The use of pigs vocalisation structure to assess the quality of human-pig relationship

Authors

Avelyne S Villain* [1,2], Carole Guérin [1], Camille Noûs [3], Céline Tallet* [1]

Affiliations

[1] PEGASE, INRAE, Institut Agro, 35590 Saint Gilles, France

[2] Behavioural Ecology Group, Section for Ecology & Evolution, Department of Biology, University of Copenhagen, 2100, Copenhagen Ø, Denmark

[3] Cogitamus laboratory, 75000 Paris, France

* Corresponding authors : avelyne.s.villain@protonmail.com, celine.tallet@inrae.fr

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Abstract

Studying human-animal interactions in domestic species and how they affect the establishment of a positive Human-Animal Relationship (HAR) may help us improve animal welfare and better understand the evolution of interspecific interactions associated with the domestication process. Understanding and describing the quality of an HAR requires information on several aspects of the

animal biology and emotional states (social, spatial and postural behaviours, physiological and cognitive states). Growing evidence shows that acoustic features of animal vocalisations may be indicators of emotional states. Here, we tested the hypothesis that vocal structure may indicate the quality of HAR. At weaning, 30 piglets were positively handled by an experimenter who talked to and physically interacted with them three times a day, while 30 other piglets only received the contact necessary for proper husbandry. After two weeks, we recorded the behaviours and vocalisations produced in the presence of the static experimenter for 5 min. We repeated this test two weeks later, after a conditioning period during which human presence with additional positive contacts was used as a reward for all piglets. We hypothesized this conditioning period would lead to a positive human-piglet relationship for all piglets. As expected, piglets that were positively handled at weaning expressed a higher attraction toward the experimenter, and, after the conditioning, piglets that were not positively handled at weaning expressed a similar level of attraction than the positively handled ones. Piglets positively handled at weaning produced shorter grunts than the other ones, regardless of the context of recording, which may indicate a more positive affect. During reunions with the static experimenter, a more positive HAR was associated with a decrease in vocal reactivity to human proximity. However, during reunions with the experimenter providing additional positive contacts and over the conditioning, proximity to the human systematically triggered shorter and higher pitched grunts, indicator of positive a emotional state. Results first show that changes in vocal structure are consistent with indicators of positive states in the presence of a human. Second, these changes are stronger when the human positively interact with the piglets, supposedly emphasizing a higher positive arousal state during these interactions. We show that vocalisation structure may be a promising indicator of the quality of human-pig relationship.

Introduction

The process of domestication was conducted to shape physiology and morphology of domestic animal species, but also their behaviour. It notably has shaped interspecific interactions between human and non-human animals, by improving animals' capacity to use human signals to adapt their behaviour both decreasing fearfulness toward humans and increasing attention toward humans (Mignon-Grasteau et al., 2005). In farms, the relationship that domestic animals form with humans is important for animal welfare. Therefore, studying human-animal interactions and their consequences to understand the mechanisms of emergence and maintenance of a positive human-animal relationship (HAR) directly applies to welfare (Rault et al., 2020). Animal welfare consists of three major aspects: the ability of an animal to control its mental and physiological stability (Broom, 2011), the decrease of experiencing negatively perceived contexts and the increase in experiencing

positively perceived contexts and species-specific behaviors (Peterson et al., 1995; Weerd & Day,
2009). A positive HAR is thought to be established through repeated positive interactions between
the human and the non-human animal. Some of the mechanisms involved in this process are:
accumulation of positive experiences through positive associative learning, modifications of
cognitive biases, shaping expectations from the non-human animal toward the human. A positive
HAR can be appreciated through behavioural and physiological measures, for example by assessing
the expression of positive emotions [reviewed in (Rault et al. 2020)]. Several behavioural measures
may help to define a positive HAR such as: short latency to approach and spatial proximity (Boivin
et al., 2000; Schmied et al., 2008), body postures (Villain, Lanthony, et al., 2020) or play behaviour
(Jerolmack, 2009). Contacts from a human such as stroking, may induce changes in body postures
and exposition of body areas by the animal to the human, supposedly vulnerable [central neck area in
cattle (Schmied et al. 2008), abdominal area in pigs (Rault et al., 2019)]. Such grooming solicitation
may be markers of engagement, trust and motivation to interact with the human. In most cases, these
behaviours are similar to those shown during intraspecific socio positive interactions, although there
are some species specific behaviours [e.g., dog vs. wolf (Gácsi et al., 2005)]. Vocal behaviour may
also help defining the quality of an HAR. First, some vocalisations type have been associated with
positive interactions with humans, for example the cat – human communication : purring is thought
to be derived from mother pup communication during nursing and is observed associated with care
solicitation from humans; meowing, which is not observed during intra specific interactions is
thought to emerge from associative learning during cat – human interactions (Brown & Bradshaw,
2014). This shows that HAR may elicit specific vocalisations from the non human animal toward the
human. Second, vocalisation structure is known to carry markers of the emotional states in several
bird and mammal species (Briefer, 2012, 2020) and markers of emotional valence (positive versus
negative) has been studied in domestic farm animals [reviewed in Laurijs et al. (2021)]. Since
positive or negative HAR is likely to affect the emotional state of animals, it is likely that it may be
reflected in the structure of the produced vocalisations.

In pigs, diversified evidence attest the possibility of a positive HAR. Animals may be handled by
humans providing regular additional positive contacts, leading to the expression of a positive
perception of humans, with evidence from behavioural and physiological studies. Cognitive bias tests
showed a positive judgment bias in piglets that had received gentle contacts with humans (Brajon et
al., 2015b). Pigs may recognise a human providing positive contacts compared to an unfamiliar one
and adapt their behaviour accordingly (Brajon et al., 2015c). Pigs may be sensitive to human voice
and respond accordingly (Bensoussan et al., 2019, 2020). Pigs vocalisations are diverse and linked to
their emotional states, attested by the use of positive or negative call types (Briefer et al., 2019, 2022;
Tallet et al., 2013). In addition, even within a call type, spectro-temporal changes are closely related

to the valence or the arousal a situation may trigger for the animal. For example, the grunt, a contact
call, is used in various contexts and is now known to be a flexible call. Positive situations have been
associated with shorter grunts compared to negative ones (Briefer et al., 2019, 2022; Friel et al.,
2019), as well as higher formants (which are frequency peaks containing more energy than others)
and a lower fundamental frequency during positive situations (Briefer et al., 2019, 2022). Grunt
structure may also change according to the arousal of a negative situation: the higher the arousal in
the negative state the higher the frequency range and bandwidth (Linhart et al., 2015) and the longer
(Puppe et al., 2005) the grunts. Variation in grunt spectro-temporal structure in positive situations of
different arousal is still unknown.

In order to determine to what extent vocalisations structure could be used as non invasive indicator
of the quality of human-pig relationship, we tested whether varying the degree of familiarity and the
quality of the human-pig interactions could modulate the spectro-temporal structure of vocalisation,
through the vocal expression of emotional state. Because it was suggested to study vocal markers of
emotions within the same call type (Briefer, 2020) and because grunts are the most commonly
produced call in various contexts, we studied the spectro-temporal structure of grunts. We predicted
that if grunts reflect the quality of the human-pig relationship, then 1. A period of positive handling
given by a human should modulate piglets vocal expression in presence of the human, leading to
grunts exhibiting markers of positive states (higher pitched and shorter grunts), 2. Spatial proximity
toward the human should influence the spectro-temporal structure of grunts (higher pitched and
shorter grunts).

Methods

Ethical note

The study was approved by the ethic committee CREEA and received the authorization no.
APAFIS#17071-2018101016045373_V3 from the French Ministry of Higher Education, Research
and Innovation. UE3P, where the experiment was carried out, is an experimental unit authorized by
the French Ministry of Agriculture to breed animals for experimentation under the number D35-275-
32. This authorization includes a derogation to follow the directive 2008/120/EC relative to the
protection of piglets and its regulations.

Subjects and housing conditions

Sixty weaned female pigs (in two replicates from January to April 2019), *Sus scrofa domesticus*, bred from crosses between Large White and Landrace females and Piétrain males were used for this study from 28 to 62 days after birth. Animal housing and experiments took place at the experimental unit UE3P (UE 1421, INRAE France).

One piglet had to be excluded from our sample size to receive care/medication due to health issues independent from the experiment. From weaning at 28 days of age, piglets from the same litter and having similar weight (<1 kg difference) were housed by three in a 1.2 x 1.3m pen on plastic duckboard. Wooden panels were used to visually isolate pens. One metal chain per pen was used for enrichment. Food and water were available *ad libitum*. Artificial lights were turned on from 8:00 to 17:00 and temperature was maintained between 26 and 27 ºC. The experiment was carried out in two replicates and two identical rearing rooms were used (5 pens per room per replicate).

Treatment: positive handling at weaning

From day 28 (day of weaning) to day 39 of life, piglets were separated into two groups that experienced a different post-weaning period as follows:

- Non positively handled piglets (H piglets): Control piglets from 10 rearing pens, housed in the same room, received the minimal amount of daily contact with a stockperson (a 1.70m tall male who did the feeding, cleaning and health checkups). The stockperson wore a dark green shirt and pants and brown shoes.

- Positively handled piglets piglets (H+ piglets): Experimental piglets from the 10 other rearing pens, housed in another room, received the same daily care given by the same stockperson as for H piglets. They additionally received repeated sessions of additional human contacts. Each pen of three piglets received 29 sessions of 10 min , from day 28 (weaning) until day 39, occurring five days a week. Three sessions per day were performed (except on the day of weaning during which only two were done with a two-hour break in between). Each session took place in the rearing pen and the order of the interventions in the pens was balanced across days. The handling procedure, using gentle tactile contacts is described in supplementary material of Villain et al. (2020) and was similar to Tallet et al. (2014). Briefly, the behaviour of the human toward the piglet was adapted to the reaction of each animal and included four steps: (1), the handler hold out the hand towards the animal; (2) if the piglet did not move away, the handler tried to touch it; (3) if the piglet accepted being touched, the handler softly stroked it along the body with the palm of her hand; and (4) once it accepted being

stroked, the handler scratched it along the body with her fingers. Scratching consisted in rubbing the

skin of the piglets with the finger tips and applying more pressure than stroking. No specific body

part of the piglets was more considered that another. Two experimenters ('AV' and 'AH') performed

these sessions (both women, both between 1.70-1.73 m tall, with a balanced number of pens

attributed to each of them). The experimenters wore the same blue overalls and green boots each

time they interacted with the piglets. The experimenters tried to imitate each others behaviours

(remote video monitoring) to decrease variability.

This intense period of additional positive contacts for half of the piglets after weaning constituted the

treatment of positive handling at weaning: positively handled piglets are referred to as H+ piglets and

non positively handled piglets are referred to as H piglets to describe the early experimental

treatment they experienced regarding a human, prior to the conditioning.

Conditioning: sessions of additional positive contacts with (un)familiar human

The conditioning took place between day 42 and 62 of age and lasted twelve days, with two trials

per day and at least three hours between trials on the same day. Piglets were habituated to the test

room for 10 min, by pen, two days before the start of the conditioning. All piglets (H and H+) were

subjected to the same conditioning. The experimental design of the conditioning is already published

in an article dedicated to the study of anticipatory behaviour (Villain, Hazard, et al., 2020).

Briefly, all piglets were individually trained to learn to associate two different stimuli with the

arrival of two different (pseudo)-social partners: either two pen mates (partner = Conspecifics) or a

familiar human (partner = Human). When entering the room, the piglets and the partner(s) would

remain in the room for 2 min. Specifically, when the human was the partner, the human entered, sat

on a bucket and positively interacted with the piglet for 2 min, in the same manner as additional

contacts was provided to the H+ piglets during the previous period (see above section) (figure 1).

Therefore, at the beginning of the conditioning, H+ piglets were already familiar with the human and

procedure, whereas H piglets were unfamiliar with the human. During the conditioning, the same

sessions occurred in both treatment groups (H and H+). After the conditioning, all piglets were

familiar with the human, but treatment groups had a different time of exposure to them. Sessions of

reunions with social partners were not studied here because they were part of an analysis on vocal

expression of positive anticipation reported earlier (Villain, Hazard, et al., 2020).

For every second trial, the 2 min reunions with the human were analysed by the same person: trials

number 2, 4, 6, 8, 10 and 11 (see behavioural analyses section).

Standard Isolation/Reunion Tests with a static and silent human

At 40 or 41 (before conditioning) and then 63 or 64 (after conditioning) days of age, piglets were
subjected to a standard Isolation/Reunion test in order to assess their perception of the human. The
test consisted of two phases. The piglet was brought individually in a trolley to the experimental
room. It was left alone for 5 min, which defined the 'Isolation' phase. Then, the human entered the
room, remained stand up for 30 seconds and they sat on a bucket, remaining silent and not moving
for 4.5 min (figure 1).

Location of the microphone
Virtual zones for behavioural scoring
Location of the human
Proximal area   Distal area

Figure 1: Design of the room used during the Isolation/Reunion tests and the additional positive contacts sessions of the conditioning. The room was split into 16 virtual zones. A proximal area (zones 10, 11, 14, 15) and a distal area (zones 1, 2, 3, 4) were defined, suing the location of the human as reference. Behavioural monitoring and analyses

Sessions and tests were recorded using a camera (Bosh, Box 960H-CDD) and behaviours were
scored *a posteriori* on videos using *The Observer XT 14.0* (Noldus, The Netherlands) software. The
room was split into 16 virtual equally-dimensioned zones to assess the mobility and exploratory
behaviour of the piglet. A proximal area, around the human was defined by merging four zones, a
distal area was defined merging the four most distant zones from the human (figure 1).

The behaviours scored during the reunion of the Isolation/Reunion test and the sessions of
additional positive contacts of the conditioning are available in table 1. Every time the shoulders of
the piglet crossed a zone, a zone change was scored. Looks and watching behaviours were scored as
point events, all other behaviours were scored as state events. Behavioural scores were then
calculated to quantify global responses (see Table 1).

Table 1: Ethogram.

Behaviour	Description
No. zones crossed [1,2]	The number of times the piglet crossed a virtual zone
No. approaches H [1]	Number of times the piglets entered the proximal area
Time watching H [1,2]	The amount of time the piglet spent watching the human
Latency to contact H [1,2]	The latency to the first contact of the human by the piglet
No. looks toward H [1,2]	The number of times the piglet turned its head toward the human
No. looks at walls and doors [1]	The number of times the piglet looked at walls and doors
Time watching walls and doors [1]	The amount of times the piglet watched walls and doors
Time in proximal area [1,2]	The amount of time the piglet spent in the proximal area
Time in distal area [1,2]	The amount of time the piglet spent in the distal area
Time in contact H [1,2]	The amount of time the piglet investigated the human
Time investigating floor [1,2]	The amount of time the piglet investigated the floor
No. contacts H [2]	Number of times the piglet was in contact with the human (initiated by the piglet or the human)

[1]: Scored during reunions of Isolation/Reunion tests. [2]: Scored during reunions of conditioning sessions

Acoustic monitoring and analyses

Vocalisations were recorded with an AKG C314 microphone placed in the center of the room and
one meter above the ground, connected to a Marantz MD661MK2 recorder. Vocalisations produced
during each phase of the trial were manually annotated according to vocal type (grunt, squeal, bark,
scream and mixed calls (Kiley, 1972)), after visual inspection of spectrograms using the 'Annotate'
function of the Praat software (Boersma & Paul, 2001), version 6.0 from http://www.praat.org/.
Checking the occurence of each call type in the several contexts of the study, we confirmed that
'grunt' was the call type used in all contexts and by most of the piglets in each context. So only the
spectro-temporal structure of grunts was further analysed. For information, a table of the number of
each call types recorded in each context as well as the number of individuals involved in the count is
presented in the electronic supplementary material. We could not conduct a robust statistical analysis
on call type utterance, due to the rarity (per subject and tests) of other vocalisations than grunt. (table
S5).

A spectro-temporal analysis was performed with custom-written codes using the Seewave R
package (Sueur et al., 2008) implemented in R (R Core Team, 2022). We first studied the spectral
properties of the remaining background noise of the experimental room (electric noises and
remaining low frequency noises from the rest of the building), using 20 examples of 0.5 second
fragments and compared it with the general frequency range of the grunts. To avoid measuring
masking effect of the background noise, grunts were filtered using a 0.2-8 kHz bandpass filtering
('fir' function). As a consequence, all results presented in this study are on a 0.2-8 kHz frequency
range, and no conclusions on frequency components of grunts below this 200 Hz threshold are drawn

here. Several acoustic parameters were then extracted from each grunt. To measure grunt duration, a
5% to maximal amplitude threshold was used ('timer' function). After normalisation, the following
spectral parameters were calculated using the 'specprop' function (FFT with Hamming window,
window length = 512, overlap = 50%): mean (Q50), first (Q25) and third (Q75) quartiles,
interquartile range (IQR), centroid and standard deviation (all in Hz). The grunt dominant frequency
(in kHz) was also calculated ('dfreq', 50% overlapping FFTs, window length = 512), which is the
mean over the grunt duration of the frequencies of highest energy of each window. Frequency peaks
were detected and the minimal and maximal peaks were kept as descriptors ('fpeaks' function,
window length = 512, peak detection threshold = 10% of the normalized amplitude). Measures of
noisiness and entropy of the grunts were assessed using: Shannon entropy (sh), Spectral Flatness
(Wiener entropy, sfm) and Entropy (H) [combining both Shannon and Temporal envelope entropy,
length = 512, Hilbert envelop). Two vocal scores were used: the logarithm of grunt duration and a
built-in spectral vocal score with all spectral parameters (see below). A table describing mean and
range of variation of each acoustic parameter in the relevant contexts of the study is available in the
supplementary material (table S4).

Statistical analyses

Behavioural and vocal response scores

All measures extracted from videos or sound analysis are named parameters throughout the text.
The symmetrical distribution of parameters (behavioural on the one hand and acoustic on the other
hand) was visually inspected, and linear transformations were computed when necessary to reach
symmetrical distribution (see tables 2, 3, 4). When this criteria was reached, Principal Component
Analyses (PCA, one for the behavioural analysis and one for the spectral acoustic analysis) were
performed using several parameters to build scores ['dudi.pca' function from 'ade4' R package
(Dray & Dufour, 2007) and 'inertia.dudi' function to extract the loadings]. These scores were then
used as statistical variables. Indeed, PCAs are generally used to reduce the number of variables
included in statistical models. It also generates quantifiable global descriptors of behaviours or
acoustic structure, since correlated parameters usually load on the same PC (McGregor, 1992). All
PCs having an eigenvalue above one were kept and constituted response scores of behavioural
('ReuPCs' and 'CondPCs' in table 2 and 3 respectively) and vocal ('VocPCs', table 4) parameters.
Only the duration of grunts was kept separated from the spectral parameters to keep it as a temporal
parameter.

Table 2: Percentage of explained variance and relative loadings of parameters on PCs, following the Principal

Component Analysis computed on the behaviours scored during the reunion of the Isolation/Reunion test. The first three PCs, having an eigenvalue above 1, constituted three behavioural scores: ReuPC1, ReuPC2, ReuPC3. Parameters that explain the most each PC are bolded (|loading|>0.4).

	ReuPC1	ReuPC2	ReuPC3
Cumulative variance explained %	38.3	60.8	74
No. of zones crossed (sqrt)	24.177	**-55.843**	-0.435
No. approaches H (ln)	**47.748**	-30.163	0.578
Time watching H (ln)	**-52.914**	-7.422	25.585
Latency to contact H (ln)	**-64.232**	-0.464	1.688
No. looks toward H (sqrt)	-7.787	**-43.721**	31.633
Time watching room	-32.048	-13.581	-6.238
No. looks at walls and doors (sqrt)	3.524	**-72.408**	-2.027
Time in proximal area (sqrt)	**69.96**	-0.156	9.584
Time in distal area (sqrt)	**-46.416**	-12.437	-1.215
Time in contact H (sqrt)	**61.041**	3.586	24.183
Time spent investigating floor	11.868	-7.503	**-42.265**

Table 3: Percentage of explained variance and relative loadings of parameters on PCs, following the Principal Component Analysis computed on the behaviours scored during the sessions of additional positive contacts of the conditioning. The first three PCs, having an eigenvalue above 1 constituted three behavioural scores: CondPC1, CondPC2, CondPC3. Parameters that explain the most each PC are bolded (|loading|>0.4).

	CondPC1	CondPC2	CondPC3
Cumulative variance explained %	41	68.5	80.7
Time in proximal area (ln)	**80.23**	2.542	-0.112
Time in distal area (ln)	-33.826	8.547	30.789
Number of contacts H (ln)	**78.55**	6.476	2.288
Time in contact H (ln)	**86.625**	0.715	-0.369
No. looks toward H (ln)	-2	**79.898**	-0.745
Time watching H (ln)	-6.757	**65.67**	-10.325
No. of zones crossed (sqrt)	0.129	33.599	**48.457**
Time spent investigating floor	0.006	**-49.286**	14.205
Latency to contact H	**-81.01**	-0.248	-2.83

Table 4: Percentage of explained variance and relative loadings of parameters on PCs following a Principal Component Analysis on spectral parameters of the grunts recorded in the entire dataset (including both types of tests, N=17 546 grunts). The transformations used to reach symmetrical distribution before the PCA are indicated in parentheses. The first three PCs, having an eigenvalue above 1 constituted three vocal response scores: VocPC1, VocPC2, VocPC3. Parameters that explain the most each PC are bolded (|loading| >0.4) .

	VocPC1	VocPC2	VocPC3
Cumulative variance explained %	59.769	76.807	87.712
Mean Dominant Frequency[1]	-13.558	**53.557**	2.220
Min frequency peak[1] (ln)	-0.349	**58.758**	24.236
Max frequency peak[1]	**-43.023**	8.760	-9.537
Mode[2] (ln)	-0.522	**66.248**	19.268
Mean[2] (ln)	**-95.092**	-2.295	2.028
Q50[2] (ln)	**-85.278**	0.280	-0.093
Q25[2] (ln)	**-52.360**	19.327	0.985
Q75[2] (sqrt)	**-88.925**	-4.645	2.309
Centroid[2] (ln)	**-95.092**	-2.295	2.028
Sd[2]	**-64.484**	-11.303	7.680
IQR[2]	**-87.981**	-5.851	2.640
Sfm[3] (sqrt)	**-94.344**	-3.189	0.962
Sh[3] (sqrt)	**-96.087**	-0.785	-0.175
H[3]	**-88.205**	-1.059	-1.063
Skewness[4]	28.032	-18.010	**48.652**
Kurtosis[4]	22.973	-16.241	**50.615**

[1]: parameters related to the pitch of the vocalisation; [2]: parameters related to the frequency distribution descriptors; [3]: parameters related to the noise component of the vocalisation; [4]: parameters related to the shape of the frequency distribution

Statistical models

All statistics were carried out on R (R Core Team, 2022). Linear mixed effect models ['lmer' function, 'lme4' R package (Bates et al., 2014)] were built when tested variables were linear (behavioural and vocal scores, grunt duration) and one binomial generalized mixed effect model was built for binary parameters (occurrence of missed contacts initiated by human during the conditioning). The following subsections describe how models were built for each type of tests. In all models described below, the identity of the replicate ('1' or '2') was used as an interacting fixed

factor, since the experiment was run in two identical replicates on two independent groups. The

identity of the human ('AH' or 'AV') was used as interacting fixed factor in all models described

below, since two experimenters were involved in the positive handling at weaning and in the session

of additional positive contacts of the conditioning (but always the same human was attributed to a

given piglet). The piglet was used as random factor to take into account the within-subject design.

All explanatory variables used in the models and interactions between them were built in respect to

the experimental design and to allow biological interpretations. As a consequence, not all

interactions between all variables were made. They are fully explained in the subsequent sections.

Isolation/Reunion tests

The aim of this part was to test the effect of the positive handling at weaning treatment (H vs. H+

piglets) and additional human contacts during sessions of the conditioning on the piglet's reaction to

human presence. Since the same Isolation/Reunion test was repeated before and after the

conditioning, we used the variable 'Conditioning time' as a two level interacting factor ('before' or

'after' conditioning, referred as "Time" in the models) to test the effect of the conditioning. Piglets

spacial behaviour and proximity to the human was studied only during the reunion phase with the

human that followed the isolation phase. Model_1 was computed:

```
274    Model_1 <- lmer (ReuPCs ~ Treatment*Time + Treatment*Replicate +
275    Treatment*HumanID + Time*Replicate + Time*HumanID + (1 | pigletID), data=
276    data_Behaviour_Reunion).
```

Concerning the analysis of vocal behaviour, the isolation phase represents a negative social context

for the piglets and may be used as a negative control when monitoring the effect of human presence

on vocal expression of emotional states (Villain, Lanthony, et al., 2020). So, the two phases of the

test were used to study the three way interaction between treatment (H vs.. H+), phase of the test

(isolation vs.. reunion) and time of the conditioning (before vs.. after). The following model_2 was

computed:

```
283    Model_2 <- lmer (VocPCs ~ Treatment*Phase*Time + Treatment*HumanID + Time*HumanID
284    + Treatment*Replicate + Time*Replicate  + (1 | pigletID/Time/Phase) , data=
285    data_Vocal_Isolation + data_Vocal_Reunion).
```

286   To go further, only the reunion phase was kept and a proximity variable was added. Indeed, the

287   piglet could vocalise either when close to human or away from them and this spatial proximity was

288   demonstrated as an important factor of changes of vocal features (Villain et al. 2020b). Thus, a two

289   level proximity factor was built: either '1' when the piglet was in the proximal area (figure 1) or '0'

290   when it was elsewhere in the room. The following model_3 was computed:

```
291   Model_3 <- lmer (Vocal response score ~ Treatment*Time*InProxArea +
292   Treatment*HumanID + InProxArea*HumanID + Treatment*Replicate +
293   InProxArea*Replicate + Time*Replicate + Time*HumanID + (1 | pigletID/Time), data
294   = data_Vocal_Reunion).
```

295   **Conditioning trials**

The aim was to study the evolution of human-piglet relationship over the conditioning [the variable

'Trial number', used as a continuous variable, referred as "Trial" in the models]. The effect of

treatment (positively handled at weaning H+ piglets or non handled H piglets) was tested as an

interacting factor with Trial. Trial was also used as a random slope to take into account individual

trajectories (Schielzeth and Forstmeier 2009). The following model_4 was built to test the

behavioural response scores CondPCs (lmer) and the occurrence of missed contact initiated by the

human during a session (presence/absence, binomial model, glmer):

```
303   Model_4  <- (g)lmer (CondPCs / Missed contact ~ Trial*Treatment + Trial*HumanID +
304   Trial*Replicate + Treatment*Replicate + Treatment*humanID + (1+ Trial |
305   pigletID), (family=Binomial), data= data_Behaviour_Conditioning).
```

306   For the analysis of vocal response scores, similarly to the Isolation/Reunion test, the piglet could

307   vocalise either when close to the human or away from them. We thus added the proximity factor in

308   the analysis of vocal response variables. The following model_5 was built :

```
309   Model_5  <- lmer (VocPCs ~ Trial*Treatment*InProxArea+ Trial*HumanID +
310   Trial*Replicate + Treatment*Replicate + Treatment*HumanID + HumanID*InProxArea +
311   Replicate*InProxArea  + (1+ Trial | pigletID), data= data_Vocal_Conditioning).
```

312   **Model validation and statistical tests**

All linear models were validated by visual inspection of the symmetrical and normal distribution of

the residuals. Anovas ('car' R package (Fox & Weisberg, 2011)) were computed on models to test

for significant effects of explanatory variables. Following the Anova, when interactions were found

significant, post hoc test were run on model interactions, correcting for multiple testing with Tukey

contrasts ('emmeans' or 'lstrends' functions from 'emmeans' R package (Lenth, 2016), for

categorical or continuous variables respectively). Considering the conditioning time (before or after

conditioning), when involved in a significant three-way interaction, this factor was fixed to allow

pairwise comparison within each time period as it was not considered relevant to assess the effect of

time only. Results of the Anova, model estimates and pairwise post hoc comparisons are reported in

the supplementary material (tables S1 and S2 for tests, table S3 for model estimates).

Results

Effect of positive handling at weaning and conditioning on piglets' reaction to human presence (Isolation/Reunion tests)

Piglets that were not handled at weaning express a similar behavioural proximity to a human after a positive conditioning as the positively handled ones.

Figure 2: Effect of conditioning and treatment on spatial behaviour and proximity toward the human during the reunion of the Isolation/Reunion test. Mean ± SE per group is indicated, different letters indicates significantly different groups. Significant interaction between treatment (H : grey squares and H+ : black circles) and time (Before the conditioning: empty elements and After the conditioning: filled elements) on behavioural ReuPC1 (letters a to c) and ReuPC3 (letters z and y). Full statistical report is available as supplementary material (tables S1 S2 for statistical tests and S3 for model estimates)

The interaction between the treatment (positively handled piglets at weaning (H+) or not (H) and the conditioning time (before or after the conditioning) was significant for both ReuPC1 and ReuPC3 (χ^2_1 = 28.0, p < 0.001, and χ^2_1 = 3.7, p = 0.05 respectively, figure 2) but not for ReuPC2 (χ^2_1 < 0.001, p = 0. 99, supplementary table S1). Post hoc tests on ReuPC1 showed that ReuPC1 was higher after the conditioning than before (H: after – before, t.ratio = 12.1, p <0.001 , H+: after – before t.ratio = 11.0, p < 0.001) and that before the conditioning, piglets that were positively handled at weaning had significantly higher ReuPC1 than non handled piglets (Before, H – H+: t.ratio = -2.1, p < 0.001), but not after (After, H – H+: t.ratio = 0.02, p = 1.0). According to the loadings, this means

that piglets that were positively handled at weaning had a lower latency to contact the human,

approached them more often and spent more time close to and investigating the human (ReuPC1)

than non handled piglets, before the conditioning. This score increased after the conditioning and no

evidence of a difference between treatments after the conditioning was found (figure 2). Post hoc

tests on ReuPC3 showed a significant effect of the conditioning time only in piglets that were

positively handled at weaning (H+: after – before, t.ratio = 5.2, p < 0.001, H: after – before, t.ratio =

2.6, p = 0.06). No difference in ReuPC3 was found between treatments before the conditioning

(Before: H – H+, t.ratio = -0.75, p = 0.87), whereas positively handled piglets had a higher -ReuPC3

after the conditioning than before (After : H – H+, t.ratio = -3.2, p = 0.009). According to the

loadings, this means that after the conditioning, piglets that were positively handled at weaning

expressed more investigation of the room after the conditioning than non handled piglets. No

evidence of any effect on ReuPC2 was found (table S2).

Piglets positively handled at weaning produce shorter grunts even when no human

is present.

Using the isolation phase as a negative control we could compare the effect of the phase of the test

(Isolation vs. Reunion with the human), taking into account the conditioning time (before or after the

conditioning) and the treatment. No evidence of any effect of neither the three way interaction ($\chi^2_1 <$

0.62, p > 0.43) nor two way interactions of interest was found (treatment: phase, conditioning

time:phase, conditioning time: treatment interactions : χ^2_1 <3.5, p > 0.06, table S2) in any of the

scores.

Regardless of the treatment, single effects of the phase of the test were significant for grunt

duration and all AcPCs (χ^2_1 > 6.6, p < 0.01, table S1). During the reunion phase with the human,

grunts were shorter (estimates of log(duration)[95% CI] : -1.32[-1.37;-1.26] vs. -1.06[-1.12;-1.00]),

had a higher frequency range, higher bandwidth and a higher noise component (-VocPC1:

0.78[0.48;1.08] vs. 0.34[0.03;0.66]), were higher pitched (VocPC2: -0.18[-0.36;0.01] vs. -0.46[-

0.65;-0.28]) and their spectrum had a higher skewness and kurtosis (VocPC3: -0.25[-0.37;-0.14] vs. -

0.11[-0.23;0.01]), compared to the isolation phase.

Regardless of the phase of the test, single effects of treatment were found for grunt duration and -

VocPC3 (χ^2_1 = 5.5, p = 0.02 and χ^2_1 = 4.9, p = 0.03 respectively, table S2). Grunts produced by

positively handled at weaning piglets were shorter (estimates of log(duration)[95% CI]: -1.25[-1.32;-

1.19] vs. -1.12[-1.2;-1.1], table S3), and differed in -VocPC3 scores, describing the shape of the

frequency spectrum (estimates of -VocPC3[95% CI]: -0.29[-0.43;-0.14] vs. -0.07[-0.22;0.08], table

S3), than grunts produced by non handled piglets.

Positive handling and conditioning affect vocal reactivity to human proximity.

Figure 3: Acoustic structure of grunt during the reunions with a silent and static human (Isolation/Reunion test). Effect of conditioning (before or after), treatment (H or H+) and location of the piglet relatively to the human (close: dark blue or away from them: light blue). Violin plots representing the median and the density of data distribution in the considered groups. (A, B) Results of post hoc tests following significant three way interaction between treatment, conditioning time and location on grunt duration (A) and on the first vocal score -VocPC1 (B). (C,D) Results of post hoc tests following significant two way interactions between conditioning time and location (C) and between treatment and location (D) on the second vocal score VocPC2. Values with no common letters differ significantly. When no letters are present, no significant difference between groups was found. Stars (*) between two groups represent a statistical trend (p< 0.10). Full statistical report is available as supplementary material (tables S1 S2 for statistical test and S3 for model estimates).

During the 5 min reunion, the piglet was scored either as close to the human or away from them.
The three way interaction of the conditioning time, the treatment and the location was significant for
grunt duration, -VocPC1 and VocPC3 ($\chi^2{}_1 > 4.9$, $p < 0.03$). Post hoc tests revealed that grunts
produced closer to the human were shorter than the ones produced further away, but only in piglets
that were not positively handled at weaning, effect being stronger before the conditioning than after
it (H piglets: away – close, z.ratio = 6.3, $p < 0.001$ before and z.ratio = 4.1 $p < 0.001$ after the
conditioning; H+ piglets: away – close z.ratio < 1.98 $p > 0.19$, figure 3A). -VocPC1 was higher, i.e.
grunts had a higher frequency range, bandwidth and were noisier when produced closer to the human
than further away, but only in non handled piglets and before the conditioning (H piglets: away –
close, z.ratio = -3.34, $p = 0.005$ before and z.ratio = -1.23 $p = 0.61$ after the conditioning; H+ piglets:
away – close, z.ratio < 0.36 $p > 0.21$, figure 3B). For VocPC2, the three way interaction did not
reach significance ($\chi^2{}_1 = 3.3$, $p = 0.07$), so only subsequent two way interactions were considered
(post hoc tests on the three way interaction can be found in supplementary, tables S1 to S3). For
VocPC2, significant two way interactions were found between the conditioning time and the location
($\chi^2{}_1 = 10.3$, $p = 0.001$) on the one hand, and between the location and the treatment ($\chi^2{}_1 = 4.2$, $p =$
0.04) on the other hand. Post hoc tests revealed that grunts produced closer to the human had a
higher VocPC2, meaning they had a higher pitch, effect being stronger before the conditioning than
after (before: away – close, z.ratio = -6.12, $p < 0.001$; after: away – close, z.ratio = -2.88, $p = 0.004$,
figure 3C). The increase in VocPC2 with the location was greater for non handled piglets than
positively handled piglets (H piglets: away – close, z.ratio = -5.54, $p < 0.001$; H+ piglets: away –
close, z.ratio = -3.82, $p = 0.001$, figure 3D). The last two-way interaction of interest between the
conditioning time and the treatment did not reach significant level ($\chi^2{}_1 = 0.80$, $p = 0.37$). For
VocPC3, post hoc tests did not reach significant levels (|z.ratio| < 2.3 $p > 0.09$ for any comparison) .

396 Emergence of positive perception of human (effect of additional
397 positive contacts sessions over the conditioning)

398 *The conditioning increases behavioural proximity to the human in all piglets.*

Figure 4: Behavioural variation of responses of piglets according to the sessions of additional positive contacts of the conditioning (A), and to the treatments (B, C). (A,B) Mean ± SE per group. Numbers in (A) refers to the trial number of the conditioning. Higher CondPC1 and lower CondPC2 over time (single effect of trial number, A). Higher CondPC2 in H piglets than H+ piglets regardless of time (single effect of treatment, A). Higher CondPC3 and lower CondPC2 in H+ piglets than H piglets (single effect of treatment, B). (C) Mean estimates ± 95% confidence interval from generalized mixed effect model. Lower probability of occurrence of missed contact by the human in H+ than H piglets (single effect of treatment). Full statistical report is available as supplementary material (tables S1 et S2 for statistical tests, table S3 for model estimates).

No evidence of any effect of the interaction between the treatment [positively handled piglets
before the conditioning (H+) or not (H)] and the trial number was found for all behavioural scores
(CondPC1, CondPC2 and CondPC3, table 3). Independently from the treatment, the higher the trial
number the higher CondPC1 (χ^2_1 = 59.3, p < 0.001, slope estimate [95% confidence interval]: 0.20
[0.15 : 0.25]) and the lower CondPC2 was (χ^2_1 = 48.6, p < 0.001, slope estimate: -0.17 [-0.22 : -
0.12]). According to the loadings, over the conditioning, piglets decreased the latency to contact the
human, made more contacts, spent more time in the proximal area and in contact with the human
(condPC1), decreased the number of looks to the human, spent less watching the human and more
time investigating the room (CondPC2) (figure 4A). Independently from the trial number, positively
handled piglets had a lower CondPC2 and a higher CondPC3 than the non handled ones (χ^2_1 = 12.8,
p < 0.001 and χ^2_1 = 7.0, p = 0.008 respectively), meaning that piglets that were positively handled at
weaning expressed a fewer number of looks to the human, spent less time watching them and more
time investigating the room (CondPC2) and crossed more virtual zone during the test (CondPC3)
(figure 4B). The probability of having at least one missed contact by the human during a session was

lower for positively handled piglets than non handled ones ($\chi^2_1 = 9.57$, p = 0.002, figure 4C), with no

interaction with the trial number ($\chi^2_1 = 0.22$, p = 0.064).

Additional positive contacts trigger shorter and higher pitch grunts in all piglets.

Figure 5: Vocal scores over the conditioning, during the 2min sessions of additional positive contacts. (A, B) Violin plots representing the median and the density of data distribution in the group. Interacting effect of location (in proximal area of the human 'close': dark blue) or elsewhere in the room ('away' from the human: light blue) and treatment (H vs. H+ piglets) on grunt duration (A) and VocPC2 (B). (C) Mean ± SE per group, interacting effect of trial number and location of piglets on VocPC2. Values with no common letters differ significantly (difference between groups: A, B or slopes: C). Full statistical report is available as supplementary material (tables S1-S3).

During the sessions of additional positive contacts of the conditioning, the three-way interaction

between the trial number, the treatment and the location was not significant for any of the vocal

scores ($\chi^2_1 < 0.18$, p > 0.67), allowing the analysis of the two way interactions of interest. The

interaction between treatment and the trial number was not significant for all vocal scores ($\chi^2_1 < 2.5$

p > 0.11). Grunt duration decreased over time and independently from the treatment (trial

number:replicate interaction, $\chi^2_1 < 5.3$ p = 0.02, slope estimate -0.03[-0.04;-0.01] for the lower slope,

table S1 and S3). However, independently from the trial number, grunt duration was lower when

piglets were located close to the human and this effect was stronger in non handled piglets than

positively handled piglets (treatment:location interaction: $\chi^2_1 = 15.8$ p < 0.001, away vs.. close, H

piglets: z.ratio = 10.2 p < 0.001, H+ piglets: z.ratio = 6.86 p < 0.001, figure 5A). -VocPC1 and

VocPC2 decreased over time but remained higher when piglets were located close to the human (trial

number: location interaction, $\chi^2_1 = 3.97$ p = 0.05 and $\chi^2_1 = 6.1$ p = 0.01 respectively for -VocPC1 and

VocPC2). According to the loadings, this means that the frequency range, bandwidth and noisiness

of grunts (-VocPC1) as well as the pitch (VocPC2) decreased over the conditioning when piglets

were located away from the human but remained high when piglets were close (slope comparison

away – close, -VocPC1 : z.ratio = -1.80 p = 0.07, VocPC2 : z.ratio = -2.34 p = 0.02, figure 5C).

Additionally, VocPC2 was higher when piglets were close to the human in non handled piglets

(treatment:location interaction, χ^2_1 = 7.6 p = 0.005, pairwise comparisons away vs. close, in H:

z.ratio = -4.9 p z 0.001 and in H+: z.ratio = -2.0 p = 0.21), meaning that non handled piglets

produced higher pitched grunts when closer to the human (figure 5B).

Impact of human identity on piglets behaviour and grunt structure

Figure 6: Effect of human identity on spatial behaviour and proximity during the reunion of the Isolation/Reunion test. Violin plots representing the median and the density of data distribution in the group. Values with no common letters differ significantly. Full statistical report is available as supplementary material (tables S1 and S2 for statistical tests, table S3 for model estimates).

Since half of the piglets had been assigned to one human experimenter and the other half to another

one, the identity of the human was included in the model. This allowed to test interactions between

the identity of the human and the treatment of positive handling at weaning on the one hand and the

conditioning time on the other hand.

During the reunions of the Isolation/Reunion test, the interaction between treatment and human

identity was significant for the first behavioural proximity score (ReuPC1, χ^2_1 = 6.01, p = 0.01) but

not the others (ReuPC2 and ReuPC3 (χ^2_1 < 1.98, p > 0.16, table S1). The effect of treatment on

ReuPC1 was higher when piglets were handled by the human 'AH' (H vs. H+, AH: t.ratio = -4.77, p

< 0.001, figure 6). When the human 'AV' handled the piglets, for which ReuPC1 scores exhibited

intermediate values, treatment was not significant (AV, H vs. H+: t.ratio = -1.33, p = 0.56). These

interacting effects of the human identity and treatment on behaviour were not found when

considering the reunions of the conditioning ($\chi^2{}_1 < 1.32$, p > 0.25 for all CondPCs, table S1).

Interactions between the human identity and conditioning time were not significant, neither

considering the reunions of the Isolation/Reunion test (ReuPCs, $\chi^2{}_1 < 0.642$, p > 0.42, tables S1),

neither the trial number during the session of additional positive contacts of the conditioning

(CondPCs, $\chi^2{}_1 < 0.11$ p > 0.74, table S1).

Table 5: Significant effects of human identity on vocal response score (VocCP1 and VocPC2) during the reunion of the Isolation/Reunion test and during the sessions of additional positive contacts of the conditioning. Only significant effect are presented here but a full statistical report is available as supplementary material (tables S1 and S2 for statistical tests, table S3 for model estimates). When single effects were interpretable, the Chi-squared statistic are reported. When significant interactions were significant, post hoc pairwise comparisons were performed with Tukey corrected and are thus reported. The estimates correspond either to the group estimate and comparisons of groups (categorical fixed effect) or slope estimates and comparison of slopes (continuous fixed effect, 'Trial number').

Vocal response score	Fixed effect	Levels	Estimate	Lower.95%CI	Upper.95%CI	Statistic	P-value
Reunion of the Isolation/Reunion test							
VocPC2	humanID	AH	0.154	-0.119	0.427	$\chi^2{}_1=$ 4.94	P = 0.03
		AV	-0.292	-0.571	-0.012		
Sessions of additional positive contacts of the conditioning							
-VocPC1	humanID	AH	0.091	0.037	0.144	$\chi^2{}_1 =$ 4.69	P = 0.03
		AV	0.076	0.021	0.132		
VocPC2	humanID*InProxArea	AH – away	0.317	0.110	0.524	Z-ratio = -1.23	P = 0.60
		AH – close	0.402	0.161	0.643		
		AV – away	0.027	-0.182	0.236	Z-ratio = -5.77	P < 0.001
		AV – close	0.462	0.212	0.712		
VocPC3	humanID * Trial number	AH – Trial number	-0.048	-0.070	-0.026		
		AV – Trial number	-0.007	-0.031	0.016	Z-ratio = -2.82	P = 0.005

Considering the vocal scores, no effect of human identity was found on VocPC1 during the

Isolation/Reunion tests but -VocPC1 was higher when the human 'AH' was in the room during the

reunion periods of the conditioning (table 5), meaning the frequency range and the bandwidth of the

grunt were higher when the human 'AH' interacted with the piglet compared to the human 'AV'.

VocPC2 was higher when the human 'AH' was in the room during the Isolation/Reunion tests (table

5), meaning that the pitch of grunts was higher and this effect was also found during the sessions of

additional positive contacts of the conditioning in interaction with the location of the piglet (χ^2_1 =

11.9, p = 0.001): VocPC2 increased when piglets were located close to the human but this increase

was significant only for the human 'AV' and not for 'AH' (table 5). VocPC3 was not different

between humans during the reunions of the Isolation/Reunion tests but, over the conditioning,

VocPC3 changed differently when piglets were handled by the human 'AH' or 'AV', as showed by

the significant interaction between trial number and human identity (χ^2_1 = 8.0, p = 0.005): the

skewness and kurtosis of grunts decreased over the conditioning when 'AH' was interacting with the

piglets, but not 'AV' (see slope estimates, table 5). No evidence of any effect of human identity was

found on grunt duration neither during the Isolation/Reunion tests nor during the sessions of

additional positive contacts of the conditioning (table S1).

Discussion

In this study, familiarity to a human and human-animal interactions were experimentally modified

in weaned piglets to study the establishment of a positive HAR and test whether grunt structure could

reflect a positive HAR. A positive conditioning paradigm, using additional positive contacts from a

human as a reward, allowed to compare the behavioural changes over time in piglets previously

positively handled at weaning or not. Two types of sessions were studied: a standard

isolation/reunion tests with the human, carried out before and after conditioning, during which the

human remained silent and did not interact with the piglet, and sessions of the conditioning, during

which the human interacted with the piglets, providing additional positive contacts, as long as the

piglets stayed close to the seated human. Behavioural data were collected to describe the positive

HAR. Grunts produced during the tests and sessions were collected and their spectro-temporal

structure confronted to the behavioural data, with the hypothesis that vocalisation structure may

reflect the quality of HAR, though vocal markers of positive emotions. Firstly, the discussion will

focus on the behavioural validation of the establishment of a positive HAR. Secondly, behavioural

and vocal expression will be confronted to discuss grunt spectro-temporal structure as indicator of

the quality of HAR. Last, we will discuss perspectives regarding the effect of human identity on the

establishment of a positive HAR.

Behavioural evidence of a rapid establishment of interest and proximity toward a human providing additional positive contacts

The standard reunion test with the human before the conditioning showed first that the treatment of

positive handling at weaning succeeded in creating two different levels of human-piglet relationship

(H and H+), as positively handled piglets expressed a higher attraction toward the human than non

handled piglets (ReuPC1), parameters considered as indicators of a positive HAR (Rault et al.,

2020). Second, this test showed that the conditioning increased the behavioural proximity toward the

human of both positively handled and non handled piglets so that non handled piglets expressed a

similar attraction toward the human as positively handled piglets. These results are in line with the

behavioural results of the sessions of additional positive contacts. The analysis of piglets' behaviour

every second sessions of the conditioning showed that, although positively handled and non handled

piglets started with different degree of proximity toward the human (trials 2 and 4, CondPC1), then,

over time and for both treatments (H and H+), piglets expressed a higher attraction toward the human

(CondPC1) and avoided less the human when the latter attempted to interact with them. So it seems

that the conditioning process allowed non handled piglets to compensate the lack of positive

handling before the conditioning and develop a similar proximity toward the human. Two minute

daily sessions of additional positive contacts changed positively the perception of the human for the

piglets, and thus their willingness to interact with them. Since no evidence of any interaction

between time and treatment was found, no conclusion on differential developmental trajectories

between treatments can be drawn, but a parallel development of the human-piglet relationship in

both groups, when considering the proximity.

Beside behavioural proximity, piglets that were positively handled at weaning expressed more

exploratory behaviours than non handled piglets after the conditioning (ReuPC3). This was also

observed during the sessions of additional positive contacts of the conditioning: positive handled

piglets started with a higher score associated with investigation than non handled piglets (CondPC2)

and it held over the conditioning. Piglets that were positively handled at weaning also expressed a

higher mobility than non handled piglets (CondPC3). These observations may be interpreted as an

expression of natural foraging and disinterest from human contact, which may be a sign of positive

welfare (Weerd & Day, 2009). In addition, this could also be interpreted in terms of attachment to

the human. Indeed, attachment to a human may facilitate exploration of novel environments or

objects, as shown in dogs (Palmer & Custance, 2008). A period of positive handling at weaning may

provide an environment secure enough for the piglets to explore their environment in the presence of

the human. Attachment has also been hypothesised in the lambs-human relationship (Tallet et al.,

2009).

Overall, the behavioural monitoring showed that 2 min sessions of positive additional contacts per

day are sufficient to increase proximity to a human to similar levels as when piglets were previously

familiarised for 2 weeks, even when piglets experienced social isolation. But it did not allow the non

handled piglets to express natural exploratory behaviours as the positively handled piglets. We

hypothesize a sequential establishment of a positive HAR over time: firstly with a decrease of

attentive state and an increase in proximity and accepted contacts, and secondly with a disinterest of

human contacts and the expression of natural foraging behaviour. The latter may require a higher

exposure time.

In the next paragraph we discuss to what extent changes in grunt spectro-temporal structure may

reflect behavioural changes linked to the positive HAR over time.

Links between vocal expression and positive HAR

A positive HAR is reflected by shorter grunts in presence and absence of a human

The social isolation phase of the Isolation /Reunion test, before any human entered the room, was

associated with longer, lower pitched grunts with a downshifted frequency spectrum, whereas the

reunion with a static human changed grunts structure to shorter, higher pitched with an upshifted

frequency spectrum and this was observed in both handled and non handled piglets (H or H+) as well

as before and after the conditioning. In terms of emotional indicators, similar changes in acoustic

features of grunts were found in studies focusing on vocal markers of valence in pigs (Briefer et al.,

2019, 2022; Friel et al., 2019; Villain, Hazard, et al., 2020), meaning that the reunion with a human,

after a period of social isolation would be perceived as positive. However, this modulation of grunt

structure was observed regardless of piglet experience with the human. It is possible that the reunion

with an either neutral or familiar human, releasing piglets from total isolation could be perceived as

positive by the piglets, as suggested in previous studies (Villain, Lanthony, et al., 2020).

In addition, and surprisingly, positively handled piglets produced shorter grunts than non handled

piglets regardless of human presence. This was previously shown in another context (anticipation of

(pseudo)social events independently from the type of partner) in the same groups of piglets (Villain,

Hazard, et al., 2020). This may show that the period of positive handling at weaning modulated vocal

expression in the long term, as this result was found both before and after the conditioning. On the

one hand, a positive HAR establishes through successive positive experiences (Rault et al. 2020)

and, on the other hand, HAR may have long term effects on behavioural expressions, as suggested by

Brajon et al. (2015) using cognitive bias tests. We can thus hypothesize this may also be reflected in

the way piglets vocalise, in general. In that case, we may have evidence of expression of another

category of affect, moods, and not only emotional expression. Indeed, as suggested by Schnall
(2010), although emotions are short-term affects triggered by an external stimulus, moods, on the
other hand, may be experienced on a longer term and may not be attributable to a specific stimulus.
Although emotions and moods do not rely on the same time scale, they may interact with one
another, and more studies are needed to understand their effects on vocal expression.

A positive HAR affects vocal reactivity toward a static human

In a previous study, we showed that pigs vocalizing close to a human that previously had provided
repetitive additional positive contacts produced shorter and higher pitch grunts, compared to when
vocalizing away from the human (Villain, Lanthony, et al., 2020). Using the same type of test with
positively handled at weaning and non handled piglets, before or after conditioning sessions with
positive interactions, we can test the effect of positive handling on this modulation of grunt structure.
Similarly to the previous study, during the standard reunion test (no contact from the human), piglets
produced shorter and higher pitched grunts with an upshifted frequency spectrum when close to the
human. It has to be noted that this effect was 1) stronger in previously non handled piglets than
positively handled at weaning piglets and 2) stronger before the conditioning than after. In other
words, the more familiar with the human associated with positive handling, the less reactive to
human proximity.

These results may be interpreted according to the behavioural results we described earlier (fig. 2).
We described that the proximity to the human was first increasing at the beginning of positive
handling experiences (see H piglets, before vs. after conditioning) before reaching a maximum (see
H vs. H+ piglets after conditioning) and that the most familiar piglets showed more exploratory
behaviours (H+ after conditioning). The acoustic results during the standard reunion mirror the
behavioural results from the same test. The least familiar piglets would vocally express the
exploration of a neutral and static human and, as the familiarity with the human increases, the human
may become part of their environment, explaining the lack of vocal reactivity when close to the static
human.

In addition, we may also be facing ceiling effects in terms of vocal flexibility, which could also
partly explain these results. We showed that positively handled piglets generally produce shorter
grunts than non handled piglets, and that the shape of the frequency spectrum of these grunts was
different. So the structure of their calls, in general is different. According to the source-filter theory
of vocal production, vocal flexibility is constrained by the dimensions and functioning of the vocal

apparatus (lung capacity, characteristics of the vocal folds, length and shape of the vocal tract, see

(Taylor & Reby, 2010) and (Titze & Martin, 1998)). It is possible that the positive HAR developed

by the positively handled piglets may have change their grunts structure to an extent that vocal

flexibility is no longer quantifiable in the experimental design of this study.

Providing rewarding additional positive contacts triggers short and high pitched grunts

Contrary to the standard reunions with a static human, the human actively interacted with the

piglets during the sessions of the conditioning, providing contacts and producing speech as long as

the piglets remained close to the human. During these sessions and contrary to the standard reunions,

grunts produced close to the human were shorter and higher pitched, regardless of the trial number of

the conditioning and treatment. Although these effects were stronger in non handled piglets than

positively handled piglets, they remained over time. We describe here two types of vocal reaction to

human proximity, depending on the human behaviour. On the one hand, time decreased vocal

reactivity to human proximity during a standard reunion with a static human. On the other hand, no

evidence of a decrease in vocal reactivity to human proximity was found during sessions of

additional positive contacts. This would mean that positive interactions with piglets consistently

triggers the production of shorter and higher pitch grunts. These changes may be explained by the

expression of a higher arousal state experienced by the piglets while being positively handled.

Indeed, in the context of these sessions, the piglet could choose to approach and stay close to the

human, which will provide positive contacts systematically. So the piglet may anticipate to receive

positive contact and systematically being rewarded. When close to the human, observed changes in

frequency distribution of grunts (increased pitch and upshifted frequency spectrum) are known to be

markers of arousal (in the negative state in multiple mamalian species (Briefer, 2012, 2020) and pigs

(Linhart et al., 2015)). In addition, these spectral changes were also associated with shorter grunts.

Although the duration of grunts is associated with the valence of a situation, the duration may also be

an indicator of positive arousal. This hypothesis has to be taken precociously since no additional

control of arousal could be done in the present study.

This working hypothesis may explain the decrease in vocal reactivity to human proximity observed

during the standard reunion test as the HAR becomes more positive. Indeed, before the sessions of

the conditioning, positively handled piglets were habituated to a human interacting positively when

present whereas non handled piglets were not, hence, during the first standard reunion test, when the

human is present but do not interact with the piglet, positively handled and non handled piglets may

have diverging expectations regarding the presence of the static and silent human. As positively

handled piglets received positive contacts every time they were in the presence of the human, they

may have expected positive contacts when approaching and experienced an absence of reward during

the test. This has already been hypothesised in piglets deprived from human voice during interactions

after a period of habituation to it (Bensoussan et al. 2020). On the contrary, piglets that were not

positively handled at weaning never experienced additional positive contacts and being close to a

human, having the possibility to investigate them may be some kind of reward after the period of

total isolation. After the conditioning, piglets from both treatments were conditioned to receive

additional positive contacts and both groups had experienced a first standard reunion test, so they

may both experience an absence of reward during the test, which may explain a lower reaction to

human proximity, and thus fewer changes on grunt spectro-temporal features.

Last, we can raise the question whether changes in grunt structure in reaction to rewarding positive

contacts may also be associated with a specific human-pig communication. In other domestic

species, owner directed vocalisations has been shown (in cats, reviewed in (Turner, 2017); in dogs

(Gaunet et al., 2022)). In addition, studies have found similar socio-communicative behaviours

toward a human in socialized pigs and dogs (Gerencsér et al., 2019). Hence, we may profit from

testing the existence of human directed vocalisations in pigs, as consequences of their socio

communicative abilities.

Effect of human identity on piglets' perception: perspectives on HAR

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We found that the identity of the human had effects on behavioural and vocal response scores.

Piglets that were handled by the human 'AH' had higher values of behavioural proximity (ReuPC1)

than piglets handled by the human 'AV' during reunion test after a period of isolation. This effect

was not found during conditioning sessions. The effect of the human did not interact with the

conditioning time, leading to the conclusion that the difference between the two experimenters may

have established during the period of positive handling at weaning, prior to the conditioning.

Additionally, when the human 'AH' was in the room, piglets produced grunts with a more upshifted

frequency spectrum and a higher pitch than when the human 'AV' was in the room. If upshifted

grunts may be a indicator of positive higher arousal, then we may conclude that 'AH' was more

likely to trigger higher positive states than 'AV'. Interestingly, the human identity and the spatial

proximity had different effects on piglets grunts during sessions of additional positive contacts but

not when the human was static during the standard reunion test. Hence, it is possible that the way one

human interacts (behavioural and vocally) with a piglet may be more or less effective at triggering

positive emotions and thus modifications of grunt structure. Several evidence exists in the literature

that pigs discriminate humans visual and auditory cues (Bensoussan et al., 2019; Brajon et al.,

2015c). Pigs may also show behavioural changes hearing human voice (Bensoussan et al., 2020). We

may question the efficiency of different human features to generate a positive HAR. In our study,

both humans that interacted with the piglets wear exactly the same clothes and standardized their

tactile interactions toward the piglets before starting the study, and agreed on the rhythm and types of

sounds (words, intonation) to use, to minimise generating variability although no systematic controls

of the human behaviour or spectral feature of voices were performed here. It thus remains unclear

whether experimenters interacted differently or if they were initially perceived differently by piglets.

Our results show that the identity of the human may modulate piglet proximity and vocal behaviour

but the design of this experiment does not allow to find the causes of these observations (behaviour,

voice characteristics, or even odour profile). Thus, more studies of human features that are most

likely to generate a positive HAR are needed and may be of interest regarding animal welfare. In

addition, studying human-piglet relationship in a more systematic way, as in other domestic species,

for example the play behaviour in dogs (Horowitz & Hecht, 2016) or the pet directed speech

(Jeannin et al., 2017; Lansade et al., 2021), may shed light on the evolution and converging strategies

of interspecific relationships. However, the influence of human identity did not modify the general

outcomes of our study, but only decreased some effects, suggesting that this variability does not

modify the main results, but should be considered in future studies.

To conclude, we showed that degrees of familiarity toward a human could be reflected in the way

piglets vocalise in their presence, and out of it. We also showed that the spatial proximity toward a

human providing additional care could change the acoustic structure of piglet grunts. These changes

are likely to be linked to positive and more intense emotional states than when piglets are further

away from the human. However, it is still unclear whether the changes in grunt structure could also

be linked to human-animal communication and more studies are needed to determine it. We did also

show that the identity of the human may be of importance, and may generate vocal changes during

additional positive contacts that were not associated with changes in behaviour of the human. More

systematic studies of human behaviour along with pig behaviour during the human-animal

interactions would be needed to have a better understanding of the evolution of HAR, especially

interactive interspecific communication as well as providing new procedures to promote positive

welfare. We suggest that analysing vocalisations structure may be a good tool to assess the quality of
human-pig relationship and help monitor the establishment of a positive HAR.

Authors contributions

685   Conceived and designed the experiment (A.V., C.T., C.N.). Performed the experiment (A.V., C.G.).
686   Collection and processing of the acoustic and behavioural data (A.V., C.G.). Statistical analyses
687   (A.V.). Contributed to the writing of the manuscript (A.V., C.T., C.N.).

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Data availability

696   The datasets used for the study are available at (Villain et al., 2022). The folder contains all datasets
697   and a readme to match the type of analysis to the proper dataset. We have made sure to report in the
698   main text of the article which R libraries and which functions in these libraries we used. All formulas
699   of the statistical models are explicit in the text to facilitate transfer of information and replicate the
700   analysis. All libraries are open source as well.

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↔ 65.5cm

■ Location of the microphone

⬚ Virtual zones for behavioural scoring

🔴 Location of the human

▭ Proximal area ┆ Distal area

A

During conditioning

Duration (s) (log)

c
b a
a

Treatment

H   H+

B

During conditioning

VocPC2

a b
ab ab

Treatment

H   H+

C

VocPC2

Trial number

2   4   6   8   10   11

a

b

Location of piglet   away from human   close to human