility and systolic tension which could not be associated with heightened levels of CK and AST – even though sub-chronic khat consumption, as these can lead to multi-organ damage. This is in line with findings of human liver and kidney damage reported by Chapman et al (2010) and Fallatah and Akbar (2010). Khat-induced ischaemic strokes have been identified in a number of case-reports (Bruno, 2003; Vanwalleghem et al., 2006; de Ridder et al., 2007). Such cardiac events can clearly
lead to fatal outcomes, as evidenced by a recent study in the Middle East which found that Yemeni khat-chewing patients with Acute Coronary Syndrome had higher in-hospital complications including Stroke and death (Al-Suwaidi, 2011).

Deaths and haemiplegia (paralysis of half of the body) because of meningeal haemorrhages are reported. The causal mechanism is ruptures of aneurisms following circulatory “coups de fouet” (rupture of the planteris muscle accompanied by sharp disabling pain) engendered by the use of khat (Le Bras and Frétillère, 1965). A similar reported mechanism is cerebral haemorrhage, cardiac arrest and pulmonary oedema (Health Canada, 2007). Cerebral haemorrhagic softening has also been described, although such cases are relatively rare (Le Bras and Frétillère, 1965). Le Bras and Frétillère (1965) also note that when an individual is experiencing hyperthermia caused by khat consumption, death can occur following strenuous physical effort (such as playing sport).

Administration of *Catha edulis* extracts has demonstrated a capacity to induce cytotoxic effects in liver and kidney cells, as well as cell death in various human leukaemia cell lines and peripheral human blood leukocytes (Al-Habori, 2005). Studies on animals have shown that khat is hepatotoxic with increases in liver enzymes and there has been histopathological evidence of acute hepato-cellular degeneration (Al-Mamary et al., 2002). However, it has only been recently that clinical human case reports have been published (Brostoff et al., 2006; Saha and Dollery, 2006). One case report suggested that hepatitis was probably due to right heart failure which in turn was possibly due to direct khat toxicity - as it did not respond to withdrawal of statins or proton pump inhibitors (Saha and Dollery, 2006).

Until recently, there have been few documented cases of khat causing liver failure. However, evidence is now emerging of an association between khat consumption and liver problems, sometimes necessitating transplant, that result in death (D’Souza et al., 2006; McCune et al., 2007; Chapman et al., 2010; Peevers et al., 2010; Stuyt et al., 2011).
A study of adults in Yemen with acute sporadic hepatitis with jaundice found that no viral cause was identified in half (51%) of the cases. Whilst there may be an unknown virus responsible or environmental toxins such as pesticides in khat leaves (Gunaid et al., 1997), liver function test abnormalities resolve when khat chewing is stopped. Young Somali males who chew khat appear to be particularly at risk of autoimmune hepatitis (D’Souza et al., 2006; McCune et al., 2007) and liver disease (McCune et al., 2007).

Chapman et al (2010) report that five out of six patients (four male, two female) had a history of previous unexplained hepatitis, all chewed khat and had been advised not to it all but resumed chewing before presenting a second time with liver problems. Having been thoroughly screened, other causes of liver injury were excluded. All patients had multi-lobular necrosis and two had a background of chronic liver disease. These findings suggest that long-term use of khat may be associated with repeated episodes of sub-clinical hepatitis, with development over time into chronic liver disease. One patient subsequently died. There were no significant features to suggest an autoimmune or immunoallergic process in these cases, it was concluded that khat consumption was the aetiological cause of the liver injury in these individuals. Whilst the mechanism of khat-related hepatotoxicity remains uncertain, the presence of a high concentration of cathinone being detected in the liver of one patient three weeks after their last use of khat, suggests that the accumulation of the drug may be of significance.

Stuyt et al (2011) report 6 cases of long-term khat chewers who migrated from East Africa to the Netherlands that presented with liver injury. Four presented with jaundice and hepatitis; one of these and two others had variceal bleeding and hepatic encephalopathy; two had clinical ascites. Three had auto-antibodies and elevated serum IgG. Other causes of liver disease were excluded. Three patients died from spontaneous bilateral peritonitis, sepsis and one case receiving a liver transplant died from pulmonary hypertension apparent during and after transplantation. These findings support the argument
that long-term khat chewing can produce repeated episodes of hepatitis (probably due to immuno-allergic or idiosyncratic) leading to fibrosis and cirrhosis.

np-SAD is aware of five deaths involving liver failure due to exposure to long-term khat consumption, three of which have been notified to np-SAD (Corkery et al, 2010). Of these, one case developed jaundice diagnosed as khat-induced hepatotoxicity which developed into recurrent khat-induced hepatitis, with pulmonary oedema and an enlarged heart; his condition developed into hepato-renal syndrome and sub-dominant liver failure (Gadelrab, 2006). A second case was admitted to hospital with jaundice, night sweats, and pyrexia which led to sub-acute liver failure; this was due to auto-immune hepatitis caused by his khat use. A third case was admitted to hospital suffering from fulminant hepatic necrosis due to khat toxicity; he had a liver transplant which was unsuccessful, and died of a pulmonary haemorrhage and multi-organ failure. Another case has been reported in the UK (McCune et al., 2007; Stein, 2007); and a further death was registered in October 2010 (personal communication to lead author from Frank Dixon, General Register Office for Scotland on 3 May 2011).

Cancer related to khat use

Keratotic white lesions are present in the mouths of about one-fifth of khat-chewers (Ali et al., 2004). This change in pathology is recognised as pre-cancerous and may develop into oral cancer (Goldenberg et al., 2004; Ahmed et al, 2010). Buccal epithelial cells experience genotoxic effects in a dose-related way in khat-chewers; this suggests that oral malignancies can be contributed to by khat (Kassie et al., 2001).

There is an increased risk of oral carcinoma amongst khat chewers (Drake, 1988; Soufi et al., 1991; Gunaid et al., 1995), especially when associated with alcohol and tobacco consumption (Soufi et al., 1991; Kassie et al., 2001). Khat has been found to be a significant contributory factor in the high prevalence of head and neck squamous cell carcinoma in the Yemeni population (Nasr and Khatri, 2000).
There may be a synergistic effect created by the consumption of alcohol, khat, and tobacco, leading to the development of changes in cells that can become cancerous (Kassie et al., 2001).

Although oral cancers due to khat chewing occur and do lead to death, there do not appear to be any published statistics on such deaths. The few studies that have examined head and neck cancers and possible associations with khat-chewing have small sample sizes and the effects of using other substances are not controlled for. There is a need for large-scale case-control studies to gauge the strength of this possible association.

**Endocarditis associated with khat chewing**

A death from endocarditis caused by infection by a streptococcal bacterium *Gemella morbillorum* in a Djibouti male has been reported (Massoure et al., 2010). The bacterium had been ingested through the patient chewing khat. Despite treatment with anti-biotics, he developed pulmonary oedema requiring assisted ventilation, became asystole, and died 8 days after being hospitalised.

**Reproductive health issues**

The effects of khat consumption on the reproductive system, pregnancy and neonates have also been noted. Khat is genotoxic, having teratogenic effects on the foetus if regularly consumed by pregnant mothers. Neonates have low birth weights – a risk factor for perinatal and young infant death (Ghani et al, 1987; Eriksson et al, 1991; Mwenda et al, 2003). Currently chewing lactating women have been found to excrete norpseudoephedrine in their breast milk, and traces were found in the urine of a breast-fed infant (Kristiansson et al, 1987). Cardiovascular complications amongst pregnant khat-chewing females have been reported: chest pains, tachycardia and hyperthermia (Kuczkowski, 2005).
Psychopathological

Khat use may be less psychotogenic when integrated within its original cultural context than when taken in an unfamiliar setting after emigration. Many of the published reports relating to khat-induced psychoses concern individuals who have emigrated to Europe or North America, become culturally isolated and are in vulnerable socio-economic situations. However, this pattern may reflect a publication bias (Manghi et al., 2009). This is due to several factors. The ethnic populations in this region using khat have increased substantially over the past two decades, particularly because of the civil war in Somalia. In traditional khat producing and consuming areas, where health care services were/are lacking, those with psychotic symptoms would be confined to the home by family members (Cox and Rampes, 2003). Increasing reports of cases of khat-related psychosis in Europe and Australia (Stefan and Mathew, 2005), have led to more familiarity with ‘Khat syndrome’ and consequent concern being expressed about it. In part, this may be attributable to the influx of refugees from war-torn countries such as Somalia, many of whom are suffering from trauma and who use khat (for example, see Bhui et al., 2003). It is likely that refugees from strife-torn parts of Somalia and elsewhere are likely to experience greater use of khat than previously realised (Odenwald et al., 2007). This may be due to the added pressures experienced in living in a refugee community, distant from traditional patterns of life, leading to increased changes in behaviour (Hughes, 2000). In most European countries such individuals would be presenting to health services and their symptoms detected and they would be referred for treatment. The stressful life (and death) situations which many of these individuals had escaped have left psychological scars and damage that are self-medicating by means of khat.

Murders have been committed by individuals suffering from psychotic states aggravated or caused by khat. For example, there have been reports in the UK of individuals who, having experienced a psychotic illness as a result of khat consumption or where such consumption exacerbated an existing mental illness or depression, caused suicidal ideation, killed others and/or themselves (see, for example, Busby, 1987;
BBC, 2007; Pantelis et al., 1989; Alem and Shibre, 1997; *The Sun*, 2007). There has also been at least one further case of murder where the assailant was believed to be under the influence of alcohol and khat and inadvertently killed an innocent bystander instead of the intended victim (Nettleton, 2001a, 2001b). The use of khat can lead to aggressive behaviour (Pantelis et al., 1989; Alem and Shibre, 1997; Arnold et al., 2008; Gelaye et al., 2008; Philpart et al., 2009) and impaired judgement or making mistakes (Walter, 1996).

Khat has had an effect on the fighting in the Somali civil war and combat related to military operations in that country (Randall, 1993). Aggression can be increased and notions of personal supremacy enhanced by khat use. Fighting over the delivery of khat at airports and children being shot over a 7kg bag of khat have been reported (Stevenson, 1992).

The np-SAD has been notified of three cases of suicide, including one following a homicide and an accidental drug overdose (BBC, 2005; Edwards, 2005; *Hackney Gazette*, 2006; Lefley, 2005; Meehan, 2006; *Somaliuk*, 2006; Turner, 2006; Corkery et al, 2010), where the individual had a history of psychiatric problems, typically schizophrenia aggravated by khat use or a khat-induced psychosis, or was suffering from depression. The Programme is also aware of another such suicide that has not been notified.

**Impaired judgement/accidents**

There is concern that fatalities are high among tired lorry drivers due to accidents - especially those from Ethiopia, who chew khat to stay awake over the course of several days whilst they travel extremely long distances in Eastern Africa (Ehrenkranz and Kushner, 2008; Harden, 1987; Mutongi, 2006). An assessment of road traffic accidents in Yemen during 1978-95 found a high correlation between road traffic fatalities and consumption of khat, as measured by the area under cultivation (Ameen and Naji, 2006).
Although khat has a stimulatory effect during the first few hours of consumption, the study also found that its use can increase the risk of road traffic accident fatalities for two reasons. First, anxiety and tensions that occur several hours after use can affect perception and risk assessment. Second, often those taking khat whilst driving will be busy drinking water, smoking cigarettes, or preparing and cleaning the leaves. These divert the attention of the driver and are likely to cause accidents especially on rural roads where drivers tend to drive fast. Le Bras and Frétillère (1965) note the neurological effects that khat can have, particularly on gross motor co-ordination and impaired vision, leading to accidents. Adverse effects on perceptual-visual memory and decision-speed have also been noted (Khattab and Amer, 1995). It has also been suggested that khat may have similar effects to amphetamine on driving ability, i.e. reduced alertness and awareness coupled with increased risk-taking (Al-Samarraie et al., 2007).

The np-SAD has been notified of two cases of individuals dying in accidents involving motor vehicles. In both cases the deceased had consumed khat and alcohol (Corkery et al, 2010). One was a taxi-driver returning home after work, who lost control of his vehicle, collided with a lorry and sustained fatal injuries. The other was a pedestrian who attempted to cross a road, stumbled in the middle of the street, and was run over by a taxi (Daily Echo, 2008a, 2008b, 2008c). A lack of motor co-ordination and impaired vision (Le Bras and Frétillère, 1965) may also have contributed to these accidents.

In these two cases, khat had been consumed with other psychoactive substances. This led to impaired judgement resulting in traumatic road traffic accidents. It may be difficult to disentangle the individual contributions of each substance, but the potential contribution of khat cannot be excluded. In a further case, the decedent died from heroin intoxication, although he had also consumed khat. This fatality may also have involved impaired judgement or perception of risk, as well as demonstrating the possible synergistic effect of combined khat and other psychoactive substance consumption. The important feature here is not that the individual died of heroin use, but that he had also consumed khat.
The role of khat in a death reported to the np-SAD that resulted from injuries sustained in a fall from a height is unclear (Corkery et al, 2010). An adult Somali male, was found dead on a concrete surface below a fifth-floor flat at an address which he had just left after purchasing khat. The post-mortem examination showed no natural disease to account for death. There were no marks of restraint or of classical defence-type injuries. Khat was found in the mouth, oesophagus, and stomach but it is unclear as to its role, although it is possible that his khat-induced psychosis may have contributed to or influenced his fall from a height. The police could find nothing else to suggest it was anything other than suicide. Alternatively, his judgement may have been impaired by consuming khat. In 2009 a 32 year-old Somali female fell or jumped from a block of flats in an English town during a khat-chewing session with her husband and a friend. She had a history of mental health issues, as well as drug and alcohol abuse. The coroner concluded that the khat's psychotic effects or hallucinations could have led to an error of judgement by the decedent (BBC, 2011).

We noted earlier that violence amongst khat sellers, who were also chewing it themselves whilst retailing it, can break out and lead to death, contributed to in part by the intoxicating effects of the plant. Violence can be demonstrated not only by khat consumers but also towards them as a result of their behaviour whilst under its influence. These two aspects can be illustrated by two recent UK cases (Corkery et al, 2010).

A 22-year old Somali male, who had been drinking alcohol and chewing khat, attended a party in Bristol and assaulted the host. When the latter tried to be reconciled with the young man, he was again assaulted and threatened in a restaurant where his assailant was drinking. The young man became aggressive and started acting strangely, demanding more alcohol. He suddenly lost his balance, collapsed, striking his head. He started having difficulties breathing. On the way to hospital by ambulance he suffered a cardiac event that led to death. His khat consumption could not be ruled out as a contributory factor (thisisbristol, 2010).
By contrast in London, a 20-year Somali youth got on a bus with a relative, after taking alcohol, cannabis, and khat. He began spitting on the floor of the vehicle, and a passenger remonstrated with him about it. When the youth got off the bus he spat at this passenger. A short while later, the victim of the spitting turned up in two cars with family members and assaulted the youth. He received severe head injuries (consistent with impacts from heavy blunt instruments) and died in hospital. His assailants were subsequently convicted of his murder (Ealing Times, 2008; Met Police, 2008). It is thought that the substances the youth had consumed may have impaired his judgement and risk assessment, increased his aggressiveness, reduced his self-control, and led to his anti-social behaviour which was sufficiently offensive for his victim to seek retaliation – with fatal results.

Discussion

This paper has demonstrated that a wide range of associations can be posited legitimately to exist between khat and death. Some of these are more clear-cut and obvious, as with medical causes associated with consumption, as well as less direct or not so obvious, such as those associated with its cultivation and trade. For the purpose of this paper and its principal aim of creating a framework for discussion around the topic, a ‘khat-related death’ is defined as meeting one or more of the following criteria (which are not necessarily mutually exclusive):

- ante- and/or post-mortem toxicology positive for cathine, cathinones, or other metabolites indicating consumption of khat;
- acute death occurring as the direct consequence of consuming khat;
- death occurring as the consequence of long-term use of khat leading to adverse physiological conditions;
- death occurring as the result of khat use aggravating a pre-existing or triggering mental health conditions leading to fatal self-harm;
• death resulting from impaired judgement caused by khat consumption;
• death associated with the cultivation, transportation or trade in khat – howsoever caused.

However, this definition is not carved in stone and is open to future refinement, as and when more objective evidence and scientific data are collected. It has to be recognised that this working definition, although based on a phenomenological approach, is nevertheless a social construction like all other drug-related deaths (Cruts, 2000).

The contribution of khat to deaths is varied; and in some cases, can be difficult to categorise. Deaths associated with trade, marketing, wholesale and retail, can equally occur with any other illicit substance. Their inclusion here is deliberate: they are part of the wider pattern of mortality associated with the consumption of khat. The association of khat with psychiatric disorders or psychopathological factors is evident in some cases, contributing to suicide and even homicide.

Apart from poisoning from insecticide, traumatic deaths can be a feature of the cultivation, transportation, and trading activities undertaken by distributors and sellers. Violence can also occur in other settings. For example, psychopathological effects – (i) impaired judgement leading to accidents and violence, (ii) causing or exacerbating psychoses or causing depression leading to suicide and even homicide.

Physiological effects encompass (a) mechanical problems e.g. choking on pieces of the plant.; and (b) toxicity (i) causing heart problems leading to fatal heart attacks, and (ii) liver failure. There is now a fairly well developed literature on the cardiac and circulatory problems caused by khat, especially with regard to myocardial infarction. There is now evidence that such consequences can be triggered by high levels of the principal psychoactive substances in khat i.e. left ventricular failure and acute pulmonary oedema.
The physiological effects of khat consumption are quite clear in a number of cases. The np-SAD cases referred to above are thought to be the first reported (as distinct from recorded) cases of death due to khat toxicity since the case reported by Heisch (1945). In one case the role of norephedrine/norpseudoephedrine producing adrenaline-like actions led to an increased likelihood of myocardial infarction (Saha and Dollery, 2006), leading to left ventricular failure and pulmonary oedema. Khat chewing is a recognised risk factor for myocardial infarction (Al-Motarreb et al. 1997, 2002a and b, 2005; Croles et al., 2009; Health Canada, 2007). Khat can also cause cerebral haemorrhage (Bentur et al., 2008), cerebral infarction (Mujlli et al., 2005) and contribute to cardiomyopathy (Saha and Dollery, 2006); both of which have high associated risks for morbidity and mortality.

Until recently, there have been few documented cases of khat causing liver failure. However, evidence is now emerging of an association between long-term khat consumption and liver problems, sometimes necessitating transplant, and now of death. It is possible that some deaths that are in fact attributable to khat consumption are going undetected because symptoms such as myocardial infarction and liver failure are not recognised as being associated with the substance. The cases reported here from the np-SAD of “khat-induced” liver failure [the Corkery khat liver condition] are consistent with recent findings in respect of hepatitis (Brostoff et al., 2006; D’Souza et al., 2006; Peevers et al., 2010) and liver disease (McCune et al., 2007). Such cases highlight that khat is a potential hepatotoxic agent and that this phenomenon needs further investigation and documentation. The determination of the aetiology of hepatitis should take into account the use of recreational drugs in all socio-cultural groups (Peevers et al., 2010). Patients presenting with such symptoms from ethnic groups associated with khat use should be asked for a full substance use history. The administration of nephrotoxic herbal remedies, including khat, should also be borne in mind when considering the origin of renal failure – again in all cultural groups. This awareness is also important in the assessment of coronary ischaemia amongst young male urban khat chewers with few other risk factors for coronary disease.
Impaired judgement or poor risk perception may play a part in fatalities (Walter, 1996), especially where violence may also be a factor. These factors may be made worse by the consumption of other psychoactive substances in conjunction with khat.
Conclusions

The information presented here illustrates some of the key issues related to the trade in and consumption of khat consumption. This paper is believed to be the first attempt to identify and classify mortality associated with or related to Catha edulis. Death can occur at any stage from cultivation to consumption. Some of the factors can occur in combination, and it can be difficult sometimes to disentangle them. This paper not only seeks to establish potential causality between khat and death but also the type of associations that can be posited both theoretically and drawn from observation. However, this is more than a mere taxonomical exercise. A complementary paper examines in more depth causality and other associations in specific case studies and reflects on the quality of available evidence (Corkery et al., 2010).

The use of case-studies and reports of individual incidents mean that the findings presented here are methodologically limited in quantitative terms, with consequent implications for analysis and interpretation. However, the research was undertaken systematically and provides a basis for the development of concepts and theories that can be refined by further, preferably population-based, research. This process could also identify other possible categories and dimensions which would need to be assimilated or incorporated into the model presented here. A similar approach to taxonomy could be applied to other substances. These would also need adaptation and refinement. For example, if applied to tobacco, it would need to take account of those manually handling the crop being poisoned by the plants because of nicotine being absorbed through the skin giving rise to toxic and even fatal results.

There are probably more deaths that have occurred in the past and which are currently happening but that are not identified as being related to the use of khat. This has meant that there is a demonstrable lack of documented cases in the literature, as well as nothing by way of quantitative data. These gaps in knowledge need to be filled employing systematic and scientifically-based approaches. This will lead to a
much better understanding of the potential risks of death associated with the use of khat, based on empirical observation.

In the meantime, the dangers arising from the use of khat and its psychoactive constituents need to be brought to the attention of those in producing/growing countries, and those countries/regions that have become hosts to ever-increasing communities from these countries. This will need to be specifically focused on target groups using a variety of media, including a variety of languages. For example, khat is a major risk factor for acute heart attack (Al-Motarreb et al., 2005) and needs to be considered in any ischaemic assessment of individuals from khat-consuming communities presenting to physicians with cardiac problems.

The fatal case studies detailed above need to be borne in mind in future reviews of khat and the physiological and mental health risks associated with its use. These risks appear to be greater than previously realised. There is need for routine and systematic research of khat-using populations in respect of morbidity and mortality arising from the use of pesticides in the cultivation of the crop.

Khat research is still in its infancy, and there is little robust information on the subject (Odenwald et al., 2011). Furthermore, the lack of negative health results for khat in the literature should not lead to complacency or an assumption that khat use is free from toxic consequences (Carvalho, 2003). This absence of negative reports is due to a lack of population-based studies (Odenwald et al., 2005), particularly in respect of the toxicology of Catha edulis. Whilst anecdotal reports are informative, systematic investigations are needed to determine the incidence and prevalence of ill-effects of khat use. Only then can the best methods of supplying preventative and therapeutic interventions be considered in an informed way.
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Conflict of interests
We are unaware of any conflict of interests.
Ethical approval

No ethical approval was required for this review
References


Accessed on 19 April 2009.


Table 1: Taxonomy of ‘khat-related’ mortality

<table>
<thead>
<tr>
<th>Directness</th>
<th>Type of association</th>
<th>Mechanism</th>
<th>Example</th>
</tr>
</thead>
<tbody>
<tr>
<td>Trade-related</td>
<td>Cultivation/production</td>
<td>Poisoning</td>
<td>Disagreement over wages =&gt; homicide – possibly aggravated effects of khat chewing clouding judgement and triggering aggression</td>
</tr>
<tr>
<td></td>
<td>Transportation</td>
<td>Fatigue</td>
<td>Long hours and driving at high speed =&gt; loss of concentration (can be in association with khat use to keep awake) leading to accidents</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Loss of concentration</td>
<td>Distraction whilst preparing khat for chewing whilst driving =&gt; accidents</td>
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<tr>
<td></td>
<td>External factors</td>
<td></td>
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<tr>
<td></td>
<td>High speed</td>
<td></td>
<td>High speed trying to escape from law enforcement officers =&gt; collision =&gt; fatal injuries</td>
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<tr>
<td></td>
<td>Distribution/marketing/wholesale</td>
<td>High speed &amp; mechanical failure</td>
<td>High speed, over-loaded vehicle, faulty brakes =&gt; failing to negotiate bend in road =&gt; fatal accident</td>
</tr>
<tr>
<td></td>
<td>Retail</td>
<td>Disputes over price</td>
<td>Homicide – possibly aggravated effects of khat chewing clouding judgement and triggering aggression</td>
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<tr>
<td></td>
<td></td>
<td>Violence in khat markets/terrorism</td>
<td>Detonation of hand grenade in khat market; militia firing guns indiscriminately into crowds; suicide bomber exploding explosives immediately after chewing khat</td>
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<td></td>
<td>Law enforcement agencies</td>
<td>Execution</td>
<td>Beheading for possession and use of khat</td>
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<td></td>
<td>Drug control</td>
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<td></td>
<td></td>
<td>Opposition to imposition of ban on khat importation</td>
<td>Protestors against ban fired on by militia enforcing ban</td>
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<tr>
<td></td>
<td></td>
<td>Breach of drug controls</td>
<td>Trader killed by militia enforcing khat-selling restrictions</td>
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<td></td>
<td>Consumption-related (medical)</td>
<td>Physiopathological</td>
<td>Choking on leaves/twigs or airway obstruction =&gt; asphyxia or cardiac arrest</td>
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<td></td>
<td>Neurological</td>
<td>Lack of motor coordination, shaking</td>
<td>Reduces control =&gt; accidents</td>
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<tr>
<td></td>
<td>Psychopathological</td>
<td>Causing and/or exacerbating psychosis and/or depression</td>
<td>Suicide and/or homicide, accidental overdose</td>
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<td></td>
<td></td>
<td>Increases aggression and notions of personal supremacy</td>
<td>Violence in Somalian civil war =&gt; homicide; fighting over khat air-freight deliveries</td>
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<tr>
<td></td>
<td></td>
<td>Impaired judgement/assessment of risk =&gt; accidents</td>
<td>Road traffic accident</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Impaired judgement/intoxicating effects</td>
<td>Misunderstanding/misinterpretation of others’ motives/actions =&gt; offending others =&gt; retaliation involving fatal assault</td>
</tr>
<tr>
<td></td>
<td>Other medical contributory factors</td>
<td>Used with other psychoactive substances and/or positive toxicology</td>
<td>Role may not be clear</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>Cerebral haemorrhage</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>Ischaemic stroke</td>
<td></td>
</tr>
</tbody>
</table>

Note: This is a theoretical framework based on the findings of this research, and is subject to revision as new information is reported. Its aim is to provide a starting point for debate and refinement.
Table 2: Main characteristics of np-SAD ‘khat-related’ deaths, UK, 2004-9

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>Case 1</th>
<th>Case 2</th>
<th>Case 3</th>
<th>Case 4</th>
<th>Case 5</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Age (years)</strong></td>
<td>39.1</td>
<td>32.5</td>
<td>40.2</td>
<td>47.0</td>
<td>27.3</td>
</tr>
<tr>
<td><strong>Gender</strong></td>
<td>Male</td>
<td>Male</td>
<td>Male</td>
<td>Male</td>
<td>Male</td>
</tr>
<tr>
<td><strong>Ethnicity</strong></td>
<td>Somali</td>
<td>Somali</td>
<td>Somali</td>
<td>Somali</td>
<td>Somali</td>
</tr>
<tr>
<td><strong>Marital status</strong></td>
<td>Separated</td>
<td>Divorced</td>
<td>Married</td>
<td>Married</td>
<td>Single</td>
</tr>
<tr>
<td><strong>Living arrangements</strong></td>
<td>Alone</td>
<td>With sibling</td>
<td>With partner &amp; children</td>
<td>Alone</td>
<td>Alone</td>
</tr>
<tr>
<td><strong>Employment status</strong></td>
<td>Unemployed</td>
<td>Invalidity/sickness benefits</td>
<td>Unemployed</td>
<td>Unemployed</td>
<td>Unemployed</td>
</tr>
<tr>
<td><strong>UK residency (years)</strong></td>
<td>11</td>
<td>15</td>
<td>Not known</td>
<td>11</td>
<td>3</td>
</tr>
<tr>
<td><strong>Significant medical history</strong></td>
<td>No</td>
<td>Yes</td>
<td>No</td>
<td>Yes</td>
<td>No</td>
</tr>
<tr>
<td><strong>Known psychiatric history</strong></td>
<td>Yes, at least 11 years</td>
<td>Yes</td>
<td>No</td>
<td>Yes</td>
<td>No</td>
</tr>
<tr>
<td><strong>Known khat using history</strong></td>
<td>Yes</td>
<td>Yes</td>
<td>Yes</td>
<td>Yes</td>
<td>Yes</td>
</tr>
<tr>
<td><strong>Evidence of using khat prior to death</strong></td>
<td>No</td>
<td>Yes</td>
<td>Yes</td>
<td>No</td>
<td>Yes</td>
</tr>
<tr>
<td><strong>Place of death</strong></td>
<td>Underground station</td>
<td>Car park</td>
<td>At home</td>
<td>At home</td>
<td>At home</td>
</tr>
<tr>
<td><strong>Role of khat in death</strong></td>
<td>Paronid psychosis associated with history of khat use =&gt; traumatic suicide (none in body at death)</td>
<td>Possible suicide/accidental fall whilst judgement impaired (found in body)</td>
<td>Possible history of excessive use =&gt; traumatic murder and traumatic suicide</td>
<td>Paronid schizophrenia exacerbated by khatting =&gt; accidental overdose</td>
<td>Ingestion of khat =&gt; high norephedrine levels =&gt; left ventricular failure =&gt; pulmonary oedema</td>
</tr>
<tr>
<td><strong>Toxicology done</strong></td>
<td>Yes</td>
<td>Yes</td>
<td>Yes</td>
<td>Yes</td>
<td>Yes</td>
</tr>
<tr>
<td><strong>Toxicology results</strong></td>
<td>Blood – negative; urine - negative</td>
<td>Blood - cathinone 104µg/L; norephedrine/norpseudoephedrine (total) 154µg/L; zuclopenthixol 123.8µg/L</td>
<td>Blood - cathinone &quot;low&quot;, cathine &quot;moderate&quot;; consistent with &quot;recreational use&quot;. Urine – Cathine and cathinone detected.</td>
<td>Blood – negative. Urine – negative.</td>
<td>Blood - cathinone &lt;50µg/L; norephedrine/norpseudoephedrine (total) 1000µg/L. Urine – norephedrine/ norpseudoephedrine detected</td>
</tr>
<tr>
<td><strong>Cause(s) of death</strong></td>
<td>1a Severe chest, abdominal and pelvic injuries</td>
<td>1a Multiple injuries [severe multiple injuries, impact onto a firm surface, fall from a height]</td>
<td>1a Hanging 1b Stab wounds to neck</td>
<td>1a olanzapine toxicity</td>
<td>1a Acute pulmonary oedema [left ventricular failure, high norephedrine levels] 1b Ingestion of khat</td>
</tr>
<tr>
<td><strong>Verdict</strong></td>
<td>Act of self-harm</td>
<td>Open</td>
<td>Suicide</td>
<td>Accidental</td>
<td>Narrative</td>
</tr>
<tr>
<td><strong>Coroner’s area</strong></td>
<td>West London</td>
<td>Inner West London</td>
<td>Inner North London</td>
<td>Cardiff &amp; Vale of Glamorgan</td>
<td>Northern London</td>
</tr>
</tbody>
</table>

Note: These cases illustrate a range of deaths linked to khat consumption – from purely circumstantial to clear and unambiguous physiological causes.
<table>
<thead>
<tr>
<th>Characteristic</th>
<th>Case</th>
</tr>
</thead>
<tbody>
<tr>
<td>Year of death</td>
<td>6</td>
</tr>
<tr>
<td>Age (years)</td>
<td>7</td>
</tr>
<tr>
<td>Gender</td>
<td>8</td>
</tr>
<tr>
<td>Ethnicity</td>
<td>9</td>
</tr>
<tr>
<td>Marital status</td>
<td>10</td>
</tr>
<tr>
<td>Living arrangements</td>
<td></td>
</tr>
<tr>
<td>Employment status</td>
<td></td>
</tr>
<tr>
<td>UK residency (years)</td>
<td></td>
</tr>
<tr>
<td>Significant medical history</td>
<td></td>
</tr>
<tr>
<td>Known psychiatric history</td>
<td></td>
</tr>
<tr>
<td>Known khat using history</td>
<td></td>
</tr>
<tr>
<td>Evidence of using khat prior to death</td>
<td></td>
</tr>
<tr>
<td>Place of death</td>
<td></td>
</tr>
<tr>
<td>Role of khat in death</td>
<td></td>
</tr>
<tr>
<td>Toxicology done</td>
<td></td>
</tr>
<tr>
<td>Toxicology results</td>
<td></td>
</tr>
<tr>
<td>Cause(s) of death</td>
<td></td>
</tr>
<tr>
<td>Verdict</td>
<td></td>
</tr>
<tr>
<td>Coroner’s area</td>
<td></td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>6</th>
<th>7</th>
<th>8</th>
<th>9</th>
<th>10</th>
</tr>
</thead>
<tbody>
<tr>
<td>36.5</td>
<td>21.9</td>
<td>41.5</td>
<td>37.8</td>
<td>29.9</td>
</tr>
<tr>
<td>Somali</td>
<td>Eritrean</td>
<td>Somali</td>
<td>Somali</td>
<td>White (Polish)</td>
</tr>
<tr>
<td>Married</td>
<td>Single</td>
<td>Not known</td>
<td>Not known</td>
<td>Not known</td>
</tr>
<tr>
<td>With partner &amp; children</td>
<td>Psychiatric unit</td>
<td>Self &amp; children</td>
<td>Alone</td>
<td>With friends</td>
</tr>
<tr>
<td>Unemployed</td>
<td>Student</td>
<td>Not known</td>
<td>Mini-cab driver</td>
<td>Employed (manual)</td>
</tr>
<tr>
<td>7</td>
<td>8</td>
<td>7</td>
<td>8</td>
<td>8</td>
</tr>
<tr>
<td>Yes</td>
<td>No</td>
<td>No</td>
<td>Not known</td>
<td>Not known</td>
</tr>
<tr>
<td>No</td>
<td>Yes</td>
<td>No</td>
<td>Not known</td>
<td>Not known</td>
</tr>
<tr>
<td>Yes</td>
<td>Yes</td>
<td>Yes</td>
<td>Not known</td>
<td>Not known</td>
</tr>
<tr>
<td>Yes</td>
<td>No</td>
<td>Yes</td>
<td>Yes</td>
<td>Yes</td>
</tr>
<tr>
<td>Long term khat use =&gt; hepatic necrosis =&gt; sub-fulminant liver failure</td>
<td>Drug-induced psychosis/psychosis exacerbated by use of ‘skunk’ and khat over long period =&gt; traumatic suicide</td>
<td>Ingestion of khat (possibly no longer active), alcohol consumption =&gt; intoxication, impaired judgement/lack of co-ordination =&gt; traumatic road traffic accident (pedestrian)</td>
<td>Alcohol and khat in system =&gt; impaired judgement/loss of control of vehicle =&gt; traumatic road traffic accident (driver)</td>
<td>Overdose of injected heroin, but khat also in system</td>
</tr>
<tr>
<td>Not applicable</td>
<td>Blood – negative, Urine – negative.</td>
<td>Blood - ethanol 185mg/100ml. Urine - ethanol 235mg/100ml; cathinone detected; norephedrine/norpseudoephedrine detected</td>
<td>Vitreous humour- alcohol 210mg/100ml; cathinone 110µg/L; cathine 510µg/L</td>
<td>Blood - alcohol 47mg/100ml; 6-MAM, Morphine 0.24mg/L; codeine 0.01mg/L. Cathinone &lt;25µg/L; norephedrine/norpseudoephedrine (total) &lt;50µg/L. Urine - alcohol 73mg/100ml; codeine detected; cathinone detected; norephedrine/norpseudoephedrine detected</td>
</tr>
<tr>
<td>1a Sub fulminant liver failure 1b Hepatic necrosis 2 Khat toxicity</td>
<td>1a Multiple injuries</td>
<td>1a Chest injuries</td>
<td>1a Multiple injuries</td>
<td>1a Morphine intoxication</td>
</tr>
<tr>
<td>Accident</td>
<td>Act of self-harm</td>
<td>Narrative</td>
<td>Accident</td>
<td>Non-dependent abuse of drugs</td>
</tr>
<tr>
<td>Inner North London</td>
<td>West London</td>
<td>Southampton &amp; New Forest</td>
<td>Northern London</td>
<td>Northern London</td>
</tr>
</tbody>
</table>

Note: These cases illustrate a range of deaths linked to khat consumption – from purely circumstantial to clear and unambiguous physiological causes.
<table>
<thead>
<tr>
<th>Characteristic</th>
<th>11</th>
<th>12</th>
<th>13</th>
<th>14</th>
<th>15</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Year of death</strong></td>
<td>2008</td>
<td>2008</td>
<td>2008</td>
<td>2009</td>
<td>2009</td>
</tr>
<tr>
<td><strong>Age (years)</strong></td>
<td>34.5</td>
<td>22.2</td>
<td>19.3</td>
<td>41.0</td>
<td></td>
</tr>
<tr>
<td><strong>Gender</strong></td>
<td>Male</td>
<td>Male</td>
<td>Male</td>
<td>Male</td>
<td>Female</td>
</tr>
<tr>
<td><strong>Ethnicity</strong></td>
<td>Somali</td>
<td>Somali</td>
<td>Somali</td>
<td>Somali</td>
<td>Somali</td>
</tr>
<tr>
<td><strong>Marital status</strong></td>
<td>Not known</td>
<td>Not known</td>
<td>single</td>
<td>Not known</td>
<td>Married</td>
</tr>
<tr>
<td><strong>Living arrangements</strong></td>
<td>With partner</td>
<td>With cousin</td>
<td>Not known</td>
<td>With partner &amp; children</td>
<td>With partner</td>
</tr>
<tr>
<td><strong>Employment status</strong></td>
<td>Unemployed</td>
<td>Unemployed</td>
<td>Student</td>
<td>Cab driver</td>
<td>Unemployed</td>
</tr>
<tr>
<td><strong>UK residency (years)</strong></td>
<td>Not known</td>
<td>15</td>
<td>Not known</td>
<td>Not known</td>
<td>4.5</td>
</tr>
<tr>
<td><strong>Significant medical history</strong></td>
<td>Not known</td>
<td>No</td>
<td>No</td>
<td>Not known</td>
<td>None</td>
</tr>
<tr>
<td><strong>Known psychiatric history</strong></td>
<td>Not Known</td>
<td>Not known</td>
<td>No</td>
<td>Not known</td>
<td>Depression, mental health issues, on prescribed medications but not being used</td>
</tr>
<tr>
<td><strong>Known khat using history</strong></td>
<td>Yes</td>
<td>Yes</td>
<td>Not known</td>
<td>Yes</td>
<td>Alcohol and drug abuse; chewed 2 bundles of khat, 2-4 times/week</td>
</tr>
<tr>
<td><strong>Evidence of using khat prior to death</strong></td>
<td>Yes</td>
<td>Yes</td>
<td>Yes</td>
<td>No</td>
<td>Yes</td>
</tr>
<tr>
<td><strong>Place of death</strong></td>
<td>Hospital</td>
<td>Hospital</td>
<td>Hospital</td>
<td>Hospital</td>
<td>Outside block of flats</td>
</tr>
<tr>
<td><strong>Role of khat in death</strong></td>
<td>Abused khat =&gt; jaundice, night sweats, pyrexia =&gt; sub-acute liver failure</td>
<td>Ingestion of alcohol &amp; khat =&gt; intoxication =&gt; fighting =&gt; collapse =&gt; cardiovascular event. Khat may have triggered an infarction or electrical instability (arythmia) causing death</td>
<td>Had consumed alcohol, cannabis &amp; khat which may have contributed to his aggressive behaviour, leading to incident which triggered events leading to his assault and death</td>
<td>Excessive use of khat =&gt; fulminant hepatic necrosis =&gt; required liver transplant (failed)</td>
<td>Its psychotic effects or hallucinations may have led her to jump or its effects could have led to an error of judgement</td>
</tr>
<tr>
<td><strong>Toxicology done</strong></td>
<td>No</td>
<td>Yes</td>
<td>Yes</td>
<td>No</td>
<td>Yes</td>
</tr>
<tr>
<td><strong>Toxicology results</strong></td>
<td>NA</td>
<td>Blood – alcohol 240mg/dl; cathine present; cathinone present. Urine – alcohol 309mg/dl; cathine present; cathinone present; norephedrine present. Blood – alcohol 170mg/dl; cathinone 0.019mg/l; carboxy-THC 0.011mg/l; phenylpropanolamine 0.1mg/l</td>
<td>NA</td>
<td>Peripheral blood cathine 1447ng/ml; cathinones 122ng/ml</td>
<td></td>
</tr>
<tr>
<td><strong>Cause(s) of death</strong></td>
<td>1a Sub acute liver failure</td>
<td>1a Unascertained</td>
<td>1a Severe head injury [consistent with impacts from heavy blunt object]</td>
<td>1a Pulmonary haemorrhage and multi-organ failure 1b Liver failure due to khat toxicity (orthotopic liver transplant) 2 Steatosis of the orthotopic (donor) liver transplant</td>
<td>1a Multiple injuries consistent with a fall whilst under the influence of khat</td>
</tr>
<tr>
<td><strong>Verdict</strong></td>
<td>Narrative</td>
<td>Open</td>
<td>Unlawful killing [attackers sentenced to life imprisonment]</td>
<td>Narrative</td>
<td>Accidental</td>
</tr>
<tr>
<td><strong>Coroner’s area</strong></td>
<td>Northern London</td>
<td>Avon</td>
<td>West London</td>
<td>Inner North London</td>
<td>Northamptonshire</td>
</tr>
</tbody>
</table>

*Note: These cases illustrate a range of deaths linked to khat consumption – from purely circumstantial to clear and unambiguous physiological causes.*