Taphonomy in the kitchen: culinary practices and processing residues of native tuberous plants of the south-central Andes

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ABSTRACT

We present comparative material for the identification of culinary residues of cooked tubers of Solanum sp., Oxalis tuberosa and Ullucus tuberosus. We use a broad concept of taphonomy that includes the study of plant modifications resulting from the preparation of food, in this case the boiling and cooking al rescoldo of fresh tubers. We undertake a number of controlled cooking experiments and compare the results with fresh samples. We discuss morphological and optical modifications of tissue fragments and intracellular particles resulting from our cooking experiments. Finally, we discuss the possibility of recognizing cooking techniques from microscopic analysis of tuber remains.

Keywords: Taphonomy; Culinary techniques; Plant processing; Starch; Tubers.

RESUMEN

TAFONOMÍA EN LA COCINA: PRÁCTICAS CULINARIAS Y RESIDUOS DEL PROCESAMIENTO DE PLANTAS TUBEROSAS NATIVAS DE LOS ANDES CENTRO-SUR. Presentamos material comparativo para la identificación de residuos culinarios de tubérculos cocidos de Solanum sp., Oxalis tuberosa y Ullucus tuberosus. Partimos de un concepto amplio de tafonomía que incluye el estudio de las modificaciones de las plantas resultantes de la preparación de alimentos; en este caso, aquellas que se deben al hervido y cocción al rescoldo de tubérculos frescos. Realizamos experimentos de cocción controlados y comparamos los resultados con muestras frescas. Describimos las modificaciones en los atributos morfológicos y ópticos de tejidos y partículas intracelulares resultantes de nuestros experimentos de cocción. Finalmente, discutimos la posibilidad de reconocer las técnicas de cocción a partir del análisis microscópico de vestigios de tubérculos.

Palabras clave: Tafonomía; Técnicas culinarias; Procesamiento de plantas; Almidón; Tubérculos.
A TAPHONOMIC APPROACH OF CULINARY TRANSFORMATIONS

This work is part of a series of studies initiated over a decade ago (Babot and Korstanje 2001; Babot 2003), aimed at generating comparative material for identifying plant processing techniques from microscopic analysis of preserved residues of plants native to the south-central Andes. It is worth noting that the transformations and agents involved in such preparations (e.g., cooking) go beyond the scope of food, as medicinal or ritual uses can also be important. Thus, although the perspective of this work is culinary, this is only circumstantial; the emphasis is on how techniques, as modes of transformation, affect matter (Pazzarelli 2012).

This study deals with the cooking of tuberous plants of the genus *Solanum* L. (Solanaceae), “potatoes”; *Oxalis tuberosa* Mol. (Oxalidaceae), “oca” and *Ullucus tuberosus* Caldas (Basellaceae), “ulluco” or “papa lisa”. We began with an ethno-botanical approach that considers the traditional uses of plants in the south-central Andean region and the various means by which tubers become food. We developed experiments to obtain tubers cooked *al rescoldo* and boiled tubers, which may themselves constitute simple foods, based on a single ingredient, or be part of complex preparations with other plant, animal and mineral ingredients.

We rely on a broad concept of “recipe” “[...] as a more or less flexible or open formula to achieve a preparation” (Babot et al. 2012: 242). The components of a recipe are its ingredients in relative amounts and combinations, the modes and techniques of preparation and service, the circumstances of its use, the implements required throughout the process, and the spatio-temporal context and the actors involved in the culinary performance (Babot et al. 2012). Of all these components, this paper deals with tubers as ingredients in recipes, and with methods of tuber preparation. Preparation sequences are not rigid or linear because a food element can be associated with multiple recipe pathways of different length and complexity (Babot 2009). “So, when we refer to old recipes, we refer to approximations to the modes of preparing foods, rather than to strict and closed models of the ingredients, their preparation and presentation [...]” (Babot et al. 2012: 241).

The taphonomic study of food residues gives us a unique perspective on how plants, animals and minerals were transformed into food and their subsequent history to the point of archaeological recovery and analysis. According to the “unrestricted taphonomy” approach (Borrero 2011) we use a broad definition of the concept as the study of the decomposition “or modification” of organisms, their parts or products, and of the processes leading to their accumulation and differential preservation in archaeological contexts, so that they can be genuine sources of archaeological information (Lewarch and O’Brien 1981; Borrazzo 2006). We are particularly interested in assessing how such a perspective can inform our interpretation of past human practices. Thus, we consider the modifications of plants’ useful parts associated with different methods of food preparation, which condition the appearance and integrity of such food remains as recovered from archaeological contexts. Such modifications can be explained in terms of different physico-chemical processes that affect the structure and the properties of starch and other vegetable intracellular particles, and histological elements (Babot 2003). Previous work has demonstrated the utility of taphonomic and experimental approaches to the macroscopic and microscopic analyses of the modes and procedures for preparing animal or vegetable foods (e.g., González and Frère 2004; Gong et al. 2011; López et al. 2011; Lovis et al. 2011; Raviele 2011; Babot et al. 2012 and references therein; Lantos et al. 2012). In particular, our approach focuses on the micro-morphological characterization of such modifications.

In this study, we characterized the anthropic/culinary modifications of micro-particles and tissue fragments in different varieties of tubers of three genera, which allowed us to formulate expectations for archaeological cases. Thus, the various stages of investigation aimed to: 1) characterize fresh tissues of different varieties of potato, oca and ulluco, since these are ingredients in traditional culinary preparations and recipes; 2) observe how aspects of the fresh tissue of potato, oca and ulluco change or disappear as a consequence of two traditional food preparation methods (boiling and cooking *al rescoldo* until completely cooked), and identify which new elements appear in processed foods, explaining these on the basis of the physico-chemical processes involved; and 3) compare the two sets of observations to identify taphonomic signatures of these culinary practices in archaeological situations.

THE TAPHONOMY OF MICROFOSSILS

Current Perspectives

The taphonomic study of plant microfossils and their archaeological associations is a versatile approach. Some taphonomic researches focus on the ways various biological (e.g., soil fauna and roots) and environmental agents (e.g., pH, moisture and temperature conditions, salt precipitation and transport) alter the microfossil record and affect its survival and integrity (Therin 1994; Hart 2003; Humphreys et al. 2003; Haslam 2004; Barton and Matthews 2006; Osterrith et al. 2013; Musaubach and Babot 2014, among others).
These studies also consider humans, through the manipulation of the natural environment, as an agent of transformation, since they can modify plants and their microfossil remains through residential use of certain places, agriculture, or burning (Therin 1994; Parr et al. 2008; Osterrieth et al. 2009, among others). These kind of taphonomic researches have focused on analysis of biosiliceous particles, calcium phytoliths and starch from the perspective of paleoecological, pedological and archaeological problems. Previous work has also included study of the effects of different laboratory manipulations on the integrity and composition of microfossil assemblages that are ultimately observed under the microscope. Many of these recent studies also involved revision of sample extraction and treatment protocols with particular attention to archaeological applications (Babot and Korstanje 2001; Coi et al. 2003; Korstanje 2003; Babot 2007; Korstanje and Babot 2007; Torrence and Barton 2007, etc.).

Other taphonomic studies closely related to the analysis of use residues on archaeological artifacts assess the survival rates and prospects of starch –one of the main microfossils useful for these purposes– under different conditions of entrapment, depth, sediment compaction and exposure, and considering the passage of time according to the nature of the processed substance (Lu 2003; Babot and Bru de Labanda 2005; Barton and Matthews 2006; Langejans 2010, among others). The potential for post-depositional contamination has also been discussed (Barton et al. 1998). Related to taphonomic studies but following different approaches are archaeologically or bromatologically inclined histological studies designed to understand the feasibility and reliability of identifying plant tissues exhibiting different degrees of fragmentation and preservation (Pochettino and Scattolin 1991; Cortella and Pochettino 1994).

We are optimistic about the possibility of obtaining archaeological information from taphonomic and contextual approaches and we consider residues (meaning remains in general, rather than a priori use residues) as dynamic systems that can provide data about use, the contemporaneous context for such use, processes subsequent to deposition, and the manufacture of associated artifacts (the latter when microfossils are included in the artifact raw materials) (Babot and Haros 2008; Hart 2011; Babot et al. 2012). Thus, when considering plant residues located on an artifact or in an archaeological matrix, it would hypothetically be possible to identify interfaces. In the first case, such interfaces would be artifact/residue/sediment matrix after deposition, where the microfossil “signal” attributable to the raw materials would decrease from the artifact mass to its surface; the “signal” related to use would decrease from the surface of the artifact to the sediment matrix, and the “signal” of the matrix would follow an opposite tendency to that of use. Hence, precautions taken during sample extraction and the criteria used to interpret manufacture, use and context are very important (Babot and Haros 2008; Zucol and Loponte 2008; Hart 2011). The exchanges, gains, losses and modifications of material that occur at the interfaces between object, use residue and matrix must be considered, including other processes that may occur between the time archaeological remains are extracted and analytical data are obtained. Sometimes, burial conditions can promote the preservation of residues, degradation byproducts and new products resulting from interactions with the sedimentary matrix (Jones 2009). This applies to different types of elements contained in the residues, whether chemical, histological or intracellular.

In this regard, we consider that interpreting the nature of a residue requires knowing the particular history of the object under analysis. In other words, a residue could indicate 1) the last recorded use of an artifact (Haslam 2006), or 2) successive use episodes recorded in the stratification of the remains (Musaubach and Beron 2012), or 3) random averaging (in a broad sense rather than referring to statistical significance) resulting from repeated use, partial cleaning between successive uses, differential decay of components and exchanges between the environment, the object’s mass and the residue, plus laboratory treatments (Babot 2007; Babot et al. 2012).

A taphonomic approach to plant microfossils and anthropogenic manipulations

A small number of studies have discussed the transformation of tissues and intracellular particles resulting from anthropogenic manipulation of plants. Examples include the identification of striae on the surface of silica phytoliths obtained during threshing of silica-rich taxa (e.g., European grains); the transformation of calcium oxalate crystals into calcite pseudomorphs, which occurs during combustion (Juan-Tresserras 1992); the fracture of biosiliceous particles as a result of activities such as grinding (Checa et al. 1999); and the tearing and disarrangement of plant fibers during mastication (Musaubach and Babot 2014). The formation of new products during cooking (e.g., micro-charcoals and particle clumps) has also been documented (Babot 2003; Tassara and Osterrieth 2008, among others).

Starch is one of the most studied intracellular particles, both in archaeology and other fields. Analyses of the physico-chemical processes that can alter starch have a long history, linked initially to biological and food industry interests (e.g., Radley 1943; Whistler et al. 1984). These studies have addressed aspects including starch grains’ loss of structure and birefringence properties through contact
with caustic chemicals, heat in the presence of water, the action of enzymes during seed germination, and the sprouting of underground organs. The first applications of this knowledge to archaeological situations were related to questions about food, particularly brewing, grinding and boiling of starchy plants (e.g., Juan-Tresserras 1992; Loy 1994; Checa et al. 1999). These studies, along with the work of Babot (2003), Beck and Torrence (2006), Fullagar (2006), Samuel (2006), Henry et al. (2009), Gong et al. (2011) and Crowther (2012), are directly related to our research, as they consider the effects past anthropogenic manipulation of plants had on their starches. Babot’s (2003) research aimed to develop modern comparative standards for application to archaeological cases (Babot 2009; Babot et al. 2012, etc.), and has focused on the optical and morphological effects of various processing techniques on starch (e.g., aeration, sun drying, toasting, ashing, freezing, desaponification/washing –i.e., removal of saponins–, and grinding). Babot’s research also includes recognition of various cooking practices from starchy residues. Beck and Torrence (2006) considered the paths of starch when used for different purposes in various cultural contexts. Fullagar (2006) evaluated the role of experiments in the functional assessment of archaeological artifacts, and Samuel (2006) discussed various ways in which modified starch has been preserved, including bread and yeast residues. Henry et al. (2009) documented the progression of starch damage due to the baking, boiling, drying, and fermentation of domesticated legumes and grass seeds, and found boiling to be the most harmful cooking technique for starch. Gong et al. (2011) studied food remains from exceptionally well preserved mortuary offerings, and developed experiments to aid identification of culinary preparations. Crowther (2012) discussed the influence of moisture on starch during cooking.

These studies of cultural manipulations have shown that, although they are modified, starch and other intracellular elements can survive multiple food preparation processes. Drying, heating, the breaking down of tissues by trampling and friction, and the formation of ice crystals, all generate physico-chemical changes in those particles, modify their completeness and degree of crystallinity, and produce optical and morphological changes in some grains (Babot 2003). While the damage caused by different food processing techniques may look similar, and although sometimes the same process can alter the starch from different biological sources in different ways (Babot 2003; Henry et al. 2009), it has been shown that, in general, distinct damage patterns seem to result from different processes (Babot 2003). Also, it has been documented that the more intense the process, the greater the severity of the damage. In addition, it is sometimes possible to infer the consecutive stages of processing, which are present as damage patterns superimposed on the same individual starch granule or on different granules within a sample.

Damaged starch grains are more susceptible to hydrolytic agents and fungal and bacterial activity than well-preserved starch. This is attributed to infiltration of the grains’ interior via cracks associated with the damage (Radley 1943; Cortella and Pochettino 1994) and to structural weaknesses resulting from physico-chemical changes to the granules. However, research on ancient starch has shown that it is possible to recover damaged grains from both modern and archaeological contexts, as this carbohydrate’s pseudo-crystalline structure survives the passage of time even when damaged.

Ancient starch analyses with a taphonomic perspective have increasingly focused on damaged starch grains (alongside those recovered intact), as well as the natural and cultural causes of the observed changes. In addition, appropriate and specific terminologies for the description of alteration processes and damage have been adopted (Babot 2003; The International Code for Starch Nomenclature (ICSN 2011)), referring to the so-called modified starch (sensu Samuel 2006; ICSN 2011). This category and one referred to as resistant starch have been distinguished from native starch, which refers to unaltered grains. Such studies have recognized certain traits that have a taphonomic rather than taxonomic origin, and have proposed that damaged starch is a source of information regarding cultural practices and post-depositional processes. Damage patterns have also been used to evaluate contamination by assessing the coherence between the types of damage and the archaeological context from which the starch was recovered (Babot 2003).

Previous publications have proposed descriptors for modified starches, but when studying human practices we suggest it is possible to rank those descriptors by grouping them following hierarchical categories: a) techniques: modes of transformation implemented by humans; b) agents other than humans –the latter included in a–; c) physico-chemical processes triggered by human activities and their results or the final state of the mass of material processed, and, finally, d) morphological and optical damage and modifications or changes that are verified in starch grains within the mass (Table 1). Following Lyman’s (1994) terminology, applied techniques and resulting physico-chemical processes constitute taphonomic processes, where the taphonomic agents are the physical, chemical and biological factors involved in the transformation of grains. In this case humans are the default participants in technical manipulation. Finally, the taphonomic effects are the results of these processes and the specific patterns of damage and changes that may be identified histologically and
described in morphological and optical terms (Table 1). Thus, for example, what we observe as a paste or a mixture of cooked food that includes resistant and modified starch “damaged in various ways” is a modification due to the retrogradation that occurs when boiled dough is left to rest, or a modification due to the melting by heating in a medium with low water content (Radley 1943; Johnson et al. 1990).

The two main physico-chemical processes we address in this paper, gelatinization and melting, are described in previous works. The effects of heating in a wet medium have been studied extensively, especially for the generation of baseline information on the properties of starch for industrial purposes. It leads to a non-reversible process called gelatinization (Radley 1943). For dry heating or for heating with low moisture content, we consider the “dry-cooking series”, toasting/roasting al rescoldo/cooking on embers/toasting/charring2, wherein the relative intensity of the heat and, therefore, the thermal alteration of the granules, increase from toasting to charring (Radley 1943; Babot 2003). Direct contact with the heat source differentiates roasting, charring and cooking al rescoldo from toasting and cooking on embers, with heat contact being indirect in the latter.

**Table 1.** Techniques, agents, physico-chemical processes and results, and morphological and optical modifications relevant to the study of human manipulation of starchy substances.

<table>
<thead>
<tr>
<th>Techniques (Modes of transformation)</th>
<th>Agents</th>
<th>Physico-chemical Processes and Results</th>
<th>Morphological and optical modifications</th>
<th>Taphonomic effects</th>
</tr>
</thead>
<tbody>
<tr>
<td>Heating in humid medium (chemical)</td>
<td>Liquid water + temperature (heat)</td>
<td>Gelatinization (Radley 1943). Irreversible – Gelatinized material</td>
<td>Loss of defined shape and structure manifested in the swelling, bursting, hilum opening and projection of grains, loss of birefringence (Babot 2003), and the presence of exudates (Messner and Schindler 2010), collapsed (Williams and Bowler 1982) or emptied grains (ghosts) (Radley 1943).</td>
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<tr>
<td>e.g.: boiling</td>
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<td>Rest of the boiled mass after cooking</td>
<td>Chemical forces</td>
<td>Retrogradation (Jacobson et al. 1997) – Retrograded material or paste</td>
<td>Formation of pastes (Biliaderis 2009), in which damage and modifications related to gelatinization can be observed</td>
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<tr>
<td>Heating in medium with low moisture content (chemical)</td>
<td>Temperature (heat) + liquid water (residual moisture)</td>
<td>Melting (Biliaderis 2009), Reversible – Melt or paste</td>
<td>Loss of the structure manifested in the formation of pastes (Biliaderis 2009), projection and/or opening of the hilum in grains, loss of birefringence (Babot 2003)</td>
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<tr>
<td>e.g.: airing or roasting</td>
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<tr>
<td>Hydration, dehydration in cold (chemical)</td>
<td>Liquid water (or relative humidity) + temperature (cold)</td>
<td>Hydration, dehydration (Radley 1943). Reversible – Hydrated or dehydrated material</td>
<td>Swelling (Williams and Bowler 1982) or shrinkage (Radley 1943) of the grains, increase or decrease of the relief (e.g., flat relief) and of the birefringence and the visibility of lamellae (Babot 2003)</td>
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<tr>
<td>e.g.: soaking, sun drying, airing</td>
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<tr>
<td>Absorption, percussion (mechanical)</td>
<td>Breakage mechanical forces</td>
<td>Disaggregation (Babot 2003), mechanical breakage, irreversible – Disaggregated material</td>
<td>Grains in different states of disaggregation or disarrangement, with presence of physical discontinuities such as fracture, truncation, cracking, surface damage such as depressions or dents (Babot 2003) and collapse (Williams and Bowler 1982)</td>
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<td>e.g.: milling, mashing and pounding</td>
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<td>Freezing (mechanical) e.g.: making chuno, a dehydrated form of potato obtained by freeze-drying (Parado and Pizarro 2008)</td>
<td>Solid water + temperature (cold to the point of freezing)</td>
<td>Dehydration, mechanical breakage, irreversible – Dehydrated material</td>
<td>Presence of physical discontinuities of the grains such as breakage and cracking, fragmentation (Babot 2003), collapse, emptying (Williams and Bowler 1982), cavities (ghosts) (Radley 1943), deep grooves and corrosion (Rechert 1913)</td>
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</tr>
<tr>
<td>Enzymatic attack (bio-chemical) e.g.: malting</td>
<td>Amylolytic enzymes</td>
<td>Amylolyisis, Irreversible</td>
<td>Loss of structure manifested in the presence of pitting (French 1984; Juan-Tresserras 1992), deep grooves and corrosion (Rechert 1913)</td>
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**PROCEDURES AND MATERIALS**

Tuber samples were obtained from traditional vending stalls in Jujuy Province (Argentina) in 2010 and 2012 and Villazón (Bolivia) in 2012. We obtained 19 ethno-varieties of potato (Solanum L.), 4 of oca (Oxalis tuberosa Mol.) and 3 of ulluco (Ullucus tuberosus Caldas), all from rural areas. Here we selected 10 varieties of potato, 2 of oca and 1 of ulluco, which were sampled and processed for culinary purposes. The purpose of using different ethno-varieties was to assess whether there were variations if different kinds of potatoes were subjected to different processing techniques, particularly since they could have been used in different traditional culinary practices (Castro 2008).

In all ethno-varieties, cortex and periderm were sampled separately from the vascular parenchyma and medullary tissue. Cortex and periderm were separated under the assumption that they were part of the “shell” that was eventually removed or peeled prior to consumption, while the parenchyma and the medullary tissue constitute the soft part intended for consumption. Three types of samplings were used for the fresh tubers to obtain reference material: 1) soft scrapings and histological cuts done freehand with a scalpel to obtain thin sections (Babot 2007), 2) diaphanization (a technique in which the tissue is treated with an oxidizing agent to make it transparent whilst retaining fabric), and 3) dry ashing (Piperno 1988). Some washing of the ash material was incorporated as well to assess tissue loss by this procedure. The first two sampling techniques provided references regarding tissue appearance and various elements of tissues, such as ergastic substances (starch, cellulose and crystalline calcium salts), of interest to archeology because they survive as microfossils. These procedures allowed us to observe intact tissue and fragments in different stages of disintegration, as well as free cells and intracellullar elements. Controlled diaphanization at temperatures below 10°C thinned the material for better observation.
Dry ashing allowed for recovery of the silicified tuber tissues. Due to the low proportion of silica produced by these plants, after the first trials, we avoided to wash ash samples because doing so results in a significant loss of material.

Cooking experiments included: a) boiling until cooking was complete, transforming the starch-rich parenchyma and medullary tissue by heating it to boiling in a humid medium with sufficient water; b) cooking al rescoldo (in direct contact with embers and ashes) until cooking was complete, transforming tissues by heating them at moderate temperature in a dry medium or in the presence of tuber residual moisture. Cooking enables consumption, increases energetic value of tubers and eliminates the causes of indigestion (Wandsnider 1997).

Boiling was performed using a stove; tubers were placed in a metal container filled with potable water to avoid contamination. Unpeeled tubers were boiled separately, reaching an average of 99°C during the time needed to soften the parenchyma and medullary tissue (between 15-20 minutes). Tubers started to lose their peel and stain water before completely cooked. Pink oca stained the water and started to lose their peel after 15 minutes in the boiling water. Nine minutes later it was completely cooked. White oca lost their peel at 17 minutes and started to smell like boiled sweet potato. Eight minutes later it was completely cooked. Ulluco took 19 minutes to begin staining the water, losing a minimal amount of peel. Two minutes later was completely cooked. Potatoes took 24 minutes to cook completely.

Cooking al rescoldo was done in a backyard fire. The fire was started with quebracho blanco (Aspidosperma quebracho-blanco Schltr.) wood. Red hot embers were produced and gradually covered with ash, at which point the charcoal was not as hot as in its red-hot state. A layer of embers and ashes was dispersed on the firebrick floor. Whole, unpeeled tubers were set over the embers and ashes and exposed to the heat source until the parenchyma and medullary tissue were softened, simulating the effect of placing tubers on the periphery of an active fire. This is a dry-cooking technique different from others such as toasting, cooking on the flame or on embers with a grate, or in ovens. Cooking on the flame involves direct contact of food with the heat source; placing a grate between the food and flame or red-hot embers separates food from the heat source. Toasting requires an intermediate object (e.g., a stone or vessel) to avoid direct contact with fire and buffer heat (Pazzarelli 2012). Cooking al rescoldo is a moderate-temperature procedure traditional in northwestern Argentina, where food is put into embers and ashes, coming into direct contact with the heat source. In our experiment, cooking al rescoldo took between 10-15 minutes depending on the type and size/form of the tuber and degree of exposure to the heat source. Ocas cooked most quickly, followed by ullucos and longer/thinner potatoes and, finally, thicker potatoes. During this process the tubers were not covered, so the periderm was carbonized where directly in contact with embers. Differences in the humidity of the cooked mass were observed in areas near carbonized and non-carbonized periderm. Oxygen and temperature might have fluctuated between the periphery and the center of the ember and ash layer within the range of 100-200°C. Quebracho blanco provided a strong, slow, constant and non-sparking charcoal combustion, with low ash production.

Boiled tubers and those cooked al rescoldo were wrapped in aluminum foil and refrigerated until sampling was complete. Sampling followed the same procedures used for fresh specimens (see above). Fresh, dry-ashed and cooked specimens were mounted on slides for viewing and photographing using a polarizing microscope (200X to 630X). Assemblage analysis focused on the various histological elements and intracellular particles present.

Characteristics recorded for fresh samples include: cell shape in two- and three-dimensions, arrangement of tissue cells (tiled, linear, concentric, etc.), color with and without polarizer, birefringence, cell size, presence of conduction elements and their characteristics, presence of cellulose in the cell walls; presence, kind, shape, color, presentation and location of calcium salt bodies within the tissues (isolated, grouped); and the occurrence of starch in the reserve parenchyma and the medullary tissue, and its disposition (isolated, in clusters, massively filling the tissue). The same characteristics were observed in food samples, with special attention to the features preserved, modified, and originated from culinary practices. In the case of native and modified starches, the specific variables summarized in the ICSN (2011) and previous studies were recorded (Korstanje and Babot 2007 and references therein; Henry et al. 2009). As mentioned, rather than a detailed description of individual starch grains, the presence of various attributes was confirmed (Table 1). Observations proceeded from questions such as: What is the general state of preservation of the intracellular tissues and starches in our samples? What is the degree of alteration? How does alteration vary with cooking techniques and their relative “aggressiveness”? Have the integrity, visibility, shape, size, color, spatial arrangement, optical properties or textures of tissues, cells and ergastic substances been altered? If so, how? Have any elements or distinctive features of fresh tissues disappeared? Have any new elements arisen as a result of processing? Finally, we compared the two sets of observations, which allowed us to generate expectations for documenting foods made from potatoes, oca and ulluco based on archaeological residues (Tables 2 and 3).
Table 2. Characteristic features of the periderm and cortex from tubers

<table>
<thead>
<tr>
<th>Aspect of tissue</th>
<th>Fresh specimens</th>
<th>Specimens cooked al rescoldo</th>
<th>Boiled specimens</th>
</tr>
</thead>
<tbody>
<tr>
<td>Turgid tissue, transparent, with mosaic-like fabric, tight or dense with straight-sided polygonal cells, occasionally rounded</td>
<td>General preservation of the original fabric of the tissue. Deformation and fragmentation by sections, with presence of net cracks and folds. Thermal alteration varies between complete carbonization and mild impact by heat. Dehydration and shrinkage: loss of relief and aging.</td>
<td>Preservation of original tissue structure by sections. Tissue fragmentation and intense cracks in the form of tears and joined cell fragments; presence of folds in tissue fragments. Distension, loss of turor of the tissue with increased apparent size. Thinning of the tissue and loss of relief.</td>
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<tr>
<td>Transparent tissue</td>
<td>Opaque tissue, slightly translucent or transparent according to the effect of heat.</td>
<td>Transparent and diaphanous tissue.</td>
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<tr>
<td>Non-birefringent and colorless tissue, or heterogeneously colored, depending on variety</td>
<td>Staining of the tissue in brown-reddish tones until reaching black in carbonized sections. Loss of the original color. In oca, all tissues appear birefringent under polarized light.</td>
<td>It can vary between color preservation, partial loss of color in parts of the tissue, or preservation of parts with watery or diluted coloration (brown-reddish tones). In oca, all the tissues appear birefringent.</td>
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<tr>
<td>Thickened cell walls with thick birefringent cellulose enrichments</td>
<td>The birefringence of cellulose is preserved or emphasized</td>
<td>The birefringence of cellulose is preserved or emphasized.</td>
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<tr>
<td>Sections of the tissue with intracellular deposits of amorphous silica (polyhedral or globular cells and vessel elements) stand out due to their higher profile</td>
<td>Colorless silicified tissues (silia phylostilis) with polyhedral crystals, sometimes attached to the tissue, or detached from it. This element highlights the fabric of the tissues</td>
<td>Colorless silicified tissues (silia phylostilis) with gray-black hues detached as a result of the thermal alteration. If not detached, they stand out from the surroundings due to their higher relief.</td>
<td></td>
</tr>
<tr>
<td>Polyhedral, tabular druses, crystal sand and subrounded discoid or globular particles. Scattered in the tissue or grouped in the intracellular space</td>
<td>Crystals very clearly observed in situ or detached from the tissue. They look unaltered; their birefringence is retained or accentuated. Eventual partial melting of the crystalline surface</td>
<td>Crystals very clearly observed in situ or detached from the tissue, located close to the cell walls or completely expelled from the tissue. They may look unaltered. Evidence of partial melting. New clumps of melted crystals are formed.</td>
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<tr>
<td>New products</td>
<td>- Massively carbonized by sections (outer areas of the tubers), and presence of micro-charcoal abundantly distributed throughout the tissue.</td>
<td>Micro-charcoals or carbonized tissues were not observed.</td>
<td></td>
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</tbody>
</table>

Table 3. Characteristic features of the parenchyma and the medullary tissue of fresh and boiled tubers, and tubers cooked al rescoldo. Note: CE= Cellulose; CR = Crystals; VR = Vessel elements; NP = New products.

<table>
<thead>
<tr>
<th>Aspect of tissue</th>
<th>Fresh specimens</th>
<th>Specimens cooked al rescoldo</th>
<th>Boiled specimens</th>
</tr>
</thead>
<tbody>
<tr>
<td>Turgid tissue, transparent and dense with polygonal cells with thin walls</td>
<td>Tissues remain structured; cells remain close but not attached to each other, or completely detached and with a globalular appearance. Dented and rough surface of cells, contracted, cracked, with lower relief. Although dehydrated, some degree of turor persists; some cells are partially or fully collapsed.</td>
<td>Tissues remain structured; cells remain close but not mutually attached, or are disaggregated and dispersed with a globalular appearance. Distension, loss of turor or firmness. Cells with cracks and fractures, partially or totally collapsed, depending on the preservation of their content.</td>
<td></td>
</tr>
<tr>
<td>Colorless to slightly brown-grayish tissue</td>
<td>Colorless to slightly brown-grayish tissue is maintained.</td>
<td>Colorless to slightly brown-grayish tissue is maintained. Tissues became diaphanous. A staining of the tissue may occur in watery or diluted brown-reddish hues.</td>
<td></td>
</tr>
<tr>
<td>Cells with thin cellulose walls</td>
<td>The cellulose thickenings are preserved, and their birefringence stands out in some cases. This element highlights the fabric of the tissues.</td>
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<td></td>
</tr>
<tr>
<td>Closely packed cells, filled with birefringent starch grains</td>
<td>The starch content is completely absent, giving the tissue an empty appearance, or the starch content is in different stages of the melting-gelatinization process. The birefringence varies depending on the situation. These are exuabales of amorphous starch within and outside of the tissues.</td>
<td>The starch content is completely absent, giving the tissue an empty appearance, or the starch content is in different stages of the melting-gelatinization process. The birefringence varies depending on the situation. These are exuabales of amorphous starch within and outside of the tissues.</td>
<td></td>
</tr>
<tr>
<td>Supernumerary simple starch grains</td>
<td>Amorphous starch masses dominate in the interior of the tissue. Occasional clumps or isolated individuals of modified starch with damage due to heating (open hilum, darkened sections, peripheral cracks and damage to the extinction cross and the birefringence). Little to no presence of unaltered resistant starch.</td>
<td>Amorphous starch masses dominate in the interior of the tissue. There is unaltered resistant starch as isolated and scarce grains, and grains of modified starch in individual gelatinization process of (open hilum, damage to the extinction cross and birefringence, longitudinal cracks) forming clumps associated with starch exudates.</td>
<td></td>
</tr>
<tr>
<td>Calcium crystals dispersed in the tissue</td>
<td>Crystals preserve their location in the tissue. Expulsion events of crystal clusters and microcrystals into the intercellular space</td>
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<td></td>
</tr>
<tr>
<td>Dense clusters of vessel elements, birefringent, silicified or not</td>
<td>They are preserved unaltered within or outside the tissue.</td>
<td>They are preserved unaltered within or outside the tissue.</td>
<td></td>
</tr>
</tbody>
</table>
RESULTS

Description of fresh tuber specimens

Periderm and cortex tissues are turgid and present a tight or dense mosaic-like fabric of polyhedral cells (Figure 1; Table 2). The tissue is non-birefringent—with the exception of periderm cell walls that are cellulose enriched—and either colorless or colored by sections (Figure 1a-b). The presence of calcium salt crystals (primarily calcium oxalate) is variable (Figure 1). They occur as polyhedra in oca and ulluco (Figures 1i-n, p-r); tabular druses and crystal sand in oca, ulluco and *Solanum* (Figure 3o-p); and flat, rounded or globular particles in *Solanum* (Figure 1h).

Parenchymal and fresh medullary tissue cells are polyhedral and globular, with thin and less cellulose-enriched membranes than the periderm. They are closely packed and contain numerous starch grains of the type described previously (e.g., Korstanje and Babot 2007 and references therein), completely filling reserve tissue (Figure 5u, Table 3). Abundant calcium crystals are observed in ulluco and oca parenchyma (Figure 7i-n); in *Solanum*, crystal abundance depends on the variety (Figures 4c, e and 5e-f). Dense clusters of birefringent conducting elements are observed, especially in the section corresponding to the vascular ring (Figures 5a-b and 7f-g).

Description of tuber specimens cooked *al rescoldo*

Carbonized or highly thermally altered sections of the periderm and cortex appear opaque to slightly translucent under the microscope. The internal structures and content of some tissues are difficult to observe, but in other sections, birefringent elements stand out well (Figures 2d-f, o-r and 3k-m; Table 2). Parts of the...
tissues that were not in direct contact with fire are less thermally altered (Figure 2a-c, h-l). Silicified parts stand out from the tissues around them due to a higher relief than the periphery. Occasionally, they appear as isolated polyhedral or globular siliceous particles (Figure 3k, g-h). Some deformation, fragmentation, crack and folds are observed, but this kind of damage is less pronounced among tubers cooked al rescoldo than among those that were boiled (Figures 2c-d, p). Tissues significantly affected by heat become brittle, dehydrated and shrunken. They are stained reddish-brown to black (in carbonized areas), which accentuates surface irregularities regardless of the original periderm color. Micro-charcoals are abundant throughout the tissue; some of them are still recognizable in the inner layers (Figures 2a-c, h-i, k-n, g and 3d-f, j, q). Typically, it is possible to observe isolated clumps, or clumps attached to the tissue, within an unctuous, reddish-brown matrix, against which crystals and occluded micro-charcoals stand out. This pattern occurs in both extensive and restricted sections of tissue (Figures 2g-h, k-n and 3d-f, g-h, q). Cell wall cellulose birefringence is retained or increased. In the cells of oca, birefringence is widespread, not restricted to cellulose accumulation (Figure 3n, r). Conducting tissues remain unaltered. Calcium crystals also remain largely unaltered, although their birefringence stands out (Figures 2r and 3n, p). In a few cases, partial melting of the druses’ crystallites is observed (Figure 3i).

Parenchyma and medullary tissues may retain their original structure. In other cases, cells remain close without completely attaching to each other, or are entirely detached. Their surfaces are dented, rough, contracted, cracked, and have lower relief than fresh internal tissues (Figures 4-5; Table 3). Despite being

Figure 2. Appearance of the periderm and cortex cooked al rescoldo of: a-o) Solanum sp. and p-r) Ullucus tuberosus. a-c) Desireé, d-i) Abajeña, j-n) Malcacha, o) Huaico potato, p-r) Ulluco. The contiguous twin micrographs correspond to views with parallel (left) and crossed (right) nicols of the same sample.
dehydrated, they retain a degree of turgor, though some are partially or fully collapsed (Figure 4a). Starches are in the process of melting/gelatinization (Figures 4e-g, j, o-r and 5a-p, s-t) or absent, giving the tissue the appearance of being almost totally empty (Figure 4c, i). Clumps (Figure 5s-t) and isolated grains of modified starch with damage from dry-cooking are occasionally observed (Table 3) (Figure 5g-j, m-n), and unaltered, resistant starch is scarce or virtually absent (Figure 5c, k-l). The birefringent cellulose of cell walls is retained. Although the crystals can maintain their location within the tissues (Figure 4c, e), clusters of multiple crystallites are often found in the intercellular spaces due to their expulsion from the interior of the cells (Figure 5e-f). Vessel elements are preserved unaltered, with high birefringence, appearing isolated or in clusters (Figure 5a-b).

Description of boiled tuber specimens

Periderm maintains its configuration and original morphology in some portions, although there is more fragmentation of the tissue than when cooking al rescoldo (Figure 6, Table 2). Highly fragmented, torn or folded sections are observed (Figures 6c, m-r). In general, boiling causes distension, loss of tissue turgor, an increase in size, and loss of relief. Tissues become transparent and noticeably thinner (Figure 6a-f). Colorless to gray silicified tissue and isolated polyhedral cells can be seen. If not detached, they stand out from surroundings tissues due to their higher relief (Figure 6e-f). A watery or diluted pigmentation remains unaltered or lost by sections in periderm cells (Figure 6e-f, l, m). Brown to reddish staining can appear by sections (Figure 6p). Birefringence of thickened cell wall cellulose is retained (Figure 6j, l, n, r). In pink oca, this phenomenon is widespread, as with cooking al rescoldo (Figure 6g). Calcium crystals may be unaltered and retain their position in the tissue (Figures 6m, q-r) or be released into the inter- or extracellular space (Figure 6k, q-r).
Occasionally, crystals are found melted, formed into clumps in which the individuals are not discernable (Figure 6j-k). No micro-charcoals or carbonized tissues are observed in the boiling specimens.

Globular or polyhedral parenchymal and medullary tissue cells with rounded edges are present in clusters -although not attached to each other-, or disaggregated and dispersed, and there are empty net spaces between them (Figures 7-8, Table 3). In general, there is a loss of turgor, and cells appear more distended than when cooked *al rescoldo* or fresh. In sections more affected by boiling, cells appear collapsed due to the partial or total loss of starch (Figure 7a-e and 8j-k). There are cracks and fractures in the cell walls; the surfaces are rough or dented, resembling those of tissues cooked *al rescoldo* (Figures 7a, e and 8a, r). Increased transparency and areas with watery or diluted reddish-brown staining are observed (Figure 7i and 8b-e). Some nodular particles in the same tones are attached to tissue (Figures 7i-j and 8a, c). Discontinuous masses of gelatinized, amorphous starch are observed in the interior or exterior of the tissues (Figure 8g-h, j-k). Resistant starch occurs as scarce, isolated individual grains (Figures 7c-e, h and 8g-i, l). Modified starch is in the process of gelatinization (Table 3), isolated (Figure 8n-o, r-s), forming clumps of individuals that have lost their original shape (Figures 7a-b and 8p-q), and associated with starch exudates within the tissues or expelled from them (Figure 8j-k). The survival of resistant starch and of partially gelatinized individuals is greater in the close proximity of cortex and periderm than in the parenchymal and medullary tissue. Vessel elements within the vascular parenchyma preserve their birefringence and structure forming dense packages with a persistent linear orientation (Figures 7i-g and 8a, c-d, l). In some cases the crystals maintain their original location within the tissues (Figures 7i-j and 8f-h) or are expelled from them to the inter- or extra-cellular spaces (Figures 7k-n and 8m). Some are partially solubilized (Figure 7n).
DISCUSSION AND CONCLUSIONS

Experimental results indicate the potential for differentiating two cooking techniques: heating in humid and dry (or low moisture) environments. Based on previous studies, we expected these techniques to be similar in their effects on tubers since both involve the same agents of modification: heat and moisture. In the case of boiling, water is abundant in the cooking environment. By contrast, water is scarce but not totally absent in the “dry-cooking series” techniques. Among the latter, there is an increase in both the temperature gradient and exposure of the matter to temperature. While there are similarities in the effects of the cooking techniques reviewed, the abundance of water clearly leads to gelatinization, while its absence leads, in theory, to melting due to loss of starch humidity, although some degree of gelatinization is also possible.

Generally speaking, certain reactions are common among tubers cooked al rescoldo and those heated in a moisture-rich medium, the latter being a more aggressive transformation technique that compromises the integrity of plant tissues and particles to a corresponding degree. However, damage to the morphology and optical properties of starch caused by cooking al rescoldo is greater than that caused by toasting, as reported elsewhere (Babot 2003). Such damage includes denaturation and a very low specimen count. In the case of cooking al rescoldo, this is influenced by the fact that plant matter is in direct contact with the heat source and subjected to higher temperatures than in toasting. Although both groups of cooking techniques analyzed here cause similar modifications and damage, our results indicate that certain characteristics can be used to differentiate them and also, to distinguish cooked from fresh samples. For them to be applicable in archaeological contexts, we need to evaluate the survival over time of the characteristics reported here.

The periderm and cortex generally survive and retain their original fabric. They react differently to cooking al rescoldo (characterized by reddish-brown staining of tissues; unctuous clumps –isolated, or attached to tissue– with crystals and occluded micro-charcoals; contraction and tissue aging) versus boiling (characterized by diaphanous and thin material, partial preservation of the original watery color, tissue distension). However, the natural color of tuberous plants’ shell could be confused with the effects of cooking on color and mistakenly attributed to anthropogenic modification. In the external tissues, massive carbonization occurs by sections, and the presence of micro-charcoals distributed throughout the tissue occurs only with cooking al rescoldo.

Figure 5. Starch grains, calcium crystals and conduction elements in parenchyma and medullary tissue cooked al rescoldo of: a-n) Solanum sp. and o-t) Oxalis tuberosa. Appearance of the fresh parenchyma and medullary tissue of Solanum sp. (Huaico potato) is in (u). a-c) Abajeña, d) Huaico potato, e-f) Desireé, g-n) Pentaoca, o-t) White oca. The contiguous twin micrographs correspond to views with parallel (left) and crossed (right) nicols of the same sample.
Results obtained for the most external tissues are important for supplementing the data regarding the parenchyma and medullary tissue, where the starches are primarily located. While the latter are distinctively altered by particular processes and, therefore, play an important role in identifying anthropogenic processes, under certain circumstances they can be denatured or absent, making information about the periderm and cortex particularly relevant. Additionally, the periderm and cortex tissue can be preserved in archaeological contexts, both inside cooking containers where whole tubers were cooked and as discarded peelings (particularly those of bitter tuber varieties). The parenchyma and medullary tissue suffer greater damage than the periderm and cortex. Contraction occurs in cooking al rescoldo, whereas distension is observed with boiling. In the latter case, a diluted reddish-brown staining of the interior tissues may appear, as well as the occurrence of some nodular particles in the same tones. Observed starch alterations are consistent with those reported elsewhere, but a higher intensity of the modifications was verified when cooking al rescoldo, with respect to what was previously documented for toasting (Babot 2003). Despite the poor survival of starch as isolated, identifiable grains, both inside or outside the cooked mass, resistant starch specimens are more common in boiled tubers than in those cooked al rescoldo. Both techniques result in modified starch grains. Calcium crystals, conducting tissues and cellulose survive both techniques. Melted or diluted crystals and crystals expelled from the tissues generally indicate manipulation in a medium with heat. Nonetheless these changes are more intense in the presence of water. Wet cooking is also more aggressive to cellulose than cooking al rescoldo. After cooking, cellulose allows excellent visualization of the original tissue fabric. Silica deposits, located in conduction and peridermal tissues and usually of little diagnostic value when taken alone, are not affected by the processes studied; on the contrary, they tend to stand out from the rest of the tissue, either in situ or detached from it.

Figure 6. Appearance of the boiled periderm and cortex of: a-d, g-l) Oxalis tuberosa, e-f) Solanum sp. and m-r) Ullucus tuberosus. a-d) White oca, e-f) Pentaoka, g-l) Pink oca, m-r) Ulluco. The contiguous twin micrographs correspond to views with parallel (left) and crossed (right) nicols of the same tissue.
No significant differences in changes produced by the processing techniques considered here were observed between the three types of tubers and their varieties. Thus, the expectations generated should apply to a large number of tuberous plants and potentially other resources as well.
Figure 8. Appearance of boiled parenchyma and medullary tissue of Oxalis tuberosa. a-q) White Oca, r-s) Pink Oca. The contiguous twin micrographs correspond to views with parallel (left) and crossed (right) nicols of the same tissue.
Finally, we must acknowledge that the daily character of kitchens and food preparation methods suggests intermediate instances that turn black and white results from controlled experiments into grays which cannot be ignored. For instance, tubers and other items might be cooked in their own liquid, without additional water, which, strictly speaking, is neither boiling nor toasting. Such exceptions and the integrity of the archaeological remains, determine the “grain” of the inferences that can made using experimental data. Despite these caveats, experimental work continues to contribute positively to our understanding of past practices.

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NOTES
1.- Resistant starch is defined as “[...] starch that is not digested in the small intestine of humans (Champ 2004; Biliaderis 2009). Useful term for explaining why some starches survive and others do not” (INSC 2011). In this paper, we used the term generically to refer to any starch grain preserved virtually unchanged.

2.- We consider charring as a special cooking technique used in ritual performances (e.g., in feeding Pachamama) (Pazzarelli 2012).