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Abstract: The objective of this study is to explore the current strategies available to monitor and detect the economically and criminally motivated adulteration of food, identifying their strengths and weaknesses and recommend new approaches and policies to strengthen future capabilities to counter adulteration in a globalized food environment. Many techniques are used to detect the presence of adulterants. However, this approach relies on the adulterant, or means of substitution, being "known" and an analytical method being available. Further techniques verify provenance claims made about a food product e.g. breed, variety etc. as well as the original geographic location of food production. These consider wholeness, or not, of a food item and so do not need to necessarily identify the actual adulterant just whether the food is complete. The conceptual framework developed in this research focuses on the process of predicting, reacting and detecting economically and criminally motivated food adulteration

Highlights

Discussion of economic and criminally motivated food adulteration

Reviews challenges that exist in the supply chain using food supply examples

Reviews techniques for determining food adulteration product wholeness

Conceptual framework developed focuses on the process of predicting, reacting and detecting economically and criminally food adulteration

Developing systems to control food adulteration

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Keywords: economically, criminally, motivated, adulteration, substitution

Ms. Ref. No.: FOODPOLICY-D-13-00249R1 Title: Developing systems to control food adulteration Food Policy

Reviewers' comments:

Reviewer #1: The paper has been greatly improved with this revision and I do recommend publication after some minor revisions suggested below. Response: Thank you

General comments:

Abstract & highlights: change "economically and criminally food adulteration" to "economically and criminally motivated food adulteration"

Response: Thank you for the suggestion. We have added "motivated" into the text (Line 19, pg. 1).

Whole paper: Suggest replacing the term "wholeness" with integrity throughout the entire paper. The concept of food integrity is in principal the same as wholeness, but the former is a more established term in food policy.

Response: Thank you for the comment this has been changed Line 31, pg.2; and as required later in the paper

Page 3: I suggest toning down "It was determined that global anti-counterfeiting activities for the food and drug sector are projected to be worth \$79.3 billion by 2014 (Li 2013)." to "It has been suggested that..." All numbers in literature on the economic cost of EMA (although I haven't reviewed this particular reference) are based on anecdotal evidence and not science, so this should be toned down. Response: The word determined has been toned down to suggested (line 58, pg. 3).

Page 3: A better definition for the specific type of adulteration (EMA) discussed in this paper should be used and referenced (see below refs). It is helpful to use the term "Economically Motivated Adulteration" instead of just "adulteration" since the later in some countries like the USA has a different meaning in regulations.

Moore, Jeffrey C., John Spink, and Markus Lipp. "Development and application of a database of food ingredient fraud and economically motivated adulteration from 1980 to 2010." Journal of food science 77, no. 4 (2012): R118-R126.

Everstine, Karen, John Spink, and Shaun Kennedy. "Economically motivated adulteration (EMA) of food: common characteristics of EMA incidents." Journal of Food Protection® 76, no. 4 (2013): 723-735.

Spink, John, and Douglas C. Moyer. "Defining the public health threat of food fraud." Journal of food science 76.9 (2011): R157-R163.

Response: Term adulteration has been amended to food adulteration and specific focus on economically and criminally motivated adulteration. This has been refocused throughout the paper. A definition of ENA has been inserted. (Spink and Moyer, 2011:32). Line 75; pg 3

Page 12: The following phrase is no longer accurate: "a comprehensive database about known problematic ingredients and detection methods does not currently exist" and should be replaced by "a comprehensive database about known problematic ingredients and detection methods did not exist until 2012 when the USP Food Fraud Database was established"

Response: The word determined has been toned down to suggested (line 238, pg. 10).

Page 14: Replace "Moore et al. (2012) reviewed and collected over 1000 records of food frauds and analytical methods in the US Pharmacopeia Food Chemicals Codex" with "Moore et al. (2012) reviewed and collected over 1000 records of food frauds and analytical methods published in the USP

Food Fraud Database" Response: Changed to USP Food Fraud Database (line 296, pg. 12).

Page 20: References and discussion on FDA's Vulnerability Assessment Software and Carver + Shock tool is not accurate. These tools are not designed to assess vulnerabilities in a food supply for EMA issues, but rather, intentional food defense issues. Would suggest re-framing this point to an argument that tools are needed to assess the likelihood or probability of food fraud/adulteration occurring and that current tools like FDA's Vulnerability Assessment Software and Carver + Shock are not suitable for this purpose. You can reference US FDA's recent proposed rule on Intentional Adulteration part IV-F on EMA.

Response: The authors have re-framed the CARVER + Shock and VAS Tools as focused on predicting attacks (from a food defense point of view) (lines 462-471, pg. 18-19).

Figure 1 is very difficult to understand. I suggest revising this figure or explaining more clearly in the text. For example the box "earliest time before food and feed is adulterated..." does not make sense to me as it appears to fall before the beginning of the food chain.

Response: The authors have placed the text box "earliest time befor food and feed is adulterated..." to the start of the food chain (on the right) and explained in detail the reactive and predictive systems from lines 541-564 (pg. 21-22).

Reviewer #2: I would recommend that this paper undergo another revision to reduce the history and provide more support for the theories that make up the framework. The figure demonstrating the framework was the strength of this paper and should become a greater focus. As it currently read, the history and legislation regarding food adulteration is the focus. The author's point can be made that there is a long history and much current activity in this field, and there is movement toward enhanced legislation, but there are many issues that cannot be solved with this approach. A framework to enhance investigation is required.

Response: History element has been reduced – Line 72 onwards – pg. 3 Wording changed to reflect comments on framework

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Abstract

9 The objective of this study is to explore the current strategies available to monitor and detect the 10 economically and criminally motivated adulteration of food, identifying their strengths and weaknesses and recommend new approaches and policies to strengthen future capabilities to 11 12 counter adulteration in a globalized food environment. There are many techniques used to detect 13 the presence of adulterants, however this approach relies on the adulterant or means of substitution 14 being "known" and no food item can ever be declared truly free of adulteration on that basis. 15 Further techniques will verify the provenance claims made about a food product e.g. breed, variety etc.as well as techniques to identify original geographic location of food production. These consider 16 wholeness, or not, of a food item and do not need to necessarily identify the actual adulterant. The 17 18 conceptual framework developed in this research focuses on the process of predicting, detecting and 19 reacting to economically and criminally motivated food adulteration.

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Introduction

Food adulteration is an age-old problem especially where there is a challenge between the physical availability of, and the market demand for, a food item. This is further impacted if there is juxtaposition between the cost of production, say of meat or meat-based products, and the price the supply chain customer (at a supplier/customer interface) or the end user is prepared to pay for the product. The objective of this study is to explore the current strategies available to monitor and detect the economically and criminally motivated adulteration of food, identifying their strengths and weaknesses and recommend new approaches and policies to strengthen future capabilities to counter adulteration in a globalized food environment. This paper begins by discussing the context of economically and criminally motivated food adulteration and then reviews the evolving techniques used to detect the presence of known adulterants, to identify product integrity, or otherwise, of foodstuffs as well as techniques to identify original geographic location of food production. A conceptual framework is developed and then its application discussed.

Whilst there is much focus in the literature, quite rightly, on the definitions of food safety and the 34 agents that render food unsafe there is less emphasis on the nature of product integrity or 35 wholeness. Adapting the term for "wholeness" in the Collins Dictionary (2013), the term product 36 integrity can be described as the inherent quality of containing all the component parts necessary to 37 38 form a total; i.e. completeness. Product integrity in this context could be further described as meeting the agreed specification that has been laid down in terms of expressing the total 39 40 completeness of the item that is "undiminished, without removal of part" (Adapted from Sykes 41 1976). By inference, failure to meet this specification indicates, to the limits of the testing methods, that a food may have been contaminated, have undergone substitution or has been adulterated. This 42 approach does not require the party undertaking the testing to identify the specific contaminant 43 rather just to identify that the specification of integrity for that commodity has not been met. As 44 analytical techniques become more accurate the depth of the specification of "what described 45 integrity" for a given food item will change and develop as discussed later in this paper. Defra 46 (2013) states that food standards legislation sets out specific requirements for the labelling, 47 composition and, in some cases, safety parameters for specific high value foodstuffs that are 48 49 potentially at risk of being misleadingly substituted with lower quality alternatives. This is as opposed to food safety that addresses food that is injurious to health (Food Safety Act, 1990). In 50 51 their Food Law Enforcement Plan 2010/2011, the London Borough of Tower Hamlets (2010: 3) 52 states that "standards inspections are seen as a second priority" to that of food hygiene and as a result, far less sampling for composition, labelling, claims, allergens, etc. is done. It is food standards that this research particularly focuses on and dependent on the adulterant or substitution concerned this may, or may not, also be a food safety problem.

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Adulteration in a globalized food environment

It has been suggested that global anti-counterfeiting activities for the food and drug sector are projected to be worth \$79.3 billion by 2014 (Li, 2013). In order to outline the context of this statistic this section compares and contrasts a number of food adulteration and fraud cases in both developed and developing countries.

62 United Kingdom / European Union

Scally (2013) argues that the lengthening of food supply chains, accompanied by the increased 63 industrialization of the food business, has had a profound effect on the food culture of developed 64 countries. Indeed he proposes that modern food processing has created the opportunity to practice 65 consumer fraud on a truly massive and international scale. The fraud can be undertaken in one 66 67 country and then the actual impact can be in countries far removed from the perpetrators especially so as the globalization and consolidation of food procurement increases further (Manning et al. 68 2005). Therefore, it is possible to contaminate food in a country where regulatory and market 69 70 controls are limited and cause major human health consequences and economic disruption in 71 another where on the surface such controls appear stringent.

Food adulteration can be described as the actions that are taken to add or adjust a food item or composite food product by the use of extraneous, substandard, or inferior ingredients. Food fraud may be carried out intentionally for economic gain, with the associated actions undertaken to avoid detection by regulatory bodies or consumers (Grundy *et al.* 2012). Economically motivated adulteration (EMA) has been described as "*The fraudulent, intentional substitution or addition of a substance in a product for the purpose of increasing the apparent value of the product or reducing the cost of its production, i.e. for economic gain.*" (Spink and Moyer, 2011:32). Economically and

criminally motivated food adulteration is nothing new. Accum (1820) identified that at that time 79 80 that there had been a range of successful prosecutions in the United Kingdom (UK) for counterfeiting and adulteration of tea, coffee, bread, beer, and pepper. These were both a concern 81 82 with regard to food safety as well as being of a food standards issue. Accum determined that adulteration was a widespread practice involving a number of food items and also exposed the 83 culinary fraud practices in London and detailed how bakers cut their flour with alum, chalk, plaster 84 85 and sawdust to make them heavier. Other fraud cases at the time involved brewers adding bitter substances such as strychnine to beer and the use of lead, copper or mercury salts to make bright 86 coloured sweets and jellies. 87

88 In April 2013, the European Commission reported on testing that had been carried out in the wake of concern over meat product adulteration (EC, 2013). The results indicated that, for the products 89 90 tested for the presence of horse DNA (n=4144), 4.7% revealed positive traces of horse DNA. For the products tested for the presence of phenylbutazone (n=3115) 0.51% showed positive traces of 91 92 the drug. In addition, Member States (MS) reported tests performed by food business operators 93 (producers, processors and distributors; n=7951) for the presence of horse DNA; 1.38% had horse DNA present. The UK Food Standards Agency (FSA) also identified products labelled as "Halal" 94 95 that contained pork (FSA, 2013). Beef adulteration in Europe highlights not only the continued problem with food fraud, but also the potential for unwitting cross-contamination at "micro levels" 96 97 during standard meat processing activities where multi species meats are processed/prepared in the 98 same vicinity and using the same equipment. This means that products (that would have previously been declared as "free from" or "whole" in terms of being suitable for a certain cultural or religious 99 100 group) as analytical methods develop, and as limits of detection reduce, may not indeed be found to 101 meet that specification. The discrepancy may be at the level of parts per million (ppm) or parts per billion (ppb) but this may not be acceptable to consumers e.g. in terms of pesticide residues or the 102 103 presence of DNA from other animal species. This creates a current and future challenge that the industry will need to address both in practical terms in trying to reduce these minimal levels furtherand also with meeting cultural expectations.

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107 United States

There is much work from the United States (US) that focuses on food fraud and food adulteration 108 (Everstine et al., 2013; Spink and Moyer 2013; Moore et al., 2012; Spink and Moyer 2011) As an 109 110 example of the types of incidents identified, a 2012 report on food fraud in US restaurants and retail outlets (Warner et al. 2012) concluded that 58% of the eighty-one retail outlets sampled, sold 111 mislabeled fish with small markets having a higher incidence of fraud (40%) than national chain 112 grocery stores (12%). Furthermore, all of the sushi bars (n=16) tested sold mislabeled fish and 94% 113 of the "white tuna" tested was not tuna at all. As previously discussed this type of adulteration 114 115 could be caused for a variety of reasons e.g. by accidental means due to a failure in either process or supply chain controls or as a result of premeditated criminal activity. 116

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118 **India**

One of the key problems in India is the intentional contamination of food with look-alike 119 substances. The look-alike substances were substituted in items like incidents of brick powder in 120 red chillies, lead chromate in turmeric and vegetable oil contamination with milk fat (Shukla et al., 121 2014). A 2011 survey in India of adulteration in liquid milk found that 68% of the randomly 122 collected samples tested (n=1791) were non-conforming (FSSAI, 2011). In some states the level of 123 non-compliance was 100%. The non-conformity of samples in rural areas was found to be 31% of 124 which 81% were loose (unpacked) samples. In urban areas 69% of samples were non-conforming 125 126 (67% loose samples). Detergent was found (8%); skimmed milk powder (45%) and glucose (27%) of the samples. In seven Indian states all samples taken were found to be impure. This demonstrates 127 the level of milk adulteration being practiced in India. The biggest dairy food fraud incident to date 128 129 using melamine, that also had serious implications for public health, was in China.

131 **China**

132 Melamine is rich in nitrogen and contains 67% nitrogen per mass unit (Merck Research Laboratories, 2001). Due to the high nitrogen content, melamine was added, as an adulterant, to 133 food commodities such as milk and wheat gluten to "increase" the perceived protein content and 134 135 avoided detection as milk was tested for protein using a method based on total nitrogen content (Schoder, 2010). In 2006 dairy production in China faced rising feed prices so 40% of dairy farmers 136 were losing money and a further 30% were just breaking (Jia et al. 2012). Whilst dairy processing 137 138 firms were demanding increased milk supply as a result of consumer demand some farmers were culling their herds due to the lack of profitability. This aggravated the already tight milk supply in 139 China. In early 2007 the new shortage of milk supplies threatened to push up the price of milk 140 141 products (Jia et al. 2012). The use of protein powders in milk was prohibited; such powders could be sourced from ground animals" parts, soy and other food sources. Later, manufacturers of plastics 142 started seeing a demand for melamine, but there was no connection made between the two 143 144 supposedly separate incidents.

An increased incidence of kidney stones and renal failure among infants was identified in China in 145 December 2007 and Sanlu Customer Service Department received consumer complaints about their 146 147 products (Xiaojing, 2011). [Concurrently there was a pet food recall for melamine contamination of pet food ingredients in the US due to contamination of wheat gluten.] In June 2008 complaints 148 149 appeared on the State Council Administration for quality, supervision, inspection and quarantine 150 (AQSIQ) website. Official inspectors then assessed the commodities produced by Sanlu, and once 151 adulteration was identified all batches produced up to December 2007 were recalled. In August 152 2008 melamine was reported as being detected in 15 out of 16 lots tested, but a recall was not 153 instigated until the government ordered Sanlu to stop production and distribution of product in September 2008 (Xiaojing, 2011). In that month it was announced that 59 infants had developed 154 155 kidney stones and one child had died. In September 2008, the WHO (2008) identified that there had

been 6240 cases of kidney stones in China with three deaths. The WHO reported that at least 22 dairy manufacturers across China were found to have melamine in some of their products (the levels varied between 0.09mg/kg and 2.560 mg/kg). Gossner *et. al.* (2009) determined that kidney and urinary tract effects, including kidney stones, affected about 300,000 Chinese infants and young children, with six reported deaths.

Further forty-seven countries received the melamine-contaminated products and sixty-eight countries banned or recalled foods suspected of containing melamine (Gossner *et. al.* 2009 citing Bhalla *et al.* 2009). Food fraud, as in this example, can occur in commercial circumstances when there is an issue with the bridging of the supply of and demand for a food commodity. Substitution can arise as a result of an illegal activity to fill the "supply gap" or to meet the cost structure at the stages of the food supply chain where there is a reticence or inability for increasing operational costs to be passed through to the end consumer.

As a result of this incident, the Chinese government was forced to react to ensure the safety and 168 169 quality of Chinese food products through the implementation of food safety laws, increasing 170 penalties for illegal practice and by instituting a system of risk evaluation that included monitoring 500,000 companies (Ramzy, 2009). It should be stressed that within the diverse and complex global 171 food supply chains there are constraints to addressing food safety, food standards and corruption at 172 local, national and international levels. Furthermore, maintaining confidence in a food supply chain 173 in order to ensure continued economic growth is not an issue localized only to China. The Chinese 174 case study merely serves as an example of the challenges presented with regard to control of food 175 adulteration. As Accum (1820) identified such activities were evident in a developing UK food 176 177 culture and the examples given in this paper highlight they continue to be prevalent today.

Although the use of melamine in China as a food adulterant gained attention from 2007, adulteration continues to be a problem with further arrests and prosecutions in China in 2011 (Coghlan, 2011). Melamine contamination has also been identified in milk purchased in twelve out of fourteen samples from markets in Iran (Hassani *et al.* 2013). These examples highlight the 182 continued use of this adulterant and why routine product testing for melamine is so critical to verify continued product compliance and to seek to prevent contaminated materials from being used in the 183 food supply chain and/or consumed. However, often food fraud is undertaken with the full 184 knowledge and understanding of the systems of surveillance and control and the analytical tests that 185 are currently used at borders and within countries. The constituents used for emerging and re-186 emerging food fraud are targeted on this basis either for the reason that they are not currently 187 188 routinely tested for in surveillance and verification testing and food import control protocols or that the adulterant used will pass existing analytical tests without identification. 189

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Economically motivated adulteration

Contamination maybe accidental or unintentional particularly when farmers or processors are 192 193 unaware of that a set of circumstances they put in place could potentially lead to contamination of food. However, when food adulteration becomes intentional, this is when criminal and 194 195 economically driven factors can come into play. Practices of deliberate contamination of food and 196 drug ingredients may be widespread and also avoid detection in poorly regulated markets where 197 surveillance is minimal. For example, in China there are over 500,000 food processing businesses and slim profit margins drove some owners to cut cost by substituting food with cheaper ingredients 198 (Zach et al. 2012). Substitution may include diluting infant formula (Xiu and Klein 2010), using 199 diethylene glycol as a substitute for glycerin (FDA 2008), using illegal red dyes in duck eggs (Du 200 and Sun, 2007) and relabeling of seafood products (D"Amico et al. 2014). If deliberate 201 contamination is motivated by financial gain, the practices are likely to be concealed and if 202 203 undiscovered, to recur (Brown and Brown 2010).

Due to their high market value, meat products are often targets for species substitution and adulteration (Cawthorn *et al.* 2013). A study undertaken in South Africa on processed meat products (n=139) identified that 68% of samples contained species that were not declared on the product labelling, with the incidence being highest in sausages, burger patties and deli meats i.e.

processed foods rather than carcass meats. Soya and gluten were identified as undeclared plant proteins in a large number of samples (28%), whilst pork (37%) and chicken (23%) were the most commonly detected animal species. Cawthorn *et al.* (2013) also reported that unconventional species such as donkey, goat and water buffalo were discovered as species that had been substituted for another origin. They conclude that mislabeling of processed meats is commonplace in South Africa and this not only violates food labeling regulations, but also poses economic, religious, ethical and health impacts.

In the EU, syndicates took advantage of the price-support structure of the European Common 215 Agricultural Policy for financial gain. For example, butter produced within the EU receives a 216 subsidy payment because of lower market prices when exported to a "third" (non-EU country). Then 217 the same consignment of butter was re-labeled as produce of the third country before being re-218 219 imported back into the EU. The re-labeled butter was subjected to income tax at a lower rate than the original subsidy paid on the export. Hence, by re-labeling the origin of the butter, syndicates 220 were able to make illegal profit of up to £30,000 per 25,000 kg consignment of butter (Kelly et al. 221 222 2005). Spink and Moyer (2011) identified seven types of food fraud (Table 1) namely adulteration, counterfeit product, diversion of products outside of intended markets, over-run, simulation, 223 tampering and theft. Each type of food fraud generates different potential levels of monetary gains 224 and the degree of gain is dependent on how well the "fraud" has been carried out and if detection of 225 the crime occurs. For example, when white sturgeon caviar is substituted with beluga caviar, 226 consumers pay five times more than they should for the product (Cohen 1997). 227

228

229 Take in Table1

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Everstine *et al.* (2013) argue that EMA incidents reveal voids in quality assurance testing methodologies that can be exploited for intentional harm. Indeed gaps in traceability, quality assurance programmes or interfaces between different certification schemes will be exploited where 234 they occur by some individuals for economic benefit. Everstine et al. (2013) suggest in their study 235 that 137 documented and distinct EMA incidents had been identified. The food product categories ranged from protein products to spices and sweeteners. Moore et al. (2012) determine that whilst 236 food ingredient fraud and EMA are emerging risks, a comprehensive database about known 237 problematic ingredients and detection methods did not exist until 2012 when the USP Food Fraud 238 Database was established. The proliferation of potential adulterants demonstrates that any 239 "screening based" approach needs to be diverse and wide reaching in its scope. Product testing can 240 be costly and introduce time delays, especially at border inspection points, in a food supply chain 241 that is both highly price sensitive and continuously driving towards a just in time approach to 242 minimize the costs of holding/storing stock. Organizations will vary in the extent to which they 243 use/undertake risk-benefit evaluations such as hazard analysis critical control point (HACCP) for 244 245 food safety and a threat or vulnerability analysis critical control point (TACCP or VACCP) assessment to determine the risk of vulnerability to fraud or bioterrorism incidents. These 246 approaches identify the process controls and product testing that is deemed necessary to minimize 247 248 risk to the organization, their customers and the final consumer (FDA, 2013a).

The WTO/SPS agreement (WHO, 1997) introduced the term "appropriate level of sanitary or 249 phytosanitary protection" (ALOP) i.e. the level of protection deemed appropriate by a Country or 250 Member State establishing a Sanitary and/or Phytosanitary (SPS) measure to protect human, animal 251 or plant life or health within its borders. By setting a food safety objective (FSO), competent 252 authorities can determine a risk-based limit that should be achieved operationally within the food 253 chain, while providing flexibility for different production, manufacturing, distribution, marketing, 254 and preparation approaches (CAC, 2007). Furthermore, a performance objective (PO) can be 255 256 determined i.e. the maximum frequency and/or concentration of a food safety hazard in a food at a specified step in the food chain before the time of consumption that provides or contributes to an 257 FSO or ALOP (CAC, 2011). However, the FSO and PO can only be determined if the food safety 258 259 hazard or contaminant is "known" and there has been a scientific risk-based determination of the

acceptable level of the hazard within a food. In the case of "unknown unknowns" this risk 260 assessment approach falls down. By its nature EMA is often within this category as the food 261 adulteration or substitution has the potential to cause harm if ingested. In instances of food fraud 262 only the fraudsters know how the food has been manipulated and to what extent the substitution is a 263 labelling or a food safety issue and also how it was introduced into the food supply chain. However, 264 265 the fraudsters may neither care nor have the knowledge, the expertise, or the resources to determine if the substitution or manipulation undertaken poses any acute or chronic risk to consumers. Hence, 266 the public health risks of adulterated food are often unknown until it is too late (Moore et al. 2012). 267 268 Spink and Moyer (2011) also state that the public health risks from adulterated food are more risky than traditional food safety threats because the contaminants are often unconventional. There are a 269 non-exhaustive number of potential EMA contaminants and a risk-based approach requires a high 270 271 degree of knowledge or expert opinion in order to appropriately quantify the level of risk. However such expert knowledge will be lacking or non-existent with some EMA, since this is the very reason 272 why they were chosen in the first place. Economic influences will create a situation where 273 274 alternative ingredients or materials are sought by supply chain partners that are "cheaper" than standard ingredients and can go largely undetected in the current product monitoring and 275 verification regimes. Food analysis is often at the accuracy level of ppm or ppb and this has led to 276 277 the development of techniques often described as food forensics. This particular field will need to develop strongly in order to meet the global challenges of food fraud. 278

279

280 Food forensics

The use of nonspecific analytical tests in routine product testing is one of the risk factor for the incidence of EMA (Everstine *et al.* 2013). The wide range of substances that can be used in food fraud coupled with the impossibility to analyse them all, make conventional testing unsuitable for food adulteration problems. In order to cover the widest range of adulterants usually requires sophisticated analytical equipment such as mass spectrometry (Di Stefano *et al.* 2012). It could be argued that the melamine adulteration incident occurred because the analytical method used to determine protein content was non-specific and thus by adulteration a "false" reading could be obtained. Kjedahl or combustion (Dumas) method measures the protein content based on total nitrogen content and do not differentiate between protein nitrogen or non-protein nitrogen (Moore *et al.* 2010). As a result of this, individuals took advantage of their ,,misused" food chemistry knowledge to enhance the determined level of the protein content of milk, knowing that the tests were of non-specific nitrogen tests.

293 The US Pharmacopeia (2012) advocates a proactive approach i.e. the testing of food ingredients for authenticity rather than testing for the absence of specific adulterants (Moore et al. 2012). Moore et 294 al. (2012) reviewed and collected over 1000 records of food frauds and analytical methods 295 published in the USP Food Fraud Database. The database is useful to identify trends and 296 developments and provide stakeholders with information on methods to detect food frauds. 297 According to Primrose et al. (2010), determining the description of food in terms of its total 298 composition, processing or origin is challenging, but there are a number of techniques that have 299 300 been successful in verifying the authenticity of food. This includes stable isotope analysis, 301 genomics and proteomics.

302 In 2005 a code of practice was developed for the control of basmati rice sold in the UK (BRC, 2005). If a product is identified as "basmati rice" then the non-basmati rice element cannot exceed 303 7% of the packed product. It is difficult to differentiate between basmati and non-basmati grains 304 305 based on visual test or physicochemical tests but research has been undertaken to identify adulteration of basmati rice as low as 1% in a sample through the use of tests that focus on variety-306 specific allele profiles (Archak et al., 2007). In the Uonuma district of Japan, high quality rice has 307 been bred with a specific genetic marker. The genetically distinctive rice sold under licence to 308 Uonuma farmers will prevent inferior rice from being falsely sold under the district"s name 309 310 (Ravilious 2006; Kitaoka et al. 2010). Kitaoka et al. (2010) suggested that the method would be able to identify food from a particular location. This is also of importance when considering 311

312 provenance i.e. the country of origin or geographic indication claims associated with food products. 313 Grundy et al. (2012) citing Kelly (2003) and Kelly and Bateman (2009) argue that analysis of stable 314 isotopes in foods can reveal EMA such as addition of cheap sugar syrups to extend honey and maple syrup; watering down of wine; preparation of fruit juice described as "freshly squeezed" 315 from concentrate; verification that chicken has been "corn-fed"; determination of whether ethanol 316 and vinegar and flavorings are natural or synthetic; and differentiation between organic and 317 conventional farming methods. All food and drink contains hydrogen and oxygen elements that 318 originate from where the animal or plant received water from the local water sources. Both 319 hydrogen and oxygen have heavy and light isotopes and the ratio of light to heavy isotopes is a 320 unique marker for climate and geographical area. Carbon isotopes can be used to differentiate plant 321 groups. Kelly et al. (2005) suggested that as a first approximation, natural abundance measurements 322 would provide information on plant "type" or diet (carbon and nitrogen isotope ratios), and 323 geographical origin (hydrogen, oxygen, sulphur and strontium isotope ratios). Therefore local 324 agricultural practices and animal diet can affect ¹⁵N/¹⁴N and ¹³C/¹²C ratios respectively. Indeed, the 325 326 geographic origin (rearing location) of animals used in meat production can be determined (Heaton et al. 2007). Beef reared in the US (n=23) and Brazil (n=10) was found to be isotopically different 327 from northern European beef (n=35), mainly because of contrasting proportion of plants with C3 328 and C4 photosynthetic pathways in the cattle diets (Schmidt et al., 2004). Isotopic maps of Europe 329 are being developed so that prized, regional products such as Champagne, Gloucestershire cheese 330 and Scottish salmon can be confidently matched with their places of origin (Ravilious 2006). More 331 332 recent research has utilized stable isotope techniques in reviewing egg authentication schemes (Rock, 2012); geographic origin of beef (Liu *et al*.2013); and authenticity and quality of food of 333 334 animal origin (Vinci et al. 2012).

One of the drawbacks of using purely chemical analytical techniques in seeking to detect food adulteration is that as previously described there is a finite number of analytes that have been determined and thus methods developed to determine their presence/absence at a defined limit of 338 detection. Utilising spectral or chromatographic techniques can identify patterns that can be 339 compared with standards for unadulterated foods and anomalies to be identified even if the exact constituent that is causing the variability is unknown. However in some instances such as the 340 adulteration of foods with Sudan 1 targeted analysis is required. This is true of spectral methods 341 such as near infra-red spectroscopy (IR) and nuclear magnetic resonance (NMR). Fingerprinting 342 refers to the spectrum or the image generated by certain analytical tools and the types of 343 fingerprinting can be classified into three categories (Table 2): spectral fingerprinting and 344 chromatographic fingerprinting and electrophoresis fingerprinting (Zhang et al. 2011). The use of 345 such fingerprinting technology has seen the detection of source, materials and components in food 346 such wines (Casale et al. 2010), cereals (Valeria et al. 2005) and fish protein (Hubert et al. 2008; 347 Serge et al. 2007). Table 3 shows the application of the different kinds of food fingerprinting in 348 food detection analysis. 349

350 Take in Tables 2 and 3

351

Additionally, DNA barcoding is a powerful method in determining morphologically unidentifiable 352 fish or meat product samples as long as the DNA is preserved in the sample (Maralit et al. 2013). It 353 is effective in determining the origin of raw materials and the detection of adulteration e.g. by 354 355 mixing products from different taxonomy such as rice and ginseng (Galimberti et al. 2013: Niu et al. 2011). The primary goal of DNA barcoding is to assembly reference libraries of code sequences 356 for known food species in order to develop reliable, molecular tools for identification (Hubert et al. 357 358 2008). DNA tests, sequencing and databases can be developed for all meat types and will make it possible to trace the meat to the individual animal type, breed and locality of origin along with 359 360 isotope analysis. In the UK, such tests are not part of routine surveillance and DNA sampling can cost £200 to £500 per food sample (Thomson 2013). This prohibits its use as an on-line quality 361 assurance and process test method. Having outlined the role of both product verification activities 362 what is the value of process verification in addressing EMA? 363

365 **Process vs. Product verification**

Food standards assessment activities focus on both product and process verification. Process 366 367 verification through the assessment of documentation, certification and traceability data is less costly than destructive product inspection and testing, but such verification rests on the ability to 368 assess valid evidence in terms of documentation, records, labelling and evidence of certification. 369 370 Fraud prevention and anti-counterfeiting tools can be used to track and trace movements of food products through the supply chain. Machine readable devices (barcodes, QR codes, data matrix) 371 allow a number of checks to be enhanced and the electronic data can be shared (Dabbene, Gay and 372 373 Tortia, 2013). Information shared between the different partners in the supply chain can decrease potential food frauds as the number of traceable units are documented and monitored for suspicious 374 transactions. 375

It is important that the traceable resource unit (TRU) or distinct batch must be uniquely identified 376 377 (Moe, 1998 citing Kim et al., 1995). Over time, product traceability methods have been developed 378 that are based on the ability to identify products uniquely as a result of physical marking on the product or its package or by the use of associated records (Moe, 1998). Moe argued that a 379 380 traceability system could be split into two elements firstly the "route" of the product and the sequence of steps that it passes through so it is traceable through manufacturing, distribution and 381 the retail system and the "scope" of the traceability in terms of the inherent nature of the product. 382 383 This has been built on in more recent years with the introduction of "mass-balance" traceability checks for a TRU. Mass balance traceability is an essential pre-requisite within the food supply 384 chain for assuring extrinsic quality. This process assures that identity preserved products are indeed 385 386 what they purport to be. Mass balance checks routinely determine an organization"s ability to identify, locate and "contain" a specific TRU of ingredient, part-processed or final product. The 387 388 capacity to do this is critical in the event of a product withdrawal or a full product recall from the 389 supply chain. It is also important to determine that the volume of product being sold as a specific

TRU where provenance, production method (organic, free range or Fairtrade) or cultural claim e.g. slaughter method (halal) and whether this could have indeed been produced in that quantity from the resources that were claimed to have originally been made available. This is largely an electronic record and/or a paper-based exercise especially if the "stock" has left the production premises. This is problematical when the reliability and authenticity of data is subverted in the event of food fraud. Therefore process verification alone is of limited value in determining or identifying EMA.

The UK Independent Farming Regulation Task Force in their 2011 report (IFRTF, 2011) 397 recommended that industry engage "fully with Government and third party assurance bodies to 398 develop a workable system of "earned recognition"". Third party certification schemes cover the 399 certification of the management of the production, storage and handling of the products at a discrete 400 401 point in the supply chain and are not, in the main, product specific certification schemes, although the generic product types are identified in the scope of certification for each organization. This 402 means that in their current form, third party certification schemes have limited impact on the control 403 404 of product verification only in as much as there was compliance with supply chain specifications on the day of the audit. This form of verification is more about the process and generic controls. 405 Furthermore, verification of process and product through review and auditing provides the auditor 406 with a range of evidence, or audit observations, which can be both qualitative e.g. interviews, 407 observations and records or quantitative based on measurement and test. However, it is important to 408 consider whether third party certification of organizations against management system standards 409 can either guarantee increased compliance with statutory food standards product requirements or 410 411 that such certification activities will address covert fraudulent behaviour which by its nature 412 involves the falsification of product, labelling and/or documentation at one point or several points in 413 the supply chain. If the records or labelling verified was:

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• falsified outside of the discrete bounds of the scope of the certification, and/or

- the processes being undertaken do not include re-confirmation of the validity of such
 documentation and labelling with the product batch delivered, and
- 417

•

there is no analytical or organoleptic evidence available of fraudulent activity when the product is being inspected,

then the fraud will not be readily identified or prevented by this type of third party certification. 419 Indeed, fraudulent behaviour, by its criminal nature, is unlikely to occur during a timetabled third 420 party certification audit. The Elliott Review Interim Report (HM Government, 2013) suggests that 421 the food industry moves to reducing the number of announced certification audits undertaken and 422 replacing them with unannounced audits. However unless the certification standards contain 423 specific elements that will be assessed with regard to EMA and food fraud this will have limited 424 425 benefit. The effectiveness of the certification activity depends upon the cooperation of the 426 organization being audited, which in the event of criminal activity may well mean the auditor will face limited disclosure. It should also be considered that if an auditor discovers criminal activity 427 during a certification audit, by the illegal nature of the issue the auditor"s well-being and safety 428 should be assured. 429

The process sampling activities used within such certification audits are constrained by the time 430 available, planned frequency of verification activities, volume of data to be assessed, any planned or 431 unplanned sampling bias, and the potential for deviation from the scope of the audit (Manning, 432 433 2013). Martz (2010) suggested that "evaluation myopia", the inability of the auditor to identify side effects or side impacts due to the rigid application and non-reflective use of a certification standard 434 or a "checklist" may also occur. This can lead to an auditor only verifying the effectiveness of the 435 436 control of food safety and food management standards criteria that have been defined in the certification or audit standard or are already "known". As already discussed the checklist does not 437 implicitly address food standards, but instead focuses on food safety and food quality, then the 438 potential for EMA, or its actual practice, might go unverified. The Elliott Report (HM Government, 439 2013) recommends that third party accreditation bodies should collect and analyse food surveillance 440

samples as this would act as an additional deterrent to food businesses knowingly trading in fraudulent food. This has potential to address known types of fraudulent activity; however emerging hazards or "unknown unknowns" are outside the scope of a biannual or triennial updating of a certification scheme and associated product sampling so emerging issues cannot be addressed by this approach and still pose an issue unless regular revision activities take place within the certification body and by the "standard owner" e.g. the British Retail Consortium. Therefore this approach has limitations in addressing EMA and food criminality.

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449 Role of food policy in minimising food adulteration

Food fraud that results in public health risk is often unknown until it is too late and the product is 450 already in circulation and has potentially been ingested. Even then the illegal activity may only be 451 452 identified by chance or as a result of a horizon scanning activity rather than from a formal riskbased approach or an annual third party audit. Predicting types of adulterants and ways of 453 manipulation can be carried out using the Rational Choice Theory (assuming rational choices by the 454 455 fraudsters which may not be the case) or indeed in terms of food bioterrorism where irrational behaviour may well underpin the behaviours that occur. The CARVER + Shock tool is a food 456 defensive tool to assess how vulnerable a food system or infrastructure is to an attack (Manning and 457 458 Soon, 2013). It allows food regulators to think like the attackers. This methodology has led to the development of Vulnerability Assessment Software (VAS) tool (FDA, 2013a). This has been 459 designed to be a prioritization tool that can be used to assess the vulnerabilities within a system or 460 infrastructure in the food industry in order to build an effective food defense system. Carver + 461 Shock and VAS tools focused on predicting attacks, but are not designed to assess vulnerabilities in 462 463 the food supply chain for EMA issues. The attacker(s) of a food system ultimately wants to hurt consumers, cause economic losses and/or reputation and to generate chaos. It is carried out with the 464 goal that the attack will be revealed within a period of time. Since food fraud or EMAs are carried 465 466 out for economical gains, fraudsters will conceal their act in order to gain as much profit as

467 possible. Similar systems can be developed to assess the likelihood of food fraud or EMA occurring in the food chain. In this case, the critical points for food adulteration are points where fraudsters 468 have the opportunity to use/substitute/addition different ingredients (i.e. agricultural/veterinary 469 inputs / processing stage) and different packaging/labeling (i.e. at packaging or distribution stage) 470 (Figure 1). In future, after incorporating food fraud methodology into certification standards, supply 471 chain assurance and product verification, it may be equally difficult to remember a national or 472 organizational food standards control programme without there being a food fraud preventive 473 system in place as it would be now a food safety system without the use of HACCP plans (Spink 474 475 and Moyer, 2013). The following section discusses the policy initiatives in the US, and UK/EU that address food adulteration including EMA. 476

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478 United States

479 The US Federal Food and Drugs Act 1906 was introduced to prevent the manufacture, sale, or transportation of adulterated or misbranded or poisonous or deleterious foods, drugs, medicines, and 480 481 liquors, and for regulating traffic therein (FDA, 2013b). The Meat Inspection Act (1906) was passed on the same day. This was superseded by the Federal Food, Drug, and Cosmetic (FDC) Act 482 of 1938, and then the Public Health Security and Bioterrorism Preparedness and Response Act of 483 2002 with Section 302 specifically addressing protection against the adulteration of food (FDA, 484 2013c). Section 302 gives high priority to increasing the number of inspections of food offered for 485 import with the greatest priority given to inspections to detect intentional adulteration. The US 486 487 passed the Food Safety Modernization Act (FSMA) in January 2011. This is considered a landmark law that shifts the food safety focus from reactive to preventive thus more in line with the European 488 489 approach. The FSMA addresses imported food safety under the Foreign Supplier Verification section where importers have the responsibility to verify inspection, testing and trace back systems 490 (FDA 2013d). In the US, there are three main federal agencies that have primary responsibility for 491 492 the safety of imported foods (Zach et al. 2012):

- Bureau of Customs and Border Protection (CBP);
- USDA Food Safety Inspection Service (USDA/FSIS); and
- US Food and Drug Administration (FDA)

496 Under the FSMA, these three agencies (CBP, FSIS, FDA) enforce, collaborate and communicate
497 between each other to reduce the risk of unsafe food.

498

499 United Kingdom / European Union

500 The UK introduced the Preventing the Adulteration of Articles of Food or Drink Act into law in 1860 and it was revised by the Adulteration of Food and Drugs Act 1872. This led to the formation 501 502 of the Society of Public Analysts in 1874. The advent of the "due diligence" defense in the UK Food Safety Act 1990 meant that organizations had to then prove that they were proactive in 503 ensuring the food they had been supplied was not injurious to health and was of the nature, 504 505 substance and quality demanded by the purchaser. The legislation differentiated between food that was sold at retail stages that was "branded" or "own-label" i.e. sold under the retailers" brand. 506 Under the Food Safety Act 1990, any supplier of a branded product was responsible for the safety 507 508 of that product, and enforcement could be taken against a wholesaler or retailer even if the offense was caused by other parties in the food chain (Lee, 2006). Whilst major multiple food retailers in 509 the UK gained commercial advantage from increased sales of own-branded food products, it also 510 exposed them to greater risks in the event of product failure. This encouraged retailers to institute 511 512 stringent private assurance programmes with their suppliers (Fearne, 1998). This so called "field to 513 fork" or "plough to plate" approach led to systems that were complex and very costly elements of 514 the procurement of own-label products (Henson and Northern, 1998). As a means to mitigate this cost the food retailers initiated the development of third-party inspection and then third-party 515 516 certification of their suppliers, as previously described in this paper whilst still seeking to maintain an acceptable level of risk with regard to product failure in terms of their own verification activities. 517

European legislation (EC Regulation 178/2002) lays down the general principles and requirements of food law, the establishment of the European Food Safety Authority (EFSA) and it also defined procedures in matters of food safety. Article 8 addresses protection of consumers' interests in the European Union (EU) and states that food law shall aim at the protection of the interests of consumers and "*shall provide a basis for consumers to make informed choices in relation to the foods they consume. It shall aim at the prevention of:*

524 *(a) fraudulent or deceptive practices;*

525 (b) the adulteration of food; and

526 (c) any other practices which may mislead the consumer".

The requirements of Article 8 also differentiate between food safety and food standards criteria. 527 This led onto the development of the Rapid Alert System for Food and Feed (RASFF) in Europe for 528 identifying non-conformance within the MS. The Emerging Risk Exchange Network (EREN) is the 529 principal body for exchanging information on emerging risks between the EFSA, MS, the EC and 530 also international organisations. The network consists of national experts and allows information 531 532 exchange through the facilitation of access to and exchange through sharing of databases (Randles, 2012). In the UK, the intelligence from the EREN network along with data from other sources feeds 533 into the Food Fraud Database. The data from these sources will feed into the predictive element of 534 535 the systems to address EMA and food crime on a global scale, however localised EMA and food crime also needs to be considered. 536

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Developing a conceptual framework

The conceptual framework developed as a result of this research focuses on the process of predicting, reacting and detecting economically and criminally food adulteration and builds on the work of Ribble *et al.* (2013) (Figure 1). At the beginning of the chain, integrity can be assured at a specific point that is before any potential attacks or substitution is possible. As the food and/or feed is utilized, produced or processed within the supply chain, or supply network, opportunities arise for 544 criminals and fraudsters to add/extract/substitute/mix/dilute the material with any substance that diminishes the integrity of such food. If EMAs were to take place at any point in the food chain, the 545 546 food safety and food standards system relies solely upon the reaction / detection protocols and system that have been developed. These protocols and systems may work through a process of 547 either passive or reactive surveillance activity. The use of supply chain intelligence needs to feed 548 into these protocols to enhance their ability to react to potential attacks or to suspicion of EMA 549 activity. Inspection protocols and product testing programmes are developed through a risk 550 assessment process that might only be undertaken on an annual basis and such attacks may occur 551 552 much more frequently. Further product testing has been focused historically on looking for specific "known" adulterants rather than determining the degree of product integrity. However as shown in 553 Tables 2 and 3 fingerprinting technologies are developing and their more widespread use will assist 554 to determine product integrity. Furthermore compliance, or not, with an integrity fingerprint does 555 not require the test to determine the actual agent used in an EMA, just that an attack has taken place 556 and that product integrity is now uncertain. If the food adulterant manages to bypass passive 557 558 mechanisms of control, the adulterated food may ultimately cause acute or chronic illness in the population or the concern over such illness cause substantial economic loss. 559

560 Concurrent risk assessment studies on economic and social factors (e.g. pressure on food prices, 561 animal disease outbreaks, or weather events causing crop loss) together with associated predictive 562 modeling can be utilized to predict the potential for EMA and wider food crime. Policy measures 563 introduced require the implementation of both predictive measures and also reaction and detection 564 methods.

565 Take in Figure 1

Prediction of food adulteration rests upon the appropriate analysis of intelligence through the use of predictive tools and expert knowledge. Cassidy and Buede (2009) argued that expert accuracy is, in general, no better than that achieved by chance as increased experience is often accompanied by an unjustified increase in self-confidence. They assert that there is a strong general tendency for 570 overconfidence when making predictions or statements of uncertainty, i.e. the predicted probability 571 of an event is often not calibrated with its actual likelihood of occurring based on the work of 572 Koehler *et al.* (2002), Yates *et al.* (1998) and Litchtenstein *et al.* (1982). Whilst this research was 573 looking at the ability to determine risk associated with issues such as whether it could be suggested 574 that this factor of expert accuracy is the same when qualitatively, or semi-qualitatively determining 575 the risk associated with food adulteration or food crime too. Koehler *et al.*, (2002) identified five 576 areas for calibrating expert judgment:

• Overprediction: always assigning probabilities that are high;

• Underprediction: Always assigning probabilities that are low

• Overextremity: overestimating high probabilities and underestimating low probabilities

• Underextremity: Underestimating high probabilities and overestimating low probabilities and

• Overconfidence: being either overprediction or overextremity.

Angner (2006) in his work on overconfidence with economic experts highlighted that 583 overconfidence increases with difficulty i.e. the more unknown a factor the more likely that 584 overconfidence occurs. Whilst this may in part lie within the requirements of the precautionary 585 586 principle associated with European food policy there is potential concern when considering EMA and food fraud that the expert assessment will be incorrect and then the resultant decision on the 587 actions to take. Anger (2006) further argues that in their role as "experts", individuals may not 588 589 receive adequate outcome feedback i.e. they will never know what would have happened in the absence of the implementation of their recommendations. It is equally important that the actual 590 591 outcomes of the implementation of their advice is fed back into the expert analysis of the future. 592 However it is important in this case in hindsight not to exaggerate the predictability of past events. 593 Therefore, how can the bias of overconfidence be mitigated in frameworks such as Figure 1? Angner (2006) suggests: 594

- Accepting that overconfidence will occur and if possible eliminating it over time by requiring experts to give arguments against their view and the reasons why they may be wrong and providing feedback on decisions that is frequent, prompt, and unambiguous;
- Require clarity in predictions and decisions so that they are not ambiguous and ensure predictions are on the public record; and
- Minimise interpersonal differences between experts.

In predicting EMA and food crime it is important to consider the contributing factors that influence 601 602 the incidence of food crime such as the motive, ability to detect the adulterant (known/unknown) the ability of the fraudster/criminal to cheat existing analytical tests, the strength of regulatory and 603 604 market controls at the point of adulteration/criminal activity and at the point of consumption, the economic or supply chain factors (pressure on food prices, factors impacting on balance between 605 supply and demand) and the complexity of supply chain and the influence of cross-border activity. 606 607 Databases and risk assessment measures as well as predictive modelling and intelligence gathering will be undertaken in order to identify the potential for EMA and food crime. Reaction and 608 detection measures will depend on the agents of adulteration/substitution and the type of food fraud. 609

- Table 1 identified seven different types of food fraud and the reaction/detection measures will vary.
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Conclusion

Activities to predict the potential for adulteration or even bioterrorism have an inbuilt weakness because the quantification of risk is usually based on historical data that may, or may not be available or may/may not reflect the actual risk now at any given time in the future. Food fraud that results in public health risk is often unknown until it is too late and may only be identified by chance rather than from a formal risk-based approach; however there is a need to develop such predictive models for the future.

Historically, analytical screening techniques were used to identify EMA, and wider food crime, butthis is only of value if the nature of the adulterant is known. There are evolving food forensics

techniques that will be able to determine food integrity through techniques such as isotope analysis or spectroscopy that do not require the contaminant to be known rather that food integrity or purity, to the level of detection, cannot be shown. This investigative framework is valuable as a means to fight food fraud/EMA. However, these tests are costly and will by and large, in the short term anyway, be used as a tool of verification and not as a form of analysis for routine batch release. Therefore they cannot be used as either a preventative control, or an on-line, real-time monitoring activity within an established quality plan.

The objective of this study was to explore the current strategies available to monitor and detect the 628 EMA and their relative strengths and weaknesses and recommend new approaches and policies to 629 strengthen future capabilities to counter adulteration in a globalized food environment. The 630 conceptual framework developed in this research focused on the process of predicting, reacting and 631 detecting economically and criminally food adulteration, with specific emphasis on calibrating the 632 confidence of experts as this underpins the horizon scanning, risk assessment and predictive 633 processes as well as informing the requirements to ensure effective reactions and detections are 634 635 undertaken.

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983 984 Table 1: Types of food fraud (Adapted from Spink and Moyer, 2011)

Туре	Definition
Adulteration	A component of the finished product is fraudulent
Counterfeit	All aspects of the fraudulent product and packaging are fully replicated
Diversion	The sale or distribution of legitimate products outside of intended markets
Over-run	Legitimate product is made in excess of production agreements
Simulation	Illegitimate product is designed to look like but does not exactly copy the legitimate
	product
Tampering	Legitimate product and packaging are used in a fraudulent way
Theft	Legitimate product is stolen and passed off as legitimately procured

Table 2: Classification of fingerprinting technologies (Adapted from Zhang *et al.*2011)

Methods	Electrophor	presis fingerprinting	Spectral fingerprinting	Chromatographic fingerprinting
	Biochemical fingerprinting	Protein electrophoresis, isoenzyme electrophoresis		
	DNA fingerprinting	Restriction fragment length polymorphism (RFLP) Random Amplified Polymorphic DNA (RAPD) Amplified Fragment Length Polymorphism (AFLP) Pulsed-field gel electrophoresis (PFGE)	Nuclear Magnetic Resonance (NMR), Infrared (IR) Ultraviolet and visible spectroscopy (UV) Mass spectrometry (MS)	Gas chromatography (GC) High performance liquid chromatography (HPLC)

Table 3: Application fields of fingerprinting in food detection (adapted from Charlton, 2010; Niu*et al.*, 2011; Sefc*et al.* 2000; Woolfe and Primrose 2004; Zhang *et al.* 2011)

Application domain	Products	Detection indicators	Detection Technology
Origin	Tea, beer, mutton,	Microelements, water,	NMR, IR, PCR
	olive oil, wine	lipid, protein,	
		carbohydrate, aromatic	
		compound, isotope	
		indicators	
Material/species	Bird's nest, aquatic	Protein, DNA	SDS-PAGE,
	product, poultry,		Isoenzyme
	vegetables, Basmati		electrophoresis, RFLP,
	rice, Genseng		RAPD, AFLP, small
			sequence length
			polymorphism
			(SSLPs)
Component	Milk, fruit, edible oil,	Protein, lipid, lecithin,	SDS-PAGE, NMR, IR,
	tea, beef, ham, health	vitamins, sugars,	UV, MS
	products	organic acid,	
Additive	Meat, milk, juice,	Nitrite, sufan,	UV, GC, LC, MS
	processed food,	melamine, clebuterol	
	carbonated beverages,	hydrochloride,	
	ice-cream	colorants, antiseptic	
Objectionable	Fried starch products,	Acrylamide, trans-fatty	UV, GC, LC, MS
constituent in	margarine, barbeque	acids, benzopyrene	
processing			

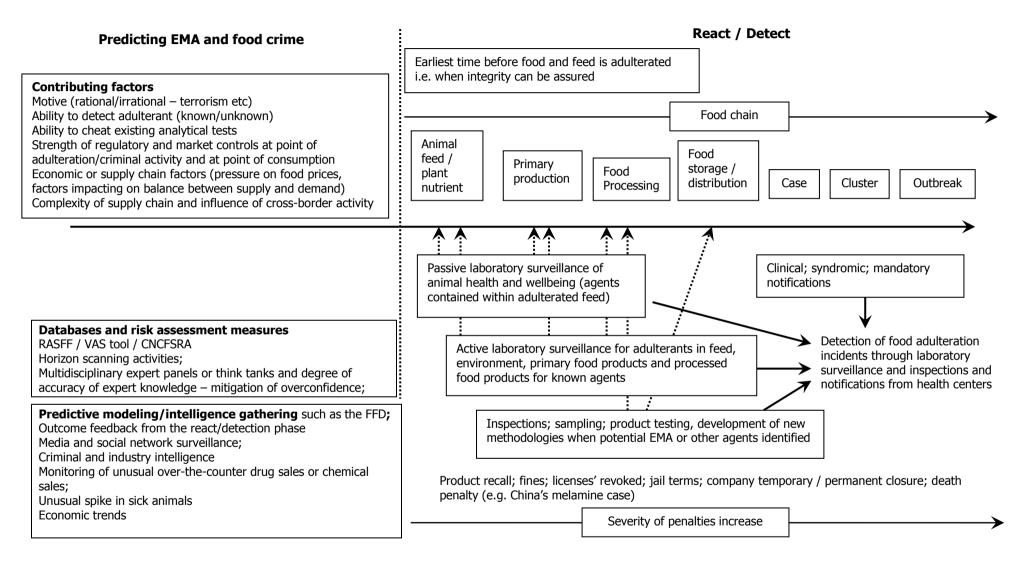


Figure 1. Predictive and reactive systems for food adulteration – role of food policy and risk assessment centres (adapted from Ribble et al. 2013) (Note: RASFF: Rapid Alert System for Food and Feed; VAS – Vulnerability Assessment Software; FFD – Food Fraud Database; CNCFSRA: China National Center for Food Safety