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⁸⁷Sr/⁸⁶Sr isotopes in grapes of different cultivars: a geochemical tool for geographic traceability of agriculture products

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15 Highlights

- 16 ⁸⁷Sr/⁸⁶Sr in fresh grapes from different genotype vines (cultivar)
- 17 ⁸⁷Sr/⁸⁶Sr differences between white and red grapes and among harvest years
- 18 ⁸⁷Sr/⁸⁶Sr in bioavailable fraction of soils, soils, and rocks beneath vineyards of production
- 19 Geologic heritage in food stuff as a fingerprint for geographic traceability

20 Abstract

⁸⁷Sr/⁸⁶Sr was determined on fresh red and white grapes, soils and rocks from three selected vineyards to verify the isotopic relationships between the fruit of the vine and geologic substrata of vineyards. ⁸⁷Sr/⁸⁶Sr were determined on sampled grapes of four different harvest years and different grape varieties, on bioavailable fraction of soils, on whole soils, and on bedrocks from the geo-pedological substratum of the vineyards. The vineyards chosen for the experimental works belong to an organic farming winery and thus cultivation procedures were strictly controlled.

Grapes were sampled during the harvests of four different but consecutive years with ⁸⁷Sr/⁸⁶Sr that does not change reflecting the values of the soil bioavailable fraction. No variations among grapes from different vine cultivars were observed. A strict isotope relationship with soil bio-

30 available fraction was observed. These findings demonstrate the reliability of ⁸⁷Sr/⁸⁶Sr, even at a

31 very small scale, for food products geographic origin assessment.

Key words: ⁸⁷Sr/⁸⁶Sr of fresh grapes, white and red grapes, geographic traceability, geologic and
 pedologic fingerprints, Pitigliano area, Vulsini Mountains, Central Italy.

34 1. Introduction

The increasing demand of high quality food products promoted the development of rigid regulations for certification of authenticity and protection from frauds (e.g., http://eurlex.europa.eu/legal-content/EN/ALL/?uri=CELEX:32002R0178). This is leading to an increasing request of suitable scientific protocols able to confirm the authenticity of food products by tracking their geographical origin. These studies aim to find scientific parameters inherited by the area of production of the agricultural products, which may be considered a fingerprint for geographic traceability of food (Kelly, Heaton & Hoogewerff, 2005).

42 In the last decades, geochemistry of light (H, C, O, N, B) and heavy (Sr, Pb) isotopes, sometimes combined with multi-elemental analysis and chemometrics, have been applied to the 43 authentication and the tracking of geographic provenance of foods (e.g., Evans, Pashley, Richards, 44 Brereton & Knowles, 2015; Medini, Janin, Verdoux & Techer, 2015; Techer, Lancelot, Descroix, 45 Guyot, 2011) and processed beverages such as wine, one of the most investigated product derived 46 from fermentation of fresh fruit (e.g., Di Paola-Naranjo et al., 2011; Durante et al., 2013, 2016; 47 Petrini, Sansone, Slejko, Buccianti, Marcuzzo & Tomasi, 2015; Rummel, Hölzl, Horn, Roßmann & 48 49 Schlicht, 2010; Vinciguerra, Stevenson, Pedneault, Poirier, Hélie & Widory, 2016).

A relevant aspect in isotope geochemistry studies is the variation of stable isotopic 50 composition of light elements by climatic variation (e.g., Christoph, Roßmann, & Voerkelius, 2003; 51 Christoph et al., 2004). On the other hand, isotopic composition of radiogenic heavy elements, 52 53 such as Sr and Pb, show the advantage of correlating directly with the geological and pedological 54 substrata with no effects related with climatic conditions, which allows to identify possible unambiguous association between the agricultural product and the geological setting of the 55 production area (e.g., Horn, Schaaf, Holbach, Hölzl & Eschnauer, 1993; Marchionni et al., 2013; 56 Tommasini et al., 2018, and references therein). 57

58 Several authors, however, showed that Sr and Pb uptaken from soil with nutrients is 59 differentially enriched in different parts of the vine (e.g., roots, branches, leaves, grapes; Amorós 60 Ortiz-Villajos et al., 2017; Bravo et al., 2017; Censi, Saiano, Pisciotta & Tuzzolino, 2014; Tommasini, 61 Davies & Elliott, 2000).

The experimental studies on the distribution of ⁸⁷Sr/⁸⁶Sr in wines available in the scientific 62 63 literature focused mainly on the evaluation of the consistency of the Sr-isotopic value through the 64 winemaking process from the soil to the wine (e.g., Almeida & Vasconcelos, 2004; Durante et al., 2013, 2016; Marchionni et al. 2016). These papers, however, did not check in detail the Sr-isotopic 65 traceability of grapes from different vines, in comparison with the Sr-isotopic variability of the 66 substrata of the vineyards (e.g., soils and rocks). Indeed, grapes are not only important for the 67 wine consumers, but also for the consumers of the fresh fruit itself. Presently, there are no 68 69 detailed Sr-isotopic studies on grapes in general, notwithstanding grape production represents the 70 world's most important fresh fruit crop with the highest total value of production (FAO-OIV Focus, 2016), with half used for wine production, and the greatest quantity of the remaining part 71 72 destined to the global market to be consumed as fresh fruit.

In this paper we report the first detailed research on the ⁸⁷Sr/⁸⁶Sr distribution in grapes from 73 different cultivars from three different geologically well-constrained vineyards. This experiment is 74 75 an extensive grape inter-varietal and site-specific study designed for evaluating the possible contribution to the ⁸⁷Sr/⁸⁶Sr of grapes of the cultivar versus the different geology/pedology of the 76 vineyards substrata. Samples of grapes from different cultivars of four different harvest years and 77 samples of soils and bedrocks were analysed for ⁸⁷Sr/⁸⁶Sr isotopic composition to shed some lights 78 on the following issues: i) the consistency of ⁸⁷Sr/⁸⁶Sr isotopic ratio in different harvest years; ii) 79 the influence of ⁸⁷Sr/⁸⁶Sr in grapes from vineyards with different geological substrata; iii) the 80 possible influence of different cultivars on the ⁸⁷Sr/⁸⁶Sr composition of grapes. The findings of this 81 study will bring further insights in the issue of food geographical traceability using radiogenic 82 83 isotope of heavy elements of geological interest.

84 **2.** Materials and Methods

In the present pilot study a restricted area with vineyards characterised by young volcanic rocks with different ⁸⁷Sr/⁸⁶Sr was selected to minimise the effects of age variability among rocks of different ages and ⁸⁷Sr ingrowth due to time integration (Faure, 1986). In addition, because the

pyroclastic volcanic rocks of the study area are rich in the glass fraction rather than in the crystalline one (Conticelli, Francalanci, Manetti, & Peccerillo, 1987) the possible ⁸⁷Sr/⁸⁶Sr bias between the geological substratum, the soil, and its bioavailable fraction is minimised.

91 The grape samples collected for this study are from a single winery in southern Tuscany, 92 Central Italy. The Sassotondo farm, managed as a high quality organic farm, is located between the 93 small towns of Pitigliano and Sorano on 72 hectares, 10 of which are destined to the vine (Vitis 94 vinifera L.) cultivation (ESM-Fig. 1, Electronic Supplementary Materials 1). The vineyards produce both red grape (Ciliegiolo, Sangiovese, Merlot, and Teroldego cultivars) and white grape 95 96 (Trebbiano, Greco, and Sauvignon cultivars). The trial was carried out in three different vineyards: 97 Pian de' Conati, Piana San Lorenzo, Crucignano, which are located in the NW and in the S, and SE of Pitigliano, respectively (ESM-Fig. 1, Electronic Supplementary Materials 1). Each vineyard is 98 99 characterised by different geological substratum.

The Pitigliano area is underlain by volcanic rocks belonging to the Latera volcano (Vulsinian volcanic district), which is made up by a succession of seven ignimbrites (e.g., Conticelli, Francalanci, Manetti, & Peccerillo, 1987; Conticelli, Francalanci, & Santo, 1991) each of them characterised by different Sr-isotopic compositions (e.g., Conticelli, Avanzinelli, Ammannati & Casalini, 2015).

105 2.1 Sampling

For this study, we collected 46 samples, 29 of red (14 Ciliegiolo, 4 Sangiovese, 4 Merlot, and 7 106 107 Teroldego cultivars) and 17 of white grapes (6 Trebbiano, 6 Greco, and 5 Sauvignon cultivars) 108 varieties. They have been taken within four harvest years from 2013 to 2016, and collected in 109 three different vineyards, each of them characterised by different bedrock and soil (a = Pian de' Conati, b = Piana di San Lorenzo, c = Crucignano vineyards; ESM-Fig. 1, Electronic Supplementary 110 Materials 1). Each sample consists of a bunch of grapes picked from a single plant of the vineyard. 111 112 Grape samples of the same cultivar and harvest year were collected from different plants of the same vineyard. 113

We collected whole soil samples from *Pian de' Conati* (4 samples), *Piana di San Lorenzo* (2 samples), and *Crucignano* (1 sample) at a mean depth of 20-30 cm (Marchionni et al., 2016) and of a unitary weight of about 500 g. Among bedrocks (Vezzoli et al., 1987), two samples are from *Pian de' Conati* (*Grotte di Castro Formation*), one sample is from *Piana di San Lorenzo* (*Pitigliano*

formation), and one sample from *Crucignano* (*Pitigliano formation*) vineyards (ESM-Fig. 1, Electronic Supplementary Materials 1). In addition ⁸⁷Sr/⁸⁶Sr data from scientific literature and from an unpublished Sr-isotopic Authors' database are also used for investigating the relationships between grapes and vineyard of provenance (Conticelli et al., 2015).

122 A detailed scheme of the sample strategy adopted is reported in the Electronic 123 Supplementary Material 1 (ESM-Tab. 1).

124 2.2 Sample preparation and analysis

All the collected samples have been prepared and measured in the Radiogenic Isotopes Lab of the Earth Science Department at the *Università degli Studi di Firenze*. The preparation protocols were different depending on the kind of material (grape, soil, rock, etc.); the treatment and preparation of the samples followed the procedure reported by Marchionni et al. (2016) and were performed in a clean chemistry laboratory "Class 1000" environment. High purity chemical reagents and Milli-Q[®] water were used during sample treatment to reduce the level of contamination.

Grape samples were first washed and rinsed with Milli-Q[®] water for three times to be sure to 132 133 have not any dust during further sample preparation. Then grapes were crushed with skin and 134 seeds. A fraction of the collected juice was treated as follow: 5 ml were evaporated to dryness in a 135 PFA beaker on a hot plate at 90°C, then 3 ml of H₂O₂ 30 vol.% UpA were added to the dried sample and left overnight at room temperature and later evaporated to dryness at 80°C. This step of the 136 137 digestion procedure was repeated twice. To assure a complete digestion of the organic matter 2 138 ml of HNO₃ 65 vol.% were added, the beaker was covered and left overnight on the hot plate at 139 170°C and evaporated to dryness. This step was repeated twice, too.

140 Soil samples underwent two different treatments: a first portion of the sample was prepared following the procedure for rocks described below to determine the isotopic composition of the 141 142 bulk soil; the remaining part of the sample underwent to an extraction treatment using Milli-Q® water and Unibest[®] resin capsules (Dobermann et al., 1994) in order to determine the chemical 143 composition of bioavailable fraction in soil solution. Unibest® resins are able to simulate the 144 mechanism of nutrient uptake by the plant roots from soil (Skogley & Dobermann, 1996). The 145 Unibest[®] resin capsule was immerged in a muddy mixture of 200 g of soil, without any pre-146 treatment, and Milli-Q[®] water for 10 days; subsequently the Unibest[®] resin capsules was 147

extracted, rinsed, to remove any residues, treated three times with 20 ml of 2N HCl in a PFA beaker, dried to get the accumulated ions from the soil solution, and then dissolved in 0.3 ml 3N HNO₃ for Sr chromatography (Marchionni et al., 2016).

Rock samples were first mechanically crushed and then pulverised with an agate ball mill; an amount of 50 mg of homogenised sample were digested in a solution of 1:4 HNO₃ + HF at 140°C for 1-2 days, then they were brought to dryness. Two further additions of HNO₃ and dissolution in 6 N HCl at 120°C and eventually evaporation to dryness followed.

All the digested samples underwent to Sr-purification by cation exchange chromatography using Sr-Spec[®] resins in 140 μ l pure quartz micro-columns with 3N HNO₃ as eluent and Milli-Q[®] water to collect Sr. Sr purification was performed using a cation exchange chromatography in a "Class 100" vertical HEPA-filtered laminar flow hood and high-purity chemical reagents (Avanzinelli et al., 2005). The extracted Sr (about 100-200 ng) was dissolved in 2N HNO₃ and loaded on single Re filaments using 1 μ l of TaCl₅ and 1 μ l of H₃PO₅ as activator and fractionation suppressor, respectively.

Sr-isotope abundance (⁸⁸Sr, ⁸⁷Sr, ⁸⁶Sr, ⁸⁴Sr) was measured by thermal ionisation mass-162 spectrometer (TIMS), in dynamic mode, using a Thermo FinniganTM Triton-Ti[®] magnetic sector 163 field equipped with nine moveable collectors (Avanzinelli et al., 2005). Each Sr-isotope ratio 164 reported in Table 1a is the average of about 120 sets of cycles, with each cycle itself representing 165 166 the average of three measurements performed during triple-jumping dynamic measurement. The 120 sets of cycles were collected in 6 blocks, each consisting of 20 cycles with 8 seconds 167 168 integration time. An idle time of 3 seconds was set before the start of the collection after each jump, to eliminate possible memory effect due to the decay of the signal in the faraday cups. 169

Procedural blank was <200 pg resulting in negligible sample correction. The external precision of NIST SRM987 international reference sample for the time of this study was 87 Sr/ 86 Sr = 0.710251 ±0.000010 (2 σ , n = 20), while the long-term mean value was 87 Sr/ 86 Sr = 0.710248 ±0.000016 (2 σ , n = 173, equivalent to an error of 23 ppm). The within run precision (i.e., $2\sigma_m$: internal precision) of 87 Sr/ 86 Sr measurements has been typically ≤10 ppm.

The reproducibility of the analytical method we used in this study is reported in Marchionni et al. (2013), where different aliquots (n = 31) of the same sample of wine (i.e., a similar organic matrix with respect to the samples of this study) were processed and measured for 87 Sr/ 86 Sr

178 composition, yielding a $2\sigma = \pm 0.000017$ (i.e., 23 ppm), which is consistent with that of the 179 international reference standard.

180 3. Results and Discussion

The values of ⁸⁷Sr/⁸⁶Sr in fresh grapes, soil bioavailable fractions, bulk soils, and whole rocks are reported in Tables 1a and 1b. The results for each cycle of measurement performed are reported as Electronic Supplementary Material (ESM-Tab. 2, Electronic Supplementary Materials 2).

185 3.1 The ⁸⁷Sr/⁸⁶Sr does not vary with time

In order to evaluate the consistency of the ⁸⁷Sr/⁸⁶Sr values over different years of harvest, first 186 of all we tested the reproducibility of the isotopic measurements on multiple sampling of bunch 187 grapes from the same cultivar (6 grapes of *Ciliegiolo* and 4 of *Teroldego*), of the same harvest year 188 189 (i.e., 2015), from different sampling points of the same vineyard (i.e., *Pian de' Conati*). The results are reported in figure 1, showing a significant constancy of ⁸⁷Sr/⁸⁶Sr values in both cases under 190 consideration. This is strongly suggestive that each bunch of grapes collected is representative of 191 the whole vineyard and cultivar of provenance. Thus, in absence of any external cause (e.g., soil 192 193 nourishment, additive addition to soil, use of chemical improver, etc.; Marchionni et al. 2016), the constancy and conservation of the isotopic composition over different harvest years is expected. 194

In figure 2 the ⁸⁷Sr/⁸⁶Sr of the grapes of the most representative cultivars sampled in the 195 different years of harvesting and from the Pian de' Conati vineyard are reported. In these plots we 196 observe for the majority of samples a fairly good conservation of the Sr-isotopic composition 197 through the different harvest years, with values within the measurement uncertainty level (1σ) . A 198 first exception is shown by the Greco and Sauvignon grapes of the 2013 harvest year, which 199 display 87 Sr/ 86 Sr values of 0.709749 ± 0.000036 (1 σ) and 0.709738 ± 0.000064 (1 σ), which are 200 201 slightly off with respect to the Sr-isotopic composition of the 2014-2016 period. This discrepancy 202 has no apparent explanations, neither analytical nor related to the farming practice. On the other hand, the Teroldego grapes display a perfect conservation of the ⁸⁷Sr/⁸⁶Sr values through the 203 204 harvest period considered.

A further notable variation is observed for *Ciliegiolo* grapes (Fig. 2) with samples from *Pian de' Conati* vineyard showing consistent higher 87 Sr/ 86 Sr values [average value 0.709957 ± 0.000046]

(1 σ)] than samples from *San Lorenzo* vineyard [average value 0.708986 ± 0.000121 (1 σ)]. This bias depends upon the different ⁸⁷Sr/⁸⁶Sr shown by the substrata of the two vineyards (ESM-Fig. 2, Electronic Supplementary Materials 1) as discussed in the next paragraph.

In summary, excluding the *Greco* and *Sauvignon* grape samples harvested in the 2013, which display lower ⁸⁷Sr/⁸⁶Sr values than other grapes from the same vineyard, the grape samples collected for this study display no significant ⁸⁷Sr/⁸⁶Sr variations through time (Fig. 2), as shown in previous experimental studies for musts and wines of different vintage years (Marchionni et al., 2013, 2016). On the other hand, systematic large ⁸⁷Sr/⁸⁶Sr variations are observed among grapes from vineyards farmed on geological substrata with different chemical composition and ⁸⁷Sr/⁸⁶Sr signatures.

217 3.2 The ⁸⁷Sr/⁸⁶Sr of bioavailable soil fraction, soil, and bedrocks

Marchionni et al. (2013) have already shown the strong correlation existing between the 218 ⁸⁷Sr/⁸⁶Sr of bottled red wines and the ⁸⁷Sr/⁸⁶Sr of the bedrocks of the area of production, although 219 a discrepancy might be observed between the ⁸⁷Sr/⁸⁶Sr of the wine and that of the geological 220 221 substratum of the area of production. This bias increases passing from wine production areas 222 characterised by a substratum made of young volcanic rocks to wine production area characterised by older either sedimentary or granitic rocks (Marchionni et al., 2013). Other studies 223 have also shown that the ⁸⁷Sr/⁸⁶Sr of the bioavailable fraction is a better proxy of the *terroir* of the 224 225 area of provenance of the food product from agriculture (e.g., Tescione, Marchionni, Mattei, Tassi, Romano, Conticelli, 2015; Vinciguerra, Stevenson, Pedneault, Poirier, Hélie, Widory, 2015, 2016; 226 227 Petrini et al., 2015; Durante et al., 2016; Marchionni et al., 2016).

These findings are confirmed by our data when the ⁸⁷Sr/⁸⁶Sr of bedrocks are compared with 228 229 whole soils and soil bioavailable fraction of the three vineyards taken under consideration in this study (Table 2 and ESM-Fig. 2; Electronic Supplementary Materials 1). Indeed, the Pian de' Conati 230 vineyard shows slight differences among the mean values of ⁸⁷Sr/⁸⁶Sr of the bedrock [Grotte di 231 Castro Formation = 0.710213 \pm 0.000037 (1 σ)], the total fraction of whole soil [0.710114 \pm 232 233 0.000054 (1σ)], and its bioavailable fraction [0.710077 ± 0.000049 (1σ)]. On the other hand, in the Piana San Lorenzo vineyard the ⁸⁷Sr/⁸⁶Sr values of the three components of the substratum are 234 slightly to significantly lower than those observed for the Pian de' Conati vineyard. In particular 235 236 large isotopic differences among geologic bedrock [Pitigliano Formation = 0.710052 ± 0.000032

(1 σ)], whole soil total fraction [0.709595 ± 0.000225 (1 σ)], and its bioavailable fraction [0.708948 ± 0.000091 (1 σ)] are observed. Similarly, in the *Crucignano* vineyard the large difference between the whole soil [0.710265 ± 0.000041 (1 σ)] and its bioavailable fraction [0.709642 ± 0.000117 (1 σ)] does also occur (Table 2 and ESM-Fig. 2; Electronic Supplementary Materials 1).

The different amount of ⁸⁷Sr/⁸⁶Sr depletion from bedrock to soil and to bioavailable fraction in 241 the three different areas can be related to the different nature of soils outcropping in the area 242 243 (Pedological Map of the Tuscany region: http://www.lamma.rete.toscana.it/territorio/cartografiatematica/pedologia/carta-dei-suoli). Different soils may release the bioavailable fraction to the 244 plants due to their different texture and inorganic composition, leading to different relationships 245 among ⁸⁷Sr/⁸⁶Sr values in bedrock, soil, vine, and wine (e.g., Marchionni et al., 2013; Petrini et al., 246 2015). Notwithstanding the significant difference observed between ⁸⁷Sr/⁸⁶Sr values of the 247 Ciliegiolo grapes from Pian de' Conati and those from San Lorenzo vineyards depends clearly upon 248 the ⁸⁷Sr/⁸⁶Sr of the bioavailable fraction rather than any other geological (i.e., bedrock, soil) and 249 biological (e.g., vine, cultivar) component (Table 2), as previously found by other studies on 250 251 isotopes of wines (e.g., Tescione et al., 2015; Vinciguerra et al., 2015, 2016; Petrini et al., 2015; 252 Durante et al., 2016, Marchionni et al., 2016).

In the following paragraphs we only discuss the relationship among the ⁸⁷Sr/⁸⁶Sr of the bioavailable fraction of the soils and of cultivars in the three different vineyards selected, leaving the analysis of the processes, which are responsible of the different mean values of ⁸⁷Sr/⁸⁶Sr in bedrock, soil, and bioavailable fraction to future studies.

257 3.3 The ⁸⁷Sr/⁸⁶Sr does not vary among cultivars, but depend upon different vineyard substrata

In figure 3 the box plots for each grape variety under consideration in this study are reported. Each box plot was calculated using all ⁸⁷Sr/⁸⁶Sr values obtained from a single grape variety of different harvest years. In addition, in this figure the range of ⁸⁷Sr/⁸⁶Sr values of the soil bioavailable fraction is reported as hatched area in the background.

In this plot we observe, with the sole exception for the *Sauvignon* grape cultivar, that the box plot derived statistically by the ⁸⁷Sr/⁸⁶Sr of the different grape cultivars, indifferently by their variety (i.e., red or white), fall well within the range of the bioavailable fraction of the soil of their vineyards (Fig. 3). As a corollary, different ⁸⁷Sr/⁸⁶Sr values are observed for grapes from the same

cultivar but harvested by vines grown in different vineyards (i.e., *Ciliegiolo* and *Trebbiano* cultivars;
Fig. 3).

On the other hand the ⁸⁷Sr/⁸⁶Sr mean values of the whole set of grape samples fall, with no 268 exceptions, well within the ⁸⁷Sr/⁸⁶Sr range of the bioavailable fractions. In addition the ⁸⁷Sr/⁸⁶Sr 269 mean value of grapes from Pian de' Conati (vineyard a) is 0.710010 ± 0.000095 (1o), which lies 270 271 within the range of variation of the relative soil bioavailable fraction $[0.710077 \pm 0.000049 (1\sigma)]$, and it is discernible from grape collected in San Lorenzo (vineyard b) $[0.708986 \pm 0.000121 (1\sigma)]$ 272 and Crucignano (vineyard c)[0.709526 ± 0.000038 (1 σ)](Table 2). These findings consistently 273 support the hypothesis that ⁸⁷Sr/⁸⁶Sr of grape depends on the ⁸⁷Sr/⁸⁶Sr of the-soil solution from 274 275 which the vine roots absorb nutrients as bioavailable substances.

276 Regarding the discrepancies observed in the box plot of *Sauvignon* these are caused by the 277 lower ⁸⁷Sr/⁸⁶Sr values of the 2013 harvest year, shown in figure 2. The same was observed for the 278 ⁸⁷Sr/⁸⁶Sr of *Greco* in the same harvest year but that did not affect greatly the box plot. Tentatively 279 the sole explanation that we can find for these 2013 samples can be found in the low number of 280 cycles during the experimental runs (i.e., 55-75; Table 1a).

281 3.4 ⁸⁷Sr/⁸⁶Sr as a tool for geographic traceability at different scales

The potential of ⁸⁷Sr/⁸⁶Sr for food geographic origin applications has been abundantly investigated in many studies, at a wide scale, on different products: orange juice from 14 producing countries in the world were analysed by Rummel et al. (2010), and Garcia-Ruiz et al. (2007) attempted the geographical discrimination of ciders from 4 different European countries.

One of the most tested products with Sr-isotopes, due also to its high economic relevance, is wine. In fact, several studies can be found in literature, aiming to discriminate wines from a global to a regional scale, according to the nations of production (Barbaste, Robinson, Guilfoyle, Medina, & Lobinski, 2001; Horn et al., 1993), to different regions in the same country (Boari et al., 2008; Marchionni et al., 2013; Vinciguerra et al., 2015, 2016), and to different wineries in the same producing region (Durante et al., 2013; Petrini et al., 2015; Marchionni et al., 2016).

292 In order to perform an inter-varietal investigation among various grapevine genotypes, the 293 scale of the study area has been further reduced to a single producer.

The values of ⁸⁷Sr/⁸⁶Sr measured in the grapes of this study show that the relation with the geologic bedrock of the vineyard of provenance is not relevantly affected by the plant genotype

(Fig. 3) and by the harvest year and, despite the reduced study area, the grape samples collected in the three different vineyards of production are isotopically well defined and differentiated by each other. This implies that the geographic provenance for grapes can be tracked at a very fine tuning, being possible to discriminate among single vineyards of the same geographic area. On the other hand, no direct information about the variety and the vine genotype can be obtained with Sr-isotopes.

The possibility of checking the origin of grapes at such high level of geographic detail has relevant implications in terms of geographic traceability control in agreement with the European regulation for Protected Denomination of Origins (PDO) and therefore the Sr-isotopic method represents a valid and robust support for the product authenticity assessment.

The robustness of this tool at the local scale of the vineyard has been verified and, as the geology of the bedrock is the only discriminant factor, its applicability at a regional scale to check the provenance of grape at the national or even global market is also encouraged.

309 4. Summary and Conclusions

310 In this study we have shown that grapes inherit their Sr isotope composition exclusively from the geologic substratum of the vineyard of production and that it does not depend on the variety 311 of the cultivar. It is worth noting that the correspondence between ⁸⁷Sr/⁸⁶Sr in vine products and 312 geologic bedrock can be limited due to the selective absorption of chemical elements by vine 313 roots. This concern can be encompassed by correlating the ⁸⁷Sr/⁸⁶Sr of grapes with that of the 314 bioavailable soil fraction. In addition, our data show that for harvest years 2013-2014-2015-2016 315 the Sr-isotope remains fairly constant, arguing for the lack of influence of the climatic variations 316 on its value. Therefore, this study clearly shows that ⁸⁷Sr/⁸⁶Sr can be a useful analytical tool to 317 check the geographical provenance of fresh grape fruits, using the ⁸⁷Sr/⁸⁶Sr of the bioavailable 318 319 fractions, which is somehow related to the geological substratum of the vineyard.

The study was conducted at the small scale to demonstrate the reliability and reproducibility of data within a well-controlled operational farming practice and well constrained geological and pedological environments to prevent any kind of food fraud and to guarantee the final product origin. In addition the reduced scale of the study shows the potential application of this tool, on a wide regional scale, and on the vineyard scale, with the possibility of confirming the authenticity of food products grown by different producers in the same viticultural region.

Our work demonstrates that Sr isotopes can be used to assess the geographical origin of grapes and may also be used to label wine production. Furthermore, they might be useful to guarantee the observance of the regulations and the control on the production chain of high quality wines.

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466 **Table headings**

- 467 **Tab. 1.a** $-{}^{87}$ Sr/ 86 Sr of analysed grapes from different harvest years
- 468 **Tab. 1.b** 87 Sr $/ {}^{86}$ Sr of substrata of the vineyards
- 469 Tab. 2 Descriptive statistics of the grape, bioavailable soil fraction, soil, and rock samples

470 Table captions

- 471 **Tab. 1** Values here reported are the mean values of one single squeezed grape batch for the
- 472 2013, harvest year, three squeezed grape batches for the 2014 harvest year, and two
 473 squeezed grape batches for the 2015 and 2016 harvest years.
- 474 Figure Captions

- 475 Fig. 1 ⁸⁷Sr/⁸⁶Sr values for the 2015 harvest year and *Pian de' Conati* vineyard (a) for different
 476 samples of *Ciliegiolo* and *Teroldego* cultivars. The error bars represent the standard
 477 deviation (1σ).
- ⁸⁷Sr/⁸⁶Sr composition of *Ciliegiolo, Greco, Sauvignon and Teroldego* cultivars collected in Fig. 2 – 478 479 the Pian de' Conati (Vineyard a) and Piana S. Lorenzo (Vineyard b) areas through the 480 2013 – 2016 harvest years interval. Ciliegiolo red grapes harvested in San Lorenzo 481 vineyard are represented with empty squares and show lower mean values. The yellow triangles and circles are related to white grape cultivars (Sauvignon and Greco 482 respectively), the red squares and diamonds represent the red grape cultivars (Ciliegiolo 483 and Teroldego), all of them grown in Pian de' Conati area. The error bars represent the 484 standard deviation (1σ) . 485
- Fig. 3 Box plot of the ⁸⁷Sr/⁸⁶Sr values distribution in different grape varieties from the three different areas compared to the soil bioavailable fraction. The hatched area represents the minimum and maximum values of the bioavailable fraction distribution. The yellow boxes are related to white grape cultivars (*Sauvignon, Greco,* and *Trebbiano*), the red boxes represent the red grape cultivars (*Ciliegiolo, Merlot, Sangiovese,* and *Teroldego*).

Grapes								
Sample Type	Harvest Year	Variety	⁸⁷ Sr/ ⁸⁶ Sr	1σ	n			
Vineyard A - Pian de' Conati								
	2012	Greco	0.709749	± 0.000036	55			
	2013	Sauvingnon	0.709738	± 0.000064	75			
		Sauvignon 1	0.709749	± 0.000036	55			
	2014	Sauvignon 2	0.710048	± 0.000039	116			
		Greco 1	0.709960	± 0.000031	112			
		Greco 2	0.710029	± 0.000036	117			
14/		Trebbiano 1	0.710069	± 0.000045	115			
vv b		Trebbiano 2	0.710049	± 0.000043	114			
n i	2015	Greco 1	0.710082	± 0.000042	116			
		Greco 2	0.710050	± 0.000035	115			
د م		Sauvignon 1	0.710065	± 0.000036	81			
C		Sauvignon 2	n.d.	n.d.	n.d.			
		Sauvignon 1	n.d.	n.d.	n.d.			
		Sauvignon 2	0.710042	± 0.000126	81			
	2016	Greco 1	n.d.	n.d.	n.d.			
	2016	Greco 2	0.710068	± 0.000172	65			
		Trebbiano 1	0.710081	± 0.000042	115			
		Trebbiano 2	0.710082	± 0.000040	91			
	2013	Ciliegiolo	0.709960	± 0.000034	113			
		Teroldego 1	0.710014	± 0.000033	113			
	2014	Teroldego 2	0.710028	± 0.000035	113			
	2014	Sangiovese 1	0.710020	± 0.000033	116			
		Sangiovese 2	0.710050	± 0.000034	115			
		Teroldego 1	0.710000	± 0.000043	117			
		Teroldego 2	0.710017	± 0.000034	113			
		Teroldego 3	0.710053	± 0.000035	117			
	2015	Teroldego 4	0.710033	± 0.000053	111			
		Ciliegiolo 1	0.709982	± 0.000041	113			
		Ciliegiolo 2	0.710008	± 0.000053	23			
R		Ciliegiolo 3	0.709949	± 0.000042	98			
е		Ciliegiolo 4	0.709942	± 0.000038	115			
d	2016	Ciliegiolo 5	0.709981	± 0.000058	76			
		Ciliegiolo 6	0.709942	± 0.000035	116			
		Merlot 1	0.710044	± 0.000041	//			
		Ivieriot 2	0.710216	± 0.000079	102			
		Sangiovese 1	0.710031	± 0.000045	114			
			0.710008	+ 0.000030	112			
		Teroldego 2	n d	± 0.000055 n d	n d			
		Ciliegiolo 1	0 709951	+ 0 000050	96			
		Ciliegiolo 2	0.709940	+ 0.000048	94			
		Merlot 1	0 710040	+ 0 000037	66			
		Merlot 2	0.710040	+ 0.000037	118			
	Vine	vard B - Piana S	an Lorenzo	- 0.0000000	110			
	2013	Ciliegiolo	0.708963	± 0.000040	117			
R	2014	Ciliegiolo	0.708834	± 0.000040	119			
	2015	Ciliegiolo 1	0.709119	± 0.000041	94			
		Ciliegiolo 2	0.709128	± 0.000034	114			
a		Ciliegiolo 1	n.d.	n.d.	n.d.			
		Ciliegiolo 2	0.708916	± 0.000035	116			
Vineyard C - Crucignano								
Pad	2016	Trebbiano 1	0.709527	± 0.000039	115			
Reu	2010	Trebbiano 2	0.709525	± 0.000038	114			

Table 1.a - ⁸⁷Sr/⁸⁶Sr of analysed grapes from different harvest years

Bedrock and Soil									
Sample Type	Formation	Fraction	Average ⁸⁷ Sr/ ⁸⁶ Sr	1σ	n				
Vineyard A - Pian de' Conati									
Rock	Gratta di Castra Ignimbrita	whole	0.710204	± 0.000033	117				
			0.710223	± 0.000039	114				
Soil		whole	0.710145	± 0.000035	114				
	Grotte di Castro Ignimbrite		0.710125	± 0.000037	114				
			0.710133	± 0.000054	116				
			0.710052	± 0.000032	112				
	Grotte di Castro Ignimbrite	bioavailable	0.710095	± 0.000050	93				
			0.710066	± 0.000039	115				
			0.710096	± 0.000035	114				
			0.710056	± 0.000056	117				
Vineyard B - Piana San Lorenzo									
Rock	Pitigliano Formation	whole	0.710052	± 0.000032	112				
Soil	Ditigliano Formation	whole	0.709371	± 0.000037	111				
		whole	0.709810	± 0.000055	116				
	Pitigliano Formation	hioavailahle	0.708884	± 0.000037	113				
		bloavailable	0.709010	± 0.000084	114				
Vineyard C - Crucignano									
Soil	Pitigliano Formation	whole	0.710265	± 0.000041	114				
	Pitigliano Formation	bioavailable	0.709642	± 0.000117	114				

Table 1.b - ⁸⁷Sr/⁸⁶Sr of substrata of the vineyards







Electronic supplementary materials - 1

⁸⁷Sr/⁸⁶Sr isotopes in grapes of different cultivars: a geochemical tool for geographic traceability of agriculture products

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Focus on the geological setting of the Vulsini Volcanic District

The Vulsinian District is a 2,000 km² widespread volcanic area formed about 600 – 100 kyr BP (Vezzoli et al., 1987; Conticelli et al., 2010). The volcanism produced a thick sequence of pyroclastic deposits and lava flows that formed the three main volcanic apparata of Bolsena, Latera and Montefiascone (ESM-FIG1). These coalescent volcanoes were characterized by similar eruptive styles with ignimbrite-forming eruptions preceded and followed by effusive and strombolian activities, usually taking place along peripheral circum-calderic fault system.

The volcanic activity in the Latera area developed from about 429 to 145 ka. Lavas were confined in the early and late stage of the volcano, whilst pyroclastic activity occurred between 278 and 166 ka, with the emplacement of six main ignimbrites inter-bedded to some events of pyroclastic fall and surges (Conticelli et al. 2010). A large polygenic caldera was formed.

The *Sassotondo* farm, the winery used for the experimental work of this study, has vineyards lying on the volcanic succession of Latera volcano in the surroundings of the town of Pitigliano (Fig ESM-1). The vineyards develop in three different areas: a) *Pian de' Conati*, which has the soil

formed on the bedrock belonging to the Grotte di Castro formation; b) *Piana San Lorenzo*, which has the soil formed on both Poggio Pinzi Tuffs and Pitigliano Formation, but the grapes were from vines on the former bedrock; c) *Crucignano*, which has a soil formed directly on the Pitigliano Formation. The different soils are formed on the above mentioned pyroclastic bedrocks (i.e., ignimbrites and falls) and they range in terms of pedology, according to the Tuscany Pedological map - <u>http://www.lamma.rete.toscana.it/territorio/cartografia-tematica/pedologia/carta-dei-suoli</u> - from "*Molli Pachic Andosols*" (autoc. *Scopetone* soil) for *San Lorenzo* vineyard, to "*Epileptic Andosols*" (autoc. *Farmacista* soil) for *Crucignano* vineyard, and "*Eutric Epileptic Andosols*" (autoc. *Aia di Tufi* soil) for *Pian de' Conati* vineyard (IUSS WGW, 2006).



ESM-Fig. 1 – Geological map of the study area showing the three vineyards of the Sassotondo farm. On the left side insets a,b,c represent the three vineyards: a = *Pian de' Conati* Area, b = *Piana S. Lorenzo*, c = *Crucignano*, whereas the polygons represent the grape variety grown. In *Piana San Lorenzo* area only *Ciliegiolo* variety is grown, while in *Pian de' Conati* area both red (*Ciliegiolo*, *Teroldego*, *Merlot*, and *Sangiovese* cultivars) and white (*Greco*, *Trebbiano*, and Sauvignon cultivars) grape varieties can be found. The inset below represents the distribution of the volcanic rocks of the Vulsinian District, Roman Magmatic province (Vezzoli et al., 1987).



ESM-Fig. 2 – Box plot of the 87 Sr/ 86 Sr value distribution of the different samples (i.e., bedrock, whole soil, and bioavailable fraction of soil) for the three vineyards analysed.

				2014	2015	2016
Vineyard A - Pian de' Conati	White Grape	Greco	1	2	2	1
		Sauvignon	1	2	1	1
		Trebbiano			2	
	Red Grape	Teroldego		2	4	1
		Ciliegiolo	1		6	2
		Merlot			2	2
		Sangiovese		2	2	
		Trebbiano				2
Vineyard B - Piana San Lorenzo	Red Grape	Ciliegiolo	1	1	2	1
Vineyard C - Crucignano	Red Grape	Trebbiano				2

ESM-Tab. 1 – Schematic representation of the sampling strategy of the bunch grapes and of related materials to the vineyards (i.e., bedrocks, whole soil, bioavailable fraction of the soil).

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