Development of an Enriched Polyphenol (Natural Antioxidant) Extract from Orange Juice (*Citrus sinensis*) by Adsorption on Macroporous Resins

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Orange (*Citrus sinensis*) juice contains a high amount of antioxidant compounds, such as polyphenols and vitamins. The aim of this work was to develop an adsorption procedure for the quantitative recovery of polyphenols from fresh orange juice. Different macroporous resins have been selected to evaluate their affinity for phenolic compound in order to purify the antioxidant compounds from the orange juice. The main compounds of orange juice were firstly characterized using an UPLC-UV-HRMS to define the metabolite profile, and subsequently three different types of adsorbent (XAD-2, XAD-4, and XAD-16N) were tested to concentrate these bioactive compounds. The time of contact was selected based on kinetic studies, and subsequently the adsorption and elution conditions were optimized in order to maximize the recovery of phenolic compounds to obtain an extract rich of bioactive compounds. Lastly, antioxidant capacity of the orange juice extract of selected macroporous resin, obtained under optimized conditions, was determined by in vitro antioxidant assays.

1. Introduction

In the last few decades, the interest for new bioactive compounds in antioxidant compounds has increased because of their capability to fight several chronic and acute diseases, including inflammatory cardiovascular and neurological diseases as well as cancers [1].

Among these new compounds, polyphenol is considered one of the most promising antioxidants of natural origin that can be used for the formulation of new drugs and nutraceuticals and as ingredient for functional foods. Antioxidants are compounds that, present at low concentration, can significantly reduce or prevent the oxidation of other molecules.

The antioxidant compounds act through several mechanisms, but the most accepted one is the scavenging of free radicals contributing to reduce the oxidative stress [2].

Antioxidants are commonly used to protect oxidizable species, such as cosmetics [3, 4], pharmaceuticals [5], and foods [6], from the deleterious effects of oxidation and are also employed as dietary supplements to neutralize the adverse effects of the oxidative stress. Moreover, they are also employed in the food industry to reduce rancidity, to protect and stabilize colours and aroma and to increase the shelf life of products [7]. Some synthetic additives with antioxidant activity, such as butylated hydroxytoluene (BHT) or butylated hydroxyanisole (BHA), are potentially toxic for humans [8]. For this reason, there has been a growing interest to replace synthetic antioxidants with natural ones in order to increase the shelf-life and improve the nutraceutical value of food products [7–9]. There is a broad range of natural antioxidant compounds, but, among them, polyphenol doubtlessly represents the most abundant and
widespread class in nature. Polyphenols are secondary plant metabolites occurring in vegetables, fruits, beverages, and other related food products with well-documented antioxidant, anticarcinogenic, antimutagenic, anallergic, and antiaging activities [10]; for these properties, particular attention is currently placed on their extraction from inexpensive agricultural food products. In this context, fruits and vegetables of the Mediterranean area can be considered an important source of polyphenol [11].

Although orange juice has lower antioxidant properties than the fruit juice of the berries (chokeberry, elderberry, and blueberry) [12, 13], orange juice can be considered a good source of polyphenol intake in the Mediterranean diet [14, 15]. Citrus genus and its varieties are few of the most cultivated crops all over the world; in particular, thanks to its geographical location and climate, Italy is one of the major citrus producing nations in Europe after Spain [16]. Numerous population studies show that regular consumption of fruit and vegetables rich in antioxidants may prevent cardiovascular diseases. The nutritional benefits of citrus consumption on human health are well demonstrated [17–22], and over the past decades, a large number of studies have been carried out with the aim of identifying the bioactive components present in different parts of citrus fruits [23, 24] in an attempt to gain a deeper understanding of the correlation between diet, health benefits, and reduced risk of diseases [17, 25]. In this context, several studies have associated the consumption of foods rich in phenolic compounds with lower risks of different types of cancer [26] and a decrease in the incidence of many diseases, as cardiovascular diseases (CVDs) [27]. Furthermore, these studies have shown that foods rich in phenolic compounds have an antioxidant, antiedematogenic, anti-inflammatory, and antiaging activity [18, 28] and may also contribute to an improvement of the blood lipid profile (e.g., triglyceride and total cholesterol reduction and increased HDL), a decrease of blood pressure and blood glucose or to the improvement of the endothelial function [29–33]. Citrus fruits are one of the richest dietary sources of flavonoids [34]. In recent years, the biological activity of citrus polyphenols and their role in the prevention and treatment of various human chronic and degenerative diseases have been extensively reported [35–37]. Given the possible increase of the commercial value of citrus juice as a natural source of antioxidant compounds, the aim of this work was to optimize and develop a method to obtain highly enriched antioxidant polyphenols and to eliminate other interfering compounds which do not possess antioxidant properties, such as proteins, sugars, fiber, or metals. In order to eliminate these interfering compounds and selectively recover the polyphenols, an easy, cheap, and green adsorption procedure has been developed. Experimental conditions affecting adsorption and elution of polyphenols from macroporous resin have been carefully studied and optimized.

2. Materials and Methods

2.1. Sample and Resin Extraction. Before being pressed, orange fruits were washed, peeled, and manually squeezed to obtain the juice. Pulps and solid material were removed by 5 minutes of centrifugation at 13000 rpm (SL 16 centrifuge, Thermo Fisher Scientific, Milan, Italy). Amberlite macroporous resins XAD-2 XAD-4 and XAD-16N, used as adsorbent material, were obtained from sigma (Sigma-Aldrich, Germany), and information on the physical and chemical characteristics of each resin was reported in Table 1.

Before performing the extraction procedure, the macroporous resins were rinsed thoroughly with distilled water in order to remove salts and impurities, followed by a drying at 70°C for 24 h, and finally immersed in ethanol for 12 h. Ethanol was replaced by water before starting the adsorption procedure. Kinetic study was performed by keeping in contact 500 mg of each resin with 10 mL of orange juice in a plastic tube of 50 mL, under planetary agitation at 300 rpm during the extraction time (30–240 min) at constant temperature (25°C). After the extraction, the resins were packed into empty SPE cartridge (6 mL), rinsed with deionized water, and end eluted with different ratios (10:0, 3:7, 7:3, 0:10 v/v) of EtOH:H2O mixture, in order to select the best elution solvent. The final extract, at concentration of 5 mg mL-1, was injected and analysed by UHPLC-UV (280 nm).

2.2. Chemicals and Standards. MS-grade solvents used for UHPLC analysis, acetonitrile (MeCN), water (H2O), and formic acid (HCOOH) were provided by Romil (Cambridge, UK); analytical-grade solvent methanol (MeOH) and ethanol (EtOH) were supplied by Sigma-Aldrich (Milan, Italy). Water was purified by using a Milli-Q system (Millipore, Bedford, USA). Gallic acid (GA), 6-hydroxy-2,5,7,8-tetramethylchromane-2-carboxylic acid (Trolox), 2,2-diphenyl-1-picrylhydrazyl radical (DPPH), and Folin & Ciocalteu’s phenol reagent were purchased by Sigma-Aldrich (Milan, Italy).

2.3. UHPLC-UV-ESI-HRMS Conditions. Separation of phenolic compounds from citrus extract was performed on a Platин Blue system (Knauer GmbH, Berlin, Germany) consisting of a vacuum degasser, an autosampler, two UHPLC pumps, a column oven compartment, and a diode-array detector (DAD). The extract was chromatographed on a Kinetex C18 (100 × 2.1 mm I.D., 2.6 μm) column from Phenomenex (Torrance, CA, USA). Water (A) and acetonitrile (B), both acidified with 0.1% HCOOH, were used as mobile phases at flow rate of 0.4 mL min-1. The gradient was programmed as follows: 0-1 min 2% B; 0-1 min 2% B; 1–3 min 2–5% B; 3–8 min 5–10% B; 8–10 min 10–15% B held for 3 min; 13–17 min 15–30% B; 17–19 min 30–50% B; 19–21 min 50–85% B; 21–23 min 85–98% B held for 4 min. Before the injection, the initial conditions were held for 5 min as the re-equilibration step. The injection volume was 10 μL, and the column was maintained at 30°C. The UHPLC system was coupled to a DAD and a LTQ Orbitrap XL (Thermo Scientific, San Jose, CA) equipped with an electrospray ionisation source (ESI). The UV signal was acquired at three different wavelengths: 250, 285, and 365 nm, whereas mass spectrometry acquisitions were carried out with the same condition previously reported [16]. Briefly,
parameters for HRMS and MS$^+$ analysis were set using negative and positive ion modes with spectra acquired over a mass range from 100–800 m/z at a resolution of 60000. The optimized value of ESI source were as follows: source voltage at 25°C, the absorbance was read at 765 nm using a Multiskan Go spectrophotometer (Thermo Fischer Scientific). The DPPH assay expressed as EC$_{50}$ was employed using a Multiskan Go spectrophotometer (Thermo Fischer Scientific). Trolox was used as reference standard, and the results were expressed as TEAC values (mmol of Trolox/g extract). These values were obtained from five different concentrations of each extract tested (between 1 and 4 mg/mL) in the assay, giving a linear response between 20% and 80% blank absorbance. All analyses were done in triplicate.

2.7. Statistical Analysis. The statistical analysis was estimated using Statgraphic Centurion XVI Version 16.1 (Rockville, USA). Data were reported as mean (SD for triplicate determinations and analysis of variance (ANOVA) using Tukey’s test were conducted to identify differences among means). Statistical significance was declared at $p < 0.05$.

3. Results and Discussion

3.1. UHPLC-UV/HRMS Analysis. The chemical identification of polyphenol compounds in orange juice was made following the same procedure of the previously published paper [16]. Table 2 shows the list of 14 compounds tentatively identified by UHPLC-UV/MS/MS experiments. In order to obtain complementary information useful to identify the main compounds in orange juice “Citrus sinensis” extract, the UHPLC-HRMS analysis was performed in both positive and negative ionisation modes. The compounds detected were tentatively characterized by means of MS data, together with the interpretation of MS/MS spectra, in comparison with those reported in the literature and with standard spectra, when available. In the identification process, the following database was used: ChemSpider (https://www.chemspider.com) and Scifinder (https://scifinder.cas.org). Figure 1 shows the representative UV (285 nm) chromatogram of orange juice extracts obtained after the elution of XAD-16 under optimal conditions.

3.2. Selection of Macroporous Resin. After the characterization of orange juice, in order to select the best sorbent material to use, three different macroporous resins were tested to evaluate the adsorption of polyphenol compounds from orange juice. In order to evaluate the best resin and
contact time able to remove analytes from orange juice, a kinetic study was performed. For this purpose, to obtain an adsorption curve of the target compounds, for each sorbent material at regular time interval, an aliquot was withdrawn and analysed by UHPLC-UV along a time of 240 min ($n \geq 3$). Pulp and solid material was removed from the juice by centrifugation. Subsequently, 10 mL of supernatant was taken by a calibrated glass pipette, placed in a 50 mL conical-bottomed test tube containing 500 mg resins, and stirred by an orbital stirrer at a speed of 200 rpm$^{-1}$. Every 30 minutes, 100 μL of sample was withdrawn using a micropipette and analysed by UHPLC in order to evaluate the adsorption kinetics of each analyte. Results of XAD-2 showed a very poor affinity (data not shown) of the analytes for the resin over the studied time (240 min). Based on these results, XAD-2 was not used for future experiments. Figure 2 shows the adsorption percentage (reduction) of each compound as function of the contact time (minutes) (0–240 minutes) for each resin (XAD-4 and XAD-16N) at a temperature of 25°C. The graphs related to XAD-4 (Figure 2(a)) and XAD-16N (Figure 2(b)) highlight a good affinity towards all compounds but with different absorption kinetics. Analysing accurately the kinetics related to XAD-16N, it is possible to notice a faster adsorption kinetics compared with those obtained using the XAD-4. When XAD-16N was used, an adsorption between 80–100% was obtained in 2 hours for all the monitored compounds. The most lipophilic analytes follow a much more rapid adsorption kinetics compared with the analytes with medium and high polarity. This behaviour was predictable, considering the hydrophobic character of styrene-divinylbenzene resin. Similar results were obtained with XAD-4 after a contact time of approximately 4 hours. Based on these results, XAD-16N and 120 minutes of contact time were selected as optimal conditions and seated for further experiments.

### Table 2: UHPLC-HRMS/MS data of compounds detected in orange juice.

| N. | m/z     | (+) HRMS | m/z     | (-) HRMS | Molecular formula | Compound                  |
|----|---------|----------|---------|----------|-------------------|---------------------------|
| 1  | 357.1165| –4.8     | 355.1041| 4.0      | C_{16}H_{20}O_{9}  | Ferulic acid glucoside    |
| 2  | 595.1627| –4.2     | 593.1515| 2.1      | C_{37}H_{80}O_{15} | Apigenin 6,8-di-C-glucoside|
| 3  | 625.1741| –3.2     | 623.1624| 3.4      | C_{28}H_{32}O_{16} | Diosmetin 6,8-di-C-glucoside|
| 4  | 673.2434| –3.9     | 649.2514| 3.1      | C_{32}H_{42}O_{14} | Limonin-glucoside         |
| 5  | 581.1838| –4.1     | 579.1722| 2.3      | C_{27}H_{32}O_{14} | Narirutin                  |
| 6  | 435.1267| –4.8     | 433.114 | 4.5      | C_{21}H_{32}O_{10} | Naringenin-glucoside       |
| 7  | 611.1941| –4.2     | 609.1831| 3.7      | C_{28}H_{34}O_{15} | Hesperidin                 |
| 8  | 465.1374| –4.7     | 463.1247| 2.0      | C_{22}H_{34}O_{11} | Hesperetin glucoside        |
| 9  | 717.2701| –3.2     | 693.2767| 2.4      | C_{34}H_{46}O_{15} | Nomilin glucoside          |
| 10 | 735.2807| –3.3     | 711.2881| 4.0      | C_{34}H_{48}O_{16} | Nomilinic acid glucoside    |
| 11 | 471.1995| –3.3     | 515.1924| 3.5      | C_{27}H_{30}O_{10} | Limonin*                   |
| 12 | 373.1269| –4.1     | 358.1033| 3.1      | C_{20}H_{20}O_{7} | Sinensetin                 |
| 13 | 403.1372| –4.7     | 388.1135| 3.1      | C_{21}H_{22}O_{7} | Nobleitin                  |
| 14 | 373.1266| –4.0     | 358.1034| 3.5      | C_{20}H_{20}O_{7} | Tangeretin                 |

*Compared with reference standards. a$m/z$ values corresponding to [M+Na]+. b$m/z$ values corresponding to [M-H+HCOOH]−.
Figure 1: HPLC-UV profiles (285 nm) of orange juice extract (XAD-16N).

Figure 2: Continued.
3.3. Selection of Elution Solvent. After the selection of the macroporous resin able to quantitatively adsorb the phenolic compounds in the orange juice, the next step was the selection of an elution solvent to desorb the analytes from resin. In this step, the efficiency of different ratios (10:0, 3:7, 7:3, and 0:10 v/v) of EtOH:H₂O mixture was evaluated. After the adsorption time, the resin XAD-16N was filtered and packed into 6mL empty cartridges and subsequently rinsed with 5mL of deionized water to remove sugars and nonadsorbed compounds. After washing the resin with 10mL of pure water, 10mL of ethanol and different ratios (7:3, and 3:7 v/v) of EtOH:H₂O mixture were passed through the cartridge. After 30 minutes, the elution solvent passed through the cartridge in order to elute and collect the analytes into a test tube. The results show a similar trend when pure water and ethanol were used as elution solvent, producing low recovery for all target analytes (Figure 3). As shown in Figure 3, the best extraction efficiency was obtained using ethanol 70%, while the extraction efficiency of target compounds decreased drastically when all the other combinations of elution solvent were used. On the basis of these results, EtOH:H₂O 7:3 v/v mixture was selected as optimal elution solvent and was used for further experiments. Another important parameter that was studied and optimized was the contact time between the selected elution solvents and the XAD-16N. As shown in Figure 4, by increasing the contact time between elution solvent and resin before performing the elution, the extraction yield increases up to 60 min and then remains almost constant for the rest of the experiment time (120 min); therefore, 60 min was selected as best contact time.

3.4. Antioxidant Capacity of Orange Juice Extract. Nowadays, the biological effects of polyphenol on human health are widely demonstrated by several studies, showing that citrus extract has a good antioxidant activity [1]. Therefore, citrus juice extract can be used as a potential source of bioactive compounds for the development of natural ingredients to be used in the nutraceutical industry or as foods additives. Before the evaluation of the antioxidant capacity of juice extracts, the total phenol content was evaluated by the Folin–Ciocalteu method [38]. The results of the Folin–Ciocalteu assay showed a good content of phenolic compounds in the XAD-16 extract (356.5 ± 4mg GAE/g of extract). The presence of a high content of phenolic compounds provided by the Folin–Ciocalteu assay suggested the possibility of a high antioxidant capacity. Whereby, in order to evaluate the antioxidant capacity of the XAD-16N extract, in vitro antioxidant activity assays DPPH and ABTS were performed. Unlike most of the other antioxidant assays used to evaluate the AOC, it is based on a single electron and/or hydrogen atom transfer reaction (SET/HAT) and the DPPH and ABTS assays are based on SET/HAT simultaneously [43], providing a more representative evaluation of antioxidant properties. In order to avoid the underestimation of the results, before the estimation of AOC, the optimization of reaction time and concentration range for both standard (Trolox) and extract were carried out. The AOC provided by the DPPH assay, expressed as EC₅₀ (9.97 ± 0.5 μg·mL⁻¹), whereas the TEAC values provided by the ABTS assay were 3.88 mmol of Trolox/g extract. Good polyphenol content shows a significant radical scavenger activity to the extract, as demonstrated by the results of DPPH and ABTS assays. This result indicates that the orange juice extract can be used as source
Figure 3: Normalized extraction efficiency of target analytes using different elution solvents. Values are means of three replicates ± relative standard deviation (RDS%).

Figure 4: Recovery percentage of target analytes using different contact times of elution solvents and macroporous resin. Values are means of three replicates ± relative standard deviation (RDS%).
of food ingredient or as natural ingredient in the nutraceutical industry.

4. Conclusions
The adsorption of phenolic compounds from orange juice on macroporous resin was studied in order to develop an easy and favourable process to obtain an extract with antioxidant properties. The main parameters affecting the extraction efficiency and elution process were optimized. The results show that, in order to obtain an excellent extraction efficiency for analytes, the adsorption needs a long contact time, with the most effective resin requiring a time of two hours to reach the equilibrium. The resin that showed the best result regarding the adsorption of phenolic compounds was XAD-16N. A mixture of EtOH: H₂O (7:3 v/v) was selected as the best elution solvent used in order to obtain a quantitative recovery of target analytes. Subsequently, the total polyphenol content and the antioxidant activity of extract were evaluated by using in vitro antioxidant assays. The results demonstrate that the orange juice extract represents a rich source of antioxidant phenolic compounds that could be employed as additives to prevent the oxidation of foodstuffs but also as functional ingredients in the nutraceutical industry.

Data Availability
The data used to support the findings of this study are available from the corresponding author upon request.

Conflicts of Interest
The authors declare no conflicts of interest regarding the publication of this study.

Authors’ Contributions
Luca Campone performed some experiments, analysed and interpreted the results, and drafted the manuscript. Rita Celano performed the HPLC-HRMS experiments and contributed to the statistical analysis. Serena Rizzo performed the antioxidant experiments and contributed to the statistical analysis. AnnaLisa Piccinelli supported the interpretation of results. Luca Rastrelli supported the interpretation of results and revised the manuscript. MariaTeresa Russo designed the overall experimental scheme and revised the manuscript. All authors approved the final version of the manuscript.

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