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An overview of the applications of total reflection X-ray fluorescence spectrometry in food, cosmetics, and pharmaceutical research

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The need for elemental analysis in quality control of food, pharmaceuticals and cosmetics, taking into account human safety, has become an essential requirement for any product destined for commercialization and targets in particular potentially toxic elements. The main objective is to obtain reliable data on elemental content that enables end-users to make informed decisions and to address problems effectively. To align with the principles of green and sustainable analytical chemistry, recent efforts have focused on incorporating new, low-cost screening and quantitative analytical tools such as total reflection X-ray fluorescence spectrometry (TXRF). This paper provides an overview of recent TXRF applications in the elemental analysis of food, pharmaceuticals, and cosmetic samples. The state-of-the-art procedures for sample preparation with details on the sample type and amount, dilution steps, mixing agents, analysed elements, and addition of internal calibration standards are presented together with analytical parameters such as limits of detection and quantification. The current challenges for applying TXRF to each of these research fields are discussed.

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1 Introduction

The determination of elemental composition in consumer products such as foodstuffs, pharmaceuticals, drugs, supplements, and cosmetics is a necessary step in industry and quality control laboratories to ensure product quality and safety.

The analytical practices involve the examination of many samples and, in most cases, the consumption and generation of high amounts of reagents and wastes, especially when atomic spectrometry techniques such as atomic absorption spectrometry (AAS) and inductively coupled plasma (ICP) based spectrometry are used. Consequently, the running costs are high, and these methods may involve time-consuming¹ sampling and sample procedures steps, when the sample cannot be directly measured. To mitigate these issues, new low-cost analytical

tools which incorporate green and sustainable analytical chemistry principles are desirable.² These tools should be able to assess the elemental composition while presenting the following characteristics: minimum sample treatment, high throughput, and adequate specificity and sensitivity for monitoring the elements of interest at the expected concentration levels.

Energy dispersive X-ray fluorescence (EDXRF) and wavelength dispersive X-ray fluorescence (WDXRF) spectrometry have been widely used in many industrial applications as screening tools and for quantitative analysis of major, minor, and trace elements in solid samples. However, the lack of matrix-matched reference materials for calibration limits their widespread use.³ Total reflection X-ray fluorescence (TXRF) is an alternative approach. TXRF takes advantage of the very low background and the higher excitation level that are achieved when a suitable sample is placed onto a reflective carrier, on which is shined an X-ray beam at an angle below the critical angle of the sample reflector material.⁴ Consequently, in comparison to other X-ray fluorescence (XRF) techniques, TXRF presents lower limits of detection (ranging from 10^{-7} g to 10^{-12} g) that may be applied to solid and liquid samples for simultaneous multi-elemental information. Reliable quantitative analysis may be obtained incorporating an internal calibration standard (ICS) to the sample, adequately prepared and deposited on a reflective carrier to comply with the thin film quantification approach. A benefit of TXRF is the possibility of

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analysing samples using simple and straightforward sample treatments, compared to acid digestion steps required by other atomic spectrometry techniques, offering thus unique screening possibility. Screening is undoubtedly a strategic way of investigation which may answer to some specific characterization demands, especially within industrial or regulatory contexts, where elemental occurrence and concentration thresholds are answers highly sought, but do not necessarily require accurate quantifications. TXRF has, however, the potential to be applied for both screening and qualitative or quantitative purposes. Qualitative and quantitative analysis aim at identifying the elements present into a sample, and at determining the concentration or mass, together with the related uncertainty, of one or more elements present into the sample, respectively. This, especially in the case of quantitative analysis, may require the development of specific sample preparation and handling procedures. In contrast, more expeditious procedures solely intending to identify, and select from a starting sample set, a group of samples that will contain one or more elements, eventually above a pre-set concentration level, is the goal of screening procedures. There, some shortcuts in the sample handling may be considered, as a yes/no answer is usually sought, without the need of thorough analysis. As TXRF unravel multi-elemental content, simultaneously, it is a promising analytical technique for consumer products. These advantages open interesting new applications for TXRF, especially in food, pharmaceuticals, and cosmetics.

In the last decade, the development and commercialization of benchtop TXRF spectrometers, offering extreme simplicity of operation in a low-cost compact design, have promoted the application of these systems in many research fields, including environment,^{5,6} geological,^{7,8} biological,^{9,10} medical,¹¹ and nuclear materials.¹² The use of TXRF in pharmaceuticals and cosmetics is very seldom, while in the field of beverages and foodstuffs it has been more widely employed.¹³ However, TXRF analysis of consumer products is still limited to research purposes and is rarely conducted as a daily routine analysis within an industrial framework (with the exception of Si wafers). Some of the limitations impairing the widespread use of TXRF, rely on the fact that TXRF is not considered as a technique of reference, due to a poor normative framework and lack of matrix-matched reference materials for calibration purposes.³ It, furthermore, lacks associated standard operating procedures, and as such is not included by regulatory bodies in the instructions or guidelines for compulsory testing.

The introduction of new techniques and methods in regulations and standards can be challenging due to the difficulty of accreditation procedures, the need to achieve a consensus between the elemental analysis experts, and consequently of worldwide recognition. Such ventures are based on a thorough assessment of the analytical performance, embracing the available guidelines of each application field and on proper validation of the developed alternative methods,¹⁴ *via* for instance proficiency tests. This review presents an overview of the published methods, protocols, sample preparation and handling procedures for the TXRF analysis, according to the

needs and regulations in the various application fields for quality and safety.

2 Elemental analysis for quality and safety of consumer products

2.1 Foodstuff

The definition of food quality is global and comprises both positive attributes, such as appearance, origin, texture, flavor, and nutritional content and value, as well as negative ones, such as contamination, discoloration, off-odors, and spoilage. Quality standards are essential in the European market to meet consumers' and regulatory bodies' requirements, and specifically target products' attributes, consequently affecting the value of the product for both producers and consumers. For industry, those attributes are evaluated *via* three key factors: conformity with the product's intended purpose and regulatory requirements, safety, and satisfaction of consumer expectations and perceptions.

Food safety refers to all hazards, whether chronic or acute, that may negatively impact consumer health according to the Food and Agriculture Organization (FAO) and the World Health Organization (WHO) (2003).¹⁵ The EU Food Safety Authorities (EFSA),¹⁶ regulates food content, ingredients, product description, chemical safety, sanitary conditions, hygiene, and other product specifications. The legislation on contaminants is based on scientific evidence and on the principle of keeping contaminants' levels as low as reasonably achievable by using good working practices. Maximum concentration levels acceptable in products have been established for specific contaminants, including heavy metals, to protect public health.

Nonetheless, the contamination of food with heavy metals remains a relevant issue for public health. Therefore, it is essential to ensure food safety by using various analytical techniques for metals identification and quantification. In particular, the EU Regulation (EC) No. 1831/2003 sets the maximum levels for potentially toxic metals in food. For example, the allowed Pb levels range from 0.02 to 2 mg kg⁻¹ wet weight for raw milk and food supplements, respectively. Cadmium limits are set between 0.005 mg kg⁻¹ wet weight for liquid infant formulae from cow milk and 3 mg kg⁻¹ for food supplements from seaweed. The permitted mercury levels are 0.1 mg kg⁻¹ wet weight in food supplements and 1 mg kg⁻¹ in muscle fish. The European Commission also established maximum levels for inorganic tin in canned baby foods, infant formulae and other canned foods, excluding beverages.¹⁷

Besides the interest in toxic elements, the determination of other elements presents in food samples, such as major essential elements, trace essential elements, ultra-trace essential elements, and non-essential trace elements, is relevant.¹⁸ Major elements are usually found in the percent range, while trace and ultra-trace elements span over a wide concentration range from ppm to ppb.¹⁹ The level of these elements varies widely in food samples, even within similar matrices. For example, the content of Ca within a group of 35 different alcoholic beverages, like beers, wines, and liqueurs, may vary from 3



to 50 mg g⁻¹; in cookies the content of Fe may fluctuate from 0.2 to 12 mg g⁻¹; and in raw apples the content of Mg may be from 0 to 4%.²⁰ Some of these elements, such as Na, Ca, Mg, K, and Cl, are macronutrients required in larger quantities for healthy nutrition, while others, such as F, V, Cr, Mn, Fe, Co, Cu, Zn, Se, Mo, and I, are micronutrients essential for plants, animals, and human nutrition. Excessive amounts of these elements can have phytotoxic and/or zootoxic effects.²¹ Fe, Zn and Mn, for instance, are toxic to humans if their daily intake level is exceeded.²² Thus, the determination of the elemental composition and concentration in food is crucial for making informed recommendations for consumption, especially for children, pregnant women, and people with specific elemental allergies or medical conditions. The interest in elemental analysis is increasing in agriculture and the food industry, with many papers focusing on the elemental composition of seasoning products,¹⁹ salts, and spices.^{23,24} Contamination due to the environment^{22,25} and food processing must be avoided²⁶ or reduced²⁷ whenever possible, but in any case controlled. Consequently, there is a strong need for fast screening tools and techniques with high sensitivity and applicable over a wide dynamic range of concentrations.

The Association of Official Agricultural Chemists (AOAC) developed the standard method performance requirements (SMPRs) for heavy metals in foods and beverages to ensure reliable monitoring through routine surveillance methods, defining the required values for concentration range, limit of quantification, repeatability, reproducibility, and recovery irrespective of the analytical techniques.²⁸ Atomic and inorganic mass spectrometries are widely used to obtain accurate information about trace levels of metals in food. Furnace atomic absorption spectrometry (FAAS), graphite furnace atomic absorption spectrometry (GFAAS), inductively coupled plasma optical emission spectrometry (ICP-OES) and inductively coupled plasma mass spectrometry (ICP-MS) are the most used analytical techniques due to the availability of good laboratory practices and standard operating procedures, which, when followed, aim to ensure analysis quality.²⁹ For a solid sample analysis, the X-ray fluorescence spectrometry is still the method most widely employed.^{30,31}

2.2 Pharmaceuticals, drugs, and supplements

The interest in the determination of elemental impurities in pharmaceuticals has increased in recent years due to greater awareness of their potential toxicity, changes in regulatory requirements, and the need for updating limit tests in pharmacopeia.³² These impurities can arise from various sources, including catalyst residues, raw materials, solvents, and container/closure systems, at different stages of the manufacturing process. Thus, their identification is required during chemistry route scouting and optimization, process development phase, or catalyst screening and loading.³³ While some elements may be intentionally added as active pharmaceutical ingredients (*i.e.*, platinum-based chemotherapeutics agents, antimicrobials containing Fe, Ag, and Au, among others),³⁴ others pose no therapeutic benefit to the patient and

must be maintained within acceptable limits. The relevant side-effects of metal impurities on the safety of drugs (adverse pharmacological and toxicological effects), or on their stability and efficacy (causes of intolerance and degradation), led to an increasing interest in their contents in drugs.³⁵ Regulatory bodies such as the European Medicines Agency (EMA) in the International Council for Harmonisation of Technical Requirements for Registration of Pharmaceuticals for Human Use (ICH) from the Q3D working group on elemental impurities, known as ICH Q3D guidelines,³⁶ and the United States Pharmacopoeia (USP) in the USP<232> chapter³² have categorized elemental impurities into three classes and established concentration limits according to a risk-based strategy and their toxicological properties, daily dosage budget fraction, and route of administration. The elements included in each class are: Class 1 (As, Cd, Hg, Pb), Class 2 (Co, V, Ni, Ti, Au, Pd, Ir, Os, Rh, Ru, Se, Ag and Pt), and Class 3 (Li, Sb, Ba, Mo, Cu, Sn, and Cr). Class 2 elements are divided into subclasses 2A and 2B based on their relatively probability of occurrence in the drug. In Table 1, concentration limits in (μg g⁻¹) for components used in oral drug products with a 10 g maximum daily dose are reported.

Recently, the USP also proposed a general chapter <232> about elemental contaminants in dietary supplements (DS)³⁷ to be monitored in finished dosage forms (*e.g.*, powders, tablets, capsules, liquids) and not in dietary ingredients.³⁸ The permitted daily exposure (PDE) endorsed by the FAO/WHO has been used to set the limits of the four significant elements of toxicological concern (As, Cd, Pb, Hg).

Previous pharmacopeia tests for heavy metals in the European Pharmacopoeia (Ph. Eur.), USP, British Pharmacopoeia and Japanese Pharmacopoeia were based on the reaction of metals with the thioacetamide reagent, resulting in the formation of coloured sulphides and their precipitation from aqueous solution.³⁹ This visual colorimetric test suffers from serious limitations such as the lack of sensitivity, specificity, and recovery to monitor properly the actual levels for some metals. For instance, elements of toxicological concern (*i.e.*, Cr, V, Ni) cannot be determined by this classical limit test, and biased results were reported for elements such as As, Cd, Mo, Pd, Pt and In with poor recovery values between 30–50%.⁴⁰

The USP chapter <233> includes two analytical procedures involving ICP-OES and ICP-MS.⁴¹ One of the major limitations of such procedures is the treatment of solid pharmaceutical samples that must be solubilized to be compatible with the sample introduction systems of ICP-OES and ICP-MS instrumentation. Furthermore, pharmaceutical products are not sufficiently soluble in deionized water or dilute acids,⁴² and solubilization using organic chemicals or concentrated acids is required.⁴³ When the sample is difficult to dissolve, a digestion method based on different acid mixtures (*i.e.*, mixtures of HNO₃ with HCl, HF, HClO₄, H₃PO₄, and H₂SO₄) is employed.³⁴ Notwithstanding, some pharmaceutical products are extremely difficult to digest even with the use of high-pressure systems and concentrated acids⁴⁴ and consequently, a universal method of sample decomposition, which applies to all dosage forms of pharmaceutical samples, does not exist.



Table 1 Concentration limits in $\mu\text{g g}^{-1}$ for components used in oral, parenteral and inhalation drug products with a 10 g maximum daily dose (according to ICH-Q3D³⁶ and USP<232>³²) and for dietary supplements (according to USP<2232>³⁷)

10	Class	Drug products, drug substances and excipients		Parenteral drug products		Inhalation drug products		Dietary supplements ^a
		ICH Q3D	USP<232>	ICH Q3D	USP<232>	ICH Q3D	USP<232>	USP<2232>
Cd	1	0.5	0.5	0.2	0.2	0.3	0.2	0.5
Pb	1	0.5	0.5	0.5	0.5	0.5	0.5	0.5
As	1	1.5	1.5 ^b	1.5	1.5	0.2	0.2	1.5 ^b
Hg	1	3	3 ^b	0.3	0.3	0.1	0.1	1.5 ^c
Methyl-Hg	1	—	—	—	—	—	—	0.2 ^d
Co	2A	5	—	0.5	—	0.3	—	—
V	2A	10	10	1	1	0.1	0.1	—
Ni	2A	20	20	2	2	0.5	0.5	—
Tl	2B	0.8	—	0.8	—	0.8	—	—
Au	2B	30	—	10	—	0.1	—	—
Pd	2B	10	10	1	1	0.1	0.1	—
Ir	2B	10	10	1	1	0.1	0.1	—
Os	2B	10	10	1	1	0.1	0.1	—
Rh	2B	10	10	1	1	0.1	0.1	—
Ru	2B	10	10	1	1	0.1	0.1	—
Se	2B	15	—	8	—	13	—	—
Ag	2B	15	—	1	—	0.7	—	—
Pt	2B	10	10	1	1	0.1	0.1	—
Li	3	55	—	25	—	2.5	—	—
Sb	3	120	—	9	—	2	—	—
Ba	3	140	—	70	—	30	—	—
Mo	3	300	300	150	150	1	1	—
Cu	3	300	300	30	30	3	3	—
Sn	3	600	—	60	—	6	0.1	—
Cr	3	1100	1100	110	110	0.3	0.3	—

^a Finished dosage forms. ^b Inorganic form: speciation is not necessary when the total mercury or arsenic is less than the limit. ^c Mercury (total).^d Methylmercury (as Hg): speciation is not necessary when the total mercury is less than the limit.

An additional inconvenience of the use of ICP-based techniques for the determination of elemental impurities in pharmaceutical products is the presence of spectral and non-spectral interferences and thus, a previous evaluation of the matrix composition should be carried out before even starting the analysis of the sample, to identify those. For instance, it is interesting to mention that almost 20% of the active pharmaceutical ingredients are hydrochlorides and spectral interferences of chlorine in ICP-MS analysis are expected.⁴⁵ For this reason, although ICP platforms are predominant for most applications dealing with inorganic impurity analysis in pharmaceutical laboratories, the use of other analytical approaches which involve a simpler sample treatment is worth of investigation. In fact, the USP chapter <233> describes the criteria for the acceptability of alternative procedures equivalent to ICP-OES and ICP-MS ones.⁴¹ Table 2 summarize those analytical characteristics and acceptance criteria, which are required in the validation of alternate quantitative approaches and test limit procedures.

In this view, the direct analysis of solid pharmaceutical samples by means of XRF technique demonstrated promising results, for instance, for the determination of Pd, Zn, Fe and Ni residues in active pharmaceutical ingredients (API).^{46,47} The major drawback reported in these studies was the lack of suitable reference materials with a matrix similar to the real

samples for validation and calibration purposes. To overcome this problem, the use of synthetic standards made of cellulose simulating the API matrix appeared to be an effective means to obtain reliable calibration curves when no suitable pharmaceutical reference materials were available. More recently, XRF potential for screening oral solid dosage drug products for elemental impurities was demonstrated.⁴⁸ The proposed method can be used as a limit test for class 1 and class 2A elements, and is fully compliant with the method validation requirements according to the ICH Q3D guidelines. It is worth to mention that in 2015 the USP included XRF as an instrumental method that can be used for both qualitative and quantitative analysis of liquids, powders, and solid materials (USP chapter <735> X-ray fluorescence Spectrometry),⁴⁹ but recommends only the EDXRF and WDXRF techniques. The same applies to the last edition of Ph. Eur.⁵⁰

2.3 Cosmetics

Cosmetics have drawn attention because of the potential negative effects of the chemicals they might contain. For cosmetic companies, proposing safe and high-quality cosmetic products become crucial, not only for regulatory purposes, but also because customers have been increasingly looking for documentation of efficacy, compliance, and quality on raw

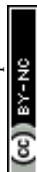


Table 2 Required analytical characteristics and acceptance criteria in the validation of alternate quantitative and test limit procedures according to the USP<233>. ⁴¹

Analytical parameter	Acceptance criteria	Quantitative method	Test limit
Accuracy	Recovery: 70–150% ($n = 3$) Levels: 50, 100, 150% limit ^a	Yes	No
Precision Repeatability	RSD $\leq 20\%$ ($n = 6$) Level: 100% limit ^a	Yes	Yes
Intermediate precision	RSD $\leq 25\%$ Repeatability study on different days, instruments and analysts	Yes	No
Specificity	Unequivocally assessment of the analyte in the presence of other target elements and matrix components	Yes	Yes
Detectability	Signal spiked sample 1 ^b $\pm 15\%$ of the average signal value for standard solution ^c Signal spiked sample 2 ^d lower than the average signal value for standard solution	No	Yes
Quantification limit (LOQ)	Demonstrated by meeting the accuracy requirement	Yes	No
Linearity	Demonstrated by meeting the accuracy requirement	Yes	No
Range	Demonstrated by meeting the accuracy requirement	Yes	No

^a Limits are reported in Table 1 (USP<232>). ^b Spiked sample 1: $1.0 \times \text{Limit}$. ^c Standard solution: $1.0 \times \text{Limit}$. ^d Spiked sample 2: $0.8 \times \text{Limit}$.

materials to finished products.¹ Potentially toxic metals may originate from various sources such as raw materials, metal-coated equipment during manufacturing, and intentional addition as pigments and ingredients.⁵¹ To mitigate against the potential toxic effects of some heavy metals, the European Union (EU) Directive prohibits their intentional use, for instance of Sb, As, Cd, Cr, Co, Hg, Ni, and Pb.⁵² The amount of metals permissible in cosmetic products is assessed based on a broad concept approach of risk evaluation because the EU regulations have not set so far specific limits. Using estimated margins of safety values that take into account both potential negative effects and systematic exposure dose, the Scientific Committee Consumer Safety (SCCS) of the European Commission has determined recommended maximum thresholds for

heavy metals in cosmetic items. As visible in Table 3, other international agencies and governments from all over the world have also established metal concentration limits for inorganic contaminants in cosmetic products. While the amount of Hg, Pb, As, Cd, and Sb are regulated, other metals such as Cr, Ni, and Co are not always considered due to their lower health risk. The strictest limitations are those enforced in Germany,⁵³ which are based mostly on levels that can be practically avoided rather than on risk. There isn't much agreement, though, on which metals or inorganic species, and in which amount, require control in cosmetics. Currently, cosmetics manufacturers are not required to disclose the metal contaminants found in their products, and consequently customers have no way to know the risks involved in using cosmetics on themselves. Several studies

Table 3 Recommended maximum levels in $\mu\text{g g}^{-1}$ for inorganic species in cosmetics according to different international administrations and governments

Element	FDA ^a (US)		BVL ^b (Germany)		FDA-CR ^c (Canada)		SCCS EC ^d (Europe)	
	Limit ($\mu\text{g g}^{-1}$)	Cosmetic type	Limit	Cosmetic type	Limit ($\mu\text{g g}^{-1}$)	Cosmetic type	Limit	Cosmetic type
Hg	65	Eye products	0.1	All	1	All	1	All
	1	Other cosmetics and colour additives	0.1	Toothpaste				
Pb	10	Externally applied cosmetics	2 ^e	All	10	All	20	All
	20	Colour additives	0.5	Toothpaste				
As	3	Colour additives	0.5 ^f	All	3	All	5	All
			0.5	Toothpaste				
Cr	50	Colour additive (FD&C Blue No.1)	—		—		100 (Cr III)	All
							25 (Cr VI)	All
Cd	—		0.1	All	3	All	5	All
			0.1	Toothpaste				
Sb	—		0.5	All	5	All	100	All
			0.5	Toothpaste				
Ni	—		10 ^g	All	—		200	All
Co	—		—		—		70	All

^a Food and drug administration (FDA). ^b Country's federal office of consumer protection and food safety (BVL). ^c Food and drug act (FDA) and Cosmetic regulations (CR). ^d Scientific committee consumer safety of the European Commission (SCCS EC). ^e For make-up powder, rouge, eye shadow, eyeliner, kajal as well as theatre, fan or carnival make-up: $5 \mu\text{g g}^{-1}$. ^f For theatre, fan or carnival make-up: $2.5 \mu\text{g g}^{-1}$. ^g Soluble Ni according to the DIN EN 71.



in a purely empirical way. Unfortunately, those are hardly verifiable and consistent with the physic processes undergoing into the sample system they represent, in particular with respect to the fundamental parameters. Alternatively, purely theoretical calibrations are determined, based on the fundamental parameters of pure elemental solutions, which consequently are not considering the sample characteristics under study. In both cases, the uncertainty budget linked to these external calibrations may be extremely important, and the results disputable.

A few commercial reference materials are available but mostly only certify the concentration of a single element in a specific cosmetic product, *e.g.*, CBW09302 cream cosmetic with a Pb concentration of $14.6 \pm 0.9 \text{ mg kg}^{-1}$. To increase confidence in the quality of results, participating in proficiency tests organized by global producers of reference materials, such as the Laboratory of the Government Chemist (LGC Ltd, UK), may be useful to at least get a feedback on the methodology employed.

3.1 Elemental quantification and figures of merit

Reflectors are typically 30 mm diameter and 3 mm thick disks made of materials like quartz glass, glassy carbon, sapphire, or acrylic glass, and which are chemically pure, smooth, and flat, and free of impurities. The carriers have to reflect X-rays efficiently, present low background levels, and be chemically inert and stable during analysis. The carrier surface needs to be cleaned prior to use to avoid memory effects, and cleaning procedures vary based on the carrier material and previously analysed samples. Cleaning can take up to 3–4 hours and involves precleaning with acetone, followed by several steps with cleaning solutions and diluted acid nitric. Usually a set of reflectors is cleaned and preserved for future use to save time. Improper handling, storage, or contamination may alter carrier properties, affecting analysis results.

The quantification approach making use of the internal calibration standard (ICS) is, and by far, the most used one as it is the simplest one. It relies on a mono-element solution of known concentration, added during the sample preparation stage to the sample to be analyzed to act as ICS. It also requires that the sample under analysis complies with TXRF requirements, *i.e.*, a controlled mass deposition, no matrix effect,

the illuminated area by the X-ray beam. Sample mass deposition must be adjusted based on the sample matrix, the analytes present and the X-ray excitation energy employed. Sample homogeneity must also be ensured, both in terms of surface and thickness, and in terms of elemental distributions, especially for the ICS.

Many procedures proved to be suitable for the analysis of solid and liquid samples by TXRF.⁶⁴ Various strategies have been used for screening and quantitative analysis of foodstuffs, pharmaceuticals and cosmetics samples, and the selection depends on the purpose of the analysis, the sample availability, and the matrix complexity. An overview is presented in Fig. 1. An internal calibration standard solution is added for quantification purposes, and a few microliters of the mixed solution are deposited on a suitable carrier for analysis. Blanks, duplicates, and replicates should be prepared and analysed at each step of the sample handling and preparation procedure.

Direct sample analysis is simple, efficient, and mostly preferred for liquid samples, either as-is or following dilution. Ultrapure water is usually employed as diluting agent to avoid contamination, and the dilution factor is kept in the range of 1–30, depending on the complexity of the sample and the analyte concentration. However, it should be avoided with samples containing a complex organic matrix, where sample solubilisation is recommended to improve accuracy and detection limits. Solid samples are typically decomposed by dry or wet digestion before analysis, as digestion produces in general a homogeneous liquid mixture. Indeed, solid particles are not present in the solution and in the deposited sample, which ensures a more homogeneous residue deposition of the sample on the reflector. Most of the digestion procedures are similar to those used in elemental analysis by other atomic spectrometric methods (mainly AAS- and ICP-based). The choice of the best decomposition procedure for a specific solid sample should be determined based on the matrix and analyte under study, and the result verified. For instance, biological samples are relatively easy to decompose *via* a mixture of nitric acid and hydrogen peroxide, whereas for some plant samples, the addition of hydrofluoric acid is mandatory to obtain a complete digestion due to their high silicon content (up to 10%).⁶⁵ The use of H_2SO_4 is not recommended due to the high temperature needed to dry the digested sample deposited on the reflector. Similarly, the use of HF should be avoided as it might damage the quartz reflector surface. To digest samples, open systems (working at atmospheric pressure) have been employed using conventional sources of heating such as sand baths, heating plates, *etc.* Microwave ovens are often used for sample digestion, as they can shorten the dissolution time and the related procedure may minimize analyte losses and cross-contamination. Block heaters for digestion of small amounts of vegetation samples (several mg) have also been successfully employed.⁶⁶

Another approach for analysing solid samples is to prepare suspensions mixing milligrams of powdered sample (10–50 mg) in an adequate disperser and depositing few μL (5–10 μL) of the resulting suspension on a reflective carrier. The amount of the analysis sample onto the carrier is highly dependent on the sample itself and its matrix, as well as of the spectrometer

$$\alpha_{C_i} = C_i \times \sqrt{\left(\frac{\alpha_{S_{\text{ICS}}}}{S_{\text{ICS}}}\right)^2 + \left(\frac{\alpha_{S_i}}{S_i}\right)^2 + \left(\frac{\alpha_{N_i}}{N_i}\right)^2 + \left(\frac{\alpha_{N_{\text{ICS}}}}{N_{\text{ICS}}}\right)^2 + \left(\frac{\alpha_{C_{\text{ICS}}}}{C_{\text{ICS}}}\right)^2}. \quad (2)$$

On a first approximation basis: $\alpha_{S_{ICS}}$ and α_{Si} are the uncertainties related to the determination of the slope value; $\alpha_{N_{IS}} = \sqrt{N_{ICS}}$ and $\alpha_{N_i} = \sqrt{N_i}$, are the uncertainties linked to the net intensity for the ICS and the analyte, respectively, and $\alpha_{C_{ICS}}$ is the uncertainty of the ICS concentration which can be roughly considered as a systematic uncertainty resulting from the ICS preparation procedure. The uncertainty α_{N_i} is a significant part of the total uncertainty (eqn (2)) for elements with a low content in the sample.

The most employed figures of merit in TXRF analysis are the Limit of Detection (LOD) and the Limit of Quantification (LOQ), which aim at describing the performance characteristics of the technique in a specific analytical process. Although their meaning is clear, a wide variety of definitions may be found across analytical techniques and application fields, and consequently various calculation procedures are applied to their determination. This lack of clarity has created and still creates confusion and lack of consistency for the scientific community, as well as for the regulators and industries. For these reasons, the LODs and LOQs values reported in the following sections cannot be compared easily, and the readers are encouraged to check in the cited literature or previous related works how the authors have considered and determined those parameters, as these procedures are unfortunately seldomly reported.

3.2 Sample preparation and handling procedures

TXRF analysis requires a small amount of sample (1–10 μL of liquids or 1–200 μg of solids) to be dropped onto a reflector and dried, while ensuring that the entire sample is contained within

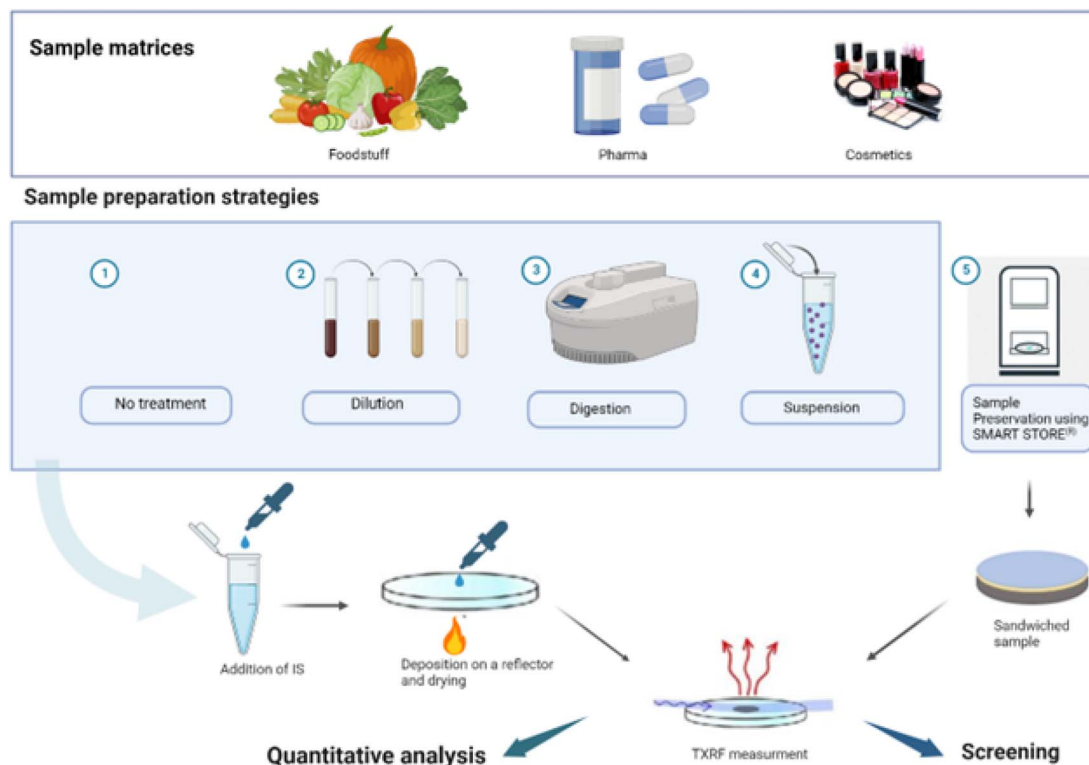


Fig. 1 Overview of sample preparation methods for TXRF analysis of foodstuff, pharmaceuticals, and cosmetics samples (created with <http://biorender.com/>).

employed (beam size, excitation energy, flux density, *etc.*), thus there is not one only volume or mass range which may be appropriate, but several of them, and as mentioned already depending on the sample type. Therefore, it is important to work in a volume/mass range adequate to obtain a thin layer on the reflector. If the deposited sample it is too thick, we will suffer from matrix effects, commonly present in EDXRF and WDXRF spectrometry.⁴

In the case of preparing the solid sample as a suspension, another interesting point is to ensure the homogeneity of the original sample and to select the appropriate amount and type of solvent and dispersing agents. The particle size of the powder material is another parameter relevant to achieve a homogeneous suspension, and reference materials usually present a particle size below 75 μm which is still too big for a TXRF analysis. Cryogenic grinders and ball mixer mills are used to obtain a good suspension homogeneity in the case of real food, pharmaceutical and cosmetic samples. While suspension offers the advantage of reduced cost and the elimination of hazardous chemicals, it may be challenging to optimise and to ensure sample homogeneity, and it may also limit the determination of analytes present in low concentration levels.

Despite digestion and suspension being the most used sample preparation strategies for TXRF analysis of solid samples, additional research is being carried out to develop faster, simpler, and more environmentally friendly sample treatment approaches, especially if elemental screening is the analytical goal. Among those, a novel device called Smart Store®

has been developed, which encloses the sample between two thin plastic films, then cuts out a 30 mm diameter disc to be stuck to a reflector and inserted into a TXRF system.⁶⁷ Promising results have been obtained using this approach for the rapid screening of heavy metals content in cosmetics⁶⁸ revealing a high potential of this device for fast and easy preparation of many types of samples to be analysed with a TXRF spectrometer.

4 Research applications of TXRF

4.1 Foodstuffs

The elemental concentration range and sensitivity of TXRF make it suitable for the analysis of food, where it can simultaneously determine trace levels of contaminants, macro, and micronutrients. Indeed, TXRF has been applied in the analysis of a wide range of liquids, like drinks and beverages (see Table 4), and solids like vegetables, fruits, dietary supplements, animal tissues and cereals (see Table 5).¹³

A direct analysis is frequently performed for the multi-element analysis of beers,⁶⁹ distilled alcoholic beverages,⁷⁰ food contact simulant,⁷¹ infusions,^{72–75} milk,⁷⁶ wine^{77–84} and whisky.⁸⁵ The recommended volume of liquids deposited on a sample carrier ranges from 5 μL to 20 μL , depending on the sample type, in order to avoid matrix effects.⁸⁶ In some cases, to reduce the mass deposition and overcome matrix effect phenomena, a minimal treatment as a sample dilution step is necessary. For example, direct quantitative analysis of honey is



Table 4 Selection of published articles based on the use of TXRF analysis of liquid foodstuff using (a) dilution and (b) digestion as sample treatments

(a) Dilution									
Sample type	Sample amount (g)	Solvent	Dilution factor (g : mL)	Elements	ICS	LOD (mg L ⁻¹)	Validation ^b	Ref.	
Honey	0.3–5	Ultrapure water	3 : 100–5:2	Multielemental: P, S, Cl, K, Ca, Cr, Ga, Mn, Fe, Ni, Cu, Zn, As, Pb, Br, Rb, Y		0.31–20.3	—	120–123	
Soy sauce	1	Ultrapure water	1 : 5	Multielemental: P, S, Cl, K, Ca, Ti, Zn, Cr, Mn, Fe, Co, Ni, Cu, As, Br, Pb, Ba, Rb, Sr		0.1–10	R, P	105	
Pineapple juice	n.a. ^a	Ultrapure water	1 : 4	Mn	Se	n.a. ^a	—	124	
Wine	1	Ultrapure water	1 : 1	Multielemental: K, Ca, Mn, Fe, Ni, Ga, Cu, Zn, Rb, Sr, Pb		0.01–0.2 (Mo-TXRF) 0.2–5 (W-TXRF)	L, P, R	79	
(b) Digestion									
Sample type	Sample amount	Digestion			Elements	ICS	LOD (µg L ⁻¹)	Validation ^b	Ref.
		Acid mixture	Energy source	Time					
Bovine milk	2 mL	HNO ₃	Muffle furnace	20 min	Multielemental: P, S, K, Ca, Ti, Cr, Mn, Fe, Ni, Cu, Zn, Se, Rb, Sr	Ga	0.08–34	A, P	98
Honey	1 g	HNO ₃ + H ₂ O ₂	MW furnace	15 min	Multielemental: Ti, Cr, Mn, Cu, Zn, As, Se, Br, Rb, Sr, Pb	Ga	n.a. ^a	—	99
	5 mL	Ashing + HCl	Muffle furnace	20 min	Multielemental: K, Ca, Ti, Cr, Mn, Fe, Ni, Cu, Zn, Se, Br, S	Ga	n.a. ^a	A	100
	6 mL	HCl	Muffle furnace	20 min	Multielemental: Ti, K, Ca, Fe, Mn, Zn, Cr, Ni, Cu, Se, Br, Sr	Ga	n.a. ^a	A	101
Soft drinks	1 mL	HNO ₃	MW furnace	15 min	Multielemental: P, S, K, Ca, Fe, Cu, Zn, As, Pb, Br, Sr	Ti	n.a. ^a	—	102
	0.8–1 mL	HNO ₃ + H ₂ O ₂	MW furnace	15 min	Multielemental: P, S, K, Ca, Fe, Cu, Zn, As, Pb, Br, Sr	Ti	2–350	—	103
Wine	10 mL	HNO ₃	Block heater	60 min	As, Pb, Br, Sr		8–1100	—	104
			MW furnace	50 min	Multielemental: Al, Cd, Mn, Cu, Fe, Cr, Ni, Pb, Zn	Mo	10	—	104
Wine	0.7 mL	HNO ₃ + H ₂ O ₂	Hot surface		Multielemental: P, S, K, Ca	Ga	60–780	P	77
Soy sauce	1 g	HNO ₃ + H ₂ O ₂	MW furnace	60 min	Multielemental: P, S, Cl, K, Ca, Ti, Cr, Mn, Fe, Co, Ni, Cu, As, Br, Pb, Ba, Rb, Sr	Ga	0.1–10	R, P	105
Sugar-cane spirits	10 mL	Ashing HNO ₃	MW furnace	n.a. ^a	Fe, Cu, Zn	Ga	8–10.9	—	106

^a n. a. – not available, MW: microwave. ^b Validation parameters: linearity (L), precision (P), accuracy (A), recovery (R).

quite problematic due to its very high concentration of sugars and high viscosity, making it also very difficult to dry and obtain a flat solid residue. In the case of wine analysis, it was demonstrated that the analytical quality of TXRF results can be improved if measurements are performed after sample dilution, especially for light Z elements like K and Ca.⁷⁹ In Table 4(a) summary of relevant publications about liquid foodstuff samples measured by TXRF after dilution is provided.

TXRF analysis of suspended solid residues (see Table 5(b)) offers a fast and straightforward way to perform screening and quantitative analysis for food safety, traceability, and quality control purposes. Non-ionic surfactants such as Triton X-100 (ref. 87 and 88) and ultrapure water^{66,89} are commonly used as dispersing agents. Polyvinyl alcohol can also be added to improve sample deposition on the reflective carrier.⁹⁰ Before deposition, the suspension mixture should be homogenized

using ultrasonic baths or vortex shakers.^{66,87,91,92} The use of polyvinyl alcohol was demonstrated to be effective in improving sample deposition for TXRF analysis of meat and cheese.⁹⁰ A novel method aiming to distinguish between mechanically separated meat (MSM) and fresh meat by monitoring K, Ca and Fe and using suspension for sample preparation and TXRF analysis is reported.^{93,94} The proposed procedure represents a powerful tool in food routine controls for consumer protection. Likewise, suspension proved to be a suitable analytical approach to check easily the presence of additives as polyphosphates (P), verify the content of salt (Cl) and detect some accidental metal contaminations (Cr, Co, Ni). In addition to the advantage of being a minimum sample treatment, suspension preparation can be of particular interest for the analysis of mass-limited samples (*e.g.*, soybean roots) since the amount of material needed to prepare the suspension (10–200 mg) is much

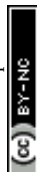


Table 5 Selection of published articles based on the use of TXRF analysis of solid foodstuff using (a) digestion and (b) suspension as sample treatments

(a) Digestion									
Sample type	Sample amount (g)	Digestion			Elements	Internal standard	LOD (mg kg ⁻¹)	Validation ^b	Ref.
		Acid mixture	Energy source	Time					
Eggs	0.5–3	HNO ₃ + H ₂ O ₂	TM heater	n.a. ^a	Multielemental: P, S, K, Ca, Cr, Cu, Fe, Ni, Mn, Pb, Se, Zn	Ga	n.a. ^a	—	108
	0.5–1.5	HNO ₃ + HClO ₄ + H ₂ O ₂	TM heater	90 min	Multielemental: Ba, Cu, Fe, Mn, Ni, Pb, Rb, Se, Sr, V, Zn	Ga	n.a. ^a	A, P	109
	1.46–1.61	HNO ₃ + H ₂ O ₂	TM heater	90 min	Multielemental: Ba, Cu, Fe, Mn, Ni, Pb, Rb, Se, Sr, V, Zn	Ga	n.a. ^a	A, P	109
Rice grain	0.5	HNO ₃	MW furnace	10 min	Se	Ga	0.05	R	125
	0.3	HNO ₃	MW furnace	15 min	Multielemental: K, Ca, Ti, Cr, Mn, Fe, Ni, Cu, Zn, Ba, Rb, Pb, Br, Sr, Ba	Ga	n.a. ^a	—	126
	1	HCl : HNO ₃ (1 : 3)	TM heater	2 h	Multielemental: As, Ca, K, Mn, P, Co Rb, S, Se, Zn		n.a. ^a	R	127
	0.5	HNO ₃ + H ₂ O ₂	MW furnace	15 min	Multielemental: K, Ca, Mn, Fe, Cu, Zn, Br, Rb, Sr	Ga	0.2–99	L, P, A	66
	0.02	HCl/HNO ₃ , (3 : 1)	TM heater	2 h					
Beans	0.15	HNO ₃ + H ₂ O ₂	MW furnace	25 min	Multielemental: K, Ca, Mn, Fe, Ni, Cu, Zn, Rb, Sr, Pb	Ga	n.a. ^a	A, P	87
Onions	0.2	HNO ₃ + H ₂ O ₂	Hot plate	n.a. ^a	As	Co	n.a. ^a	A, P	128
Porridge flour	1	HNO ₃	TM heater	n.a. ^a	Multielemental: Mn, Fe, Cu, Zn	Ga	n.a. ^a	A, P	129
Maize flour	0.35	HNO ₃	TM heater	n.a. ^a	Multielemental: Mn, Fe, Cu, Zn	Ga	n.a. ^a	—	130
Candies	0.5	HNO ₃ + HCl + H ₂ O ₂	MW furnace	n.a. ^a	Multielemental: Ca, Ti, Cr, Mn, Fe, Cu, Zn, Br, Rb, Sr, Pb	Ga	0.02–0.1	A, P	107
Chocolate	0.4	HNO ₃ + H ₂ O ₂	MW furnace	25 min	Multielemental: P, S, K, Ca, Ti, Cr, Mn, Fe, Ni, Cu, Zn, Br, Rb, Sr, Ba	Ga	n.a. ^a	—	112
Pork meat	0.3	HNO ₃	MW furnace	n.a. ^a	Multielemental: P, S, Cl, K, Ca, Ti, Mn, Fe, Ni, Cu, Zn, Se, Br, Rb, Sr, Ba, Pb	Ga	0.01–1	A, P	131
Edible Clams	0.5	HNO ₃ + H ₂ O ₂	MW furnace	15 min	Multielemental: Ti, Mn, Cu, Zn, As, Br, Sr	Y	n.a. ^a	A	92
(b) Suspension									
Sample type	Sample amount (mg)	Disperser agent	Disperser (mL)	Elements	Internal standard	LOD (mg kg ⁻¹)	Validation ^b	Ref.	
Beans root	10	Triton X 100 (1% wt)	0.99	Multielemental: K, Ca, Mn, Fe, Ni, Cu, Zn, Rb, Sr, Pb	Ga	n.a. ^a	A, P	87	
Microgreens	100	Triton X-100	5	Multielemental: P, S, K, Ca, Fe, Zn, Mn, Cu, Ni	Ga	n.a. ^a	A	88	
				Multielemental: P, S, K, Ca, Cl, Mn, Fe, Ni, Cu, Zn, Br, Rb, Sr		0.2–30	A, P, LOQ	91	
Vegetables	20	Ultrapure water	1	Multielemental: K, Ca, Mn, Fe, Cu, Zn, Rb, Sr	Ga	2–100	L, A(R), P	66,89	
Edible Clams	100	Ultrapure water	1	Multielemental: Ti, Mn, Cu, Zn, As, Br, Sr	Y	n.a. ^a	A	92	
Wheat flour	50	Triton X-100 (1% wt)	2.5	Multielemental: P, S, Cl, K, Ca, Mn, Fe, Ni, Cu, Zn, Se, Br, Rb, Sr	Y	0.13–28	A(R)	132	
			1	Multielemental: Cu, Fe, Mn, Zn	Ga	0.27–0.44	A(R)	133	
Meat	100	Triton X-100, (0.1% wt) + PVA	5	Multielemental: K, Ca, Fe, Cu, Zn	Ga	0.08–1.67	—	93	
	200		5	Multielemental: P, Ca, Fe	Ga	n.a. ^a	—	90	



Table 5 (Contd.)

(b) Suspension								
Sample type	Sample amount (mg)	Disperser agent	Disperser (mL)	Elements	Internal standard	LOD (mg kg ⁻¹)	Validation ^b	Ref.
Cheese	200	Ultrapure water + Triton X-100 (0.1% wt) + PVA (0.2%)	5	Multielemental: P, S, Cl, K, Ca, Fe, Zn	Ga	n.a. ^a	—	90
		Ultrapure water + Triton X-100 (0.1% wt) + PVA (0.2%)						
Raw meat, salami, sausages, fish, mussels	50	Triton X-100 (0.5% wt)	4	Multielemental: P, S, Cl, K, Ca, Cr, Mn, Fe, Ni, Cu, Zn, Br, Rb, Sr	Ga, V	n.a. ^a	A(R)	94

^a n.a. – not available, PVA: polyvinyl alcohol, MW: microwave, TM: thermo block. ^b Validation parameters: linearity (L), precision (P), accuracy (A), recovery (R), limit of quantification (LOQ).

lower than the one required for digestion.⁹⁵ Despite the number of benefits that there is in preparing suspension for the TXRF analysis of solid materials, only 15% of the sample treatment methods use it⁶⁴ due to calibration issues, weighing errors, and sample/suspension inhomogeneity that need to be properly addressed.^{96,97} In addition, the results for lighter elements, mainly K and Ca, are not as good as for mid-high Z elements in some food matrices because of their complicated matrix and of their effects, among other limiting parameters. In these situations, external calibration is performed to ameliorate the results utilizing a set of reference materials with a matrix resembling that of the actual samples.⁶⁶

Decomposition of the liquid to be deposited onto the reflector is also a common sample preparation method to improve the limits of detection. This pre-treatment is reported for the TXRF analysis of milk,⁹⁸ honey,^{99–101} soft drinks,^{102,103} wine,¹⁰⁴ soy sauce¹⁰⁵ and sugar-cane spirits.¹⁰⁶ As shown in Table 4(b), sample decomposition is usually performed by wet ashing procedures involving the use of concentrated acids (HNO₃, HCl) or some mixtures of acids in combination with oxidizing agents (HNO₃ + H₂O₂). To overcome problems of contamination or volatilization during the digestion process, microwave ovens are preferred as heating sources. Dry ashing by incineration of the liquid samples at 500 °C and dissolution of the resulting ashes in HNO₃ or HCl has been used as a digestion procedure for the determination of Fe, Cu and Zn in sugar cane spirits¹⁰⁶ or multi-elemental analysis of honey.¹⁰⁰ For the complete dissolution of food solid samples, a mass of about 0.1 g to 1 g is mixed with acids and other oxidating compounds. Often, concentrated HNO₃ or a mixture of HNO₃ + H₂O₂ are employed but in some cases the additional use of HCl or HClO₄ has been reported to get the total solubilisation of vegetables,⁶⁶ candies¹⁰⁷ or eggs.^{108,109} Microwave ovens and thermo block heaters are two types of heating sources employed during digestion. The main advantage of those systems is the possibility to digest a low quantity of sample, therefore using reduced acid volumes, as it was done for a very small amount of vegetables (20 mg) and only 1 mL of aqua regia (HCl:HNO₃, 3:1).⁶⁶ Compared to liquid

samples, digestion of solid foodstuffs requires a longer time due to the complexity of sample matrices.

Microextraction and selective adsorption procedures can also be considered to enhance TXRF microanalytical capability. For instance in the case of Se and Hg determination of low level (0.6 g L⁻¹) in seafood samples¹¹⁰ and in the determination of Sb in milk after nanoparticles immobilization reflectors,¹¹¹ respectively.

Promising results have been obtained using the Smart Store® system for the rapid screening of chocolate samples. In this case, chocolate was first dissolved in 3 mL of Ga solution in bain-marie conditions. Then, the total dissolved sample was placed on a polymeric foil and dried at room temperature. The plasticized sample was put onto a quartz sample carrier with a diameter of 30 mm to be measured with the TXRF spectrometer.¹¹² Though underestimation of the low Z elements occurred, the procedure demonstrated to be a fast and easy approach for screening.

4.2 Pharmaceuticals, drugs, and supplements

At present, there is a lack of general procedures in the pharmacopoeias for a reliable application of TXRF method in pharmaceutical analysis. Over the past few years, many studies have investigated the potential of TXRF for determining elemental impurities in pharmaceutical products and dietary supplements (DS). Table 6 summarizes the research published on this topic between 2000–2020, in which TXRF has been applied to analyse various dosage forms of pharmaceutical products, including APIs, drug substances, and metal-containing drugs. However, only finished dosage forms of DS have been considered. Solid samples can be dissolved by organic solvent-based diluents, such as DMF, DMSO, and methanol^{33,113–115} or suspended using an aqueous non-ionic surfactant solution.¹¹⁶ More intricate sample treatments, such as microwave digestion, have also been used, primarily for the analysis of more complex matrices.³⁸ One critical factor for determining elemental impurities in pharmaceutical samples



Table 6 Selection of publications about analytical methods published in the literature in the last twenty years regarding the determination of elements in: (A) pharmaceutical products and (B) dietary supplements by TXRF^a

(A) Pharmaceutical products								
Samples	Analytes	Sample preparation details	Internal standard	TXRF measurement	LOQ ($\mu\text{g g}^{-1}$)	Purpose of the method	Validation ^b	Ref.
API (solid)	K, Ca, Fe, Ni, Sr, Pd	10 mg dissolution (THF, DMSO), 10–20 μL deposition	Ga	Mo-TXRF (1000 s) W-TXRF (2000s)	10–20	Quantitative	S, L, P, A, D	49
API (solid)	PGMs metals (Rh, Pd, Ir, Pt)	150–500 mg dissolution (water, methanol), 10 μL deposition 500 mg MW digestion (HNO_3 + HCl), 10 μL deposition	Y	W-TXRF (1000 s)	1.9–4.1	Screening	S, P, A	113
Metal containing drugs ^c	Au, Pt	Dissolution (DMF + water + PVA), 4–10 μL deposition	Mn	W-TXRF (1000 s)	1.7–8.7	Quantitative	P, A	114
Drug Substances ^e	Ca, Fe, Zn, Sr	Dissolution (0.2–0.3 M HNO_3) 1% sample solution, 10 μL deposition	Se	Mo-TXRF (250 s)	0.2 ^d	Quantitative	P, A	114
Drug Substances ^e	Ca, Fe, Zn, Sr	Dissolution (0.2–0.3 M HNO_3) 1% sample solution, 10 μL deposition	Se	Mo-TXRF (600 s)	—	Screening	—	115
Drug Substances ^f	Pd, Rh, Pt, Cr, Fe, Cu	Dissolution (50% methanol + 2% HNO_3 + 0.5% HCL) 5 mg/1 mL, 10 μL deposition	Sr	Mo-TXRF (1000 s) W-TXRF (2000s)	<2–10	Screening platform	L, P, A, D	134
	Pd, Rh, Pt, Cr, Fe, Cu	Dissolution (50% methanol + 2% HNO_3 + 0.5% HCL) 10 mg mL^{-1} , 5 μL deposition	V	WL-TXRF (500 s) AgK-TXRF (500 s)	2–15			
(B) Dietary supplements, DS (finished dosage forms)								
Samples	Analytes	Sample preparation details	Internal standard	TXRF measurement	LOQ ($\mu\text{g g}^{-1}$)	Purpose of the method	Validation ^b	Ref.
Commercial DS (tablets) ^g	P, Ca, Mn, Fe, Cu, Zn, K, Cr, Se, I	DS without capsule: dissolution (water), 2 μL deposition DS with capsule: dissolution (HNO_3), 2 μL deposition	Ni, Ga, Y	Mo-TXRF (250 s)	—	Screening	A	33
Commercial DS (different dosage forms) ^h	Pb, As, Cr, Hg	20–50 mg digestion (HNO_3 +HCl + H_2O_2) filtration	Ga ⁱ	Mo-TXRF	0.2–3.1	Quantitative	L, P, A, D	38
Commercial DS and seaweed products	I	500 mg digestion (NH_3), 25 μL deposition	Cd, Ag, Y	W-TXRF (1000 s)	—	Screening	None	135
Commercial DS (tablets)	Se	50 mg powder suspension (2.5 mL 1% Triton X100) 10 μL deposition	Ga	Mo-TXRF (1000 s)	0.3–0.6	Quantitative	P, A	136

^a API: active pharmaceutical ingredient, MW: microwave, THF: tetrahydrofuran, DMSO: dimethylsulfoxide, DMF: dimethylformamide, PVA: polyvinyl alcohol. ^b Validation parameters: specificity (S), linearity (L), precision (P), accuracy (A), limit of detection (D). ^c Cisplatin, auranofin.

^d Liquid form (0.2 ppm). ^e Ca-folate, Levodopa, Enalapril maleate. ^f Triprolidine HCl, diphenhydramine HCl, Chlorpheniramine maleate, pseudoephedrine HCl, ephedrine sulfate, scopolamine HBr. ^g Komplivit (DS with capsule), Multifort (DS without capsule). ^h Fat burner, laxative, tejocote root, omega oil. ⁱ In the case of Hg (Ga solution contain EDTA to avoid Hg losses during drying).

with a commercial TXRF spectrometer is the instrumental setup and, in particular, the X-ray anode source determining the excitation energy. This aspect is particularly important for platinum group metals (PGMs), which are commonly used in medicinal chemistry. The determination of Pd and other PGMs has been the focus of many studies. Better results were obtained using X-ray sources equipped with W and Ag anodes that allow elemental determination through the Pd–K emission lines, free

of interferences,¹¹⁴ while with Mo X-ray sources spectral interferences overlapping with Pd–L emission lines occur.¹¹³ However, Mo X-ray sources provides better sensitivity for mid-Z elements (*i.e.*, Fe, Ni, Zn, Cr, Cu, *etc.*).³³ Therefore, a trade-off must be made between PGMs analysis *versus* sensitivity for most other elements of interest, and a thorough analysis may require the use of TXRF systems equipped with both Mo and W X-ray tubes. The quantification limits reported for TXRF



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The wide acceptance of TXRF by the analytical community is thus hindered by the lack of proper standard operating procedures and fit-for-purpose reference materials, which are both needed to claim the validity of results at a certain level of confidence. Establishing the uncertainty budget is also a priority to ensure the physical and chemical traceability of results and reliability in quantification. Such awareness has initiated already pre-normative research activities tackling the above-mentioned issues.¹¹⁹ Those ventures are essential to establish all the physical and chemical interdependencies linked to the use of TXRF, and prior to developing any normative documentation (*e.g.*, standard) under the ISO/TC201/SC10 framework. Only then, TXRF may be proposed as a standardized chemical method and labelled as the technique of reference, similar to AAS or ICPs techniques, for the elemental analysis of consumer products such as food, pharmaceutical, and cosmetic samples.

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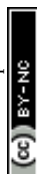
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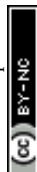
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