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### Abstract

The development of non-destructive methodology based on Near-Infrared Reflectance spectroscopy (NIRs), Hyperspectral Imaging (HSI), Nuclear Magnetic Resonance (NMR) and Magnetic Resonance Imaging (MRI) techniques to determine quality characteristics of fresh meat has been reviewed in this study, which has been mainly focused on researches published in the last decade. This review has put special attention on the instrumentation, data acquisition and main applications of each technique, finding a wide variety of possibilities of systems and methodologies as well as evidences of accurate and promising results. Most analysed samples have been pork and beef, followed by lamb and chicken, while there are few studies on fresh meat from rabbit and duck. The evaluation of the methodology exposed in the revised articles has been carried out in an experimental way but lacking real application in the meat industry. For that, these non-destructive techniques should be improved, especially regarding the speed, price and influence of external factors.

**Keywords** Near-Infrared Reflectance spectroscopy; Hyperspectral Imaging; Nuclear Magnetic Resonance; Magnetic Resonance Imaging; fresh meat; non-destructive analysis.

**Taxonomy** Meat, Analytical Method

**Corresponding Author** Trinidad Perez-Palacios

**Corresponding Author's Institution** University of Extremadura

**Order of Authors** Teresa Antequera, Daniel Caballero, Silvia Grassi, Bethany Uttaro, Trinidad Perez-Palacios

**Suggested reviewers** Xinglian Xu, Kate Kemsley, Yael Vodovotz, Da-Wen Sun

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Figure 2.docx [Figure]

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Trinidad Pérez Palacios  
Research Institute of Meat and Meat Product  
University of Extremadura

Cáceres, december 2019

Dear editor,

Please find enclosed the second revised version of the manuscript entitled “Use of Non-destructive Techniques for the Quality Evaluation of Fresh Meat: A Review” to be considered for publishing in *Meat Science*.

The present manuscript is a review aimed on published studies about the use of NIRs, HSI, NMR and MRI to determine the quality of fresh meat, paying special attention to the methodologies used (instrumentation, data acquisition and data analysis) and their applications.

We confirm that this work is original and has not been published elsewhere nor it is currently under consideration for publication elsewhere. We state that there are no known conflicts of interest associated with this publication. We declare that the manuscript is prepared strictly according to the Journal format as provided in the instruction to authors.

Please address all correspondence concerning this manuscript to me at [triny@unex.es](mailto:triny@unex.es).

Thank you for your consideration of this manuscript.

Yours sincerely,

Trinidad Pérez Palacios  
Research Institute of Meat and Meat Product  
University of Extremadura  
10003-Cáceres  
Spain

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3 1 **TITLE**  
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6 3 Use of Non-destructive Techniques for the Quality Evaluation of Fresh Meat: A Review  
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10 5 **AUTHORS AND AFFILIATIONS**

11 6 Teresa Antequera <sup>1</sup>, Daniel Caballero <sup>1,2τ</sup>, Silvia Grassi <sup>3</sup>, Bethany Uttaro <sup>4</sup>, Trinidad Perez-Palacios  
12  
13 7 <sup>1,\*</sup>.

14 8  
15  
16 9 <sup>1</sup> Food Technology Department, Meat and Meat Products Research Institute, University of  
17  
18 10 Extremadura, Av. Ciencias S/N, ES-10003, Cáceres, Spain.

19 11  
20  
21 12 <sup>2</sup> Chemometrics and Analytical Technology Section, Food Technology Department, Faculty of  
22  
23 13 Science, University of Copenhagen, Rolighedsvej 26, DK-1958, Frederiksberg C, Denmark. <sup>τ</sup>Present  
24 14 address.

25 15  
26  
27 16 <sup>3</sup>Department of Food, Environmental and Nutritional Sciences, Università degli Studi di Milano, Via  
28  
29 17 Mangiagalli 25, IT-20133, Milano, Italy.

30 18  
31  
32 19 <sup>4</sup>Lacombe Research and Development Centre, Agriculture & Agri-Food Canada, 6000 C and E Trail,  
33  
34 20 Lacombe, Alberta, Canada, T4L 1W1.

35 21  
36  
37 22 **\*CORRESPONDING AUTHOR**

38 23 Trinidad Pérez Palacios

39  
40 24 e-mail address: triny@unex.es

41  
42 25 Telephone: +34927251052 Ext. 51052

43 26 Fax: +34927257110  
44  
45 27

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29 **ABSTRACT**

30 The development of non-destructive methodology based on Near-Infrared Reflectance spectroscopy  
31 (NIRs), Hyperspectral Imaging (HSI), Nuclear Magnetic Resonance (NMR) and Magnetic Resonance  
32 Imaging (MRI) techniques to determine quality characteristics of fresh meat has been reviewed in  
33 this study, which has been mainly focused on researches published in the last decade.  
34 This review has put special attention on the instrumentation, data acquisition and main applications  
35 of each technique, finding a wide variety of possibilities of systems and methodologies as well as  
36 evidences of accurate and promising results. Most analysed samples have been pork and beef,  
37 followed by lamb and chicken, while there are few studies on fresh meat from rabbit and duck.  
38 The evaluation of the methodology exposed in the revised articles has been carried out in an  
39 experimental way but lacking real application in the meat industry. For that, these non-destructive  
40 techniques should be improved, especially regarding the speed, price and influence of external  
41 factors.

42  
43 **KEY WORDS**

44 Near-Infrared Reflectance spectroscopy; Hyperspectral Imaging; Nuclear Magnetic Resonance;  
45 Magnetic Resonance Imaging; fresh meat; non-destructive analysis.

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121 47 **1. INTRODUCTION**  
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123 48 Meat and meat products are highly appreciated, which is principally due to their sensory properties, but  
124 49 their nutritional composition is also relevant. Evaluation of meat and meat products by means of many  
125 50 physico-chemical, sensory and microbiological analysis is the subject of the industry, laboratories  
126 51 and researches, in order to i) guarantee the global quality of these kind of food, ii) assure that they  
127 52 accomplish the legal requirements and iii) give response to the demands of the consumers.

131 53 Physico-chemical characteristics, such as pH, colour, water activity, content of moisture, lipids,  
132 54 protein or salt, and sensory attributes are the most demanded parameters to be determined. However,  
133 55 traditional methodology applied to carry out these analyses require the destruction of the meat pieces  
134 56 to correctly take a representative sample. These techniques also consume solvents, take long times  
135 57 and are tedious. Besides, a trained panel is needed to evaluate the samples in the case of the sensory  
136 58 analysis (Pérez-Palacios, Caballero, Caro, Rodríguez, & Antequera, 2014).

141 59 As response to these drawbacks, several studies have been developed to evaluate the capability of  
142 60 different techniques based on images and/or spectra to analyse quality parameters of meat and meat  
143 61 products in a non-destructive way, with the final aim of proposing the evaluated techniques as  
144 62 alternative and/or complementary to the traditional methods. Couple-Charges Devices cameras  
145 63 (Cameras CCD), computed tomography (CT), Near-Infrared Reflectance spectroscopy (NIRs),  
146 64 Hyperspectral Imaging (HSI), Nuclear Magnetic Resonance (NMR) and the Magnetic Resonance  
147 65 Imaging (MRI) are some of these proposed techniques.

152 66 The present manuscript was focused on reviewing the published studies at evaluating the use of NIRs,  
153 67 HSI, NMR and MRI to determine the quality of fresh meat, paying special attention to the  
154 68 methodologies used (instrumentation, data acquisition and data analysis) and their applications.

157 69 Thus, this review article has been organized as follow: section 2 exposes the scientific searches that  
158 70 have been carried out; section 3 is about the instrumentation of each of the focused technologies;  
159 71 section 4 deals with the procedure for the data acquisition and methods for the data analysis; section  
160 72 5 presents the latest applications of NIRs, HSI, NMR and MRI techniques for the analysis of fresh  
161 73 meat; section 6 discusses the advantages and disadvantages of these non-destructive methodologies;  
162 74 and section 7 summarizes conclusions for each technology and points out some future goals.  
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168 76 **2. SCIENTIFIC SEARCHES**  
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170 77 The searches on the scientific literature were carried out by using *Scopus*, *Science Direct* and *Web of*  
171 78 *Science*. The key words used were “meat” in combination to “Near Infrared Reflectance  
172 79 spectroscopy” or “NIR” or “NIRs” or “Hyperspectral Imaging” or “HSI” or “Nuclear Magnetic  
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180 80 Resonance” or “NMR” or “Magnetic Resonance Imaging” or “Magnetic Resonance Images” or  
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182 81 “MRI”, and the areas of interest were limited to Food Science and Technology and Science  
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184 82 Technology Other Topics. In this way, around 140, 200, 170 and 150 documents were retained for  
185 83 NIRs, HSI, NMR and MRI, respectively. Then, only the publications that fulfilled the research aim  
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187 84 (research papers at evaluating the use of NIRs, HSI, NMR, or MRI to analyse fresh meat) in  
188 85 approximately the last ten years were selected (25 + 31 + 9 + 10 documents, respectively) to be  
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190 86 exhaustively analysed.

### 191 87 192 193 88 194 195 89 **3. INSTRUMENTATION**

#### 196 90 **3.1. NIRs**

197  
198 91 The NIRs method works in the region of the electromagnetic spectra from approximately 780 nm to  
199 92 2500 nm, although quite often VIS wavelengths (~400-700 nm) are also collected. Generally  
200 92 speaking, there are three formats of NIRs equipment – benchtop, portable, and miniature, all of which  
201 93 are being used in fresh meat research. Benchtop equipment, such as the FOSS family of analysers,  
202 93  
203 94 may be operated under controlled environmental conditions (temperature, humidity, airflow) and are  
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205 95 often used in laboratory situations, usually with ground meat. Portable equipment such as the ASD  
206 96 range of spectrometers may or may not be more compact, but are more tolerant to environmental  
207 96  
208 97 operating conditions, and are often used on intact meat. With the advent of microelectromechanical  
209 98  
210 98 systems (MEMS) and micro-optical electromechanical systems (MOEMS), the numbers and types of  
211 99  
212  
213 100 miniature and micro-NIR spectrometers has been increasing in recent years. Some companies  
214 101 currently offering miniature equipment that has been tested on meat include Telspec Inc., Consumer  
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216 102 Physics (SCiO™), and Viavi Solutions Inc..

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218 103 In the field of meat analysis, several labs have reported custom made equipment. Gentilin *et al.* (2016)  
219  
220 104 and Zhang, Peng, Zhao, and Sun (2017) have both pursued prediction of moisture content, with the  
221 105 former developing a hardware/software platform with rapid response and a high signal to noise (S:N)  
222  
223 106 ratio, and the latter developing equipment for use on the fresh meat conveyor line which, once  
224  
225 107 triggered, will automatically make multiple measurements, dynamically display and then preprocess  
226 108 spectra, then generate predictions. Piao, Okura, and Irie (2018) reported on small portable equipment  
227  
228 109 developed by them, then manufactured to be used under cold conditions for the prediction of fat  
229  
230 110 quality, successfully transferring calibrations from a master unit to up to 5 slave units.

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232 111 Moving towards miniaturization and/or reduced costs, Kandpal, Lee, Bae, Lohumi and Cho (2019)  
233 112 report the development of a system based on a monochrome camera and multiple LEDs spanning

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113 458-950 nm for rapid evaluation of fresh meat quality. Some researchers have been working with the  
114 reduction of full spectra wavelengths to only those of importance to meat. Habib and Ullah (2016)  
115 performed computer simulation and testing of NIRs bandpass filters, specifically for regions  
116 important to meat classification or meat quality determination, that block wavelength contamination  
117 which would otherwise occur from the Distributed Bragg Reflectors (DBR) stopband. In an  
118 alternative solution to the DBR stopband restrictions, Ullah, Butt, Fomchenkov, and Khonina (2016)  
119 has reported filters composed of indium phosphide (InP) and air-gap layers to replace the alternating  
120 InP/SiO layers of the of DBRs of the Fabry-Pérot filters in miniaturized spectrometers in the 930-  
121 1450 nm range which is commonly used for meat.

### 3.2. HSI

123 The HSI technique is based on collecting and processing information from across the electromagnetic  
124 spectra. Hyperspectral devices merge the whole advantages of spectroscopy with the advantages of  
125 spatial information of surfaces (Amigo, 2020). Thus, they can be defined as fast and non-destructive  
126 instruments able to characterize sample from a chemical (chemical interactions) and physical  
127 (properties of the surface) point of view, after the proper data collection and analysis.

128 Nowadays, there are different commercial hyperspectral cameras, and their technology is  
129 continuously advancing in terms of acquisition speed and spectral/spatial resolution.

130 Even if there are different commercial HSI instruments, their setup is based on the interaction light-  
131 matter, or better, on the interaction between photons (with a specific energy and trajectory) with  
132 molecules of the sample under study (Weisskopf, 2010). Thus, the instruments are mainly composed  
133 by a light source, a set of optical lenses, a wavelength dispersive device and a detector.

134 The photons are emitted by the light source (halogen lamps, LEDs or lasers) which should emit in the  
135 spectral range of interest, with high energy, without effects on the sample and guaranteeing  
136 illumination homogeneity (Amigo & Grassi, 2020). The latter is often achieved by light sources  
137 forming a 45° angle with the sample.

138 Most of the HSI systems take advantages of the reflection and transmission phenomenon to collect  
139 information about the chemical and physical properties of a samples. Indeed, when a photon striking  
140 a molecule, its energy is absorbed only if it has the same vibration frequency of the electron of the  
141 molecule. In case the frequencies of the photon waves are not the same than the natural frequencies  
142 of the molecules of the sample, they are reflected or transmitted. They are reflected in case molecules  
143 are “opaque” to those frequencies, which means that, when interacting with the matter, the electrons  
144 of the molecules on the sample surface vibrate just for short periods before allowing the photon with  
145 different energy to arrive to the detector. Depending on the properties of the molecules, as well as the

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146 physical characteristics of the sample, the photon can be reflected in a specular mode (the same angle  
147 than the incident) or in a scattered mode (different angle than the incident). The photons are  
148 transmitted in case they hit “transparent” molecules, a.k.a. the vibrations pass through the matter and  
149 are reemitted on the opposite side of the object and reach the detector (Abbott, 1999).

150 After the emission, before or after interacting with the sample, the radiation is dispersed into narrow  
151 wavelength bands which are recorded individually by the detector. Among the wavelength dispersive  
152 devices, the most common are variable and tunable filters, imaging spectrographs and Fourier-  
153 Transform spectrometers (Amigo & Grassi, 2020).

154 At the end of the acquisition procedure the detector collects the coming incident light and convert it  
155 into electrical signals. Even if present in the market with different variation in the architecture and  
156 composition, mainly charge-coupled devices (CCDs) and the complementary metal-oxide-  
157 semiconductor detectors (CMOS) are implemented in HSI cameras (Amigo & Grassi, 2020).

### 3.3. NMR

159 In NMR, and also in MRI, the signal is produced by excitation of the nuclei of the samples with radio  
160 waves into nuclear magnetic resonance. High (HF) or low field (LF) NMR systems can be used for  
161 analyzing fresh meat. The LF-NMR systems generate magnetic field between 0.15 and 0.50 T. These  
162 systems are cheaper than the HF-NMR ones and do not have maintenance costs. However, their  
163 obtained spectra are of lower quality and sensitivity than those from HF-NMR systems, which  
164 generate magnetic field higher than 2T. However, HF-NMR systems are very expensive and require  
165 high maintenance costs, since they need to be cooled with helium or liquid hydrogen (Feig, 2011).  
166 The radiofrequency (RF) for LF-NMR systems is between 30 MHz and 100 MHz, and in HF-NMR  
167 system, it is higher than 100 MHz. Besides the type of magnetic field, there are several types of  
168 antennas oriented to excite the spin of different isotopes of chemical elements (Hornak, 1997), being  
169  $^1\text{H}$  the most used. And the spectra can be weighted on two different relaxation time: T1 and T2. T1  
170 spin relaxation time (spin-lattice relaxation time) is the time from the longitudinal magnetization of  
171 the molecule of the sample from which the spectra will be obtained until their equilibrium value has  
172 been reduced by an “e” factor. T2 spin relaxation time (spin-spin relaxation time) describes the same  
173 process for the transverse magnetization. In this case, T2 is always smaller than T1, although both  
174 processes happen simultaneously. As acquisition sequence, most studies have applied the Carr-  
175 Purcell-Meiboom-Gill sequence (CPMG) that allows measuring relaxation times intensity of any  
176 nucleus (McIntosh, 2013).

### 3.4. MRI



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178 The MRI acquisition in fresh meat has been carried out by using HF and LF scanners. HF MRI  
179 scanners offers images with a high quality, however, they are very high-priced and involve high  
180 maintenance costs (Feig, 2011). The LF MRI scanners are cheaper and do not require maintenance  
181 costs, but they have lower S:N ratio and, consequently, their images are of lower quality than those  
182 from the HF MRI ones. Among the obtained publications on MRI for this revision, most of the studies  
183 carried out in meat with HF MRI scanners have analysed dry-cured hams, there being a low number  
184 of works focused on fresh meat. However, the use of LF MRI scanners to analyse fresh meat is higher  
185 and it has experimented an increase in the last years.

186 For MRI, besides the type of magnetic field (HF or LF), the antenna, the relaxation time and the  
187 acquisition sequence are critical parameters for the image acquisition. The antennas that excite <sup>1</sup>H are  
188 the most used in meat (Pérez-Palacios, Caballero, Antequera, Durán, Ávila, & Caro, 2017).  
189 Nevertheless, some studies have also been carried out with antennas that excite <sup>23</sup>Na (Hansen, Van  
190 Der Berg, Ringgard, Stodkilde-Jorgensen, & Karlsson, 2008; Vestergaard, Risum, & Adler-Nissen,  
191 2005), which are more specific for determining the salt content in meat and meat products. T1 is the  
192 spin relaxation time usually applied in most studies (Pérez-Palacios *et al.*, 2017). Different acquisition  
193 sequences can be selected, such as Multi-Slice (MSE), Inverse-Recovery (STIR), Gradient Echo  
194 (GE), Spin Echo (SE) or Volumetric (T3D), among others. Moreover, the echo time (TE) and the  
195 repetition time (RT) must be set for the image acquisition. TE is the time from the center of the radio  
196 frequency pulse to the center of the echo, and RT represents the length of time between corresponding  
197 consecutives series of pulses and echoes (Hendrick, 2005). SE sequences are characterized by long  
198 TE (around 20-30 ms) and short RT (less than 800 ms) and present a high S:N, while GE sequences  
199 have short TE (12 to 20 ms) and long RT (between 1200 and 2500 ms), with a lower S:N than SE.  
200 T3D sequences are a special GE sequences with 3D reconstruction, with a similar TE than GE and a  
201 very short RT (between 30 and 100 ms) (Ávila, Caballero, Antequera, Durán, Caro, & Pérez-Palacios,  
202 2018; Caballero *et al.*, 2017a). SE is the most used sequence acquisition in the MRI studies of meat,  
203 while GE and T3D have been recently proved (Pérez-Palacios *et al.*, 2017).

204 Coils are also supporting elements to receive the MR signal. They have to be placed as near as  
205 possible to the area to be scanned. Coils can be classified as a function of their shape, volume or  
206 surface coils, or of their technology, linear or quadrature coils. In the case of musculoskeletal systems,  
207 the use of volume and quadrature coils, which surround almost completely the scanned area and  
208 receive the signal through two orthogonal channels, respectively, is the optimum combination. The  
209 size of the coil is also considered in the MRI studies in meat, using body coils for hams and carcass  
210 and hand/wrist, head or knee coils for smaller samples such as pork loins or chicken breast (Perez-

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416 211 Palacios, Antequera, Durán, Caro, Rodríguez, & Palacios, 2011; Bernau *et al.*, 2015; Frelka *et al.*,  
417 212 2019).

## 419 420 213 **4. DATA ACQUISITION AND ANALYSIS**

### 421 422 214 **4.1. NIRs**

423  
424 215 Successful and reliable data acquisition must take into account equipment, environment, and sample  
425 216 condition. Firstly, a measurement method appropriate to the sample preparation must be chosen, then  
426 217 the equipment be calibrated. The most common measurement methods, shown in Figure 1, are  
427 218 reflectance (R) where most of the recorded light is reflected directly from the illuminated surface, or  
428 219 transmittance (T) where the light which travels through the sample is recorded. Both can be converted  
429 220 to absorbance for data analysis following a log transformation ( $\log 1/R$  or  $\log 1/T$ ; Cortés, Blasco,  
430 221 Aleixos, Cubero, & Talens, 2019); interactance, records the light reflected from the interior of the  
431 222 sample and often uses fiber optics for illumination and detection as they can be placed in direct contact  
432 223 with the sample; and transflectance, in which light travels through the sample, encounters a reflector,  
433 224 then travels back through the sample before detection (Alander, Bochko, Martinkauppi, Saranwong  
434 225 & Mantere, 2013). Reflectance and interactance require thick samples to accommodate the long  
435 226 travel distances of NIRs wavelengths, while transmittance and transflectance require thin samples to  
436 227 facilitate sufficient light travel through the sample. In all cases, meat fiber direction should be  
437 228 consistent among samples. Long-standing do's and don'ts for equipment and sampling (Williams &  
438 229 Norris, 2004) include keeping equipment temperature constant, and operating under similar relative  
439 230 humidity during each data collection session to keep the amount of noise in the spectra similar for all  
440 231 samples. Samples may be homogenized, minced or ground, or intact, and of a thickness appropriate  
441 232 to the measurement method. Sample temperature must be consistent to prevent alterations to the  
442 233 spectral baseline and position of absorption bands. If samples have been dehydrated, measures must  
443 234 be taken to ensure moisture level is similar in all before reading. Sampling must be done to truly  
444 235 represent the subject, thus intact samples may require multiple reading locations; fiber orientation  
445 236 may be of importance for fresh intact meat samples. A recent review by Pasquini (2018) fully  
446 237 addresses NIRs fundamentals, chemometrics, and instrumentation while Xu, Xie, and Ying (2019)  
447 238 focus on identifying error sources.

448 239 The most common approach to analysis of NIRs spectra is pre-processing followed by processing.  
449 240 The pre-processing may be accomplished in a number of different ways and generally involves  
450 241 removal of noise at spectra extremes, further noise reduction over the remaining wavelengths  
451 242 (smoothing), and scattering and slope corrections. The lowest S:N is usually found at the extremes of  
452 243 the spectra, which is often removed simply by clipping the spectra. In the remaining body of the

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475 244 spectra where S:N is higher, correction of scattering due to physical structure variations of the sample  
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477 245 is usually performed with Multiplicative Scatter Correction (MSC), while Standard Normal Variate  
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479 246 (SNV) is performed to correct spectra slope.

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481 247 Reduction of random noise is often achieved by smoothing with a least squares polynomial fit, or in  
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483 248 combination with 1<sup>st</sup> or 2<sup>nd</sup> derivative transformation, a method established by Savitzky and Golay  
484 249 (1964), although some authors have chosen to use Extended MSC (EMSC), such as Andersen,  
485  
486 250 Veiseth-Kent & Wold (2017) who explored the effect of pH-related changes in extracted pork  
487 251 myofibrils as part of an equipment comparison study. Prevolnik Povse *et al.* (2017) and Soladoye *et*  
488  
489 252 *al.* (2018) both used SNV and Detrend (SNVD), the former for quality prediction in intact pork fat  
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491 253 layers and homogenized pork lean (n=56-130), and the latter for overall pork belly firmness from  
492 254 intact pork fat layers and lean (n=198).

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494 255 By far, the most common processing approach for data exploration is principle component analysis  
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496 256 (PCA) followed by regression or classification algorithms depending on the purpose of the study. In  
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498 257 the case of regression, which is quantitative analysis, partial least squares (PLS, or PLS regression  
499 258 (PLSR)), is frequently used. All but two of the studies in Table 1 reported using this approach. Perez-  
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501 259 Palacios, Caballero, González-Mohino, Mir-Bel, & Antequera (2019), preferred to use the simpler  
502 260 algorithm, multiple linear regression (MLR), when predicting texture-related characteristics of pork  
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504 261 loin cooked sous-vide at 70°C for 1, 2, 4, 6, or 8 h. In the case of classification, Moran, Andres, Allen,  
505 262 & Moloney (2018) whose research is discussed in more detail in ‘Section 5.1 NIRs’, used PLS  
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507 263 discriminant analysis (PLS-DA).

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509 264 Some recent alternative data analysis approaches in the meats field have included pre-processing  
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511 265 synchronous 2D correlation spectroscopy to identify the key wavelengths, which were then used in  
512 266 SVM models (Wang, W., Peng, Sun, Wei, & Zheng, 2018a), and multi-index statistical information  
513  
514 267 fusion (MISIF) for variable selection, (Qu *et al.*, 2018). Wu, Zhong, and Yang (2018) chose to forego  
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516 268 preprocessing and instead establish a prediction model for freshness by using a double-layer stacked  
517 269 denoising autoencoder neural network (SDAE-NN) algorithm, which proved to out-perform PLSR  
518  
519 270 and back propagation neural network (BP-NN). Processing has been approached by using combined  
520  
521 271 stacked interval partial least squares (siPLS) and sparse partial least squares regression (SPLSR) to  
522 272 create stacked interval sparse PLSR (sisPLSR), which aims to "find favorable rotations of the  
523  
524 273 classical PLS solutions while also utilizing local information in a spectra" (Poerio & Brown, 2017).  
525 274 Harrington (2018) modified an algorithm for training a restricted Boltzmann Machine, a type of  
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527 275 neural network, to improve PLS calibrations for moisture, fat, protein.

## 4.2. HSI

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277 Before sample acquisition, HSI systems should be calibrated for spectral and spatial information. The  
278 spectral calibration is performed with a black and a background reference as for spectroscopic  
279 devices. In detail, an image of the dark response is recorded by turning off the light sources or  
280 covering the lenses with non-reflective opaque black cap, thus, obtaining a 0% of reflectance image;  
281 and an image of the background response is recorded by a high reflectance standard or a spectra with  
282 100% reflectance (ElMasry and Sun, 2010). Moreover, a spatial calibration is needed to set the ground  
283 coordinates (X-Y spatial directions) of the measuring scene by a printed checkerboard.

284 The proper acquisition mode is fundamental to obtain reliable results when performing an HSI  
285 analysis of sample, as it is for all the considered non-destructive techniques. There are different  
286 system configurations according to the procedure of image acquisition: spectral scanning (area  
287 scanning), spatial scanning (point and line scanning), and snapshot imaging (Qin, 2010).

288 In spatial scanning system, the intensity spectra of one or multiple spatial positions are acquired. In  
289 the case of one-point scanning, so called whisker-broom imaging, the spectra of each single pixel is  
290 acquired at a time by moving the sample in the measuring position which will have constant lighting  
291 path and diffusion (Figure 2.a). Actually, it will be the HSI systems more similar to normal  
292 spectrometers. The light source, the lens, the dispersive device (normally prism or optical gratings)  
293 and the line detector array remain fixed in a position, whereas the sample is moved systematically in  
294 two spatial dimensions.

295 The main advantages are the constant lighting path between the optical system and the sample and  
296 the high spectral resolution; resulting, however, in time-consuming measurements.

297 In the case of multiple spatial position scanning, so called line scanning or push-broom imaging, the  
298 intensity spectra of a portion of the sample is acquired at a time (Figure 2.b). Actually, a set of pixels  
299 dispose in a line (2-D spatial-spectral information) is acquired and then the sample is moved in just  
300 one direction, which normally is transverse to the slit.

301 The setup of those systems requires two-dimensional dispersing element (normally prism or optical  
302 gratings) and a two-dimensional detector array perpendicular to the surface of the sample.

303 Nowadays, these systems are the ones preferred for benchtop instruments applied for research  
304 purposes, but their setup is promising for industrial applications; indeed, they reach spectral resolution  
305 comparable to point scanning instruments, being faster up to one hundred times (ElMasry, Mandour,  
306 Al-Rejaie, Belin, & Rousseau, 2019).

307 The most common spectral scanning setups are area or plane scanning, which collect a global spatial  
308 information one single wavelength at a time. In this configuration, the whole system remains in a fix

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309 position; i.e. the camera, the lens, the dispersive device and the field of view of sample location are  
310 in plane parallel to the detector (Figure 2.c). The main advantage of these systems is the affordable  
311 prize, the speed of acquisition; however, they normally return poor chemical information, with  
312 exception of cameras implemented with acousto-optic tunable filters (AOTF), which allows to  
313 acquire a higher wavelengths respect to variable filter ones. Their application for HSI benchtop  
314 instruments is not common, even though they are gaining importance in microscopy (Gottschall,  
315 Meyer, Schmitt, Popp, Limpert, & Tünnermann, 2018), as biological samples could be sensitive to  
316 the heating produced by the source lamps.

317 The last step of HSI acquisition is the file storage in the so called hyperspectral datacube, i.e. a 3D  
318 matrix in which along the 2D matrix (m rows and n columns) are stored the two orthogonal spatial  
319 directions and along the third dimension is store the spectra information ( $\lambda$ ). There are different ways  
320 to ordered in a logical manner the collected spatial and spectral information, such as band interleaved  
321 by pixel (BIP), by line (BIL) and band sequential (BSQ). Dedicated software allows the proper  
322 management of the datacube, whichever is the storage format, for further data analysis.

323 As discussed previously, the result of HSI analysis is an image with spectral information combined  
324 with spectral information stored in each pixel. The large amount of data and their high correlation  
325 need a proper handling to extract the relevant results by the adaptation of the multivariate data analysis  
326 techniques. In this section the steps required previously to multivariate data analysis performance are  
327 discussed, indeed, there are different “cleaning” procedures to properly pre-process the acquired  
328 images for discarding erroneous data values and non-informative background. For more detailed  
329 information, refer to Vidal & Amigo (2012).

330 The determination of dead pixels and spikes is fundamental to get rid of spurious information which  
331 may affect the performance of multivariate data analysis techniques.

332 Dead pixels are those pixels with missing or zero values and can be isolated or grouped in a specific  
333 location (line or area) of the image as they result from anomalies of the HSI detector. Several  
334 techniques are present in software dedicated to image analysis (Mobaraki & Amigo, 2018), such as  
335 thresholding techniques from median spectra calculated from the data, or more robust methods like  
336 genetic. Once located they are normally corrected by interpolation (by mean or median) with  
337 neighbour pixels ensuring a good representation of the information, as the neighbour pixels are  
338 generally highly correlated.

339 Other failures to be detected, and corrected accordingly, are rapid and sharp rise-fall of the signal,  
340 defined as spikes, resulting from failures of the detector or of the electronic circuits. Even if it is quite  
341 simple to detect spikes by visual inspection of the spectra, there are several approaches which can be

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652 342 used to automatically correct them in huge dataset as the ones generated by HSI. For a comprehensive  
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654 343 revision refer to Vidal & Amigo (2012).  
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656 344 Most of the time the acquired HSI is not containing just the information of the sample under study but  
657 345 it covers all the scanned area; thus, this area should be discarded. The more direct, but tedious and  
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659 346 long, strategy is the manual selection of the sample area by image visualisation at one, combine or  
660 347 whole spectral variables or by visualization of the scores image obtained by after a PCA. From the  
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662 348 same visualisation (wavelength or scores) a threshold value, defined manually or by histogram  
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664 349 inspection, can be applied to discriminate the sample from the background. Anyway, it should be  
665 350 considered that thresholding is always a critic point in image analysis as it could be affected by many  
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667 351 factors, mainly the type of data and the personal experience.  
668 352 After the mentioned “cleaning” procedures the data stored in as HSI are normally pre-treated and  
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670 353 analyzed adopting the techniques of classical spectroscopy, including clustering, classification and  
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672 354 regression methods.

### 673 674 355 **4.3. NMR**

675  
676 356 Prior to analyze the spectra acquired from the NMR systems, it must be pre-processed. For that,  
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678 357 firstly, the chemical shift misalignment must be corrected by using shifting algorithms, being the  
679 358 most commonly applied the icoshift algorithm (Savorani, Tomasi, & Engelsen, 2010). This algorithm  
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681 359 optimizes by shifting of spectral intervals, aligning the peaks of the spectral intervals and the spectra  
682 360 simultaneously. For that, correlations among the spectral data are used. After that, the noisy regions  
683  
684 361 must be removed and then, NMR spectra must be normalized and scaled (Craig, Cloarec, Holmes,  
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686 362 Nicholson, & Lindon, 2006). These algorithms allow increasing the representation of lower  
687 363 concentrations and minimizing the contribution of noise. Finally, some pre-processing techniques in  
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689 364 classical spectroscopy are applied, being the most used the algorithms: Savitzky-Golay that aims to  
690 365 reduce the noise of the spectra, which is often achieved by smoothing algorithms in combination with  
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692 366 1st or/and 2nd derivative transformation; Standard Normal Variate (SNV), which aims to correct the  
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694 367 slope of the spectra in order to optimize the spectral data; Multiplicative Scatter Correction (MSC),  
695 368 which aims to correct the scattering due to external interactions with the sample, i.e., lights,  
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697 369 temperature,... (Rinnan, Van Der Berg, & Engelsen, 2009). As example, Figure 3 shows a NMR  
698 370 spectra of beef. Then, for extracting the data from the spectra, the identification of each peak is related  
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700 371 to each component of the sample. For that, the MMCA (Metabolite-Metabolite correlation analysis)  
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702 372 is commonly used to identify the component of each spectra (Craig *et al.*, 2006). Finally, the intensity  
703 373 of each peak is measured as concentration value of each component.  
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374 For the data analysis, in general, Partial Least Square (PLS), Principal Component Analysis (PCA)  
375 or Multiple Linear Regression (MLR) have been usually applied in order to obtain accurate results in  
376 NMR.

377 Other statistical tools such as ANOVA, Pearson's correlation coefficient, Naïve-Bayes, first-order  
378 and second-order statistics have been applied for analyzing the data obtained from the spectra  
379 (Delorme, Sejnowski, & Makeig, 2007). From all these statistical techniques, PCA must be noted.  
380 PCA is the most used exploratory data analysis technique, since it is usually used to identify patterns  
381 in measured data and to visualize the distribution of the data. This technique allows evaluating the  
382 relationship among the variables by using mapping and displays techniques for understanding the  
383 structure of the complex multivariate datasets (Bro, & Smilde, 2014). PLS (Bro, 1996) aims to  
384 maximize the covariance between the predictor and the response data. Its popularity can be ascribed,  
385 in part, to its speed, since the model parameters for each component can be calculated easily. Its ease  
386 of use because of the only meta-parameter to be optimized is the number of components. Its  
387 interpretation since the PLS scores, loadings and weights can be investigated in order to determine  
388 whether the model components have a meaning for the meat and meat products (Martens, & Naes,  
389 1989). MLR is used to represent linear relationship between a dependent variable and several  
390 independent variables. This technique obtains a linear regression equation, which can be used to  
391 predict future values (Hastie, Tibshirani, & Friedman 2001).

#### 4.4. MRI

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393 Once the MRI image has been acquired, it is analysed to extract numerical information. Pre-  
394 processing and/or segmentation techniques are firstly applied. The objective of the pre-processing is  
395 to improve the obtained image by outstanding certain features or eliminating noise (Sonka, Hlavac,  
396 & Boyle, 1999). The segmentation techniques extract elements of interest from the images (Maravall,  
397 1993), such as the thresholding methods (Cheriet, Said, & Suen, 1998; Otsu, 1979) or fuzzy logic  
398 (Raof *et al.*, 2008). Other applied segmentation techniques in MRI from meat are active contour,  
399 which detect the muscle of interest (Caro, Rodríguez, Cernadas, Durán, & Villa, 2001; Caro,  
400 Rodríguez, Durán, & Antequera, 2012), and the algorithms for selecting region of interest (ROI)  
401 (Molano, Rodríguez, Caro, & Durán, 2012), which is the maximum area inscribed in the muscle  
402 previously selected. The last step of the image analysis is the extraction of the computational  
403 characteristics. This process allows describing the MRI as a vector of features. For this task,  
404 algorithms based on texture features have been the most used: Gray-Level Co-occurrence Matrix  
405 (GLCM), Gray-Level Run Length Matrix (GLRLM) and Neighbouring Gray-Level Dependence  
406 Matrix (NGLDM). GLCM counts the number of times that each pair of gray levels ( $i,j$ ) occurred at a

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770 407 given distance  $d$  in all directions. GLRLM measures runs into the image, i.e., a set of consecutive  
771 pixels in the image with the same gray level value. NGLDM considers the relationship between an  
772 408 element and all its neighbouring elements at one time rather than one direction at a time. The  
773 409 application of these algorithms results on the values of different computational texture characteristics.  
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775 410 The use of fractal algorithms for analysing MRI from meat has been recently studied. They study the  
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777 411 degree of symmetry or self-similarity found in a structure at all scales, allowing the identification of  
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779 412 recurring patterns and removing the possibility of image compression (Hibbert, 1991). Three fractal  
780 413 algorithms have been used in meat: the traditional algorithm to compute the fractal dimensions, the  
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782 414 fractal texture algorithm (FTA) and the one-point fractal texture algorithm (OPFTA). The traditional  
783 415 algorithm measures the number of boxes (small fractions of the image depending of the size of the  
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785 416 original image) needed to cover an area occupied by the object as a function of the size of boxes. FTA  
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787 417 (Caballero, Caro, Ávila, Rodríguez, Antequera, & Pérez-Palacios, 2017b) is a novelty texture  
788 418 algorithm based on the number of times that a pattern is repeated in each image depending of box  
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790 419 size calculated in each case. These fractal features are gathered in a vector, and each vector was  
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792 420 computed based on second order statistics. OPFTA (Caballero *et al.*, 2018) is an algorithm based on  
793 421 features obtained from fractal properties values into smaller rectangles of 32x32 pixels. From all these  
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795 422 values, the value for the box size equal to eight is selected. After that, these values are gathered in  
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797 423 order to create a matrix, so, each cell of the matrix represents one ROI from the image. Finally, the  
798 424 features were calculated on the matrix by applying second order statistics.  
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800 425  
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802 426 Other algorithms to analyse the MRI images of meat are the mapping techniques (Zarei, & Sepyani,  
803 427 2016), the use of contrast on the images (Vala, & Baxi, 2013), the histograms (Bajd, Skrlep, Candek-  
804  
805 428 Potokar, & Sersa, 2017), and the 3-D version of the texture (Ávila, Caballero, Durán, Caro, Pérez-  
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807 429 Palacios, & Antequera, 2015). The mapping techniques is based on geometric transformation of  
808 430 images, re-locating the points in the source images on different coordinates in a destination image.  
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810 431 This allows describing some features of the images (Zarei, & Sepyani, 2016). The contrast on the  
811 432 images stands out some zones of the images. The use of this technique joins to the thresholding  
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813 433 methods allow describing some features of the images and characterize them (Vala, & Baxi, 2013).  
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815 434 The 3D version of the texture algorithms has also been applied for the analysis of 3D reconstructed  
816 435 MRI images (Ávila *et al.*, 2015).  
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818 436 Regarding the data analysis, in the case of MRI, most studies have applied usual statistical tools such  
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820 437 as Pearson's correlation coefficients, analysis of variance (ANOVA), PCA or statistics measurements  
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822 438 as the first and second order statistics. In the last years, the application of data mining in these studies  
823 439 has increased. Data mining is one of the steps of a larger process known as Knowledge Discovery in  
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829 440 Databases (KDD) (Fayyad, Piatetsky-Shapiro, & Smyth, 1996), and it is related to large volume of  
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831 441 data. Most of the data mining techniques applied in the MRI studies of meat are MLR and Isotonic  
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833 442 Regression (IR). IR estimates ordered values for an independent variable as a function of one of the  
834 443 input variables (Barlow, Bartholomew, Bremner, & Brunk, 1972), only selecting the input parameters  
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836 444 with the best adjustment results. Partial Least Square (PLS) has also been applied the MRI studies in  
837 445 meat.  
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839  
840 446 Most of the MRI studies based on regression methods have applied the common cross-validation  
841 447 methodology (Kohavi, 1995), which divides the data in two sets, training and testing, with  
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843 448 information of images from the same sample. Recently, a modified of the usual method has been  
844 449 developed. It consists of three sets (training, validation and test) and leaving one meat piece out when  
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846 450 creating the dataset (Ávila *et al.*, 2019).  
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## 848 451 **5. APPLICATIONS OF NON-DESTRUCTIVE METHODOLOGIES FOR FRESH MEAT** 849 850 452 **ANALYSIS.**

### 851 852 453 **5.1. NIRs** 853

854 454 Since the excellent and comprehensive reviews by Dixit *et al.* (2017), Kademi, Ulusoy, and Hecer,  
855  
856 455 (2019), and Prieto, Pawluczyk, Dugan, and Aalhus (2017), there have been roughly 6 types of NIRs  
857 456 studies, depending on their objective, that have analysed fresh meat in the last 4 years. The first type  
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859 457 of study used NIRs as a standard laboratory method. Konarska, Kuchida, Tarr and Polkinghorne  
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861 458 (2017) used correlation analysis to compare three approaches to measuring beef marbling on 12  
862 459 muscles: image analysis, and subjective evaluation from images of the intact sample, and NIRs on  
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864 460 the homogenized sample using the NIRFlex N-500 (Buchi, Switzerland) with its built-in prediction  
865 461 algorithm. The overall strength of correlation for % marbling between NIRs and image analysis was  
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867 462 0.60 ( $P \leq 0.01$ ) while for individual muscles it varied from 0.13 (*gluteus medius*) to 0.77 (*serratus*  
868 463 *ventralis cervicis*;  $\geq 0.56$  significantly different from 1 at  $P \leq 0.01$ ). Mínguez, Sánchez, Hernández,  
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870 464 Ragab, El Nagar and Baselga (2017) used NIRs to predict fatty acid composition and % protein from  
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872 465 crossbred ground freeze-dried rabbit *longissimus lumborum* in order to evaluate the genetic groups  
873 466 used in the crossbreeding and to estimate genetic contributions to meat quality. A difference in fatty  
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875 467 acid composition ( $P \leq 0.05$ ) was detected for one genetic line, but no differences in % protein. The  
876 468 second type of study used NIRs for authentication. Moran *et al.* (2018) tested various pre-processing  
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878 469 approaches to confirm degree of ageing of three different, intact, bloomed, beef muscles following 3,  
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880 470 7, 14 and 21 d of vacuum-packaged storage. In this preliminary study, it was found that the best  
881 471 prediction model was specific for each muscle, and was best following spectral pre-treatment such as  
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883 472 Savitsky-Golay 1<sup>st</sup> or 2<sup>nd</sup> derivatives on the full spectra (400-2400 nm). When tested, model  
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sensitivity ranged from 96.3-100%, specificity from 98.8-100%, and overall correct classification from 99.1-100%. Pieszczek, Czarnik-Matusewicz, & Daszykowski (2018) explored two class modelling techniques: one-class classifier partial least squares (OCPLS) and soft independent modelling of class analogy (SIMCA) to recognize pure minced beef, pork, or lamb from NIRs spectra. For beef, models offering the best performance, with an emphasis on specificity, were either SIMCA preceded by SNV (sensitivity = 99.8% ± 0.78; specificity = 98.98% ± 0.56) or OCPLS preceded by MSC (sensitivity = 99.11% ± 5.07; specificity = 99.9% ± 0.37). SIMCA performed best for pork and lamb, with no pre-processing for the former (sensitivity = 99.94% ± 0.49; specificity = 87.20% ± 4.00), and preceded either by MSC (sensitivity = 99.98% ± 0.20; specificity = 98.00% ± 0.58) or ISC (sensitivity = 99.48% ± 0.79; specificity = 99.06% ± 0.47) for the latter. Table 1 summarizes the 4 other types of studies along with their calibrations and performances: detecting adulteration, prediction, equipment testing, and exploration into various spectra pre-processing and processing approaches.

Prediction studies have been diverse (Table 1). Li, Z., Jia, Wang, Liu and Dong, (2016) reported predicting cooked beef texture from the raw meat. Pork studies include prediction of pork components with particular emphasis on fat composition (Richli, Kaufmann & Scheeder, 2016; Prevolnik Povse *et al.*, 2017), identifying potential PSE (Li, X. *et al.*, 2016), determining cholesterol levels (Wang, H. *et al.*, 2017), post mortem meat quality (Andersen, Wold, Gjerlaug-Enger & Veiseth-Kent, 2018a) and belly firmness (Soladoye *et al.*, 2018). Lamb studies looked at lipid peroxidation to replace TBARS (Ripoll, Lobón & Joy, 2018), and classification for eating quality (Knight *et al.*, 2019). Equipment used was a mix of benchtop and portable, and usually intact samples were tested. The wavelength range was more frequently VIS-NIRs than NIRs alone. Pre-processing commonly consisted of smoothing, 1st or 2nd derivative, and SNV, while processing was always PLS or PLSR. NIRs is a growing field, therefore new equipment is continuously under development. There have been a number of reports on testing results in recent years. Dixit and his research group have reported a group of studies centering around equipment developed in-house, and which consists of a Fabry-Pérot interferometer, a 4-point photodiode array, and collimating lenses, enabling multipoint data capture (Dixit *et al.*, 2016a; 2016b; 2016c). The equipment has been designed for rapid on-line detection and prediction of ground meat composition, thus performance has been reported for different sample stand-off distances, movement speeds, and combinations of both. Several equipment testing reports were based on technology comparisons (eg. NIRs, hyperspectral imaging (HSI), Fourier transform-NIRs (FT-NIRs), Raman, fluorescence, etc.), and the NIRs results can be found in Table 1 (Andersen *et al.*, 2017; Andersen, Wold, & Veiseth-Kent, 2018b; Nolasco-Perez *et al.*, 2019).

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946  
947 506 Piao *et al.* (2018) explored the efficacy of transferring calibration equations for beef fatty acid groups  
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949 507 developed on master equipment, to slaves. Wang, W., Peng, Sun, and Li (2017) tested custom in-  
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951 508 house equipment constructed of two spectrometers with different wavebands, operated in sequence,  
952 509 for the ability to predict pork freshness on a conveyor line. This team also explored different data pre-  
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954 510 processing and processing approaches (Wang, W., Peng, Zheng, Tian, & Wei, 2016; Wang, W., Peng,  
955 511 Sun, Zheng, & Wei, 2018b) and included reducing the spectra to key wavelengths to potentially  
956  
957 512 increase the rate of data collection (Wang W., *et al.*, 2018a).

## 959 513 5.2. HSI

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961 514 Hyperspectral imaging for the assessment of meat quality by chemical composition has been applied  
962  
963 515 mostly for moisture, fat content and composition and protein. Also, and Total Volatile Basic Nitrogen  
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965 516 (TVB-N), TBARS and K-value have been investigated as quality indexes for meat freshness  
966 517 evaluation. As concern technological and sensory attributes, WHC (water holding capacity), WBSF  
967  
968 518 (Warner–Bratzler shear force), SSF, (slice shear force) and colour by CIE - Lab\* chromatic scale  
969 519 have been considered. Thus, the works dealing with their determination by HSI are here revised  
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971 520 (Table 2), excluding reference dealing with microbiological quality determination; in case of interest  
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973 521 in this field please refers to the review by Kamruzzaman, Makino, and Oshita (2015).

974 522 Different line scanning approaches (or push-broom imaging) have been proposed for moisture content  
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976 523 determination. Kandpal, Lee, Kim, Mo, and Cho (2013) proposed a VIS/NIRs – HSI system ranging  
977 524 from 400 to 1000 nm to predict moisture content in chicken breast. 52 samples were used to calibrate  
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979 525 the PLSR model, whereas other 20 chicken breasts were used to validate the model reaching optimal  
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981 526 levels ( $R^2_p$  of 0.94 and SEP of 0.71%.) when only the NIRs (700-1000 nm) range was used. The  
982 527 laboratory-based push-broom NIRs hyperspectral imaging system (900–1700 nm) proposed by  
983  
984 528 Barbin, ElMasry, Sun, and Allen (2013) allowed the construction of good PLS regression models,  
985 529 developed from feature-related wavelengths, to predict moisture content in minced pork samples ( $R_p$   
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987 530 of 0.91 and SEP of 0.62). The same research group developed moisture content prediction models for  
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989 531 fresh minced beef samples collected from different muscles reaching prediction abilities for  $R_p$  of  
990 532 0.89 and accuracy (SEP) of 0.46 (ElMasry, Sun & Allen, 2013).

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992 533 By a larger NIRs range (880 – 1720 nm), Zhao, Esquerre, Downey and O'Donnell (2017)  
993 534 demonstrated higher prediction ability for moisture content in ground beef samples ( $R^2_p$  of 0.99;  
994  
995 535 RMSEP of 0.64 w/w). However, the samples used by Zhao *et al.* (2017) to calibrate (n=36) and  
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997 536 validate (n=9) the model were quite small compared to Barbin *et al.* (2013) and ElMasry *et al.* (2013),  
998 537 both considering around 80 samples for the training set, and 40 samples for the testing set.

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1007 538 Also lamb meat has been tested for moisture prediction by push-broom hyperspectral imaging system  
1008 539 in the spectral range of 900–1700 nm (Pu, Sun, Ma, Liu & Kamruzzaman, 2014). In their study a  
1009 540 relevant number of samples has been used, being 127 lamb meats including *semimembranosus*,  
1010 541 *semitendinosus* and *longissimus dorsi* muscles. Results comparable to those of Barbin *et al.* (2013)  
1011 542 and ElMasry *et al.* (2013) have been obtained by applying MLR models after a hierarchical variable  
1012 543 strategy (UVE-SPA-CSA), being the  $R_p$  of 0.92, the RMSEP of 0.58 and the RPD equal to 2.53.

1013 544 Fat content as total amount, intermuscular fat (IMF), or specific classes composition (SFAs, saturated  
1014 545 fatty acids; UFAs, unsaturated fatty acids and oleic acid) have been widely investigated across  
1015 546 different meat species.

1016 547 The study by Wold, O'Farrell, Høy, and Tschudi (2011) revealed how simplified HSI instrument  
1017 548 (multispectral imaging system with 15 wavelengths between 760 and 1040 nm) can be used for the  
1018 549 online estimation of fat content in beef trimming. Indeed, they reached high fat prediction accuracy  
1019 550 (RMSEP of 0.6%) in 100 batches. More recently, Lohumi, Lee, Lee, Kim, Lee and Cho (2016)  
1020 551 calibrated models for fat content in beef from different quality grades by ANOVA, spectral angle  
1021 552 measure (SAM), and Euclidean distance measure (EDM) methods, reaching  $R^2_c$  of 0.91, 0.95, and  
1022 553 0.96 however, they used a small number of samples ( $n = 24$ ) and they did not perform any validation  
1023 554 using independent samples. By revising the last decade literature, it seems that the best model on term  
1024 555 of prediction for fat in raw beef samples was develop by Zhao *et al.* (2017); they obtained  $R^2_p$  of 0.99  
1025 556 and RMSEP of 0.73% w/w of fat by applying EMCV-PLSR algorithm. Similarly, Pu *et al.* (2014)  
1026 557 developed MLR models after a hierarchical variable strategy (UVE-SPA-CSA) obtaining  $R_p$  of 0.98,  
1027 558 RMSEP of 0.36 and RPD of 4.13. The strength of their work over Zhao *et al.* (2017) is due to the  
1028 559 high number of samples and they representability as they considered 126 lamb meats including  
1029 560 *semimembranosus*, *semitendinosus* and *longissimus dorsi* muscles (84 samples used to calibrate the  
1030 561 model and 42 for its independent validation).

1031 562 The research group by Kobayashi developed models for the prediction of specific fat categories in  
1032 563 beef samples (Kobayashi, Matsui, Maebuchi, Toyota & Nakauchi, 2010 and Kobayashi, Mori,  
1033 564 Nishino, Toyota & Nakauchi, 2012). They obtained reliable models for the prediction of saturated  
1034 565 fatty acids ( $R^2_p = 0.87$ , RMSEP =1.69, RPD=2.43), unsaturated fatty acids ( $R^2_p = 0.89$ , RMP =3.41,  
1035 566 RPD=2.84) and oleic acid ( $R^2_p = 0.71$ , RMSEP =3.13, RPD=1.855) by predicting at least 32 meat  
1036 567 samples from three 25-month-old Japanese Black (Wagyu) cattle.

1037 568 Both Liu and Ngadi (2014) and Huang, Liu and Ngadi (2017) developed regression models to predict  
1038 569 IMF in porcine meat by combining different image features – from WLD to Gabor filter and improved  
1039 570 GLCM (gray level co-occurrence matrix) - with spectral information. Liu and Ngadi (2014) reached  
1040 571 an optimal model performance in prediction with an adjusted  $R_p^2$  and RMSEP of 0.93 and 0.17%,  
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1065 572 respectively; whereas the rib by rib models developed by Huang *et al.* (2017) reached a  $R_p$  of 0.90  
1066 and an RMSEP of 0.92% for the 3rd last rib even if the model was constructed with only 18 samples  
1067 573 and validated with 6.  
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1070 575 A relevant study for IMF prediction was developed by Craigie *et al.* (2017). Indeed, they performed  
1071 HSI acquisition of 2454 samples of lamb loin (*M. longissimus lumborum*) at processing plant level  
1072 576 over consecutive years. They evaluated the performance of more than ten different regression  
1073 577 algorithms reaching prediction abilities up to  $R_p^2$  of 0.72 and RMSEP of 0.45 with a Gaussian process  
1074 regression (GPR) approach.  
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1078 580 Prieto Osika, Aalhus, Lopez-Campos, Juarez, and Pawluczyk, (2018) reported that models based on  
1079 HSI to predict protein are generally less accurate than the ones for moisture or fat. Indeed, Pu *et al.*  
1080 581 (2014) only reached  $R_p$  of 0.67, RMSEP of 0.41 and RPD of 1.31 when predicting protein content by  
1081 582 the UVE-SPA-CSA-MLR model calibrated with 126 lamb samples. Certainly, the estimation of  
1082 protein content by HSI data seems highly dependent on the sample form: models developed based on  
1083 583 minced meat generally reached higher performances over intact muscles analysis. This is proved by  
1084 584 the studies of Kamruzzaman, ElMasry, Sun and Allen (2012a) and ElMasry *et al.* (2013). The latter  
1085 developed models for protein prediction in beef samples from three different muscles (*M. longissimus*  
1086 585 *dorsi*, *M. semitendinosus* and *M. psoas major*) reaching high performance ( $R_p^2$  of 0.86, SEP of 0.29)  
1087 and for pork samples reaching  $R_p^2$  of 0.88 and SEP of 0.40. It has been hypothesized that  
1088 586 homogenized samples, such as minced meat, overcome interferences of muscle fibre organization  
1089 587 and muscle physical characteristics (Prieto *et al.*, 2018) leading to better prediction capabilities.  
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1097 592 Khulal, Zhao, Hu and Chen (2017) developed models to predict TVB-N in poultry by back  
1098 propagation neural network (BPNN) algorithm. They analysed fifty chicken breast fillets by an *ad*  
1099 593 *hoc* developed hyperspectral imaging system and, by combining the spectral variables with the texture  
1100 594 ones, they calibrated a good regression model, further validated by other 25 samples, reaching a  $R_p$   
1101 of 0.75, and a RMSEP of 6.39mg/100g of meat.  
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1105 597 Great interest has been posed by the scientific community in predicting TVB-N in pork meat in recent  
1106 years. Both Li, Chen, Zhao and Wu (2015) and Guo, Huang, Zhu, Guo and Qin (2018) developed  
1107 598 methods based on NIRs-HSI systems based on line-scanning in the 400-1000 nm range to predict  
1108 599 TVB-N in fresh pork. Even if they used different regression strategies, namely Least-squares support  
1109 vector machine (LS-SVM) and Back propagation artificial neural network (BPANN), they were able  
1110 600 to obtain optimal models in terms of coefficient of determination ( $R_p^2 > 0.93$ ). However, the error  
1111 601 (RMSEP) obtained by Li *et al.* (2015) resulted considerably lower (RMSEP= 1.86mg/100g) in respect  
1112 602 to the one obtained by the BPANN model (5.52mg/100g).  
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Differently, Lee, Kim, Lee and Cho (2018) proposed the use of a hyperspectral fluorescence imaging system, based on high-intensity light-emitting diodes at 365 nm, for the determination of TVB-N contents in pork meat. In their work, 186 fresh pork *longissimus* muscles were purchased from a local supermarket, trimmed and shaped in size of 5cm x 4cm x 2.5cm to be analysed. The developed model was based on LS-SVM, leading to an optimal prediction capability ( $R^2_p$  of 0.967 and RMSEP of 1.90%).

As far as concern lipid oxidation, a research by Xiong, Sun, Pu, Xie, Han, and Luo (2015a) studied the possibility of developing a regression model based on HSI data to predict TBARS in chicken breast slices during storage at 4 °C for 0, 3, 6, 9 days. They established a simplified model by selecting 10 optimal wavelengths using the successive projections algorithm (SPA) able to predict TBARS with good performance, being the  $R_p$  of 0.80 and the RMSEP of 0.16 mg/100g.

Cheng, Sun, Pu and Liu (2016) proposed a feature level fusion of HSI spectral data and textural data to develop PLSR model for the prediction of another important freshness indicator of meat, K-value. They selected six feature wavebands (407, 481, 555, 578, 633, and 973 nm) and merged them with texture data of the grayscale images extracted by GLCM at the selected wavebands. The data fusion approach resulted in improved models with  $R^2_p$  of 0.92 and RMSEP of 4.0%.

If the success of HSI is expected for chemical composition analysis, its application for technological attributes has also been investigated - with more or less success - in the recent years.

Indirect measure of pH has been proposed for beef (ElMasry *et al.*, 2013), pork (Barbin *et al.*, 2012) and lamb (Kamruzzaman, ElMasry, Sun & Allen, 2012b). Even if applying different regression approaches models developed for beef and lamb did not performed well ( $R^2_{CV}$  lower than 0.70). The same authors hypnotised that the considered pH variation in animal flash after-post mortem process could be too small to construct a robust model. However, the model developed for pork gave  $R^2_p$  of 0.90 and RMSEP of 0.10, when considering a similar variation range was 5.31 – 6.43.

HSI systems demonstrated to be useful for colour determination intuitively when the spectral range covered also the visible part (400-700 nm). This is the case of the models developed by Wu, Peng, Li, Wang, Chen, and Dhakal, (2012) for beef and by Kamruzzaman, Makino, and Oshita (2016a) for beef, pork and lamb. Those models reached high prediction capabilities for  $a^*$  determination, which describes the colour space from green (-) to red (+), thus, the expected variation in red meat. Also, models developed for  $L^*$  - lightness from black (0) to white (100) – gave high predictive performances ( $R^2_p > 0.96$ ).

WHC modelling brought to heterogeneous results. Barbin *et al.* (2012) and ElMasry, Sun and Allen (2011) obtained good models after variable selection in prediction ( $R^2_p=0.89$ , RMSEP=0.79%) and cross-validation ( $R^2_{CV}=0.89$ , RMSECV=0.26%), respectively. The variable selection strategy (RC-

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LS-SVM optimal wavelengths) resulted successful for the models developed by Kamruzzaman, Makino, and Oshita (2016b) leading to excellent WHC predictions and clear distribution maps for beef, pork and lamb samples. Less promising results were obtained by the authors using the full NIRs spectral range (Kamruzzaman *et al.*, 2012b).

Models based on HSI for the prediction of meat texture, highly relevant for consumer acceptance, has been recently investigated. For instance, Xiong *et al.* (2015b) achieved acceptable results ( $R_p$  of 0.87 and RMSEP of 0.05) by applying PLS regression to images collected in the Vis/NIRs-HSI range (400 to 1000 nm) to predict hydroxyproline in poultry meat. For extensive details about recent application of HSI for texture prediction in meat products refer to Reis *et al.* (2018).

**5.3. NMR**

Several studies have been found in the scientific literature by applying LF-NMR and HF-NMR systems to estimate quality traits of fresh meat (Table 3).

The use of LF-NMR to analyse pork samples was firstly tested with the aim of study the relationship between cooking temperature, the water distribution and some sensory attributes (Bertram, Aaslyng, & Andersen, 2005). T2 as relaxation time and CPMG sequence for the spectra acquisition were applied, while PLS was selected for the data analysis. High correlation coefficients were found between the changes in the sensory attributes, which were caused by the temperature of cooking, and the spectral data, i.e. juiciness ( $R^2 = 0.82$ ), tenderness ( $R^2 = 0.87$ ). Moreover, Straadt, Rasmussen, Andersen, and Bertram, (2006) demonstrated the relationship between the water holding capacity and the water distribution of loins with different days of aging while are cooking and the NMR data ( $R > 0.75$ ). These authors used a LF-NMR system (23.2 MHz) with T2 relaxation time and CPMG sequence.

In the case of chicken samples, most studies used LF-NMR and T2 relaxation time. Thus, Li, Wang, Xu, Xing, and Zhou (2014) tried to determine the effect of freezing-thawing with different conditions of high pressure on water holding capacity or cooking loss of chicken by means of LF-NMR. These authors used CPMG sequence and applied correlation coefficients as data analysis technique. Thus, good to excellent but inverse relationship were found both parameters (water holding capacity  $R = -0.707$ , cooking loss  $R = -0.920$ ). Under the same analytical conditions, previous studies had also found high correlation coefficients between T2 signal intensity and water activity ( $R > 0.90$ ) (Venturi, Rocculi, Cavani, Placucci, Dalla Rosa, & Cremonini, 2007), cooking loss ( $R = 0.986$ ) and moisture content ( $R = 0.953$ ) (Shaarani, Nott, & Hall, 2006).

The use of LF-NMR has also been evaluated in beef. Pereira and Colnago (2012) determined the moisture content on beef samples by applying CPMG sequences weighted on transverse

magnetization relaxation time (T2) on LF-NMR (8.9 MHz), and using PLS and PCA as data analysis techniques. Correlation coefficients obtained between moisture content and T2 intensity values of NMR were higher than 0.96. Moreover, mixes of beef and horse were tried to be classified as a function of the quantity of each type of meat by LF-NMR (60 MHz). For that, T1 and T2 relaxation times were applied and PCA and Naïve Bayes statistics were used as classification techniques. Results showed that 106 of 107 samples were correctly classified, showing the accuracy of this methodology to differentiate mixtures of different meat (Jakes *et al.*, 2015).

The effect of thawing and post-thawing on a number of quality parameters of rabbit meat (lipid oxidation, water holding capacity, instrumental color and texture) has also been studied by applying LF-NMR (70 MHz) (Jia, Liu, Nirasawa, & Liu, 2017). For that, transverse magnetization relaxation time (T2) and CPMG pulse sequence were used on LF-NMR. ANOVA and correlation coefficients were applied as data analysis techniques. Correlation coefficients higher than 0.95 were achieved for all parameters studied in this work.

HF-NMR has not been highly used in fresh meat, with only a study being found in duck samples (Liu, Pan, Ye, & Cao, 2013). These authors evaluated the capacity of HF-NMR to discriminate meat samples as a function of the age of the ducks and to determine 22 metabolites. These authors used T1 relaxation time and CPMG pulse sequence on HF-NMR (400 MHz), and PCA and PLS as data analysis techniques. Results showed accurate classification results (percentage of correct classification higher than 60 %), differentiating among meat samples from duck of 27, 50, 170 and 500 days old, and correlations coefficients (higher than 0.70 for most metabolites).

#### 5.4. MRI

Most of the MRI studies in meat have been published after the year 2000, and they have been mainly carried out by using HF scanners, being the dry-cured ham the most sampled meat product in this kind of works. However, the interest on LF-MRI is experimenting a significant increase nowadays.

Regarding the publications about the use of MRI to analyse fresh meat in the last years, which are summarized in Table 4, the earliest works (2011-2016) used HF-scanners, whereas LF ones have been preferred in the latest researches (2015-2019). It is also noted that these recent studies on MRI and fresh meat can be fairly gathered as a function of their main objective: classifying of the meat samples, analysing the percentage of intramuscular fat or lean, optimization of the MRI methodology to predicted the quality parameters of meat and evaluation of the changes due to freezing/thawing the meat samples.

Studies centred on classification purposes have been carried out with hams (Perez-Palacios *et al.*, 2011; Caballero *et al.*, 2016). These authors aimed to discriminate between hams from pigs fed with



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different diets (acorn and grass vs. high oleic acid enriched feeding) (Perez-Palacios *et al.*, 2011), and as a function of their salt content (Caballero *et al.*, 2016). In both studies the images were acquired by using HF-H<sup>1</sup> MRI scanners with a quadrature whole-body coil and applying T1-weighted spin echo sequences. Three similar steps were carried out in both studies for the MRI analysis. Active Contours was applied on the images to recognize the *Biceps femoris*, in both studies, and the *Semimembranosus* muscles, only in the salt study. Then, the Region of Interest (ROI) was selected on each muscle. In the salt study, each ROI was divided into minor rectangles, which can be called mini-ROIs. Finally, the analysis of the ROIs and mini-ROIs was done by applying three computational texture algorithms (GLCM, GLRLM, NGLDM), obtaining a vector of 15-17 computation texture features.

Results obtained in the study of Perez-Palacios *et al.* (2011) showed visual differences between the MRI of hams from pigs fattened with acorn and grass (darker grey colour that represents the muscle) and high oleic acid concentrates (brighter white colour that represents the fat) (Figure 4). Besides, ANOVA showed statistical differences in the values of the texture features between the two batches and PCA displayed a clear separation of the two groups of hams.

In the case of the salt study, Caballero *et al.* (2016) applied classification techniques of data mining, getting a 77-79% of correct classification of ham muscles with different salt content when applying the J48 decision tree technique. These authors also achieved to predict the salt content as a function of the computational texture features with high accuracy by MLR.

As for the studies based on determining the content of fat and lean by MRI, Lee, Lohumi, Lim, Gotoh, Cho, and Jung, (2015) scanned different beef samples (ribeyes of four categories) by using a LF-H<sup>1</sup> MRI with a head coil and SE-T1 sequences. Then, the threshold method was applied to calculated total area of the image and of the intramuscular fat, as well as their number of pixels. The percentage of intramuscular fat was then calculated considering these data and the density of the fat and muscle  $= (\text{total pixel number of intramuscular fat} * \text{fat density}) / (\text{total pixel number of muscle} * \text{muscle density}) + (\text{total pixel number of intramuscular fat} * \text{fat density}) * 100$ . Results on this study were more precise for samples with high fat percentage than for that with low fat content, which authors have ascribed to thresholding method. It classifies each pixel as pure fat or not, but some pixel may contain both tissues.

In the same line, Bernau *et al.* (2015) investigated about the use of MRI to evaluate the carcass composition of boars, instead of the standard protocol. Samples were scanned in a LF-H<sup>1</sup> MRI with a large body coil and GE-T1 sequences. Images were analysed semi-automatically by the interactive segmentation function to separate muscles from fat and removing bone and cartilage tissues manually. Total volume and the volume of fat and lean were calculated on the different regions and muscles of

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1360 739 the carcass and compared with the results from the standard protocol. A stepwise regression analysis  
1361 740 was carried out on these traits, resulting in a regression equation for the percentage of lean meat with  
1362 741 a high coefficient of determination ( $R^2 = 0.88$ ).

1363 742 The principal objective of most studies using LF MRI scanners has been the optimization of the  
1364 743 procedure to achieve high accurate results (Perez-Palacios *et al.*, 2017; Caballero *et al.*, 2017a; Ávila  
1365 744 *et al.*, 2018). In these works, pork loins were scanned by using a LF-H<sup>1</sup> MRI with a hand/wrist coil.  
1370 745 Different combinations of acquisition sequence, image analysis methods and prediction techniques  
1371 746 of data mining techniques have been tested to reach the best option for predicting the quality  
1372 747 characteristics of pork loins most precisely. Perez-Palacios *et al.* (2017) focused on the texture  
1373 748 algorithms (GLCM, GLRLM, NGLDM), Caballero *et al.* (2017a) on fractal algorithms (CFA, FTA,  
1374 749 OPFTA) and Ávila *et al.* (2018) on 3D texture algorithms (3D GLCM, 3D GLRLM, 3D NGLDM),  
1375 750 whereas the same acquisition sequences (SE, GE and T3D weighted in T1) and data mining  
1376 751 techniques (MLR and IR) were evaluated in the three works.

1381 752 Perez-Palacios *et al.* (2017) observed visual differences in the MRI of loins depending on the  
1382 753 acquisition sequence, with SE offering sharper and better-defined images than GE and T3D, and also  
1383 754 found the significant influence of the acquisition sequence on the values of all computational texture  
1384 755 features. Prediction equations of the different physico-chemical parameters analysed (percentage of  
1385 756 moisture and lipid, water activity and instrumental colour coordinates) showed accurate correlation  
1386 757 coefficients when applying SE or GE in combination with any of the acquisition sequences or data  
1387 758 mining techniques tested, while the use of T3D sequence lead to less precise results. In view of that,  
1388 759 it was considered the time consumed and the required resources to choose the best option, which was  
1389 760 SE-GLCM-MLR.

1390 761 As for the research of Caballero *et al.* (2017b) with fractal algorithms, there were also found  
1391 762 significant differences in all fractal features among GE, SE and T3D acquisition sequences. The  
1392 763 prediction results were affected by the sequence acquisition, the fractal algorithm and the data mining  
1393 764 technique, finding the best prediction results with the combination SE-OPFTA-MLR. In fact, SE  
1394 765 offers a better performance in terms of signal to noise ratio than GE and T3D, OPFTA simulates the  
1395 766 texture of the MRI better than FTA and CFA (Caballero *et al.*, 2018), and MLR is recommended  
1396 767 when the values of the database are not highly correlated (Pérez-Palacios *et al.*, 2014).

1400 768 In the 3D study of Ávila *et al.* (2018), the MRI analysis initiated with the selection of a ROI of 20 x  
1401 769 20 pixels. The ROIs of all the images of a piece were reconstructed in three dimensions by linear  
1402 770 interpolation methods. Then, the 3D reconstructed MRI of loins were analysed by the three classical  
1403 771 algorithms for texture analysis adapted to work with three-dimensional images. Again, the influence

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1420 772 of the sequence acquisition on the values of the computational texture features was detected. As  
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1422 773 occurred in the study of Perez-Palacios *et al.* (2017) with two-dimensional images, combinations of  
1423 774 SE or GE with any texture algorithm and any regression technique offered precise prediction results  
1424 775 for the physico-chemical parameters of loins. However, Ávila *et al.* (2018), taking into consideration  
1425 776 the computational cost apart from the accuracy of the methodology, proposed the combination of GE-  
1426 777 3D GLCM-IR as the best option.  
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1429 778 In these studies, to validate the optimum procedure achieved, the adjustment between real and  
1430 779 predicted values of the physico-chemical parameters analysed were statistically compared, having  
1431 780 both groups of data the same performance and not being significantly different.  
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1435 781 Another recent study on pork loins has been published by Ávila *et al.* (2019), who used a LF-H<sup>1</sup> MRI  
1436 782 scanner with a hand/wrist coil and SE T1-weighted acquisition sequences to predict physico-chemical  
1437 783 characteristics of fresh loins. The authors of this study considered that conclusions of the previous  
1438 784 works at evaluating the use of MRI to analyse meat might be preliminary, due to the use of  
1439 785 conventional texture descriptors and regressors instead of stronger methods and of optimistic  
1440 786 methodology to measure the performance. In this sense, experiments of this study were developed  
1441 787 with 15 texture descriptors to analyse the MRI (such as Haralick descriptors, local binary patterns,  
1442 788 fractal features, Gabor or wavelet features), in combination to 28 regression techniques to analyse the  
1443 789 data (linear regressors, neural networks, deep learning, support vector machines, regression trees,  
1444 790 ensembles, boosting machines and random forests, among others), having 720 combinations in total.  
1445 791 To guarantee a realistic evaluation, three data partitions (for training, validation and test) were used  
1446 792 instead of the usual two sets (training and test sets), and the dataset was created leaving one meat  
1447 793 piece out, instead of the common random partitioning of the image collection. The test set was  
1448 794 composed by one meat piece, while the images of the remaining meat pieces were divided into the  
1449 795 training and validation sets at random. Good to excellent prediction results were achieved for most  
1450 796 physico-chemical parameters analysed, but there was not possible to set a common combination of  
1451 797 texture vector and regressor that provides accurate correlations for all characteristics tested.  
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1455 798 Most recent studies on MRI and fresh meat have been focused on analysing the effect of  
1456 799 freezing/thawing. Cheng *et al.* (2019) worked with beef *semimembranosus* muscles that were scanned  
1457 800 by using a LF-H<sup>1</sup> and SE acquisition sequences weighted in T1 and T2. The image analysis of this  
1458 801 study consists of measuring the relative intensity of the MRI from samples subjected to 0, 1, 2, 3, 4,  
1459 802 5, 6, 7 freeze-thaw cycles. The obtained data were analysed by ANOVA followed by the Tukey  
1460 803 procedure. The signal intensity of T1 and T2 images significantly decreased as the number freeze-  
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thaw cycles increased. Besides, it was observed a decrease in the brighter of the T1 and T2 images from external to centre regions with the increase of the freeze-thaw cycles.

The assessment of the loss of quality of chicken breast due to freezing/thawing was study by Frelka *et al.* (2019), who used a HF-H<sup>1</sup> MRI with a knee coil. Three different sequences acquisition were tested (3D T1 – rapid GE, proton density – turbo SE and T2 – turbo SE). The analysis of the images was done visually and by measuring the percentage of pixels. The water changes that take place during the freezing/thawing cycles were clearly observed in the images from the three sequence acquisitions. The quantitative analyses based on the percentage of pixels were only done in the images from proton density and T2 sequences, noting differences between unfrozen and frozen/thawed samples, especially in proton density images.

## 6. ADVANTAGES AND DISADVANTAGES OF NON-DESTRUCTIVE TECHNIQUES METHODS

Some of the greatest advantages of NIRs in meat quality evaluation are that it is non-destructive, can be non-contact, and has varying degrees of portability to meet different application needs. The typical halogen light source is easily available, and technological advances are ever increasing compactness, ruggedness, and accuracy, as well as decreasing cost. The technology also has a number disadvantages particularly related to the inhomogeneity and high moisture content of meat. The former requires that intact samples have large or multiple reading areas, and the latter that both sample and ambient temperature remain consistent. Additionally, in an industrial situation where NIRs could be used for monitoring or sorting on a moving belt, samples are not static, due to shape variations the distance from a fixed reading head is inconsistent, and data collection and analysis must be rapid. As discussed above, there has been some progress in several of these areas and the future looks promising.

The main advantage of HSI technology is the ability in predicting the concentration gradient of chemical constituents as spatial distribution, which can be especially useful for visualizing meat quality traits. HSI allows to merge the digital imaging spatial resolution with the chemical information obtainable by point spectroscopy. This is particularly relevant for fat content and distribution modelling, indeed the success in intramuscular fat determination could permit producers to exploit to the maximum level the quality of each meat cut answering the specific needs of the market.

The huge disadvantage of HSI technology is related to the long acquisition time and the large amount of produced data for each single measure. However, simplified instruments (multiband cameras) developed for specific applications could reduce the spectral range to be scanned up to few selected

wavelength, thus minimizing both acquisition time and generated data, which anyway will be fast managed with the proper *ad hoc* chemometric method.

One of the benefits of using NMR is the high sensitivity for distinguishing the different component of the meat, which is due to the strength of the magnetic field and it is dependent on the presence of the magnetic atom on the sample to analyze. Thus, the evaluation of fresh meat, which has a high content of  $^1\text{H}$ , by NMR with antennas for exciting  $^1\text{H}$  spin, achieves accurate results. It is also notable, the low time of analysis of NMR systems, being lower than 2 minutes per sample. However, this technique is quite sensitive to changes of the temperatures, being necessary low room-temperature to have accurate results. In addition, the size of the samples to be analyse should be very small, which require their destruction in some cases.

The use of MRI-computer vision as an alternative methodology to analyse different quality parameters of meat offers the advantage of being a non-destructive, non-invasive, non-radiant and innocuous techniques. Besides, it takes less time than traditional and destructive methods. However, some improvements, mainly on the image acquisition time and software, should be developed in order to work automatically, give the results on-line and fulfil the requirements of the meat industries. This is the only technique be able to take information from the inner of solid samples of medium-large volume, and consequently, it is really interesting for meat pieces such as loins or hams.

## CONCLUSIONS AND FURTHER STUDIES.

The use of NIRs in the laboratory for objective assessment of meat quality is developing, with best results for composition-related measurements. Colour, pH, and drip loss are predicted best when the VIS-NIRs spectra is used. Routine use of NIRs on a production line is still in the future, although possibly a very near future. Technological advances reducing size, cost, and data analysis time are making NIRs more easily available, although new equipment should not be blindly used, but tested for accuracy and precision before being relied upon.

Most of the reported works are feasibility studies mainly developed at laboratory scale, whereas there is a lack of studies proving model's robustness at processing plant level. Thus, more effort is desired to bridge the gap between spectroscopic devices' potential and their implementation as Process Analytical Technology tool, and this way should involve both researchers and industries. Furthermore, few cases present a data fusion approach, which could be considered a strategic way to improve quality attribute prediction by non-destructive techniques. For instance, the combination of HSI and digital image analysis could provide improved models for quality assessment at industrial level.

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868 NMR systems noted for their high sensitivity and accurate in the detection of the component of the  
869 meat and for the analysis speed. The most studies developed with NMR on fresh meat have been  
870 carried out with LF-NMR systems, with CPMG as pulse sequence and T2 as relaxation time. The  
871 results obtained show the ability of this technique to determine main physico-chemical parameters of  
872 fresh meat, and to evaluate the effects of cooking and freezing/thawing.

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873 In the last decade, most studies developed with MRI on fresh meat have been carried with low-field  
874 scanners, to optimize the methodology and predict quality parameters of loins. Nevertheless, the use  
875 of high-field scanners has been reduced in the last years, being mainly applied with classification  
876 purposes. The latest publications on MRI have been focused on evaluating of the effect of  
877 freezing/thawing on meat. The accurate results obtained on these MRI studies allow you to indicate  
878 that the combination of different acquisition parameters, algorithms for analysing the images and data  
879 analysis techniques can be proposed as an alternative methodology to analyse fresh meat with high  
880 reliability in a non-destructive way. Nevertheless, the future work will be to operate as a quality  
881 assessment system in the meat industries, by improving the time for image acquisition and to develop  
882 a software that analyses the images automatically and provides the results on-line.

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**FIGURE CAPTIONS**

**Figure 1.** Illustration of common modes of measurement employed with NIRs.

**Figure 2.** Hyperspectral systems configurations: a) One-Point, 2) Multiple spatial scanning, and c) area/plane scanning.

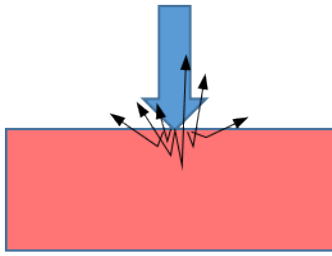
**Figure 3.** NMR spectra of a beef sample by using a HF-NMR system.

**Figure 4.** MRI of hams from pigs fattened with different diets (A: acorn and grass; B: high oleic acid concentrate).

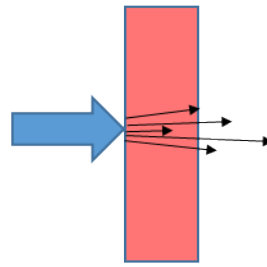


**Figure 1.**

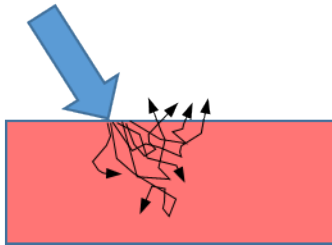
Reflectance/absorbance



Transmittance/absorbance



Interactance



Transflectance

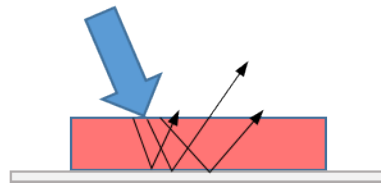
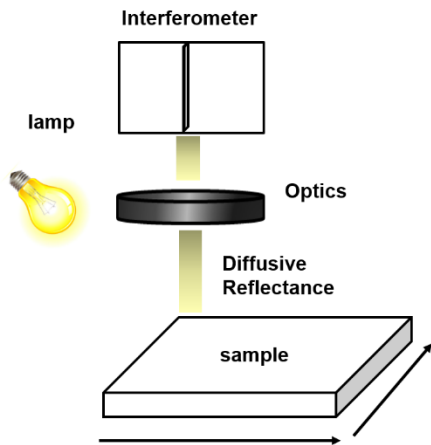
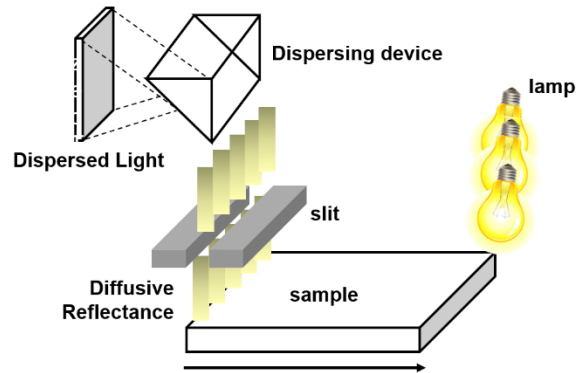


Figure 2.

**a) One-point scanning  
(whisker-broom)**



**b) Multiple spatial position scanning  
(push-broom)**



**c) Plane scan**

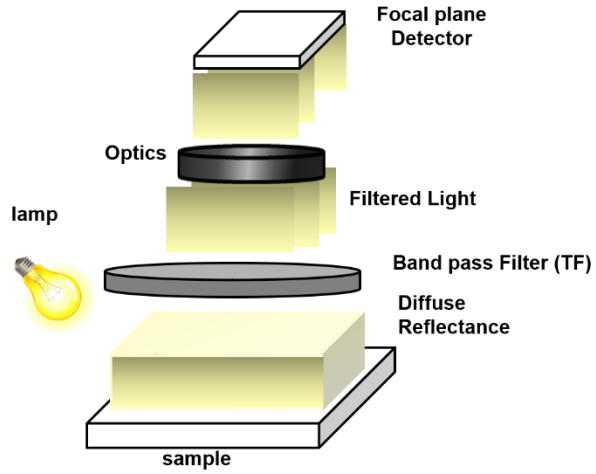
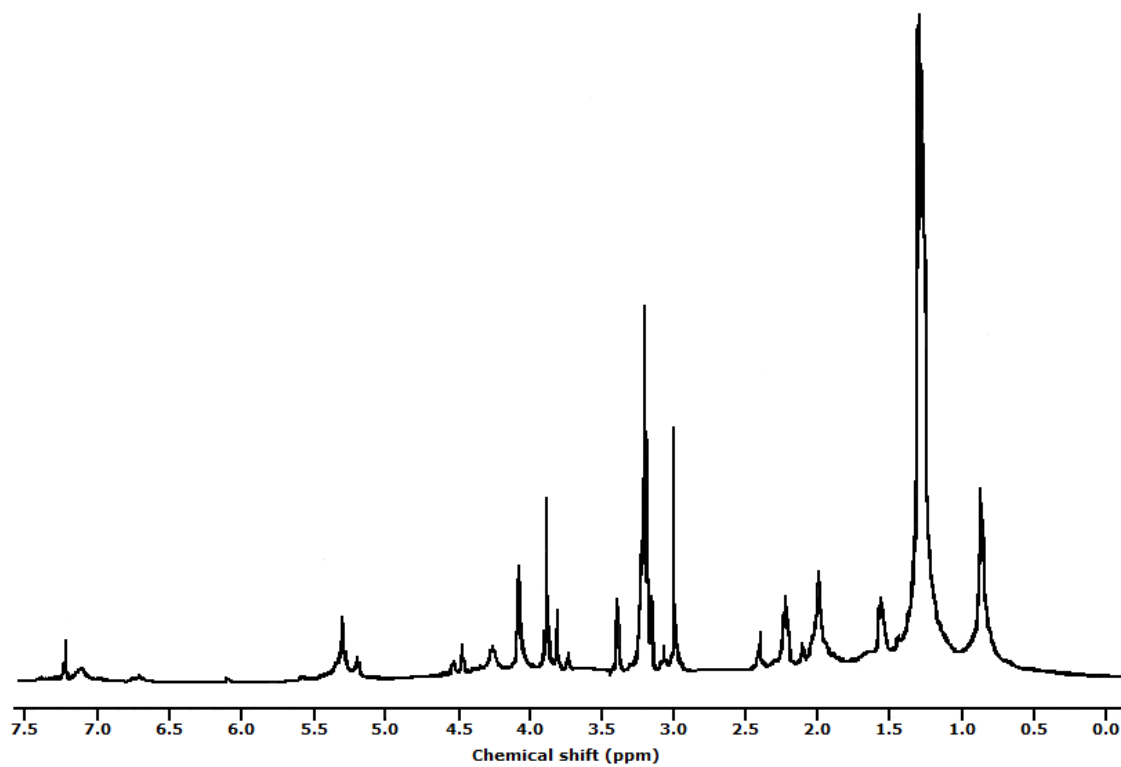


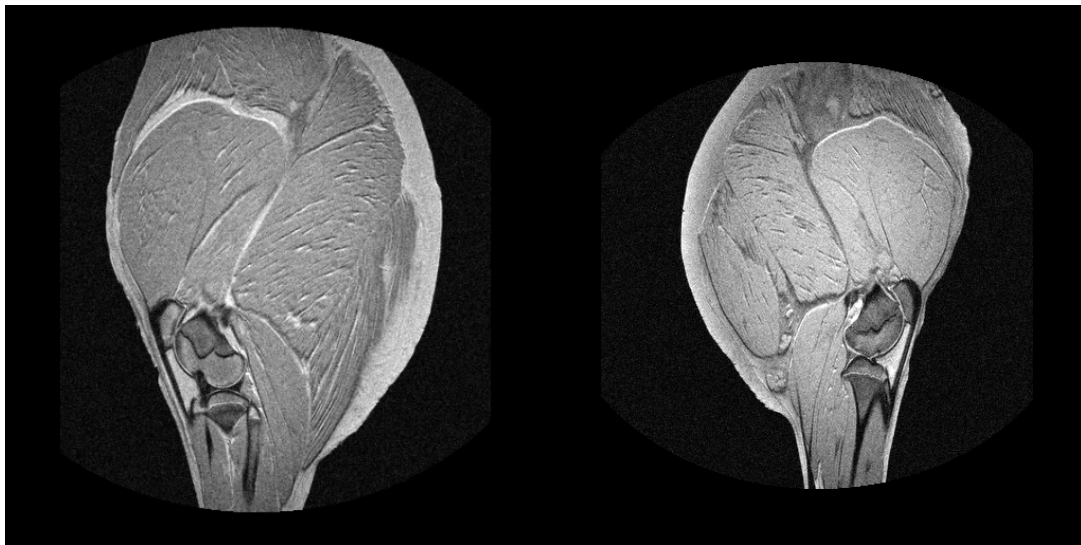
Figure 3..



**Figure 4.**

**A**

**B**



**Table 1.** Summary information about the use of NIRs to analyse fresh meat since 2016.

Meat sample	Sample preparation	Equipment type	Wavelength range (nm)	Data pre-processing	Data Processing	Variable	Calibration performance		Cross validation or Prediction performances		References
beef, pork, chicken	ground	benchtop	400-1700	included: none, smooth with 1st & 2nd der. <sup>a</sup> , normalization, SNV, MSC, median center.	PLSR	<b>Adulteration</b> Full spectrum of known species & mixes	$r_c = 57-88\%$	RMSEC = 0.23-0.51	$r_p = 53-83\%$	RMSEP = 0.24-1.36	Rady & Adedeji, 2018
						Selected wavelengths of known species & mixes	$r_c = 90-96\%$	RMSEC = 0.09-0.26	$r_p = 78-86\%$	RMSEP = 0.17-0.40	
beef, pork	ground	portable; NirScan Nano	900-1700	none, MSC, SNV, 1st or 2nd der.	PLSR	Beef	$r_c^2 = 0.91-0.93$	RMSEC = 5.48-6.05	$r_p^2 = 0.33-0.70$	RMSEP = 13.07-19.72	Nolasco-Perez et al., 2019
						Pork	$r_c^2 = 0.16-0.69$	RMSEC = 11.40-18.67	$r_p^2 = 0.01-0.28$	RMSEP = 20.32-23.90	
beef	intact	portable; ASD FieldSpec	800-2500	wavelet analysis to denoise; smooth, 1st order differential, 2nd order differential, SNV.	PLS	<b>Prediction</b> Hardness (N)	$r_c = 0.74-0.94$	RMSEC = 6.25-10.19	$r_p = 0.60-0.079$	RMSEP = 8.89-10.89	Li, Z. et al., 2016
						Springiness (mm)	$r_c = 0.54-0.93$	RMSEC = 0.69-1.11	$r_p = 0.35-0.71$	RMSEP = 0.91-1.27	
						Chewiness (mJ)	$r_c = 0.69-0.94$	RMSEC = 14.01-25.16	$r_p = 0.52-0.66$	RMSEP = 22.06-27.31	
beef	intact	benchtop; FAT-Analyzer S-7040	920-970	SG smoothing, 2nd der.	PLSR	Adhesiveness (N·mm)	$r_c = 0.44-0.72$	RMSEC = 0.30-0.40	$r_p = 0.22-0.69$	RMSEP = 0.24-0.32	Ishikawa et al. 2017
						Moisture content, over time (g/g·d·s)	-	-	$r_p^2 = 0.81$	RMSEP = 0.34	

pork	intact	benchtop	833-2500	SG smoothing, then 1st or 2nd der. & combined with SNV or MSC; COE.	PLSR	pH	$r^2_c = 69.70-78.10$	RMSEC = 0.09-0.10	$r^2_{cv} = 64.55-70.10$	RMSECV = 0.10-0.11	Li, X. et al., 2016
						CIE L*	$r^2_c = 66.31-81.90$	RRMSEC = 1.72-2.32	$r^2_{cv} = 60.02-11.18$	RMSECV = 1.91-2.51	
						CIE a*	$r^2_c = 17.26-54.55$	RMSEC = 0.96-1.30	$r^2_{cv} = 13.48-31.61$	RMSECV = 1.16-1.32	
						CIE b*	$r^2_c = 40.82-61.46$	RMSEC = 0.51-0.63	$r^2_{cv} = 38.38-45.49$	RMSECV = 0.60-0.64	
pork	intact	benchtop; NIRFlex N-500 (Buchi)		1st der., and normalization		SFA (%)	$r^2_c = 0.96$	SEC=0.52	$r^2_v = 0.92$	SEV = 0.62	Richli et al., 2016
						MUFA (%)	$r^2_c = 0.83$	SEC=0.63	$r^2_v = 0.76$	SEV = 0.67	
						PUFA (%)	$r^2_c = 0.94$	SEC=0.43	$r^2_v = 0.92$	SEV = 0.49	
						IV	$r^2_c = 0.98$	SEC=0.68	$r^2_v = 0.98$	SEV = 0.67	
						Fat (%)	$r^2_c = 0.86$	SEC=1.57	$r^2_v = 0.62$	SEV = 1.97	
						Moisture (%)	$r^2_c = 0.71$	SEC=1.71	$r^2_v = 0.64$	SEV = 1.67	
pork	lean - homogenized; fat - intact	benchtop; FOSS NIR Systems 6500	400-2500	1100-2400nm: 1st der., SNVD	PLSR	IMF (%). Lean	$r^2_c = 0.98-0.99$	SEC = 0.14- 0.23	$r^2_{cv} = 0.95-0.97$	SECV = 0.25-0.30	Prevolnik Povse et al., 2017
						Moisture (%). Lean	$r^2_c = 0.90-0.91$	SEC = 0.39- 0.45	$r^2_{cv} = 0.63-0.82$	SECV = 0.65-0.75	
						Protein (%). Lean	$r^2_c = 0.45-0.92$	SEC = 0.48- 0.92	$r^2_{cv} = 0.28-0.81$	SECV = 0.73-1.05	
						SFA (g/100g). Fat; Lean	$r^2_c = 0.95; 0.98$	SEC = 0.44; 0.26	$r^2_{cv} = 0.83; 0.58$	SECV = 0.79; 1.33	

						MUFA (g/100g). Fat; Lean	$r^2_c = 0.98; 0.18$	SEC = 0.35; 2.39	$r^2_{cv} = 0.91; 0.11$	SECV = 0.70; 2.54	
						PUFA (g/100g). Fat; Lean	$r^2_c = 0.97; 0.78$	SEC = 0.32; 1.51	$r^2_{cv} = 0.89; 0.53$	SECV = 0.57; 2.21	
						n-3 PUFA (g/100g). Fat; Lean	$r^2_c = 0.96; 0.62$	SEC = 0.04; 0.12	$r^2_{cv} = 0.83; 0.55$	SECV = 0.08; 0.13	
						n-6 PUFA (g/100g). Fat; Lean	$r^2_c = 0.97; 0.77$	SEC = 0.29; 1.43	$r^2_{cv} = 0.89; 0.52$	SECV = 0.51; 2.08	
						n-6/n-3 PUFA (g/100g). Fat; Lean	$r^2_c = 0.80; 0.12$	SEC = 0.48; 1.44	$r^2_{cv} = 0.30; 0.02$	SECV = 0.89; 1.52	
pork	intact	portable; Sup-NIR-1520	1000-1799	SG 1st der., SG smoothing, OSC	PLS	Cholesterol (mg/100g)	$r_c = 0.91$	SEC=2.56	$r_p = 0.66$	SEP = 4.99	Wang, H. et al., 2017
pork	Intact	benchtop; FOSS NIRSystem XDS rapid content analyzer.	400-2500	Divided into 400-1850 and 780-1850 then SNV applied to each region.	PLSR	pH (400-1850 nm)	-	-	$r^2_{cv} = 0.28$	RMSEV = 0.07	Andersen et al., 2018a
						EZ-Drip (%) (780-1850 nm)	-	-	$r^2_{cv} = 0.06$	RMSEV = 1.69	
						Vacuum drip (%) (780-1850 nm)	-	-	$r^2_{cv} = 0.12$	RMSEV = 1.00	
						IMF (%) (780-1850 nm)	-	-	$r^2_{cv} = 0.57$	RMSEV = 0.11	
pork	intact	portable; ASD LabSpec4	350-1900	One of: SG smoothing and 1st or 2nd order der., SNV, SNVD,	PLSR	Subjective firmness score	$r^2_c = 0.74-0.80$	RMSEC=0.41-0.48	$r^2_v = 0.44-0.72$	RMSEP = 0.45-0.66	Soladoye et al., 2018

				SNVD plus SG smoothing and 1st or 2nd order der.		Bar bend angle	$r^2_c = 0.73-0.99$	RMSEC=0.97-14.76	$r^2_v=0.51-.072$	RMSEP = 13.71-17.99	
lamb	homogenized	benchtop; FOSS NirSystems 6500	400-2500	Included scatter correction, der.s, smoothing	PLS, PCR	TBARS	$r^2_c = 0.80-0.85$	RMSEC=0.32-0.38	$r^2_{cv} = 0.77-0.80$	RMSECV = 0.37-0.41	Ripoll et al., 2018
lamb	intact	portable; ASD Labspec Pro	500-2000	SG 1st der.	PLS	24 h pH	$r^2 = 0.37-0.70$	-	-	SECV = 0.10	Knight et al., 2019
						IMF (%)	$r^2 = 0.55-0.60$	-	-	SECV = 0.50-0.70	
						Shear force (N)	$r^2 = 0.20-0.34$	-	-	SECV = 7.00-8.90	
<hr/>											
<b>Analysis Testing</b>											
pork	intact	custom; in-house	350-2500	SG smoothing, SNV	PLSR, with CARS or siPLS	Moisture (%)	$r_c = 0.95; 0.95$	SEC = 0.58; 0.60	$r_p = 0.91; 0.90$	SEP = 0.38; 0.40	Wang et al., 2016
						Cook loss (%)	$r_c = 0.95; 0.93$	SEC = 0.01; 0.02	$r_p = 0.92; 0.91$	SEP = 0.02; 0.03	
						Tenderness (N)	$r_c = 0.94; 0.94$	SEC = 3.59; 3.99	$r_p = 0.90; 0.88$	SEP = 6.90; 7.26	
						siPLS					
pork	intact	portable; custom; in-house	350-2500	Included: none, SNV, SG 1st der., or combinations; model clustering	PLSR, using selected wavelengths	L*	$r_c = 0.93$	SEC = 0.72	$r_p = 0.93$	SEP = 1.16	Wang et al., 2018b
						a*	$r_c = 0.97$	SEC = 0.46	$r_p = 0.96$	SEP = 0.64	
						b*	$r_c = 0.97$	SEC = 0.28	$r_p = 0.96$	SEP = 0.42	
						pH	$r_c = 0.95$	SEC = 0.05	$r_p = 0.95$	SEP = 0.06	
						TVB-N (mg/100g)	$r_c = 0.97$	SEC = 2.04	$r_p = 0.95$	SEP = 2.20	
						Fat (%)	$r_c = 0.96$	SEC = 0.13	$r_p = 0.94$	SEP = 0.18	



						Protein (%)	$r_c = 0.96$	SEC = 0.32	$r_p = 0.96$	SEP = 0.33	
Beef fat and lean trimmings	minced	custom; in-house; multi-point	1515-1900	SNV, SG smoothing.	PLSR	<b>Equipment Testing</b>					
						Static	$r_c^2 = 0.95$	SEC = 5.93	$r_p^2 = 0.82$	SEP = 3.05	Dixit et al., 2016a
						Slow motion	$r_c^2 = 0.9$	SEC = 5.62	$r_p^2 = 0.92$	SEP = 3.98	
						Fast motion	$r_c^2 = 0.95$	SEC = 5.99	$r_p^2 = 0.85$	SEP = 3.97	
fat beef trimmings & lean beef	minced	custom; in-house; multi-point	1515-2000	SNV, SG smoothing, MSC, 1st & 2nd order der., SNV plus SG smoothing.	PLSR	Fat (%)	$r_c^2 = 0.98-0.99$	RMSEC = 2.60-4.25	$r_p^2 = 0.95-0.99$	RMSEP = 2.79-5.67	Dixit et al., 2016b
						Moisture (%)	$r_c^2 = 0.98-0.99$	RMSEC = 1.69-2.90	$r_p^2 = 0.94-0.98$	RMSEP = 2.75-4.62	
						Protein (%)	$r_c^2 = 0.96-0.98$	RMSEC = 0.96-1.36	$r_p^2 = 0.90-0.95$	RMSEP = 1.56-2.28	
						Ash (%)	$r_c^2 = 0.98-0.99$	RMSEC = 0.02-0.04	$r_p^2 = 0.95-0.99$	RMSEP = 0.03-0.06	
Beef fat and lean trimmings	minced	custom; in-house; multi-point	1515-2050	SNV, SG smoothing, MSC, 1st & 2nd order der., SNV + SG smoothing	PLSR	Fat (%)	$r_c^2 = 0.99$	SEC = 1.94	$r_p^2 = 0.78-0.94$ ; baseline correction = 0.77-0.83	SEP = 6.75-8.61; baseline correction = 11.12-11.40	Dixit et al., 2016c
						Moisture (%)	$r_c^2 = 0.99$	SEC = 1.76	$r_p^2 = 0.76-0.94$ ; baseline correction = 0.74-0.84	SEP = 4.96-6.75; baseline correction = 8.49-9.22	
						Protein (%)	$r_c^2 = 0.99$	SEC = 0.43	$r_p^2 = 0.85-0.91$ ; baseline correction = 0.85-0.86	SEP = 1.71-2.27; baseline correction	

						Ash (%)	$r^2_c = 0.98$	SEC = 0.04	$r^2_p = 0.75-0.83$ ; baseline correction = 0.57-0.75	= 2.31-2.54 SEP = 0.11-0.13; baseline correction = 0.14-0.19	
pork myofibril extracts	extracted	benchtop - FOSS NIRSystem XDS rapid content analyzer	400-2500	divided into 400-700, 1100-1700, and 1700-2400, then EMSC on each.	PLSR	pH (1100-1700 nm)	-	-	$r^2_{cv} = 0.14$	RMSECV = 0.42	Andersen et al., 2017
pork	intact	custom; in-house	350-2500	Combinations of SG smoothing, SNV, and 1st der.		pH	$r_c = 0.84-0.94$	SEC=0.05-0.18	$r_p = 0.79-0.91$	SEP = 0.08-0.23	Wang, W. et al., 2017
						TVB-N (mg/100g)	$r_c = 0.93-0.94$	SEC=2.06-2.33	$r_p = 0.90-0.92$	SEP = 2.62-2.86	
beef	intact	in-house, portable	700-1050	2nd der., SG smoothing	PLS	MUFA	-	-	Master $r^2_{cv} = 0.79$ ; Slave $r^2_p = 0.69$	Master SECV(%) = 1.69; Slave SEP(%) = 2.03	Piao et al., 2018
						Oleic	-	-	Master $r^2_{cv} = 0.71$ ; Slave $r^2_p = 0.64$	Master SECV(%) = 1.86; Slave SEP(%) = 2.06	
						SFA	-	-	Master $r^2_{cv} = 0.81$ ; Slave $r^2_p = 0.67$	Master SECV(%) = 1.67; Slave	

SEP(%) =  
2.19

pork myofibril extracts	extracted	Benchtop - FOSS NIRSystem XDS rapid content analyzer	400-2500	Wet samples: divided into 400- 900, 1100-1700, and 1700-2350 nm, then EMSC on each. Dry samples: EMSC in 1100-2500 nm only.	PLSR	Degree of proteolysis; dried	-	-	$r^2_{cv} = 0.74$	RMSECV = 1.42	Andersen et al., 2018b
						Degree of proteolysis; liquid	-	-	$r^2_{cv} = 0.10$	RMSECV = 2.61	

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\* **1st or 2nd order der.:** First or second order derivative; **CARS:** Competitive Adaptive Reweighted Sampling; **COE:** Constant Offset Elimination; **MSC, E-:** Multiplicative Scatter Correction, Extended-; **OSC:** Orthogonal Signal Correction; **PC1:** Principal Component 1; **PCR:** Principal Component Regression; **PLS, -R, si-:** Partial Least Squares, -regression, synergy interval-; **r<sub>c,p</sub>:** coefficient of correlation for -calibration, -prediction; **r<sup>2</sup><sub>c, cv, p, v</sub>:** coefficient of determination for -calibration, -cross validation, -prediction, -validation; **RMSE, -C, -CV, -P, -V:** Root Mean Square Error of -calibration, -cross validation, -prediction, -validation; **SE, -C, -CV, -P, -V:** Standard Error of -calibration, -cross validation, -prediction, -validation; **SG:** Savitzky Golay; **SNV, -D:** Standard Normal Variate, -detrend; **CIE:** Commission internationale de l'éclairage; **GC:** Gas Chromatography; **IMF:** Intramuscular fat; **IV:** Iodine Value; **MUFA:** Monounsaturated Fatty Acids; **PUFA:** Polyunsaturated Fatty Acids; **SDS-PAGE:** Sodium Dodecyl Sulfate-Polyacrylamide Gel Electrophoresis; **SFA:** Saturated Fatty Acids; **TBARS:** Thiobarbituric Acid Reactive Substances; **TPA:** Texture Profile Analysis; **TVB-N:** Total Volatile Base Nitrogen.

**Table 2.** Use of HSI in the analysis of fresh meat in the last decade.

Meat sample	Quality Parameter	Type od HSI	Data analysis	Model evaluation				Reference
				Calibration or Cross-validation		Prediction		
				Sample	Accuracy	Sample	Accuracy	
Chicken	Moisture	Line scan system (400–1000 nm)	PLS	52	$R^2_{CV} = 0.9$ SECV = 0.55 %	20	$R^2_p = 0.94$ SEP = 0.71 %	Kanpal et al. (2013)
	TVB-N	Line scan system (450–950 nm)	BPNN	50	$R^2_c = 0.94$ RMSEC= 3.95 mg/100 g	25	$R_p = 0.75$ RMSEP = 6.39 mg/100 g	Khulal et al. (2017)
	TBARS	Line scan system (400–1000 nm)	SPA	95	$R_{cv} = 0.83$ RSECV = 0.14 mg/100 g	63	$R_p = 0.80$ RMSEP = 0.16 mg/100 g	Xiong et al. (2015b)
Pork	Moisture	Line scan system (900–1700 nm)	PLS	80	$R^2_{CV} = 0.88$ SECV = 0.76 %	40	$R^2_p = 0.91$ SEP = 0.62 %	Barbin et al. (2013)
	IMF	Line scan system (900–1700 nm)	PLS	13	$R^2_{CV}=0.94$ RMSECV=0.17 %	7	$R^2_p=0.97$ RMSEP=0.17 %	Liu and Ngadi (2014)
	IMF	Line scan system (900 -1700 nm)	MLR	18	$R_c = 0.97$ RMSEC=0.89 %	6	$R_p = 0.90$ RMSEP = 0.92 %	Huang et al., (2017)
	TVB-N	Line scan system (400-1000 nm)	LS - SVM	48	-	24	$R_p = 0.95$ RMSEP = 1.86 mg/100 g	Gou et al. (2015)
	TVB-N	Line scan system (400-1000nm)	BP NN - Adaboost	48	$R_c = 0.94$ RMSEC = 5.30 mg/100 g	24	$R_p = 0.93$ RMSEP = 5.52 mg/100 g	Li et al. (2015)
	TVB-N	Fluorescence system (400–1000nm)	LS - SVM	68	$R^2_c = 0.97$ RMSEC 1.88 %	22	$R^2_p = 0.97$ RMSEP = 1.90 %	Lee et al. (2018)
	K - value	Line scan system (328–1115 nm)	PLS	140	$R^2_{CV}=0.95$ RMSECV=3-5	70	$R^2_p = 0.92$ RMSEP = 4	Cheng (2016)
	pH	Line scan system (900–1700 nm)	PLS	50	$R^2_{CV}=0.87$ RMSECV= 0.11	25	$R^2 = 0.90$ RMSEP = 0.09	Barbin et al. (2012)
	Colour (L*, a*, b*)	Line scan system (900–1700 nm)	PLS	50	L*, $R^2_{CV}=0.93$ RMSECV= 1.36 a*, R2CV=0.75 RMSECV= 0.67 b*, R2CV=0.89 RMSECV= 0.49	25	L*, $R^2_p=0.90$ RMSEP= 1.63 a*, $R^2_p=0.72$ RMSEP= 0.78 b*, $R^2_p=0.85$ RMSEP= 0.50	Barbin et al. (2012)
Beef	Moisture	Line scan system (897–1753 nm)	PLS	54	$R^2_{CV}=0.89$ SECV=0.51 %	27	$R^2_p = 0.89$ SEP = 0.46 %	Kamruzzaman et al. (2016)
	Moisture	Line scan system (880 - 1720 nm)	EMCV-PLS	36	$R^2_{CV}=0.99$ RMSECV=0.72 %	9	$R^2_p=0.99$ RMSEP = 0.64 %	M. Zhao et al. (2017)
	Fat	Line scan system (970–2500 nm)	PLS	60	$R^2_{CV} = 0.89$ RMSECV = 4.87 %	30	$R^2_p = 0.90$ RMSEP = 4.81 %	Kobayashi et al. (2010)

	Fat	Multispectral system (15 wavelength between 760 - 1040 nm)			$R_{CV} = 0.98$ RMSECV=3.0%		$R_p=0.99$ RMSEP=0.6 %	Wold et al. (2011)
	Fat	Area imaging system equipped with a filter wheel (1000–2350 nm)	PLS	126	$R^2_{CV} = 0.85$ RMSECV =5.85	66	$R^2_p = 0.739$ RMSEP =5.15	Kobayashi et al. (2012)
	Fat	Line scan system (897–1753 nm)	PLS	54	$R^2_{CV} = 0.88$ SECV = 0.66 %	27	$R^2_p=0.84$ SEP=0.65 %	ElMasry et al. (2013)
	Fat	Line scan system (400–1000 nm)	EDM	24	$R^2=0.96$ RMSECV = not reported	-	-	S. Lohumi et al. (2016)
	Fat	Line scan system (880 - 1720 nm)	EMCV - PLS	36	$R^2_{CV} = 0.99$ RMSECV = 0.79 %	9	$R^2_p = 0.98$ RMSEP = 0.73%	Zhao et al. (2017)
	UFAs	Area imaging system equipped with a filter wheel (1000–2350 nm)	PLS (filter based method)	126	$R^2_{CV} = 0.86$ RMSECV = 4.11 %	66	$R^2_p = 0.64$ RMSEP =3.95 %	Kobayashi et al. (2012)
	Oleic acid	Area imaging system equipped with a filter wheel (1000–2350 nm)	PLS (filter based method)	126	$R^2_{CV} = 0.85$ RMSECV =3.57 %	66	$R^2_p = 0.71$ RMSEP =3.13 %	Kobayashi et al. (2012)
	SFAs	Line scan system (970–2500 nm)	PLS	61	$R^2_{CV} = 0.84$ RMSECV =1.78 %	31	$R^2_p = 0.87$ RMSEP =1.69 %	Kobayashi et al. (2010)
	UFAs	Line scan system (970–2500 nm)	PLS	62	$R^2_{CV} = 0.88$ RMSECV =3.32 %	32	$R^2_p = 0.89$ RMSEP =3.41 %	Kobayashi et al. (2010)
	Protein	Line scan system (897–1753 nm)	PLS	54	$R^2_{CV} = 0.88$ SECV = 0.31 %	27	$R^2_p = 0.86$ SEP = 0.29 %	ElMasry et al. (2013)
	pH	Line scan system (900–1700 nm)	PLS	321	$R^2_{CV} = 0.73$ RMSECV = 0.06	-	-	ElMasry et al., (2012)
	Colour (L*, a*, b*)	Line scan system (900–1700 nm)	PLS	321	L*, $R^2_{CV} = 0.88$ RMSECV = 1.21 b*, $R^2_{CV} = 0.81$ RMSECV = 0.57	-	-	ElMasry et al. (2012)
	Colour (L*, a*, b*)	Line scan system (400–1100 nm)	MLR	65	L*, $R^2_{CV} = 0.96$ SECV = 0.61 a*, $R^2_{CV} = 0.96$ RMSECV = 0.75 b*, $R^2_{CV} = 0.97$ RMSECV = 0.19	-	-	Wu et al. (2012)
<b>Lamb</b>	Moisture	Line scan system (900-1700 nm)	UVE -SPA - MLR	84	$R_{CV} = 0.91$ RMSEC=0.56 %	42	$R_p = 0.92$ RMSEP = 0.58 %	Pu et al. (2014)
	Fat	Line scan system (900-1700 nm)	UVE -SPA - MLR	84	$R_{CV} = 0.95$ RMSECV=0.38 %	42	$R_p = 0.98$ RMSEP = 0.36 %	Pu et al. (2014)
	IMF	Line scan system (550 - 1700 nm)	GPR	1628	$R^2_{CV} = 0.75$ RMSEC = 0.43 %	829	$R^2_p = 0.72$ RMSEP = 0.45 %	Craig et al. (2017)

Protein	Line scan system (900-1700 nm)	UVE -SPA - MLR	84	$R_c = 0.80$ RMSEC=0.36 %	42	$R_p = 0.67$ RMSEP 0.41 %	Pu et al. (2014)
pH	Line scan system (900-1700 nm)	PLS	201	$R_{CV} = 0.65$ RMSEC=0.085	-	-	Kamruzzaman et al. (2012b)
Colour (L*)	Line scan system (900-1700 nm)	PLS	201	$R_{CV} = 0.91$ RMSEC=1.32	-	-	Kamruzzaman et al. (2012b)

\* **IMF**: Intramuscular fat; **SFA**: Saturated Fatty Acids; **TBARS**: Thiobarbituric Acid Reactive Substances; **TVB-N**: Total Volatile Base Nitrogen; **UFA**: Unsaturated Fatty Acids; **BP NN**: back propagation neural network; **EDM**: Euclidean distance measure; **EMCV**: ensemble Monte Carlo variable selection; **GPR**: Gaussian process regression; **MLR**: Multiple Linear Regression; **PLS**: Partial Least Squares regression; **LS – SVM**: Least Square - Support Vector Machine; **UVE-SPA-MLR**: **UVE-SPA-MLR**: Uninformative Variable Elimination - Successive Projections Algorithm - Multiple Linear Regression;  $R_{c,cv,p}$ : coefficient of correlation for -calibration, -cross-validation, -prediction;  $R^2_{c,cv,p}$ : coefficient of determination for -calibration, -cross-validation, -prediction; **RMSE, -C, -CV, -P**: Root Mean Square Error of -calibration, -cross-validation, -prediction; **SEP**: Standard Error of Prediction.

**Table 3.** Summary information about the use of NMR to analyse fresh meat in recent years.

Meat sample	Objective	Type of System	Acquisition sequences	Data analysis techniques	Main results	Reference
Pork	Relationship between cooking temperature and the water distribution within cooked pork loin.	LF	T2	PLS	High correlation coefficient on some sensory attributes (Juiciness, Tenderness)	Bertram et al., 2005
	Relationship between water holding capacity and the water distribution of cooked pork meat.	LF	T2	Correlation coefficients	High correlations as a function of different days of aging	Straadt et al., 2006
Chicken	Determining the effect of the water on frozen and thawed chicken.	LF	T2	Correlation coefficients	High correlation for water activity and cooking loss	Li et al., 2014
	Determining the water activity in chicken	LF	T2	Correlation coefficients	High correlation between water activity and T2 intensity signal.	Venturi et al., 2007
	Monitoring the cooking loss and the moisture percentage during the cooking of chicken meat.	HF	T2	ANOVA and correlation coefficients	High correlation between moisture and T2 intensity signal	Shaarani et al., 2006
Beef	Determining the moisture content of beef meat.	LF	T2	PLS and PCA	High correlation between moisture and T2 intensity signal	Pereira, and Colnago, 2012
	Discriminating between beef meat and horse meat on meat mixing.	LF	T1 and T2	PCA and Naïve-Bayes	High percentage of samples with different meat mixing were correctly classified	Jakes et al., 2015
Rabbit	Monitoring the effect of thawing and post-thawing on the quality information from frozen rabbit meat.	LF	T2	ANOVA and correlation coefficients	High correlation coefficients	Jia et al., 2017
Duck	Discriminating duck samples as function of age and its quality	HF	T1	PCA and PLS	High percentage of correct classification for duck samples.	Liu et al., 2013

\* **HF**: High Field; **LF**: Low Field; **PLS**: Partial Least Square; **ANOVA**: Analysis of Variance; **PCA**: Principal Component Analysis

**Table 4.** Summary information about the use of MRI to analyse fresh meat in the last decade.

Meat sample	Main goals	Type of meat	Methodology				Main results	Reference
			Scanner - coil	Sequence acquisition	Image analysis	Data analysis		
Ham	Classifying as a function of pig diet	Ham	HF – H <sup>1</sup> , body coil	SE - T1	Contour Active ROI Texture algorithms	ANOVA Pearson`s Correlation PCA	Visual differences Statistical differences in computational texture	Pérez-Palacios <i>et al.</i> , 2011b
Ham	Classifying as a function of salting stage Prediction of salt content	Ham	HF – H <sup>1</sup> , body coil	SE - T1	Contour Active ROI Texture algorithms	Data mining (OneR, J48 decision tree MLR)	High percentages of correct classification Very good correlation coefficients of prediction	Caballero <i>et al.</i> , 2016a
					ROI Texture algorithms	Data mining (MLR, IR)	SE – GLCM – MLR (best option for prediction)	Pérez-Palacios <i>et al.</i> , 2017
				SE - T1 GE - T1 T3D - T1	Fractals algorithms	Data mining (MLR, IR)	SE – OPFTA - MLR (best option for prediction)	Caballero <i>et al.</i> , 2017a
Pork Loins	Optimization of the methodology Prediction of quality parameters	Pork Loins	LF - H <sup>1</sup> , hand/wrist coil		Interpolation and 3D reconstruction 3D Texture algorithms	MLR and IR	GE - 3D GLCM - IR (best option for prediction)	Ávila <i>et al.</i> , 2018b
				SE - T1	A collection of 15 texture features from different algorithms	A collection of 28 regressors	Any option provides the best result for all attributes tested. All combinations achieved good to excellent correlation for most parameters analysed.	Ávila <i>et al.</i> , 2019






Pork carcass	Determining the lean meat percentage	Pork carcass	HF – H <sup>1</sup> , large body coil	GE - T1	Interactive segmentation function for separating muscle	Stepwise regression	Accurate prediction	Bernau <i>et al.</i> , 2015
Beef	Visualization and prediction of intramuscular fat distribution	Beef	HF – H <sup>1</sup> , head coil	SE - T1	Thresholding	Correlation coefficients	High correlation coefficients between real and predicted data	Lee <i>et al.</i> , 2015
Chicken breast	Assessment of drip after freeze/thaw	Chicken breast	HF – H <sup>1</sup> , knee coil	3DT1 - rapid GE Proton density - turbo SE T2 - turbo SE	-	-	MRI showed freeze/thaw changes, especially when using 3DT1 - rapid GE	Frelka <i>et al.</i> , 2019
Beef	Investigating the water distribution changes in frozen/thawed samples	Beef	LF – H <sup>1</sup>	SE - T1 SE – T2	Signal intensity measure	-	T1 and T2 MRI displayed the decrease of water in frozen/thawed samples	Cheng <i>et al.</i> , 2019

\* **HF**: High Field; **LF**: Low Field. Sequence Acquisition: **SE**: Spin Echo; **GE**: Gradient Echo; **T3D**: Turbo 3D. Image Analysis: **ROI**: Region of Interest. Data Analysis: **ANOVA**: One-Way Analysis of Variance; **PCA**: Principal Component Analysis; **MLR**: Multiple Linear Regression; **IR**: Isotonic Regression. Main Results: **GLCM**: Gray Level Co-Occurrence Matrix; **OPFTA**: One-Point Fractal Texture Algorithm.

**Declaration of interests**

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

The authors declare the following financial interests/personal relationships which may be considered as potential competing interests:

				<p>PEREZ PALACIOS MARIA TRINIDAD - 760313225</p> <small>Firmado digitalmente por PEREZ PALACIOS MARIA TRINIDAD - 760313225 Nombre de reconocimiento (DN): c=ES, serialNumber=DN:CN=760313225, givenName=MARIA TRINIDAD, sn=PEREZ PALACIOS, ou=PEREZ PALACIOS MARIA TRINIDAD - 760313225 Fecha: 2019.12.17 12:03:39 +01'00'</small>
Teresa Antequera	Daniel Caballero	Silvia Grassi	Bethani Uttaro	Trinidad Perez-Palacios

## **AUTHOR CONTRIBUTION**

**Teresa Antequera:** investigation and methodology in the MRI section; review; conceptualization; administration; supervision.

**Daniel Caballero:** investigation, methodology, writing and editing the NMR section; review.

**Silvia Grassi:** investigation, methodology, writing and editing the HSI section; review.

**Bethany Uttaro:** investigation, methodology, writing and editing the NIR section; review.

**Trinidad Perez-Palacios:** investigation, methodology, writing and editing the MRI section; review; conceptualization; administration; supervision.