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**Measurement of ripening of raspberries (*Rubus idaeus* L) by near infrared
and colorimetric imaging techniques**

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ABSTRACT

This work includes the study of 168 samples of raspberries ‘Glen Lyon’, representative of the whole maturation period, by colorimetric and near infrared (NIR) imaging techniques, as well as the quantification of total phenols, total anthocyanins and antioxidant activity by chemical methods.

Samples shown significant differences depending on the maturation stage regarding CIELAB colour parameters and total anthocyanins content, mainly among the last steps. The application of Partial Least Squares Regression allowed predicting the chemical features from image analysis data, with coefficients of determination (R^2) up to 0.75. The best prediction was for total anthocyanins when including colorimetric data. The proposed methodology could become a reference method for assessing important quality attributes of raspberries. Moreover, it may be useful for industries interested in implementing rapid and accurate automatic inspection.

KEYWORDS

Raspberry ‘Glen Lyon’ (*Rubus idaeus* L); Image Analysis; Hyperspectral Image Analysis; Phenolics; Partial Least Squares Regression

1. Introduction

Raspberry (*Rubus idaeus*) is a fruit native to Europe and northern Asia. It is a good source of compounds, with recognized bioactive properties, such as phenolic compounds, vitamins and carotenoids (Kalt et al. 1999; Pantelidis et al. 2007). Raspberry 'Glen Lyon' produces very glossy orange-red fruit of medium size and excellent skin firmness and shelf life. This cultivar has good resistance to Spur Blight (*Didymella applanata* infection) and is a very heavy yielding variety when grown on fertile soils with good management (Jennings 2002).

The cultivation of berries in South-western Spain has experienced a huge development within the last 30 years, producing high-quality fruits with a great economic repercussion mainly due to exportations. This fact increases the need of developing rapid techniques for monitoring quality parameters along the maturation, when the biosynthesis of bioactive compounds such as anthocyanins occurs (Dobson et al. 2012). Anthocyanins are pigments accounting for the brilliant red, purple and blue colours of many fruits, such as berries and red grapes, as well as vegetables and derived food products (Gordillo et al. 2012).

Colour and appearance are features that define the quality of food products, being the first attributes perceived in tasting and, frequently, the only criterion for consumer's acceptance. That is why producers take this issue so seriously (Fernández-Vázquez et al. 2011). In this respect, Image Analysis has revolutionised the way of evaluating appearance in food (Sun 2004). Computing advances every day, producing faster and capable systems that broaden the scope of application in food industry. Because of the unceasing rise of hygiene and safety standards, the need for objective quality determination in food products continues growing (Brosnan and Sun 2004). Foodstuff usually have

peculiar sizes and shapes, and Digital Image Analysis is the best way to satisfy the requirements of measuring heterogeneous samples in an objective manner. Moreover, it allows measuring not only colour but also other features related to appearance, such as texture or heterogeneity, which do not vary the colour but affect how the human eye perceive it (Valous et al. 2009). Colour can be measured by Tristimulus Colorimetry and expressed in terms of CIELAB colour space (1976 $L^*a^*b^*$ colour space). It is an international standard for colour measurements, recommended by the Commission Internationale de l'Eclairage (CIE) in 1976. Within CIELAB, L^* is a qualitative attribute of relative luminosity, and ranges between black ($L^*=0$) and white ($L^*=100$). Coordinate a^* takes positive values for reddish colours and negative values for greenish ones. In the same way, b^* takes positive values for yellowish colours and negative for bluish ones. a^* and b^* become to C^*_{ab} (chroma) and h_{ab} (hue) when are converted to polar coordinates (Figure 1), which are more useful parameters with meanings easier to understand according to the human perception. Taking into account the CIE guidelines, the colour measurement includes the complete visible spectrum. However, there are several studies that uses simple RGB digital cameras for colorimetric purposes (Hutchings 2002; León et al. 2006; Fernández-Vázquez et al. 2011; Rodríguez-Pulido et al. 2013).

Beyond the colour, near infrared (NIR) hyperspectral imaging is a technique that generates a spatial map of spectral information, making it a useful tool in many applications. These images, called hypercubes, are three-dimensional data matrices where two axes (x and y) represent the spatial coordinates, while the third (λ) axis portrays the spectral dimension. Hypercubes are easier to understand if we envision a battery of images of the same scene, each one

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representing the reflectance at one wavelength in greyscale. This way, hyperspectral imaging covers the potential of NIR spectroscopy and the versatility of computer vision.

Multivariate statistic techniques are essential for assessing the relationship between instrumental data and physicochemical properties of food products. Partial least squares regression (PLSR) is a common procedure used to relate a large number of independent variables, such spectra or colorimetric attributes, to one or few response variables (Kemsley et al. 1996; Esquerre et al. 2009). It is very effective in spectral analysis since it reduces a great number of redundant information (Lin et al. 2009; Pojić and Mastilović 2012).

The objective of this work was the overall study of the Glen Lyon raspberries maturation (from very early until overripe stages) by both colorimetric and NIR imaging techniques, assessing the capability of these techniques to predict the presence of some interesting compounds (phenols, anthocyanins) in these berries. The study considered one berry per sample to avoid possible errors of considering samples that include unequal berries in each maturation level group. To the best of our knowledge, there are not previous articles studying these berries individually, from very early stages of maturation, and merging these two kind of imaging techniques.

2. Material and Methods

2.1 Samples

Description and extraction procedure

One hundred and sixty eight berries of Glen Lyon raspberry cultivar grown in South-western Spain (37.2° N, 6.5° W) and having very heterogeneous stages of

1 maturation on February 2014 were considered in this study. Clearly unripe
2 samples were small, hard, and pale green. On the contrary, overripe samples
3 showed symptoms of degradation such as feeble flesh and sticky skin.
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6 After collecting, the weight and size of every sample (it is highlighted that one
7 sample means one berry) were measured, and the visible and NIR images were
8 taken. Then, samples were stored at -20 °C until performing the chemical
9 analyses.
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15 The extraction of phenolic compounds was carried out following a modification of
16 the methodology proposed by Sheeram et al. (2006). Each berry was freeze-
17 dried (lyophiliser Cryodos 80 Telstar, Spain) and grounded to a homogeneous
18 powder (A 11 Analytical Mill, IKA, Staufen, Germany). Fifteen mg of powder were
19 extracted with 500 µL of methanol containing 0.1 % of HCl, sonicated during 15
20 min and centrifuged at 13700 g (10 °C, 10 min) (Microfuge 22R, Beckman
21 Coulter, Brea, CA), and the supernatant was collected. This procedure was
22 performed four times more, until the solid residue had no coloration. The extract
23 was concentrated at 30 °C under vacuum (Eppendorf Concentrator Plus,
24 Hamburg, Germany) until elimination of methanol; finally, made up to 2 mL with
25 HCl 0.1 M, and filtered through 0.5 µm pore-size filter. This amount of extract (2
26 mL) was enough to perform all the chemical analyses in triplicate.
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46 2.2 Image Analysis

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48 *Colorimetric analysis from digital images*

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51 The DigiEye® imaging system used (Luo et al. 2001) comprises an illumination
52 cabinet (VeriVide Ltd., Leicester, UK) with fluorescent lamps that reliably simulate
53 the CIE Standard Illuminant D65. A digital camera (Nikon D-80) attached at the
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1 top of the cabinet receives and stores the images in 48-bit TIFF format. By means
2 of the certified colour chart DigiTizer (VeriVide Ltd., Leicester, UK), the software
3 transforms from RGB to CIELAB colour spaces by applying advanced non-linear
4 multivariate models. Besides colour of samples, heterogeneity was measured by
5 using the Mean Colour Difference from the Mean (MCDM), proposed by Berns
6 (2000).
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16 *NIR hyperspectral image analysis*

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18 The acquisition of NIR hyperspectral images were made with an equipment that
19 comprises the following parts: a) Xenics® XEVA-USB InGaAs camera (Xenics
20 Infrared Solutions, Inc., Leuven, Belgium); b) spectrograph Specim ImSpector
21 N17E Enhanced, that covers a range of 884-1717 nm (Spectral Imaging Ltd.,
22 Oulu, Finland); c) two 70 W tungsten iodine halogen lamps as light source; d)
23 Mirror Scanner for scanning the sample surface (Spectral Imaging Ltd., Oulu,
24 Finland); and e) computer system to run the SpectralDAQ 3.62 acquisition
25 software (Spectral Imaging Ltd., Oulu, Finland). A more detailed description as
26 well as a scheme of this equipment are shown in Nogales-Bueno et al. (2014).
27 Reflectance of images was normalised with respect to the reflectance of a white
28 Spectralon® ceramic tile (Labsphere Inc., North Sutton, NH), and these
29 reflectance values were transformed into absorbance ($\text{Log}(1/R)$ units), better
30 related to concentration and more adequate for the prediction models (Williams
31 2001; Williams et al. 2009; Rodríguez-Pulido et al. 2014).
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2.3 Chemical analysis

Total phenolic content

The total phenolic content was determined using the Folin-Ciocalteu assay (Singleton and Rossi 1965). Briefly, 30 µL of extract, 900 µL of deionized water, 150 µL of Folin-Ciocalteu reagent, and 450 µL of a solution of sodium carbonate (20 %) were mixed, and deionized water was added to make up a total volume of 3 mL. The solution was vortexed, reading the absorbance at 765 nm with an Agilent UV-vis 8453 spectrophotometer (Palo Alto, CA) after stand for 2 hours to ensure the reaction takes place. The calibration standard used was gallic acid and results were expressed as gallic acid equivalents (mg gallic acid/g of fresh fruit).

Total Anthocyanins

Total anthocyanins content was measured by the spectrophotometric method of Wrolstad et al (2005). 200 µL of extract were added to 800 µL of buffer solution at pH 1.0. Another aliquot of 200 µL of extract was added to 800 µL of buffer solution at pH 4.5. Absorbance values at 520 and 700 nm were measured after 15 minutes to calculate the total amount of anthocyanins as follows:

$$\text{Total Anthocyanins} \left(\frac{\text{mg}}{\text{L}} \right) = \frac{A \times MW \times DF \times 10^3}{\epsilon \times l}$$

being $A = (A_{520} - A_{700})_{\text{pH}1.0} - (A_{520} - A_{700})_{\text{pH}4.5}$, MW=Molecular Weight (malvidin-3-glucoside = 493.2 g×mol⁻¹), DF=Dilution Factor, ε=molar extinction coefficient, (malvidin-3-glucoside = 20200 L×mol⁻¹×cm⁻¹), and l=pathlength.

The results were transformed to mg malvidin-3-glucoside per gram of fresh fruit.

Antioxidant capacity

The FRAP method (Benzie and Strain 1996) was applied to determine the antioxidant capacity, with a modification adapted to the analysis of food samples, (Pulido et al. 2000; Saura-Calixto and Goñi 2006), expressing the results as equivalents of Trolox, a water-soluble vitamin E analogue. The method consists on the reduction of the complex Fe^{3+} -TPTZ to Fe^{2+} -TPTZ in presence of antioxidants, and the reduced complex has a maximum of absorbance measurable at 593 nm by spectrophotometry.

2.4 Data treatment

The image processing and all the statistical treatments were performed by using Statistica 8.0 software and algorithms programmed under MATLAB R2012b.

3. Results and discussion

3.1 Sample grouping and physical analysis

The high range of maturation degree suggested us to make a pre-classification of the samples, firstly by simple visual inspection, dividing finally, the 168 samples into seven stages, from clearly unripe to overripe (Figure 2a). The assessment of uncertain samples were made based on the reflectance spectra. Regarding the physical properties, moisture and weight had almost the same behaviour during maturation (Figure 2b). Both weight and moisture increased quickly at stages 1 and 2 up to 2.5 grams and 84 % of moisture per berry, approximately. The last stage showed a decrease of weight due to the phenomenon of over-ripening.

3.2 Imaging techniques

Colorimetric Analysis

The DigiEye imaging system stored the images having the colorimetric information in CIELAB colour space (16 bits per colorimetric variable, $L^* a^* b^*$).

The segmentation criterion recognized every fruit, removing automatically the pedicel when present, and so did not compute for colour measurements. The imaging analysis also determines size and heterogeneity. Figure 1 shows the colorimetric coordinates of each sample grouped by stage of maturation. The scatterplot at a^*b^* -diagram had a horseshoe shape. It can be seen a descent of hue from 70° until 20° at the same time that chroma increases at the beginning and decreases at final stages, showing the highest chroma at stage 4, which means that the fruits had the most saturated colour even when they had not reached the optimal features for being consumed. Regarding the lightness, L^* values increased slightly at stage 3 and then, dropped up to 30-40 CIELAB units. These results are in agreement with other studies. Gordillo et al. (2012) reported that a greater amount of anthocyanins implies lower values in lightness because high concentration of pigments makes the sample absorbs more light, or rather reflects less light. Globular features of this fruits coupled with the unequal synthesis of pigments affected the overall appearance. These factors made the heterogeneity (measured as MCDM) increased quickly up to stage 4, and then, decrease when reached fully ripe or slightly before full ripeness. Table 1 shows the colorimetric results grouped by maturation stage.

Near infrared spectra (Hyperspectral Imaging)

After calibration, segmentation of hyperspectral images, and transformation into $\text{Log}(1/R)$ units, the spectra from each berry were extracted, discarding the first and the last wavelengths to avoid noise. Hence, the useful interval of spectrum ranged between 950 and 1650 nm. The most advanced stages had higher absorbance values (data not shown), being a high difference in absorbance (around 1400 nm) where the first O–H stretching overtone contributes, therefore, the moisture affects widely to this band. In this case, the influence can be attributed to the increase of water occurring along the maturation process (Osborne et al. 1993). Results also indicates that the first stages showed chemical changes more noticeable, while there were not significant changes ($p < 0.05$) between spectra belonging to stages 6 and 7.

3.3 Chemical analyses

Table 1 summarises the chemical analyses results. Regarding the phenolic content, the first stage had the highest concentration and decreased dramatically at stages 2 and 3. This can be due to the lowest moisture having at the first stages, which increases the concentration. However, the stage 1 had the highest standard deviation, so there were high variability among samples at this stage. Phenolic concentration slightly increased at stages 5, 6, and 7, probably due to the synthesis of anthocyanins, which are phenolic compounds as well, and hence, contribute to the total concentration.

The antioxidant capacity had the same behaviour (Figure 3a). On the other hand, it is well known that phenolics are one of the main compounds responsible for the antioxidant capacity in raspberries. Therefore, the direct relationship between

these variables, studied by simple linear correlation (Figure 3b), showed a value of 0.85 for the coefficient of determination (R^2). There was not a suitable distribution of values along all the range of concentrations because only samples belonging to stages 1 and 2 had the highest concentration of phenolics. The remaining samples (stages 3 to 7) appeared located near to the origin of coordinates. On the other hand, the changes of anthocyanins occurred differently to the total phenolics and antioxidant power ones. As expected, berries on stage 1 virtually had not anthocyanins, and small amounts appeared at stage 2, maintained at stage 3. From this point, the biosynthesis of anthocyanins follows an ascendant linear way until the full-ripe stage (Figure 3a). In general, these results were in agreement with those found in literature (Sariburun et al. 2010; Piljac-Žegarac and Šamec 2011). However, some results reported that they may vary depending on variety and crop conditions (Manganaris et al. 2014).

3.4 Relationship between optical features and chemical composition

The study included the assessment of the ability of predicting the chemical composition from data obtained by both colorimetric and NIR imaging techniques. For this aim, we considered the colorimetric and infrared datasets as independent variables, and the chemical result as dependent variable. The models calculated by merging both spectroscopic and colorimetric datasets did not yield better results than those obtained individually. PLSR built the models regardless the stage of maturation of berries. Table 2 shows the results. For total phenols, that ranged between 190 and 1592 mg gallic acid/100 g of fresh fruit, the prediction was almost the same regardless the data allocated as independent variables. The worst prediction were for antioxidant activity (all coefficient of determination < 0.65), which ranged between 230 and 1638 μ equiv. Trolox/100 g of fresh fruit,

1 while the best prediction were for total anthocyanins, maybe because their
2 presence is clearly noticeable even to the naked eye. Figure 4 shows the
3 scatterplot of predicted vs measured values of total anthocyanins considering
4 colorimetric data as independent variables. Except for samples having a high
5 anthocyanins concentration, all the samples appeared spread around the line of
6 equality. The coefficient of determination above mentioned may appear
7 insufficient to be a substitute of conventional chemical analysis, but they are quite
8 acceptable comparing to those obtained by similar techniques (Clark et al. 2003;
9 Downey and Kelly 2004; ElMasry et al. 2007; Nicolaï et al. 2007).

10 Based on the definition of the chemical composition for the seven proposed
11 stages, we developed a software under MATLAB in order to assess the
12 maturation stage of each berry automatically from hyperspectral images
13 (Supplementary video). The time needed to perform the loading, segmentation
14 and assessing of images is less than one second, but video has slowed down to
15 better show the process.

40 **4. Conclusions**

41 The methodology for inspecting single raspberries by colorimetric and NIR
42 imaging techniques have been established. Thus, it is possible to predict
43 important nutritional or bioactive properties by using a conventional digital
44 camera, avoiding long and destructive chemical analyses. Moreover, the
45 numerical methodology developed could become a reference method for the
46 industry and satisfy the need of the on-line and in real time quality control of intact
47 and individual berries, instead of result from chemical analysis representative for
48 a whole batch.

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FIGURE CAPTIONS

Figure 1. Scatterplot of colour data of raspberries samples regarding the stage of maturation (Lightness (L^*), a^* and b^* are expressed in CIELAB units). a^*b^* -diagram also shows the correspondence between a^*b^* and $C^*_{ab}-h_{ab}$).

Figure 2. a) Example of colour images of raspberries used grouped by stage of maturation. b) Means with error bars (standard deviation) of weight and moisture of raspberries grouped by stage of maturation.

Figure 3. a) Means with error bars (standard deviation) of total phenols, antioxidant activity, and total anthocyanins along the maturation. b) Scatterplot explaining the relationship between total phenols and antioxidant activity.

Figure 4. Scatterplot of predicted vs measured values when total anthocyanins are predicted from colorimetric data obtained from images of raspberries by PLSR

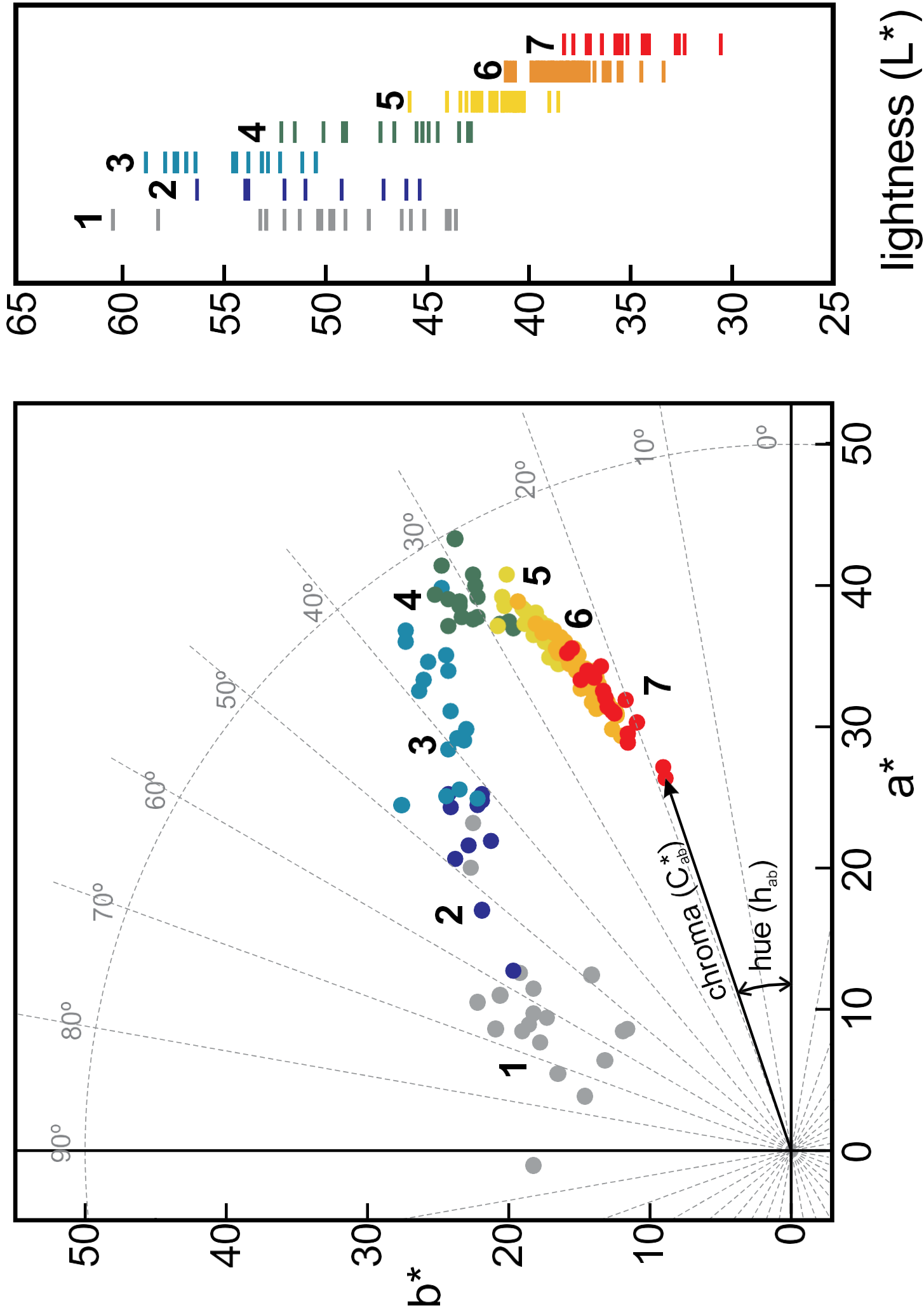
Table 1. Colorimetric coordinates and chemical composition of raspberries at each stage of maturation (mean $\pm$ SD). Lightness, chroma and MCDM are in CIELAB units. Hue values are expressed in degrees. Total phenols are in mg gallic acid/100 g of fresh fruit. Total anthocyanins are in mg malvidin-3-glucoside/100 g of fresh fruit. Antioxidant activity are in μ equiv. Trolox/100 g fresh fruit. Different letters indicate significant differences (Tukey's HSD test, $\alpha=0.05$).

Stage	1	2	3	4	5	6	7
L *	49.73 $\pm$ 4.61 ^a	50.94 $\pm$ 3.62 ^a	54.67 $\pm$ 2.67 ^b	47.06 $\pm$ 2.91 ^c	41.58 $\pm$ 1.60 ^d	38.51 $\pm$ 1.62 ^e	35.54 $\pm$ 2.06 ^f
C* _{ab}	20.98 $\pm$ 4.74 ^a	31.45 $\pm$ 3.41 ^b	39.95 $\pm$ 4.08 ^c	45.11 $\pm$ 2.04 ^d	40.99 $\pm$ 1.77 ^c	37.33 $\pm$ 2.59 ^e	34.07 $\pm$ 3.25 ^b
h _{ab}	61.66 $\pm$ 10.76 ^a	46.34 $\pm$ 5.14 ^b	39.06 $\pm$ 3.96 ^c	30.59 $\pm$ 1.66 ^d	26.24 $\pm$ 0.91 ^e	24.03 $\pm$ 1.07 ^{ef}	21.96 $\pm$ 1.74 ^f
MCDM	6.60 $\pm$ 0.80 ^a	6.71 $\pm$ 0.36 ^{ab}	7.29 $\pm$ 0.44 ^{bcd}	8.27 $\pm$ 0.54 ^e	7.87 $\pm$ 0.39 ^e	7.77 $\pm$ 0.68 ^{oe}	7.19 $\pm$ 0.69 ^{ad}
Total phenols	894 $\pm$ 333 ^a	407 $\pm$ 91 ^b	326 $\pm$ 74 ^b	251 $\pm$ 39 ^b	274 $\pm$ 43 ^b	281 $\pm$ 63 ^b	298 $\pm$ 60 ^b
Total anthocyanins	7.5 $\pm$ 6.8 ^a	14.4 $\pm$ 5.8 ^{ab}	14.4 $\pm$ 5.6 ^{ab}	22.4 $\pm$ 7.0 ^b	38.0 $\pm$ 8.2 ^c	48.9 $\pm$ 11.3 ^d	60.8 $\pm$ 16.5 ^e
Antioxidant activity	749 $\pm$ 395 ^a	324 $\pm$ 129 ^b	202 $\pm$ 117 ^{bc}	142 $\pm$ 62 ^{bc}	123 $\pm$ 64 ^c	141 $\pm$ 81 ^c	164 $\pm$ 90 ^{bc}

Table 2. Calibration and Cross Validation results for the PLS models obtained from both visible and NIR datasets. RMSE_c and RMSE_{cv} were expressed as follows: for Total phenols (mg gallic acid/100 g of fresh fruit), for Total anthocyanins (mg malvidin-3-glucoside/100 g of fresh fruit), and for Antioxidant activity (µequiv. Trolox/100 g fresh fruit).

		R ² _c	RMSE _c	R ² _{cv}	RMSE _{cv}	PLS factors
Total phenols	VIS	0.75	116.1	0.73	121.4	3
	NIR	0.75	116.3	0.70	126.8	6
Total anthocyanins	VIS	0.76	9.9	0.75	10.1	3
	NIR	0.66	11.5	0.63	12.0	4
Antioxidant activity	VIS	0.65	149.2	0.62	156.1	3
	NIR	0.64	150.4	0.61	157.3	3

Figure 1



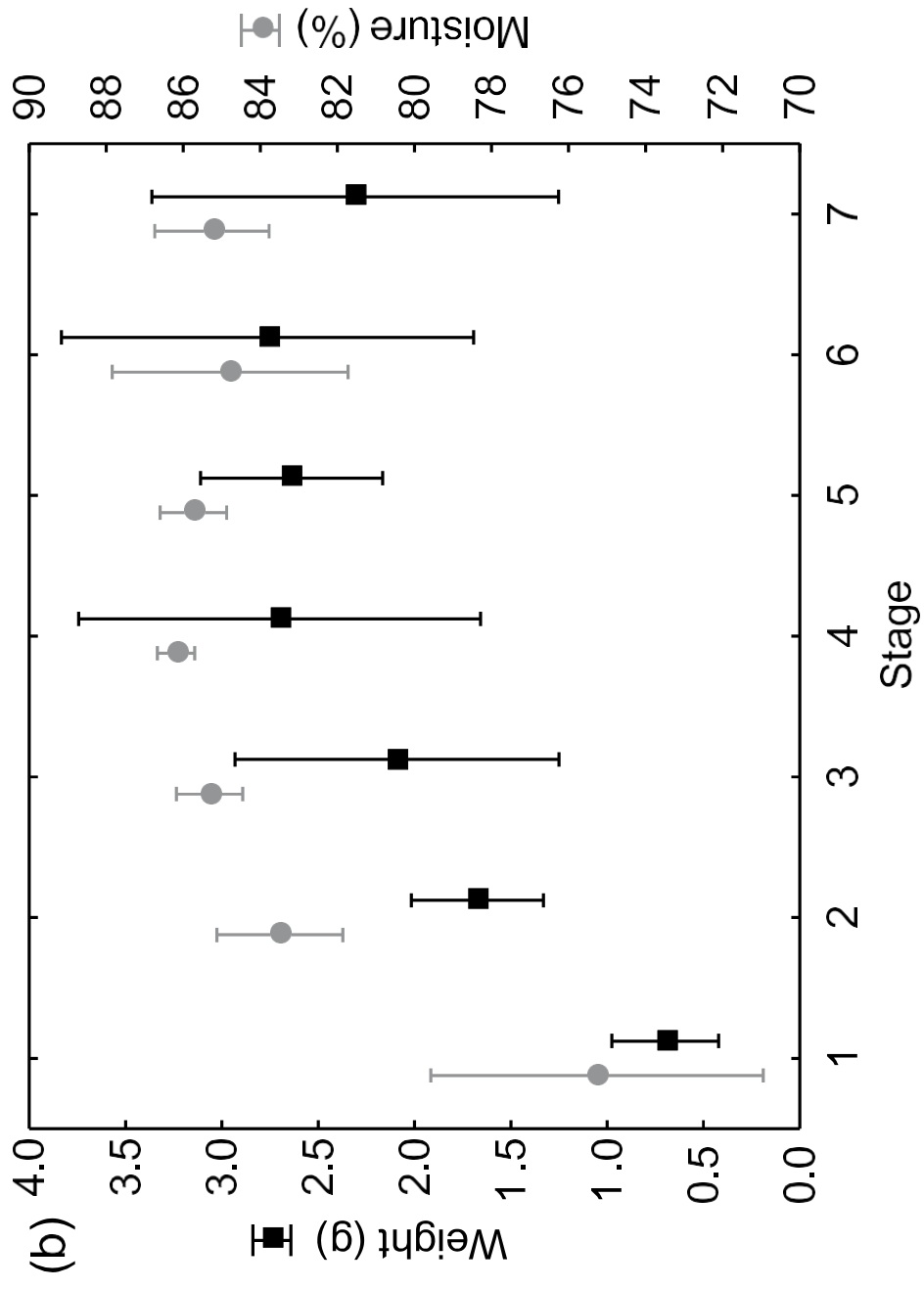
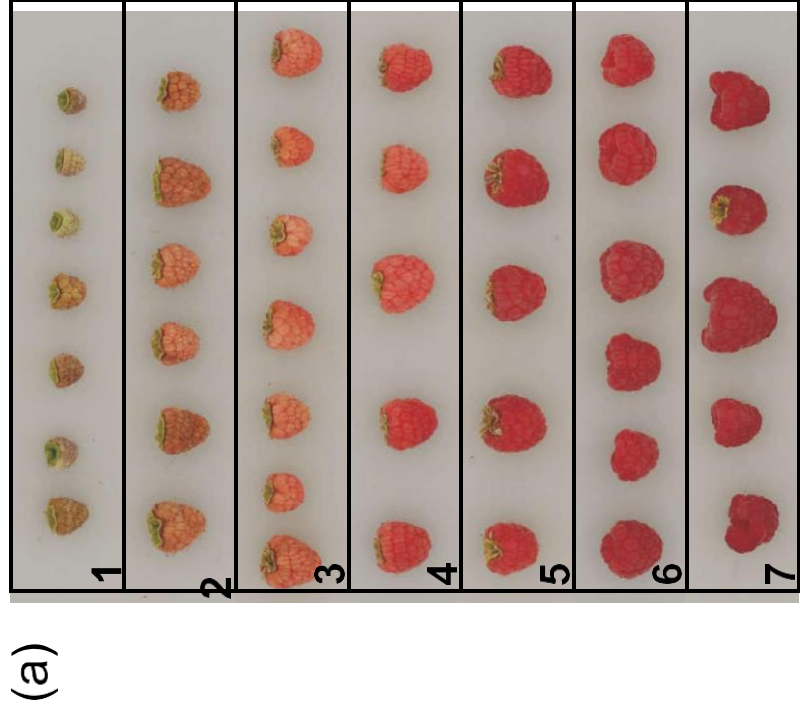


Figure 2

Figure 3

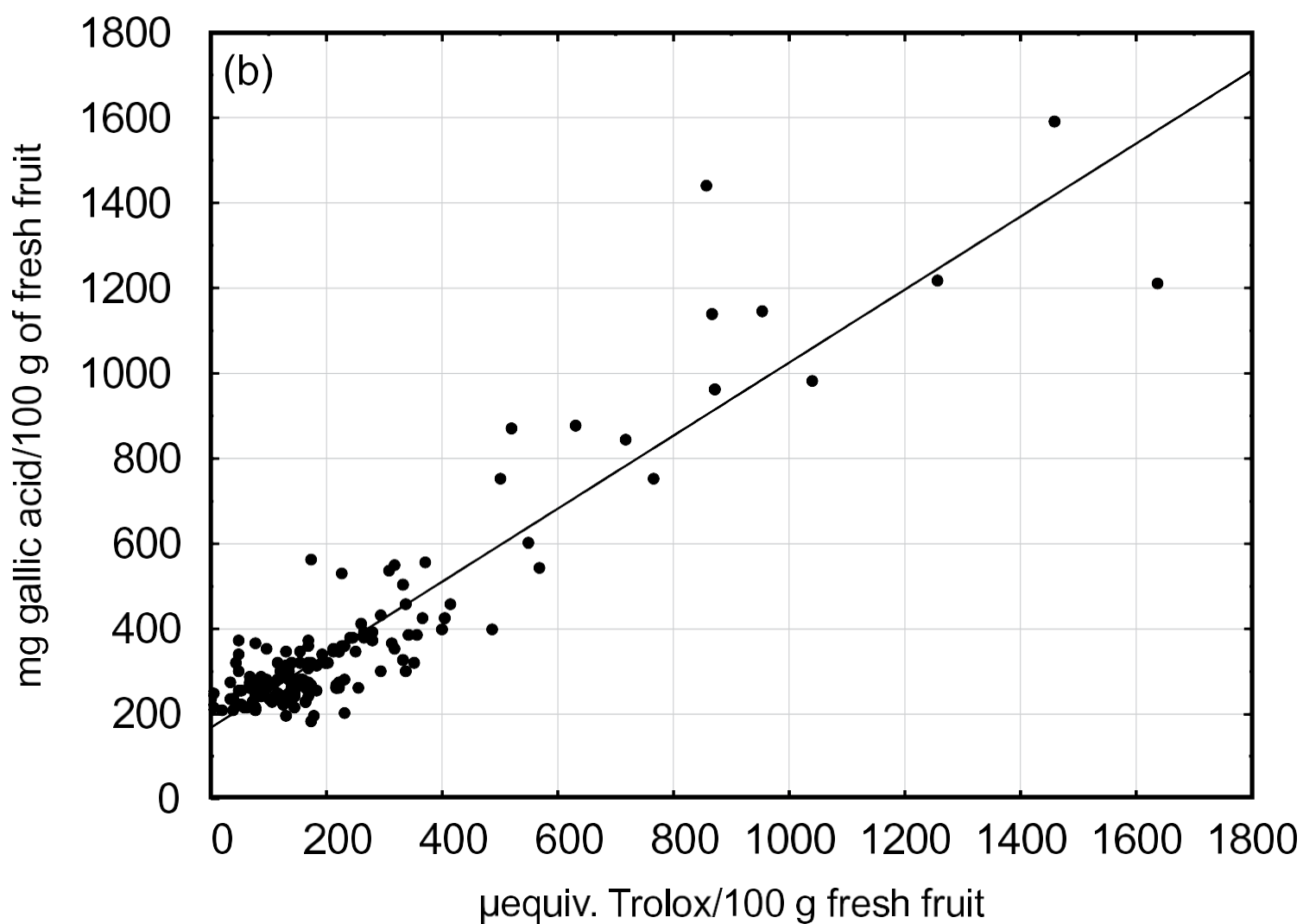
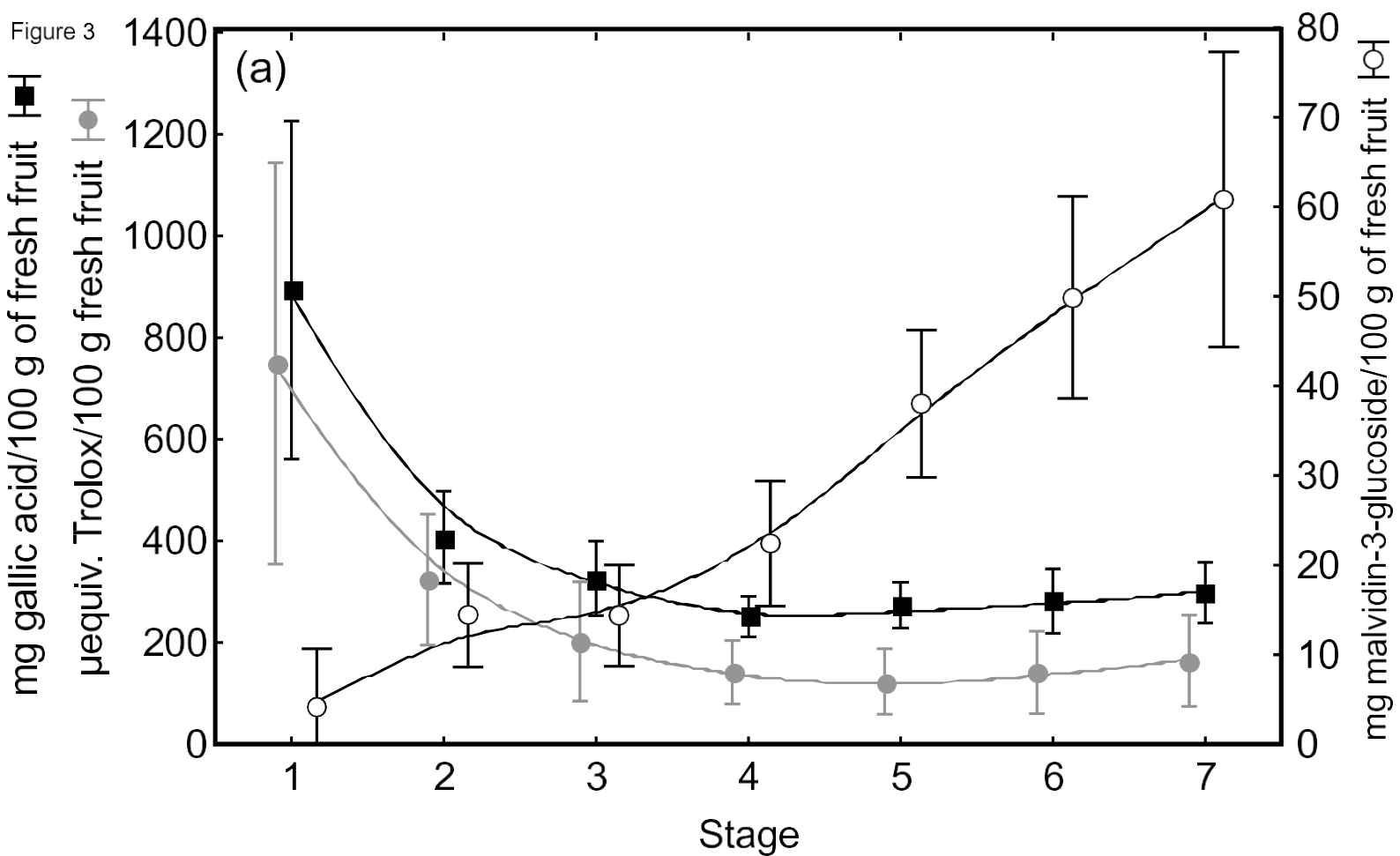
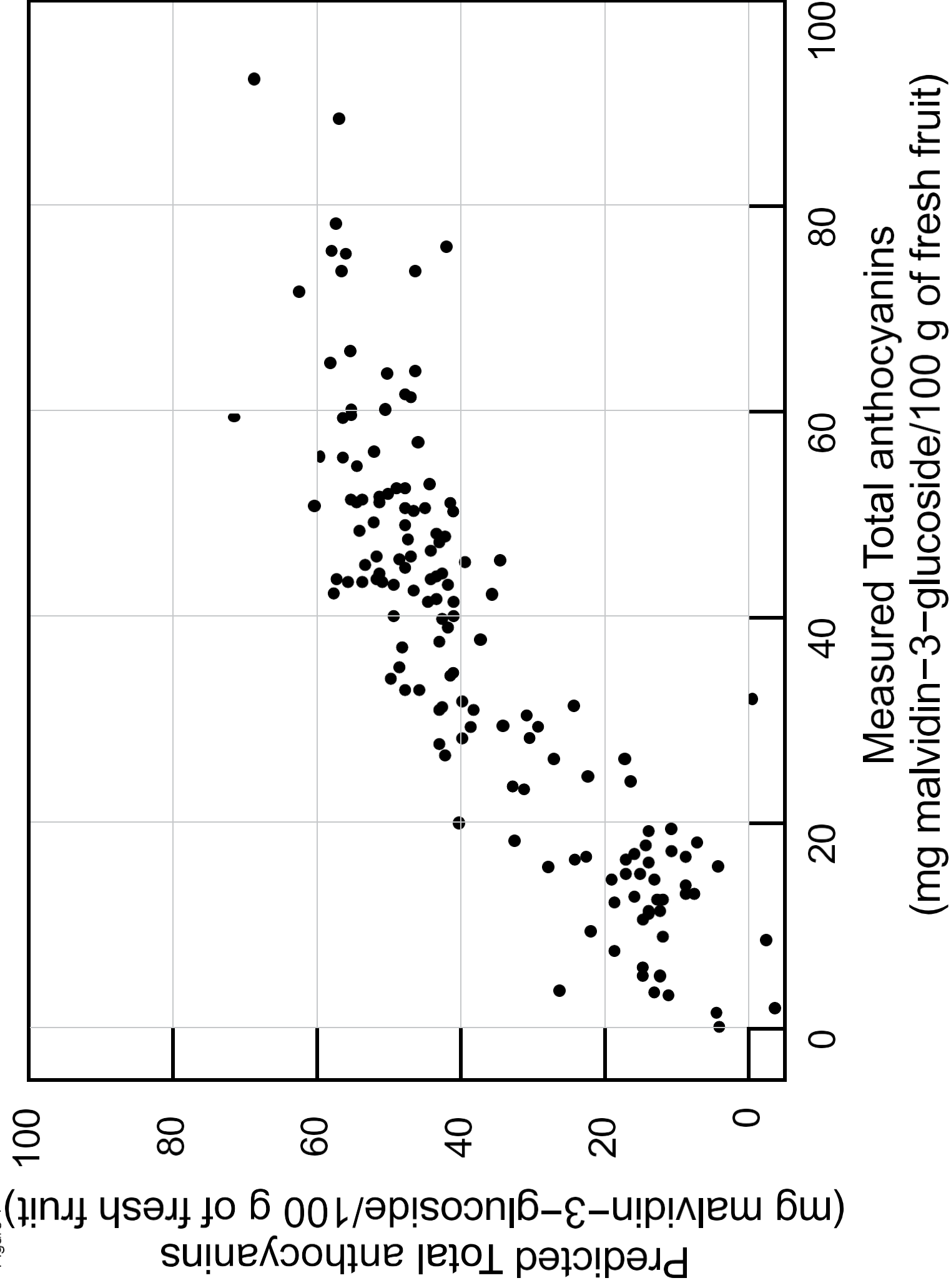


Figure 4





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by near infrared and colorimetric imaging techniques

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