



ADVANCED RELEASE

Note: In the interests of public health protection, the EMCDDA is releasing this report before formal page layout in the EMCDDA house style. The final report will be available on the EMCDDA website in due course.

**Risk assessment report on a new psychoactive substance:
methyl 2-[[1-(4-fluorobutyl)-1*H*-indole-3-carbonyl]amino]-3,3-dime-
thylbutanoate (4F-MDMB-BICA) in accordance with Article 5c of
Regulation (EC) No 1920/2006 (as amended)**

Statement regarding the United Kingdom

This report covers a reference period from January to November 2020. The United Kingdom left the European Union as of 1 February 2020. However, during the transitional period, the United Kingdom continues to participate in the European Union Early Warning System on new psychoactive substances. Unless stated otherwise, for the purpose of this report, the term 'Member States' shall include the United Kingdom.

Contents

Statement regarding the United Kingdom.....	2
Contents.....	3
Introduction	4
Background.....	6
Chemical and physical properties and the methods and precursors used for manufacture ...	7
Pharmacological and toxicological properties.....	10
Health risks	10
Social risks.....	13
Extent and patterns of use, availability and potential for diffusion	14
Commercial and industrial uses, the extent of such use and its use for scientific research and developmental purposes.....	16
Other relevant information	16
Conclusion	17

Introduction

New psychoactive substances can cause serious cross-border threats to health. In Europe, Regulation (EC) No 1920/2006 of the European Parliament and of the Council of 12 December 2006 on the European Monitoring Centre for Drugs and Drug Addiction (recast) (as amended) (hereafter the 'Regulation') and Council Framework Decision 2004/757/JHA of 25 October 2004 laying down minimum provisions on the constituent elements of criminal acts and penalties in the field of illicit drug trafficking sets up a three-step legal framework of early warning, risk assessment and control measures that allows the EU to rapidly detect, assess and respond to the public health and social risks caused by new psychoactive substances. The European Monitoring Centre for Drugs and Drug Addiction (EMCDDA) is responsible for the first two steps in this system, namely operating the EU Early Warning System on new psychoactive substances (EWS) in close cooperation with Europol, and conducting risk assessments. The European Commission is responsible for proposing control measures. Thus, the legal framework allows the institutions of the EU and the Member States to act on all new psychoactive substances (NPS) that appear on the European drug market.

In accordance with Article 5a of the Regulation, methyl 2-[[1-(4-fluorobutyl)-1*H*-indole-3-carbonyl]amino]-3,3-dimethylbutanoate (commonly known as 4F-MDMB-BICA) was formally notified as an NPS by the EMCDDA on behalf of Belgium on 2 July 2020. The notification was based on the identification of the substance in 1.5 kilograms of white powder seized by Belgian Customs at Bierset Airport, Belgium, on 31 March 2020. The seizure originated from China and was en route to the Netherlands.

Based on the information reported to the Early Warning System, and, in accordance with Article 5a of the Regulation, on 2 September 2020, the EMCDDA assessed the existing information on 4F-MDMB-BICA. The EMCDDA concluded that the assessment gave rise to concerns that 4F-MDMB-BICA may pose health or social risks at EU level, and, consequently, determined that an initial report should be produced in accordance with Article 5b of the Regulation. The initial report was submitted to the Commission and Member States on 14 October 2020. Based on the findings of the initial report, on 28 October 2020, the Commission requested that the EMCDDA carry out a risk assessment on 4F-MDMB-BICA in accordance with Article 5c of the Regulation.

This risk assessment report presents the summary findings and the conclusion of the risk assessment carried out by the Scientific Committee of the EMCDDA on 4F-MDMB-BICA. The report is intended for policymakers and decision-makers in the institutions of the EU.

The report has been prepared and drafted in accordance with the requirements of Article 5c of the Regulation as well as the conceptual framework and the procedure set out in the EMCDDA risk assessment operating guidelines. It is written as a stand-alone document, which presents a summary of the information considered during the detailed assessment of the scientific and law enforcement information available at this time. The conclusion section of the report summarises the main issues addressed and reflects the opinions held by the

members of the Scientific Committee. A list of the information resources considered by the Scientific Committee, including a detailed technical report on 4F-MDMB-BICA prepared by the EMCDDA (Annex 1), is provided below.

In accordance with Article 5c of the Regulation, the meeting to assess the risks of 4F-MDMB-BICA was convened under the auspices of the Scientific Committee of the EMCDDA with the participation of four additional experts designated by the Director of the EMCDDA, acting on the advice of the Chairperson of the Scientific Committee, chosen from a list of experts approved by the Management Board of the EMCDDA. The additional experts were from scientific fields that were either not represented or not sufficiently represented in the Scientific Committee, and whose contribution is necessary for the balanced assessment of the risks posed by 4F-MDMB-BICA. A further six experts were observers to the risk assessment: two experts from the Commission, two experts from the EMCDDA, one expert from Europol, and one expert from the European Medicines Agency (EMA). The meeting took place on 7 December 2020. Owing to the on-going response to the coronavirus (COVID-19) pandemic caused by severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2), the meeting was conducted by videoconference and hosted by the EMCDDA.

The risk assessment was carried out on the basis of information provided to the Scientific Committee by the Member States, the EMCDDA, Europol, the EMA, the European Centre for Disease Prevention and Control (ECDC), the European Chemicals Agency (ECHA) and the European Food Safety Authority (EFSA). A list of the Scientific Committee members, observers and other participants attending the risk assessment meeting is annexed to this report (Annex 2).

For the risk assessment, the Scientific Committee considered the following information resources:

- the EMCDDA technical report on methyl 2-[[1-(4-fluorobutyl)-1*H*-indole-3-carbonyl]amino]-3,3-dimethylbutanoate (4F-MDMB-BICA) (Annex 1);
- the EMCDDA initial report on the new psychoactive substance methyl 2-[[1-(4-fluorobutyl)-1*H*-indol-3-yl]carbonyl]amino)-3,3-dimethylbutanoate (4F-MDMB-BICA);
- open source information, including scientific articles, official reports, grey literature, Internet drug discussion forums and related websites;
- additional information provided during the course of the risk assessment meeting by the participants;
- EMCDDA operating guidelines for the risk assessment of new psychoactive substances;
- Regulation (EC) No 1920/2006 of the European Parliament and of the Council of 12 December 2006 on the European Monitoring Centre for Drugs and Drug Addiction (recast) (as amended); and
- Council Framework Decision 2004/757/JHA of 25 October 2004 laying down minimum provisions on the constituent elements of criminal acts and penalties in the field of illicit drug trafficking (as amended).

Background

Methyl 2-[[1-(4-fluorobutyl)-1*H*-indole-3-carbonyl]amino]-3,3-dimethylbutanoate (4F-MDMB-BICA) is a synthetic cannabinoid receptor agonist (synthetic cannabinoid) which does not appear to have a history in the scientific literature. However, it is structurally related to compounds of the indole-3-carboxamide class that feature pendant amino acid esters (methyl *L-tert*-leucinate) previously developed as part of pharmaceutical research.

Synthetic cannabinoids, such as 4F-MDMB-BICA, are functionally similar to Δ^9 -tetrahydrocannabinol (THC), the major psychoactive principle of cannabis. Like THC, they bind to and activate the CB₁ and CB₂ cannabinoid receptors which form part of the endocannabinoid system — a system that helps regulate a large number of physiological functions in the body such as behaviour, mood, pain, appetite, sleep, the immune system, and the cardiovascular system.

Many synthetic cannabinoids were first developed to study the endocannabinoid system as well as to explore their potential as therapeutic agents to treat a number of diseases and their symptoms (such as neurodegenerative diseases, drug dependence, pain disorders, and cancer).

Since around 2006, 'legal high' products containing synthetic cannabinoids have been sold in Europe as 'herbal smoking mixtures' and marketed as 'legal' replacements for cannabis. These products are made by dissolving the synthetic cannabinoids in solvents such as acetone or methanol and then mixing them with, or, spraying them on, plant material such as the herbs *Turnera diffusa* (Damiana) and other herbs such as *Melissa*, *Mentha* and *Thymus* species of the Lamiaceae family. Such products are generally referred to by a variety of names in Europe, including 'Spice' ⁽¹⁾, 'herbal smoking mixtures', 'herbal incense', and 'synthetic cannabis'. They may also be mixed with tobacco. Manufacturers of smoking mixtures frequently change the synthetic cannabinoids in the products, which means that product names are not a reliable source of information regarding the actual substances that are present. More than 210 synthetic cannabinoids have been identified on the European drug market since 2008. They are the largest group of substances that are monitored by the EMCDDA through the EU Early Warning System. In recent years, alongside smoking mixtures, new dosage forms, including e-liquids for vaping using electronic cigarettes, as well as paper (including blotters) impregnated with synthetic cannabinoids, have appeared on the drug market.

A number of cannabinoids are controlled under the United Nations Convention on Psychotropic Substances, 1971. These include: the major active principle of cannabis, Δ^9 -tetrahydrocannabinol ⁽²⁾ (Schedule I), dronabinol ⁽³⁾ (Schedule II) as well as an increasing

⁽¹⁾ 'Spice' was the most common brand name used for these types of products when they first appeared on the European market.

⁽²⁾ Including some of its named isomers and their stereochemical variants.

⁽³⁾ And its stereochemical variants.

number of synthetic cannabinoids, including 4F-MDMB-BINACA ⁽⁴⁾ and 5F-MDMB-PICA ⁽⁵⁾(Schedule II) that are closely related to 4F-MDMB-BICA.

4F-MDMB-BICA has been available on the drug market in Europe since at least March 2020. It is important to highlight that, as 4F-MDMB-BICA has emerged on the drug market only recently, no formal epidemiological studies have been conducted, which limits understanding of the frequency and patterns of use of the substance. Related to this, the presence of 4F-MDMB-BICA on the drug market and its involvement in serious adverse events (such as acute poisonings presenting to hospital emergency rooms and medico-legal death investigations) may be underreported because the substance is not routinely screened for in some forensic and toxicology laboratories in Europe. Therefore, the presence of 4F-MDMB-BICA on the drug market may be undetected in some areas. In addition, in some areas not all laboratories are part of national early warning systems; for this reason, the identification of 4F-MDMB-BICA may go unreported to the Reitox national focal points, and, as a result, to the EMCDDA. Finally, it is also important to note that, in some settings, the on-going COVID-19 pandemic may have reduced the capacity of early warning systems, including forensic science and toxicology laboratories, to detect and report events involving 4F-MDMB-BICA.

Chemical and physical properties and the methods and precursors used for manufacture

Chemical and physical properties

Methyl 2-[[1-(4-fluorobutyl)-1*H*-indole-3-carbonyl]amino]-3,3-dimethylbutanoate (4F-MDMB-BICA) is a synthetic cannabinoid receptor agonist (synthetic cannabinoid). The 4F-MDMB-BICA code name used for the substance is derived from its structural features ⁽⁶⁾: a methyl 3,3-dimethylbutanoate linked group (MDMB), a 4-fluorobutyl tail (4F and B) attached to the indole nitrogen atom of an indole core (I), and a carboxamide linker (CA).

4F-MDMB-BICA is structurally related to 4F-MDMB-BINACA ⁽⁴⁾, 5F-MDMB-PICA (5F-MDMB-2201) ⁽⁵⁾, and 5F-MDMB-PINACA ⁽⁷⁾, all of which are under international control (Schedule II of the 1971 United Nations Single Convention on Psychotropic Substances).

The chemical structures, molecular formulae and molecular weights of these compounds are provided in Figure 1.

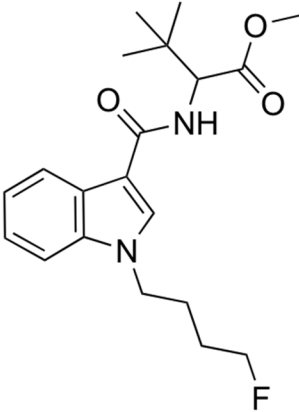
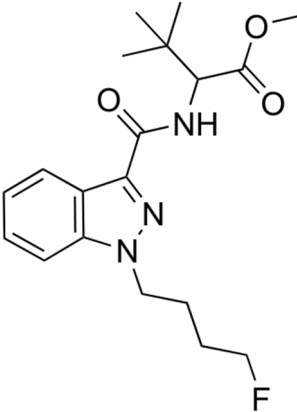
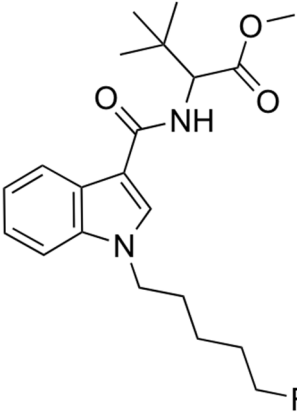
Figure 1. Chemical structures and molecular information of 4F-MDMB-BICA, 4F-MDMB-BINACA, and 5F-MDMB-PICA.

⁽⁴⁾ 4F-MDMB-BINACA: methyl 2-[[1-(4-fluorobutyl)-1*H*-indazole-3-carbonyl]amino]-3,3-dimethylbutanoate.

⁽⁵⁾ 5F-MDMB-PICA (5F-MDMB-2201): methyl 2-[[1-(5-fluoropentyl)-1*H*-indole-3-carbonyl]amino]-3,3-dimethylbutanoate.

⁽⁶⁾ Different naming systems exist and are used for applying short/code names to synthetic cannabinoids.
<http://www.emcdda.europa.eu/topics/pods/synthetic-cannabinoids>.

⁽⁷⁾ 5F-MDMB-PINACA (5F-ADB): 2-[[1-(5-fluoropentyl)-1*H*-indole-3-carbonyl]amino]-3,3-dimethylbutanoate

4F-MDMB-BICA	4F-MDMB-BINACA	5F-MDMB-PICA
		
Molecular formula: $C_{20}H_{27}FN_2O_3$	Molecular formula: $C_{19}H_{26}FN_3O_3$	Molecular formula: $C_{21}H_{29}FN_2O_3$
Molecular weight: 362.44	Molecular weight: 363.43	Molecular weight: 376.47
Monoisotopic mass: 362.2006	Monoisotopic mass: 363.1958	Monoisotopic mass: 376.2162

In its pure form 4F-MDMB-BICA has been described as a white solid. It is soluble in organic solvents such as chloroform and expected to be only partially soluble in water.

4F-MDMB-BICA contains a stereogenic centre and therefore two possible enantiomers may exist, (*S*)-4F-MDMB-BICA and (*R*)-4F-MDMB-BICA. No information is available on whether the 4F-MDMB-BICA detected in the European drug market corresponds to the (*R*)- or (*S*)-enantiomer, or a mixture of both, which may in part reflect the fact that stereochemical analysis is not routinely undertaken in forensic laboratories. Based on the literature and the precursors that are most likely to be used, an (*S*)-configuration of the stereocentre could be expected.

Information provided from seizures and collected samples reported to the EMCDDA have noted that 4F-MDMB-BICA is typically found in herbal/plant material (including as commercially-branded 'legal high' products) and as a powder. Other forms, such as pieces of paper impregnated with the substance (including blotters), liquids and e-liquids have also been reported.

The analytical identification of 4F-MDMB-BICA in physical and biological samples is possible using standard analytical techniques. These include chromatographic and mass spectrometric methods.

Analytical reference material is important for correct identification and for facilitating the quantification of 4F-MDMB-BICA in physical and biological samples. Such material is commercially available in form of the (S)-enantiomer.

Methods and precursors used for manufacture

There is no information on the actual manufacturing methods or chemical precursors used to synthesise the 4F-MDMB-BICA that has been identified on the drug market in Europe. Information from one seizure made by customs suggests that at least some 4F-MDMB-BICA on the market in Europe has been supplied from China. Although the synthetic routes reported in the scientific literature involve several steps, they are straightforward and make use of common manufacturing equipment. The starting materials for either synthetic route are readily available. The multi-step production of 4F-MDMB-BICA requires basic knowledge of and experience in synthetic organic chemistry.

Commercially available domestic or industrial chemicals which could be used for synthesis of 4F-MDMB-BICA may be potentially toxic. Use of such products as reagents or solvents may result in serious toxic effects if the resultant contaminated product is consumed. The herbal plant material which is used as a basis for smoking mixtures may also contain toxicologically relevant substances (such as pesticides that could potentially be present in the plant material).

Pharmaceutical forms

Information on the pharmaceutical forms available for 4F-MDMB-BICA is limited. There are three main types of products containing synthetic cannabinoids that are available on the drug market: smoking mixtures, where the cannabinoid is mixed with herbal plant material or tobacco that is then smoked (similar to herbal cannabis, the mixture is usually prepared for smoking as a hand-rolled cigarette ('joint')) or inhaled using a vaporiser; e-liquids, where a solution of the cannabinoid is prepared by mixing it with a solvent, which is then inhaled ('vaped') using an e-cigarette; in addition, synthetic cannabinoids are also impregnated on to paper which can then be smoked or vaped. The latter is a commonly used approach to smuggle synthetic cannabinoids into prison in some countries. To a lesser extent, people who are using these substances may also prepare their own products using powdered synthetic cannabinoids purchased from a vendor or dealer. It is expected that similar to related synthetic cannabinoids 4F-MDMB-BICA may also be administered orally and sublingually.

Products containing synthetic cannabinoids such as 4F-MDMB-BICA rarely state the correct ingredients and concentrations. Given the illicit nature of the trade in 4F-MDMB-BICA, the composition of products containing 4F-MDMB-BICA is likely to vary over time and place, as well as based on the specific location in the drug supply chain from which the product was obtained (e.g. from the manufacturer, wholesaler, retailer or at street-level illicit market). In Europe, 4F-MDMB-BICA has been seized by law enforcement in the form of powders, smoking mixtures, pieces of paper impregnated with the substance (including blotters),

liquids and e-liquids. Information on the amount of 4F-MDMB-BICA present in seized powders (chemical purity) has not been reported. In at least some of the seizures other synthetic cannabinoids were present. Information on the enantiomeric composition of seized and collected samples has not been reported.

Pharmacological and toxicological properties

Pharmacological properties

There is limited information on the pharmacological properties of 4F-MDMB-BICA. Currently available information from *in vitro* studies suggests that 4F-MDMB-BICA activates the cannabinoids type 1 (CB₁) receptor as a full agonist at reasonably high potency. These data show that 4F-MDMB-BICA has comparable potency to JWH-018 that is under international control, but is less potent than 4F-MDMB-BINACA and 5F-MDMB-PICA, which are potent full agonists that are also under international control. This suggests that 4F-MDMB-BICA may produce effects in humans similar to other cannabinoid receptor agonists, such as THC and other synthetic cannabinoids. No information is available on binding and activation of the CB₂ receptor nor the effects induced *in vivo* by 4F-MDMB-BICA.

No studies are available on the pharmacodynamics of 4F-MDMB-BICA on other pharmacological targets.

Data on the pharmacokinetics of 4F-MDMB-BICA, including information on metabolites and their effects, are unavailable.

Toxicological properties

No information is available on the toxicological properties (including pre-clinical safety data) of 4F-MDMB-BICA. Ingestion of highly potent full CB₁ receptor agonists have been associated with acute toxicity in humans; this suggests that the acute toxicity of 4F-MDMB-BICA may be much greater than THC and similar to that observed for related synthetic cannabinoids under international control. The involvement of other, non-cannabinoid toxicological targets or unexpected drug-drug interactions in the overall pharmacotoxicological effects of 4F-MDMB-BICA cannot be excluded.

There is no known antidote to poisoning caused by synthetic cannabinoids. Treatment in poisoning cases should be symptomatic.

Health risks

No studies are available on the acute and chronic health effects of 4F-MDMB-BICA. The clinical features of poisonings caused by 4F-MDMB-BICA are expected to be similar to those reported from related synthetic cannabinoids resulting in gastrointestinal, neurological, cardiovascular, renal clinical features. These dose-dependent effects appear to be much more pronounced and severe when compared to cannabis.

Acute toxicity

There is limited information on the acute toxicity of 4F-MDMB-BICA. Based on the available information, the health risks are likely to be similar to those observed with other synthetic cannabinoids. Adverse effects from overdosing 4F-MDMB-BICA might include gastrointestinal (e.g. nausea and vomiting (including hyperemesis)), neurological (e.g. hallucination, seizures, convulsions, agitation, anxiety, paranoia, confusion, delusions, catatonia, lethargy, psychosis (including susceptible individuals) and severe central nervous system depression (such as rapid loss of consciousness/coma), cardiovascular (e.g. tachycardia, hypertension, acute myocardial infarction and sudden cardiac death) and renal (e.g. acute kidney failure) clinical features, and respiratory depression.

In addition, some of the features of poisoning—particularly loss of consciousness, respiratory depression, and behavioural effects—may place users at additional risks, such as choking on/aspirating vomit, drowning, falling, hypothermia as a result of falling unconscious outside in cold weather, and self-inflicted violence/injury. The aggressive and violent behaviours reported with synthetic cannabinoids may also place others at risk of injury.

The reasons for these more pronounced and severe effects, as well as severe and fatal poisoning, are poorly understood, but at least two factors are likely to be important: the high potency of the substances and the unintentionally high doses that users are exposed to.

Firstly, studies have found that many of the synthetic cannabinoids, including 4F-MDMB-BICA, which are sold on the drug market, are typically potent, full agonists compared to THC. This means that even at very small doses they can activate the CB₁ receptor much more potently than THC, which may be responsible for the more pronounced and unexpected pharmacotoxicological effects in humans.

Secondly, in respect to smoking mixtures, the process for mixing the synthetic cannabinoids with the plant material to make the smoking mixture can lead to dangerous amounts of the substances in the products. This is because producers have to guess the amount of substances to be added, while the mixing process makes it difficult to dilute them sufficiently and distribute them consistently throughout the plant material. This can result both in products that contain toxic amounts of the substances in general, as well as in products where solid particles of synthetic cannabinoids are clumped together, forming highly concentrated pockets within the plant material. In fact, in the latter case, simply tapping a packet containing a smoking mixture can dislodge the substances from the plant material. In addition, paper (such as blotters and cards) impregnated with synthetic cannabinoids can pose a similar high risk of poisoning because the amount of synthetic cannabinoid can be unevenly distributed in different parts of the paper, sometimes forming highly concentrated sections on the paper. These issues are exacerbated as the products are typically smoked or vaped, allowing the substances to be rapidly absorbed into the bloodstream and to reach the central nervous system and other parts of the body to cause their effects.

Together, these factors, coupled to the typically high potency of synthetic cannabinoids, makes it difficult for users to control the dose that they are exposed to. This can lead them to rapidly administer a toxic dose unintentionally. Accounts from patients and people who witness poisonings suggest that in some cases a small number of puffs from a cigarette (“joint”) have been sufficient to cause severe and fatal poisoning.

Similarly to other synthetic cannabinoids, the use of 4F-MDMB-BICA together with other drugs, especially central nervous system depressants (such as alcohol, opiates/opioids, and sedative-hypnotics) is likely to increase the risk of life-threatening poisoning.

Acute poisonings

A total of five acute intoxications with confirmed exposure to 4F-MDMB-BICA have been reported to the EMCDDA by the United Kingdom. The cases occurred between June and July 2020. All cases included clinical features of poisoning similar to those reported for other synthetic cannabinoids, such as confusion, tachycardia, respiratory insufficiency, reduced conscious level, abnormal sweating, and agitation. In all cases other substances were identified including other synthetic cannabinoids, benzodiazepines, opioids, and other substances such as pregabalin, cocaine, THC, and ketamine. In all cases the poisoning was considered life threatening and required hospitalisation of the patient.

Deaths

A total of 21 deaths with confirmed exposure to 4F-MDMB-BICA have been reported by Hungary. The deaths occurred between May and August 2020. In the vast majority of cases other substances were identified including other synthetic cannabinoids, benzodiazepines, stimulants, and other substances such as THC, alcohol, and ketamine. Reported ante-mortem symptoms and clinical features included loss of consciousness, chest pain, respiratory problems, tremor, seizures, somnolence, aggressive behaviour, and foaming at the mouth. Three of the cases were found dead. One of the cases involved a prisoner and another involved a homeless person. At least some of the individuals were people known to use drugs. The reported causes of death were: cardiac arrest due to substance overdose (7 cases), acute heart failure (7 cases), traumatic shock (2 cases), strangulation (1 case), brain oedema (1 case), and asphyxiation following aspirating vomit (1 case).

Physical, mental and behavioural effects

The physical effects of 4F-MDMB-BICA have not been studied. Based on the pharmacological properties of the substance, as well as information from observations from closely related substances, it is likely that the dose-dependent effects of 4F-MDMB-BICA are similar to those commonly reported for other synthetic cannabinoids, including tachycardia, dry mouth, nausea, vomiting, balance deficiencies, and ocular effects such as reddened conjunctivae, glassy eyes and delayed or unresponsive pupil light reactions, and impaired motor performance.

The mental and behavioural effects of 4F-MDMB-BICA have not been studied. Based on the pharmacological properties of the substance, the effects of 4F-MDMB-BICA appear to share some similarities with cannabis, THC, and other synthetic cannabinoids. This may include: relaxation, euphoria, lethargy, confusion, anxiety, fear, distorted perception of time, depersonalisation, hallucinations, and paranoia.

Overall, these effects may be much more pronounced and severe when compared to cannabis. In addition, psychotic episodes, as well as aggressive and violent behaviour, have also been reported.

There are no studies on the effects of 4F-MDMB-BICA on the ability to drive and operate machines. However, synthetic cannabinoids impair the mental and physical ability required to drive and operate machines. This effect is likely to extend to 4F-MDMB-BICA.

Chronic toxicity

There is no information on the chronic toxicity of 4F-MDMB-BICA. Similar to other synthetic cannabinoids, chronic use has been associated with greater risks for developing mental health disorder than cannabis, which may include dependence. Acute and chronic use of synthetic cannabinoids has also been associated with cases displaying detrimental cardiovascular health.

Abuse liability and dependence-producing potential

No information is available on the abuse liability and dependence producing potential of 4F-MDMB-BICA. Given what is currently known about the properties of synthetic cannabinoids in general (and some similarities to THC), it is expected that 4F-MDMB-BICA may have a potential for abuse and dependence. It has been suggested that consumption of synthetic cannabinoids can produce tolerance and withdrawal-like symptoms when use is discontinued following a regular use. Withdrawal-like symptoms following cessation of synthetic cannabinoids have been described in the literature. These include: anxiety, unstable mood, crying fits, feeling of inner emptiness, spatial disorientation, hyperacusis (increased sensitivity to ordinary environmental sounds), somatic pain, shortness of breath, hyperventilation, intense sweating and sensations of motor and inner restlessness.

Social risks

There is no specific information on the social risks posed by 4F-MDMB-BICA. There is no information on whether or not criminal groups are involved in the manufacture, trafficking and distribution of 4F-MDMB-BICA within Europe.

It is possible that any such social risks may have some similarities with those associated with the use of cannabis and other synthetic cannabinoids. Such risks include negative impacts on social functioning and criminal activities, such as the involvement of organised crime in the manufacture, trafficking and distribution of the substance.

It is noteworthy that in some settings, synthetic cannabinoids are increasingly used by high-risk drug users and other vulnerable groups, such as the homeless and prisoners. In at least some cases, these people are specifically seeking out synthetic cannabinoids because the substances have developed a reputation for causing profound intoxication. They are also considered cheap, and easy to smuggle. Reports suggest that this has exacerbated existing health and social problems for these vulnerable groups, as well as creating new ones.

In prisons, alongside the adverse health effects, the market in and use of synthetic cannabinoids has been linked to an increase in aggression, violence, bullying, and debt. In some cases this has caused a serious threat to the overall safety and security of the prison environment.

Extent and patterns of use, availability and potential for diffusion

There is limited information on the extent and patterns of use, availability and potential for diffusion of 4F-MDMB-BICA in Europe. As 4F-MDMB-BICA has been on the drug market for only a short period of time, no formal epidemiological studies have been conducted, which limits the understanding of the extent and patterns of use, as well as the potential for diffusion. In relation to this, the presence of 4F-MDMB-BICA on the drug market and its involvement in serious adverse events may be undetected in some areas because the substance is not routinely screened for in some forensic and toxicology laboratories.

4F-MDMB-BICA has been available on the drug market in Europe since at least March 2020, when it was seized for the first time by customs in Belgium. As of November 2020, 4F-MDMB-BICA has been identified in twelve Member States: Belgium, Croatia, Cyprus, Finland, Germany, Hungary, Italy, Lithuania, Poland, Slovenia, Sweden, and the United Kingdom. A total of 111 seizures have been reported, which include 5.6 kg of powder and 0.6 kg of smoking mixtures. Of particular note are four large scale seizures of 4F-MDMB-BICA powder reported by Belgian customs (totalling approximately 5.5 kg) and eleven seizures that occurred in prisons in five Member States. The most recent identifications of 4F-MDMB-BICA reported to the EMCDDA are from seizures made by police in Cyprus and customs in Finland in September 2020. Of note is that 4F-MDMB-BICA has been increasingly identified in biological samples in parts of Germany during the last few months of 2020.

Based on the available information, it appears that at least some of the 4F-MDMB-BICA on the market in Europe has been supplied from China. It is imported into Europe as bulk powders and then sold and distributed in wholesale and retail amounts within Europe either as a powder for processing into products or finished consumer products. There are three main types of products containing 4F-MDMB-BICA that are available on the drug market: smoking mixtures, where 4F-MDMB-BICA is mixed with herbal plant material or tobacco that is then smoked (similar to herbal cannabis, the mixture is usually prepared for smoking as a hand-rolled cigarette ('joint')) or inhaled, for example using a vaporiser; e-liquids, where a solution of 4F-MDMB-BICA is prepared by mixing it with a solvent, which is then inhaled using an e-cigarette; in addition, 4F-MDMB-BICA is also impregnated on to paper which can

then be smoked or vaped. In some countries, the latter is a commonly used approach to smuggle synthetic cannabinoids into prisons. To a lesser extent, people who are using these substances may prepare their own similar products using 4F-MDMB-BICA purchased from a vendor or dealer.

4F-MDMB-BICA may be sought by those looking for 'legal' substitutes for cannabis. This includes individuals subject to drug testing (such as drivers, prisoners, those in drug treatment, and those subject to workplace drug testing), as commonly used drug tests may be unable to detect the compounds. In addition, reports suggest that in some areas, high risk drug users and other vulnerable groups, such as homeless people and prisoners, may specifically seek out synthetic cannabinoids because they have a reputation for causing profound intoxication. They are considered to be cheap and easy to smuggle. Similar to other new psychoactive substances, it also appears that there is interest in 4F-MDMB-BICA by people who self-experiment with a range of substances (so-called psychonauts). Overall, the available information does not suggest widespread use of the substance.

Limited information is available regarding the dose and the dose regimens of 4F-MDMB-BICA. User reports specifically about 4F-MDMB-BICA were not particularly revealing. It is not possible to discern the 'typical' dosages administered as most individuals use pre-made smoking mixtures. Synthetic cannabinoids such as 4F-MDMB-BICA may be active at less than 1 mg. Products containing synthetic cannabinoids such as 4F-MDMB-BICA rarely state the correct ingredients and concentrations. As such, people who use such products will be unaware that they are using 4F-MDMB-BICA nor will they be able to obtain accurate dosage information. 4F-MDMB-BICA can be inhaled by smoking and by vaporising e-liquid solutions ('vaping'), for example by using electronic cigarettes. To a lesser extent, other routes of administration for synthetic cannabinoids have been reported; these include oral, sub-lingual, and rectal.

The high potency of the synthetic cannabinoids, coupled to the unintentionally high doses that users are potentially exposed to, is responsible for outbreaks of mass poisonings involving smoking mixtures. Such outbreaks have ranged in size from four or five to over 800 victims, including deaths. While many of the outbreaks that have been reported so far are from the United States, they have also occurred in Russia and Europe. Mass poisonings can rapidly overwhelm emergency responders and other local healthcare systems.

Unknown to users, synthetic cannabinoids have also been sold as cannabis, ecstasy (MDMA) and other illicit drugs. In some cases, this has led to severe poisoning. Opioids have also been identified in smoking mixtures; while the overall number of detections appears to be relatively small, they could pose a risk of severe opioid poisoning, including life-threatening respiratory depression, especially in individuals with no tolerance to opioids. Users of smoking mixtures will be unaware of this risk.

Commercial and industrial uses, the extent of such use and its use for scientific research and developmental purposes

Based on the available information reported to the EMCDDA from the EMA, it appears that 4F-MDMB-BICA is not an active substance in any medicinal product for human use or in any veterinary medicinal product in Europe. However, although unlikely, the use of 4F-MDMB-BICA as an active substance in medicinal products prepared extemporaneously or in investigational medicinal products cannot be excluded in some Member States.

There is currently no information suggesting that 4F-MDMB-BICA is used for legitimate purposes, including commercial and industrial uses, other than a scientific research or forensic application.

Other relevant information

Restrictive measures

International restrictive measures

At international level, 4F-MDMB-BICA is not controlled under the United Nations Single Convention on Narcotic Drugs, 1961, nor under the Convention on Psychotropic Substances of 1971. 4F-MDMB-BICA has not been subject to assessment nor is it currently under assessment by the United Nations system.

National restrictive measures

Fifteen Member States (Bulgaria, Czechia, Denmark, Estonia, Finland, Greece, Ireland, Malta, the Netherlands, Portugal, Romania, Slovakia, Slovenia, Spain, and Sweden) reported that 4F-MDMB-BICA is not subject to restrictive measures at national level.

4F-MDMB-BICA is subject to restrictive measures in thirteen Member States. Eight Member States (Croatia, Cyprus, France, Italy, Latvia, Luxembourg, Poland, and the United Kingdom) control 4F-MDMB-BICA under drug control legislation. One Member State (Lithuania) controls 4F-MDMB-BICA under medicines legislation. Four Member States (Austria, Belgium, Germany, and Hungary) control 4F-MDMB-BICA under new psychoactive substance legislation. In addition, 4F-MDMB-BICA is controlled under medicines legislation in Norway and under drug control legislation in Turkey.

It is unknown if 4F-MDMB-BICA is controlled in China, where at least some of the substance on the European market has been sourced from.

COVID-19 pandemic

In some settings, the on-going COVID-19 pandemic caused by the severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) may have reduced the capacity of early warning

systems, including forensic science and toxicology laboratories, to detect and report events involving 4F-MDMB-BICA.

Conclusion

4F-MDMB-BICA (methyl 2-[[1-(4-fluorobutyl)-1*H*-indole-3-carbonyl]amino]-3,3-dimethylbutanoate) is an indole-based synthetic cannabinoid receptor agonist (synthetic cannabinoid). The (*S*)-4F-MDMB-BICA stereoisomer has been shown to act as a potent, full agonist at the cannabinoid type 1 (CB₁) receptor when investigated under *in vitro* conditions. This suggests that in humans, 4F-MDMB-BICA may produce similar effects both to THC, which is responsible for the major psychoactive effects of cannabis, and, in particular, closely related synthetic cannabinoids that are under international control, such as 4F-MDMB-BINACA and 5F-MDMB-PICA.

4F-MDMB-BICA is often sold as a 'legal' replacement for cannabis and other synthetic cannabinoids. It is typically administered by smoking a mixture of the substance with herbal plant material or tobacco. These are either from ready-to-use products sold as commercial 'legal high' products or made by drug dealers; less commonly they are self-prepared. Similar to herbal cannabis, the mixture is usually prepared for smoking as a hand-rolled cigarette ('joint') but it may also be smoked in a pipe or 'bong'. 4F-MDMB-BICA can also be inhaled using an e-cigarette or other vaporisation devices. In the case of impregnated papers, typically these are either smoked with tobacco or vaped using an e-cigarette.

The high potency of synthetic cannabinoids and the variable content of the substance in products such as smoking mixtures and impregnated papers constitute a high risk of poisoning, which can be life-threatening or even fatal. These factors are responsible for the outbreaks of mass poisonings seen with synthetic cannabinoids. Although no mass poisonings have been reported for 4F-MDMB-BICA, it is used in a context making it liable to the occurrence of mass poisonings. Such outbreaks have the potential to overwhelm local healthcare systems, which is of particular concern given the on-going COVID-19 pandemic and the additional burden already on healthcare systems. There is no antidote to poisoning caused by synthetic cannabinoids.

4F-MDMB-BICA has been available in Europe since at least March 2020 and has been identified in 12 Member States (Belgium, Croatia, Cyprus, Finland, Germany, Hungary, Italy, Lithuania, Poland, Slovenia, Sweden, and the United Kingdom). 4F-MDMB-BICA has been identified in smoking mixtures, powders, paper impregnated with the substance (including blotters), and liquids (including e-liquids). Of note is that the appearance of 4F-MDMB-BICA on the market appears to coincide with the recent decision to internationally control two closely related synthetic cannabinoids commonly found on the drug market in Europe, 4F-MDMB-BINACA and 5F-MDMB-PICA. As such, it is conceivable that 4F-MDMB-BICA will be a replacement for these substances. Of note is that 4F-MDMB-BICA has been increasingly identified in biological samples in parts of Germany during the last few months of 2020.

As with other synthetic cannabinoids, the available data suggests that 4F-MDMB-BICA is used by people who use cannabis, by those who are regularly subjected to drug testing procedures, and by 'psychonauts'. It may also be used by high-risk drug users and other vulnerable groups (such as prisoners and people experiencing homelessness) as synthetic cannabinoids have gained a reputation for causing profound intoxication, they are cheaper than other drugs, and easy to smuggle. No information is available on the size and demand and the characteristics of these groups of people.

Five acute intoxications with confirmed exposure to 4F-MDMB-BICA have been reported to the EMCDDA by the United Kingdom. The cases occurred between June and July 2020. While exposure to other substances was reported in most cases, including other synthetic cannabinoids, at least some of the clinical features of the poisonings were consistent with exposure to synthetic cannabinoids. In all cases the poisoning was considered life threatening and required hospitalisation of the patients.

Twenty-one deaths with confirmed exposure to 4F-MDMB-BICA have been reported by Hungary. The deaths occurred over relatively short period of time, between May and August 2020. In the vast majority of cases other substances, including other synthetic cannabinoids, were identified. In at least some of these cases, 4F-MDMB-BICA may have contributed to the death.

As 4F-MDMB-BICA has emerged on the drug market in Europe only recently, it is important to note that its presence on the drug market and as the cause of serious adverse events may be undetected because the substance is not routinely screened for in some laboratories. In addition, the on-going COVID-19 pandemic may have reduced the capacity of early warning systems to detect and report events involving 4F-MDMB-BICA.

There is no information available on the chronic health effects of 4F-MDMB-BICA, including abuse liability and dependence producing potential. The chronic health risks might share some similarities to those seen with other synthetic cannabinoids. This may include dependence.

There is no information on whether or not criminal groups are involved in the manufacture, trafficking and distribution of 4F-MDMB-BICA within the European Union. The available information suggests that at least some of the 4F-MDMB-BICA seized in Europe is manufactured by chemical companies based in China. The impact of the on-going COVID-19 pandemic on the manufacture, trafficking, distribution and use of 4F-MDMB-BICA is also currently unknown. However, seizures of bulk powders by European national customs agencies during the first wave of the epidemic highlight that it has been imported into and distributed within Europe during 2020. It is conceivable that should there be a reduced availability of cannabis and other synthetic cannabinoids in Europe, criminal groups as well as people who use drugs, may use a range of replacement substances, including 4F-MDMB-BICA.

There is no information on the actual chemical precursors or manufacturing methods used to synthesise the 4F-MDMB-BICA that has been identified on the drug market in Europe.

Although the synthetic routes reported in the scientific literature involve several steps, depending on precursor availability, they are straightforward and make use of common manufacturing equipment. The starting materials for either synthetic route are readily available. The multi-step production of 4F-MDMB-BICA requires a basic knowledge of and experience in synthetic organic chemistry.

There is no specific information available on the potential social risks posed by 4F-MDMB-BICA. The social risks might share some similarities with those posed cannabis and other synthetic cannabinoids. As highlighted, synthetic cannabinoids are increasingly used by high-risk drug users and vulnerable groups, such as prisoners and people experiencing homelessness, which has exacerbated existing health and social problems as well as creating new issues. For example, in prisons, alongside the adverse health effects, the market in and use of synthetic cannabinoids has been linked to an increase in aggression, violence, bullying, and debt. In some cases this has caused a serious threat to the overall safety and security of the prison environment. As such, it is a concern given the reports of seizures of 4F-MDMB-BICA in prisons and other custodial settings in at least five Member States.

There is currently limited information on the extent or patterns of use of 4F-MDMB-BICA in Europe, nevertheless, information from law enforcement seizures that occurred in 2020 suggest that its availability and potential for diffusion within the Union may be significant. Based on the available information, the consumption of 4F-MDMB-BICA appears to have the potential to cause harms to health associated with its acute toxicity and abuse liability or dependence-producing potential. This harm to health is considered life-threatening because it may cause death. There is also a potential for severe physical and mental impairment. In addition, although limited, information suggests potential social harms related to consumption of 4F-MDMB-BICA including from use in prisons. Overall, these effects appear to be comparable to those of other closely related synthetic cannabinoids that are under international control, such as 4F-MDMB-BINACA and 5F-MDMB-PICA.

It appears that 4F-MDMB-BICA is not an active substance in any medicinal product for human use or in any veterinary medicinal product in Europe. However, although unlikely, the use of 4F-MDMB-BICA as an active substance in medicinal products prepared extemporaneously or in investigational medicinal products cannot be excluded in some Member States. There is currently no information suggesting that 4F-MDMB-BICA is used for legitimate purposes other than research or forensic applications in Europe.

4F-MDMB-BICA is not listed for control in the Single Convention on Narcotic Drugs, 1961, nor in the Convention on Psychotropic Substances, 1971. 4F-MDMB-BICA is not currently under assessment by the United Nations system.

Eight Member States (Croatia, Cyprus, France, Italy, Latvia, Luxembourg, Poland, and the United Kingdom) and Turkey control 4F-MDMB-BICA under drug control legislation. Five Member States (Austria, Belgium, Germany, Hungary, and Lithuania) and Norway control 4F-MDMB-BICA under other legislation. It is unknown if 4F-MDMB-BICA is controlled in

China, where at least some of the substance on the European market has been sourced from.

The Committee notes that while the available information on 4F-MDMB-BICA indicates that it is a potent synthetic cannabinoid *in vitro*, overall there is limited information available on its pharmacology, toxicology, abuse liability and dependence-producing potential. Additional information may emerge with later studies but ideally would have been available in this assessment. The Committee notes that, to strengthen early warning, preparedness and response activities within the EU, such studies should be undertaken systematically prior to risk assessment.

As for any NPS, many of the questions related to 4F-MDMB-BICA that are posed by the lack of data on the risks to individual health, risks to public health and social risks could be answered through further research. Areas where additional information would be important include studies on epidemiology (frequency and patterns of use, including studies that examine the groups of people who use 4F-MDMB-BICA and risk behaviours); the market; chemical profiling; extended pharmacological profiling; metabolic pathways; behavioural effects; acute and chronic toxicity; the potential interaction between 4F-MDMB-BICA and other substances; the abuse liability and dependence-producing potential; and the public health and potential social risks associated with its use.

The Committee notes that a decision to control 4F-MDMB-BICA has the potential to bring with it both intended and unintended consequences. Potential intended consequences include reduced levels of availability and ultimately use. This may reduce the health and social risks arising from the use of 4F-MDMB-BICA. It is important to recognise that a potential unintended consequence of control may be the manufacture and availability of other substances. Indeed, pharmacologically analogous substances that may replace 4F-MDMB-BICA are already being sold on the drug market. The implementation of control measures may also lead to the criminalisation of those who continue to use this substance with the possible attendant risks of socio-economic stigmatisation and marginalisation.

Finally, the Committee notes that it is important to continue to collect accurate information on 4F-MDMB-BICA and to disseminate it to people who use the substance, as well as to practitioners, policymakers and decision-makers.