**Group-size dependent punishment of idle subordinates in a cooperative breeder where helpers pay to stay**

**Stefan Fischer1, Markus Zöttl1,2, Frank Groenewoud1,3 & Barbara Taborsky1**

1Behavioural Ecology, Institute of Ecology and Evolution, University of Bern, Wohlenstrasse 50a, 3032 Hinterkappelen, Switzerland

2Department of Zoology, University of Cambridge, United Kingdom

3Behavioural Ecology and Self-Organization, Centre for Ecological and Evolutionary Studies, University of Groningen, Netherlands;

*Corresponding author:*

Stefan Fischer

email: stefan.fischer@iee.unibe.ch

## **Summary**

In cooperative breeding systems, dominant breeders sometimes tolerate unrelated individuals even if they inflict costs on the dominants. According to the ˈpay-to-stayˈ hypothesis, (i) subordinates can outweigh these costs by providing help and (ii) dominants should be able to enforce help by punishing subordinates that provide insufficient help. This requires that dominants can monitor helping and can recognize group members individually. In a field experiment we tested whether in the cooperatively breeding cichlid *Neolamprologus pulcher* subordinates increase their help after a forced ˈidleˈ period, how other group members respond to a previously idle helper, and how helper behaviour and group responses depend on group size. Previously idle helpers increased their submissiveness and they received more aggression than control helpers, suggesting that punishment occurred to enforce help. Subordinates in small groups increased their help more than those in large groups, despite receiving less aggression. When subordinates were temporarily removed, dominants in small groups were more likely to evict returning subordinates. Our results suggest that only in small groups helpers face a latent threat of punishment by breeders as predicted by the pay-to-stay hypothesis. In large groups cognitive constraints may prevent breeders to track the behaviour of a large number of helpers.

**Keywords:** conflict; cooperation; helping; territory defence; eviction; cichlids;

## **Introduction**

Cooperative breeding, where subordinate individuals help to rear a brood produced by dominant breeders, occurs in several species of invertebrates and vertebrates, including mammals, birds and fish [[1](#_ENREF_1)]. Several mechanisms have been proposed to explain how selection can favour cooperative behaviours that directly benefit other group members at the expense of the direct fitness of the actor. Evolutionary explanations distinguish between indirect and direct benefits of helping [[2](#_ENREF_2)]. Kin selection explains altruistic behaviour of individuals helping relatives [[3](#_ENREF_3), [4](#_ENREF_4)], whereas direct fitness benefits can be gained by helpers through three main routes. (1) By group augmentation through helping [[5](#_ENREF_5), [6](#_ENREF_6)], which may increase the survival prospects of helpers. (2) By increasing the chances of own future reproduction for instance through territory inheritance [[7](#_ENREF_7)], acquisition of breeding skills [[8](#_ENREF_8)] or an increase in 'social prestige', which may aid the acquisition of dominance or mating partners [[9](#_ENREF_9)]. (3) The ˈpay-to-stayˈ hypothesis [[10](#_ENREF_10)] predicts that if subordinate group members inflict costs on dominant breeders due to competition for resources and reproduction, the ensuing conflict of interest can be resolved if helpers assume important duties such as territory defence or brood care to compensate for these costs [[11](#_ENREF_11), [12](#_ENREF_12)].

Among the above mentioned mechanisms, ˈpay-to-stayˈ has received empirical [[13-22](#_ENREF_13)] and theoretical support [[11](#_ENREF_11), [12](#_ENREF_12)]. Empirical support to date stems mostly from a single species, however. In the cooperatively breeding cichlid *Neolamprologus pulcher* evidence from several experimental studies in the laboratory and the field showed that cooperation is largely based on pay-to-stay in this species [[15](#_ENREF_15), [17-19](#_ENREF_17), [21](#_ENREF_21), [23-25](#_ENREF_23)]. Evidence in other species is scare [[13](#_ENREF_13), [22](#_ENREF_22)] and reported indications of pay-to-stay may be also explained by alternative ultimate or proximate mechanisms [[see 22](#_ENREF_22)]. Studies investigating ˈpay-to-stayˈ focused mainly on predictions regarding behavioural responses of group members towards helpers not paying in accordance with need [[13-15](#_ENREF_13), [17](#_ENREF_17)]. Others investigated how the tolerance of subordinates depends on the need for help [[18](#_ENREF_18), [26](#_ENREF_26)], how helpers adjust their cooperative effort to alternative options [[19](#_ENREF_19), [27](#_ENREF_27)], how dominant individuals benefit from subordinates [[16](#_ENREF_16)] and if the amount of helping increases tolerance in the group [[28](#_ENREF_28)]. Most studies have supported predictions derived from the pay-to-stay hypothesis [but see: [16](#_ENREF_16), [28](#_ENREF_28)].

Among the mechanisms proposed to explain altruistic helping, only pay-to-stay predicts that dominant individuals enforce help by subordinates [[11](#_ENREF_11)]. Therefore, several experimental studies attempted to test whether ˈidleˈ helpers are punished by dominant individuals [[13-15](#_ENREF_13), [17](#_ENREF_17), [29](#_ENREF_29)]. While some of these studies reported an increase in aggression on return of a temporarily removed helper [[13](#_ENREF_13), [17](#_ENREF_17)] it has so far not been shown conclusively that punishment increases the helping effort of subordinates [[30-32](#_ENREF_30)]. Furthermore, it has been argued that pay-to-stay may be unlikely to maintain cooperation because of the required cognitive skills [[31](#_ENREF_31)]. Dominant individuals need to monitor helpfulness of individual subordinates and react accordingly by punishing idle helpers. Most experimental evidence for pay-to-stay has been established in small groups of cooperative breeders, sometimes well below the average group size [[13](#_ENREF_13), [15](#_ENREF_15), [24](#_ENREF_24)], hence it remains unclear if cooperation in large groups can be maintained by this mechanism as well.

The cooperatively breeding cichlid *Neolamprologus pulcher* lives in social groups containing 1 to >25 helpers varying in relatedness, size and sex [[23](#_ENREF_23), [25](#_ENREF_25), [33-36](#_ENREF_33)]. In this species many important predictions derived from the pay-to-stay hypothesis have received experimental support. *N. pulcher* helpers have been shown to increase their helping effort in the home territory both after being temporarily removed [[field experiments: 17](#_ENREF_17)] and after being prevented from helping [laboratory experiments: [14](#_ENREF_14), [15](#_ENREF_15)]. However an important prediction of pay-to-stay models, namely that breeders increase their aggression to punish ˈlazyˈ subordinates has not yet been conclusively demonstrated in this species [[14](#_ENREF_14), [15](#_ENREF_15), [17](#_ENREF_17)].

Punishment or elevated aggression by dominant individuals often leads to the eviction of subordinate group members in cooperative breeders [[18](#_ENREF_18), [37-39](#_ENREF_37)]. Theory predicts that dominants should show an increased propensity to evict 'lazy' helpers and subordinates that directly compete with dominants for reproduction [[40](#_ENREF_40)]. In *N. pulcher*, adult subordinates can steal fertilizations or bud off parts of the dominants' territory [[18](#_ENREF_18), [41-45](#_ENREF_41)]. Eviction rates of subordinates are predicted to increase with decreasing group productivity [[40](#_ENREF_40)], which is often associated with group size [[36](#_ENREF_36), [46](#_ENREF_46), [47](#_ENREF_47)]. However, if dominants suffer cognitive constraints in large groups, eviction rates should be higher in small groups where dominants are able to monitor ˈlazyˈ helpers.

Here we exposed helpers from large and small groups to three experimental treatments in the field: (1) a temporary prevention of helping behaviour of a subordinate while being present in the territory; (2) a temporary removal of a helper from the territory; and (3) a treatment controlling for handling and manipulation procedures. We aim to answer the following questions: (1) Do *N. pulcher* helpers increase their submissiveness, helping effort and/or do they receive more aggression from other group members after being prevented from helping while being present at the territory? (2) Do breeders and other helpers reduce their helping effort on return of the ˈidleˈ helper, depending on group size? (3) Are small groups more efficient in executing punishment? (4) What is the functional context responsible for elevated aggression levels towards experimentally manipulated, 'idle' helpers? Two answers have been proposed for the fourth question; (i) punishment by dominant individuals [[17](#_ENREF_17), [31](#_ENREF_31)] or (ii) competition over rank, where lower ranked individuals try to step up the queue and inherit the position of the manipulated helper [[17](#_ENREF_17), [48](#_ENREF_48)]. To distinguish between these possibilities, manipulating both the helping effort and helper presence is necessary. If punishment is the main driving force of aggression, helpers prevented from help and helpers temporarily removed from the territory should receive similar amounts of aggression, and aggression should mainly be shown by dominant individuals. Alternatively, if a helper's absence is perceived as a higher level of cheating, aggression particularly of dominant individuals should be increased, which may result in a higher eviction probability when helpers are temporarily removed from the territory. If aggression reflects conflict over rank, helpers temporarily absent from the territory, should receive aggression mainly from other helpers.

## **Methods**

*Study species*

*Neolamprologus pulcher* inhabits rocky habitats all along the shoreline of Lake Tanganyika, from 3 to 45 m depth [[49](#_ENREF_49)]. Territories of *N. pulcher* cluster into colonies, where several social groups live in close proximity to each other [[17](#_ENREF_17), [35](#_ENREF_35), [50](#_ENREF_50)]. The groups defend distinct patches of stones containing a central breeding shelter and often additional shelters for the other group members to hide from predator attacks [[36](#_ENREF_36), [51](#_ENREF_51)]. Eggs are attached to the inner walls of the breeding shelters and are mainly cared for by the breeder females and small helpers [[23](#_ENREF_23)]. Larger helpers engage in defence against con- and heterospecific territory intruders and egg and fish predators, in digging out shelters for breeding and for protection from predators [[21](#_ENREF_21), [23](#_ENREF_23), [33](#_ENREF_33), [52](#_ENREF_52)].

*General field methods*

The study was conducted at the southern tip of Lake Tanganyika at Kasakalawe point near Mpulungu, Zambia, from September to December 2011 and 2012, by SCUBA diving. Our experimental population consisted of five colonies of different sizes. From these colonies we haphazardly allocated 14 large (range: 16–25 members, including the breeder pair) and 21 small family groups (range: 5-7 members) to experimental groups. We haphazardly caught the experimental fish using a Plexi glass tube and two hand nets. Their standard lengths (SL; from the tip of the snout to the posterior end of the vertebral column) were measured using a measuring board with a 1 mm grid. Only helpers between 34-48 mm SL were used for the experiment. We took a small fin clip from one of the last dorsal fin rays allowing us to recognise the focal helper during the following 3-4 days. As in previous studies, the removed fin rays regrew within 1-2 months [[e.g. 52](#_ENREF_52)], and all marked fish resumed normal behaviour within a few minutes after handling (S. Fischer & F. Groenewould, pers. obs.). Our study adheres to the legal requirements of Zambia and Switzerland and was conducted under the Memorandum of Understanding of 2008 between the Ministry of Agriculture and Cooperatives, Zambia, the University of Zambia, the Universities of Basel and Bern, Switzerland and the University of Graz, Austria. All experiments were conducted according to the guidelines of the Association for the Study of Animal Behaviour.

*Experimental treatments*

'Prevented-help' treatment (n=20). To prevent a focal fish from helping, it was placed in a clear plastic cylinder (height: 10cm, diameter: 8.5cm) equipped with a shelter built from two small PVC-plates. Holes in the lid of the cylinder allowed for water exchange. The plastic cylinder holding the focal helper was placed close to its original shelter within the home territory. A few minutes after start of confinement, the focal helpers started to move freely in the cylinder and did not show obvious signs of stress. One individual even returned to the cylinder deliberately after the lid had been removed to release the fish back in its home territory. Throughout this treatment the focal helper was present in the territory but did not participate in the typical helping duties such as territory defence and maintenance, and brood care. After 24h we released the focal helper from the cylinder and recorded its behaviour and the behaviours of all other group members for 10min, which is sufficiently long to obtain a representative sample of the behavioural repertoire of *N. pulcher* [see [14](#_ENREF_14), [53](#_ENREF_53)].

Control treatment (n=20). A focal helper was placed in the plastic cylinder and, as in the prevented-help treatment, it was individually marked and placed close to its original shelter at the territory. It was released after 5min; the observer returned 24h later and recorded the behaviour of the focal helper and the group members for 10min. Where possible, we performed the prevented-help and control treatments in the same group (only twice we had to use different groups), using similar sized individuals (comparison of focal SL between controls and treatments: Welch two sample t-test, t=1.24, df=36.87, p=0.22). This treatment served to control for potential effects of the catching and placement in the cylinder on the subsequent behaviour of the focal helper, and for the potential disturbance of other group members by these manipulations. It does only partly control for possible effects of the releasing procedure, because in the prevented-help treatment the behavioural observations directly followed the release. To test for the potential influence of this temporal proximity, we performed a separate series of behavioural recordings on different family groups, which is presented in "Electronic supplementary material".

Helper removal treatment (n=20). A focal fish was caught and transferred to a 0.8×0.8×0.6m mesh cage that was installed outside of the colony area. After 24h the fish was relocated to its family. After releasing the fish, the behaviours of the focal helper and of all other group members were recorded for 10min. After this observation we visited each family after 24h to confirm the acceptance status of the focal helper. For a graphical representation of all treatments see ESM Fig.A1.

*Behavioural recordings*

During the 10min behavioural observations, we recorded all submissive and aggressive behaviours of the focal fish (according to an established ethogram of *N. pulcher;* [[54-56](#_ENREF_54)]) shown towards other group members, predators and space competitors, and all aggressive behaviours of all other group members towards the focal helper, predators and space competitors. To score the behaviours we used a Psion handheld computer (Psion Teklogix Workabout Pro 7525) packed within a waterproof plastic bag and equipped with Noldus Pocket Observer 3.0. For our analysis we distinguished between aggressive behaviours towards group members and towards non-group members (intruders). Aggression or 'territory defence' against intruding con- and heterospecific individuals was combined into a single 'defence' variable for analysis.

We scored focal helpers as either evicted or accepted 24h after the release. Accepted focal helpers resumed immediately their helping duties within the group. Evicted helpers received aggression after the release, they subsequently left their group and did not return to the groups within 24h.

*Statistical analyses*

To compare the submissive behaviours, received aggression, the frequency of defence behaviours of focal helpers, the frequency of the total group defence behaviours and the frequency of the defence behaviours of breeders and other helpers between the treatments and group sizes, generalized linear mixed models (GLMM) with log link function were used to account for a Poisson error structure. Treatment and group size of the family (large or small) were always fitted as fixed effects, the SL of the focal helper was included as covariate and group identity as a random factor to account for repeated measurements within the same group. Full models always included the interaction between treatment and group size. To analyse the defence behaviour of breeders and other helpers we included the class of group member (breeder, helper) as a factor in the model. Here, all 2-way and 3-way interactions between the treatment, group size and the class factor were included in the model. To simplify the models we used stepwise backward elimination of non-significant interaction terms [[57](#_ENREF_57), [58](#_ENREF_58)]. All models were checked for over-dispersion [[57](#_ENREF_57)].

To analyse if competition over rank or punishment is the main behavioural mechanism driving elevated aggression levels in the prevented help treatment we calculated per capita aggression rates towards the focal helper from breeders and from helpers. Because *N. pulcher* groups have linear size-based hierarchies, potential competitors for a focal helper's rank can only be either of the same size or smaller than the focal helper [[55](#_ENREF_55)]. As we could not catch and precisely measure all group members, we estimated the sizes of non-captured group members to the nearest 1.0 cm, resulting in a size distribution of 1 cm length-classes. For the analysis of per capita aggression rates, we then divided helpers in two groups being either larger (all 1-cm classes larger than the focal helper) or 'similar sized' (i.e. same or a smaller 1-cm class than focal). However there was only one case where helpers 2 cm smaller than the focal participated in aggression towards the focal helper. The per capita received aggression rates were highly zero inflated, therefore we transformed the data into a binomial data structure, where focal helpers either received aggression or not. To compare the probability of receiving aggression from breeders, larger or similar-sized helpers, we used a GLMM with a logit link to account for a binomial error structure. Here, the type of group members (breeder, larger and similar-sized helpers), as well as group size (large, small) were fitted as fixed effects. All interactions of the type of group members and group size were included in the model. To compare if helpers were treated differently in large or in small groups in the helper-removal treatment, and after being removed vs. after being prevented from helping while present, we used eviction rates rather than aggression frequencies. This was necessary, because it was not possible to obtain the received aggression rates of evicted individuals, as the timespan between release and eviction was very short. Eviction rates were statistically compared using Fisher's Exact tests. For statistical analysis we used R 3.0.1 [[59](#_ENREF_59)] with the package ˈlme4ˈ[[60](#_ENREF_60)]. Data are deposited in the Dryad digital depository [[61](#_ENREF_61)].

## **Results**

*Submission, helping effort and received aggression after experimental prevention of help*

Subsequently to being prevented from helping by confinement in a clear cylinder (ˈprevented helpˈ treatment) only focal helpers in small groups increased their amount of defence compared to helpers in the control treatment (Fig. 1a, Table 1a).

After confinement in the prevented-help treatment, focal helpers showed more submissive behaviour towards other group members and received more aggression from group members than after the control treatment (Fig. 1b, c; Table 1b, c). In large groups, focal helpers showed a higher increase in submission after being confined in the cylinder compared to focal helpers in small groups (Fig. 1b; significant interaction term treatment × group size, Table 1b). In larger groups, focal helpers generally tended to show more submission and they received more aggression (Fig.1b, c; Table 1b, c).

*Influence of group size on helping and eviction rates*

After focal helpers were released in the prevented help treatment, the total group defence behaviours decreased in all groups compared to the control treatment (Fig. 2a; Table 2a). Large groups reduced their total defence behaviours in this situation more than small groups (Fig. 2a; Table 2a). When comparing the defence behaviours of breeders and of other helpers in small and large groups, we found that helpers decreased their defence significantly in large groups, whereas breeders did so in small groups (Fig. 2b, c; see 3-way interaction in Table 2b).

In the helper removal treatment, focal helpers faced a significantly higher eviction probability in small groups compared to helpers in large groups (Fisher's Exact Test, *N*=20, *P*=0.003, Fig. 2d). In the prevented help treatment all helpers were re-integrated after the release.

*Causes of change in aggression*

To disentangle whether changes in the other group members' aggression reflect punishment or rather conflicts over rank between helpers, we analysed per capita aggression rates and compared the eviction probabilities of focal helpers after being prevented from helping and after being temporarily removed from the territory. After prevention from helping, focal helpers in small groups received aggression mainly from breeders, whereas in large groups mainly similar sized helpers displayed aggression towards the focal helper (Fig. 3a; Table 3).

Helpers that had been temporarily removed from the territory were significantly more often evicted from their home territory than helpers that had been only prevented from helping but had been present in the territory (Fisher's Exact Test, *N*=20, *P*=0.001, Fig. 3b).

## **Discussion**

In this study we experimentally manipulated the helping behaviour of subordinates in a cooperative breeder living under natural conditions. In our prevented-help treatment we managed for the first time to reduce helping behaviour in a field experiment without removing the helper from the territory, thus allowing us to test for responses to 'idle' helpers. Other studies experimentally manipulating behaviours of subordinates have either varied the need for help [e.g. [62](#_ENREF_62), [63](#_ENREF_63)] or increased helping behaviours by supplementary feeding experiments [[64](#_ENREF_64)].

Our results suggest that helpers in small groups compensated for an ˈidleˈ period by increased investment in territory defence, and that they faced a higher eviction probability after temporary removal from the territory. Furthermore all helpers increased their submissiveness towards other group members, which confirms results of a previous laboratory experiment [[15](#_ENREF_15)]. After being prevented from helping, helpers in large groups received more aggression, probably because more group members were present, and they increased submission more than helpers in small groups, which may indicate an appeasement strategy [[see 15](#_ENREF_15)]. However, in contrast to previous experiments, where the breeders' aggression was not found to increase significantly after the helpers' resumption of their normal duties [[15](#_ENREF_15), [17](#_ENREF_17)], here we found that breeders and other helpers increased their aggression towards focal helpers after being experimentally prevented to help. Focal helpers increased their defence behaviours after being prevented to help only in small groups. A closer inspection of the elevated aggression rates in the prevented help treatment revealed that mainly breeders directed their aggression towards ˈidleˈ focal helpers in small, but not in large groups. This supports the hypothesis that punishment may be the main function of elevated aggression levels in small groups, where breeders may be better able to control helpers and punish them when the breeders' expectation to receive help in territory defence has been violated. In large groups, mainly helpers similar in size to the focal helper showed elevated aggression levels. This did not result in higher cooperative effort of the focal helper, suggesting that the ultimate function of increased aggression towards the focal helper in large groups might be competition over rank among group members. This is consistent with previous experimental work suggesting that competition over rank rather than coercion might be responsible for increased aggression after helper removal [[17](#_ENREF_17), [32](#_ENREF_32)].

Punishment, where dominant individuals penalize ˈidleˈ subordinates, is a crucial component of cooperation based on the pay-to-stay mechanism [[11](#_ENREF_11)]. It has been argued that unequivocal evidence for punishment is lacking in cooperatively breeding animals [[31](#_ENREF_31)]. In two other studies where helpers were experimentally removed both breeders and other helpers increased their aggression towards returning helpers [[13](#_ENREF_13), [29](#_ENREF_29)]. However, an increase of aggression by group members in 'idle' helpers has not been documented when the behaviour of helpers was manipulated while they remained in the territory. Instead in a previous experiment, such treatment caused *N. pulcher* helpers of small groups to pre-emptively compensate for experimentally induced idleness by increasing helping and submissive behaviour [[15](#_ENREF_15)]. Also, it has not been shown that dominant group members increase their aggression towards 'idle' helpers in dependence of group size.

Our results suggest that in large groups' factors other than the breeders' demand for help [[18](#_ENREF_18), [40](#_ENREF_40), [65](#_ENREF_65), [66](#_ENREF_66)] might determine aggression rates. Our results are consistent with the idea that members of large groups may be limited in their ability to reliably track the behaviours and presence of all other group members. Therefore a ˈidleˈ or absent helper may more easily remain undetected by other group members in large groups, whereas it might be possible to monitor activities of each helper in small groups. *N. pulcher* breeders are able to individually recognize subordinate group members [[67](#_ENREF_67), [68](#_ENREF_68)]. It may be more difficult in larger groups to remember all own helpers beyond prolonged times of separation. Based on a theoretical model and a simulation study, similar arguments have been put forward for banded mongooses (*Mungos mungo*) and for humans, where either reproductive restraint or monitoring of other group members does not work in large groups [[37](#_ENREF_37), [69](#_ENREF_69)]. Thus proximate constraints may influence the mutual exchange mechanism maintaining cooperation in *N. pulcher*. Alternatively, large group size may erode the benefits of punishing 'idle' subordinates for the breeders and hence they refrain from punishing despite having the cognitive ability to do so. Such an adaptive explanation has been suggested by models of the evolution of reciprocity and altruistic punishment [[70](#_ENREF_70)]. Our data do not allow to differentiate whether proximate constraints or adaptive causes explain the lack of punishment in large groups.

Remarkably, all conclusive experimental evidence for pay-to-stay results from experimental laboratory studies using small group size [[15: Group size 4](#_ENREF_15), [24: Group size 3](#_ENREF_24)]. If the pay-to-stay mechanism is limited by the ability to enforce help and to coerce subordinates also in large groups, with increasing group size the function of helping might gradually switch from pay-to-stay towards group augmentation benefits that is, a mutualism without coercion. This idea is consistent with data suggesting high fitness pay-offs for breeders and helpers in large groups [[35](#_ENREF_35), [36](#_ENREF_36)].

With the notable exception of *N. pulcher* [[15](#_ENREF_15), [17-19](#_ENREF_17), [21](#_ENREF_21), [23-25](#_ENREF_23)], among cooperative breeders there seems to be little conclusive evidence showing that dominants use aggression to elevate the work rates of helpers [[71](#_ENREF_71)]. This raises the question how social and ecological conditions in *N. pulcher* differ from other known cooperative systems and which ecological factors enable dominant breeders to enforce help from subordinates? First, *N. pulcher* groups have a low average relatedness among group members [[25](#_ENREF_25), [34](#_ENREF_34)] due to a frequent exchange of the dominant breeders of groups by predation, and therefore kin selected benefits of helping are small, particularly for older helpers in their late juvenile and early adult stage. Second, outside options for helpers are strongly constrained because of an extremely high predation risk outside of groups [[51](#_ENREF_51)]. Founding of new groups appears to be difficult, as it was observed only rarely and small groups have a low chance of persistence [[35](#_ENREF_35)]. Therefore voluntary dispersal only occurs if vacancies arise due to predation in existing groups [[19](#_ENREF_19)]. Thus, the particular relatedness structure and ecology of *N. pulcher* combined with the unique possibility to experimentally manipulate this system such that alternative mechanisms can be excluded may explain why the pay-to-stay mechanism has received its strongest support to date in this cichlid fish. We would like to propose that the potential for helper coercion should be studied in cooperative breeders, which share the prerequisites of low within-group relatedness and constrained outside options. Potential candidates may be found among cooperative breeding bird species such as pied kingfisher or white-winged trumpeters, where unrelated individuals can join in alloparental care and single pairs are unable to defend a territory to raise young [[for a review see 72](#_ENREF_72)].

In conclusion, we found that punishment of lazy helpers can explain why subordinates pay to stay in *N. pulcher*, at least in small groups. Our study shows that punishment of ˈidleˈ helpers is not dependent on the physical absence of the subordinate [e.g.: [13](#_ENREF_13), [17](#_ENREF_17)]. Instead, dominant individuals seem to directly evaluate the helping effort of subordinates in this species, in which help is traded for acceptance in the territory. Furthermore, our results suggest that group size may influence the regulation of conflicts between group members by changing the relative importance of breeders and helpers in penalizing ˈidleˈ subordinates. We propose that preventing subordinates from helping duties, without removing them from the territory should be generally adopted in studies focusing on the compensation ability of ˈidleˈ helpers [e.g.: [13](#_ENREF_13), [17](#_ENREF_17), [73](#_ENREF_73)]. Our results have important general implications for the understanding of cooperation in highly social species, where unrelated individuals help dominant breeders to raise their offspring [[24](#_ENREF_24), [74-76](#_ENREF_74)].

## **Acknowledgements**

We would like to thank the whole Tanganyika field crews 2011 and 2012, in particular, Arne Jungwirth, Joachim G. Frommen and Mathilde Bessert-Nettelbeck for helping to collect the data. We are grateful to Redouan Bshary and Michael Taborsky for discussion and comments on an earlier version of the manuscript. The manuscript was greatly improved by the comments of two anonymous reviewers. Ruben Shapola and Danny Sinyinza supported this study by logistic help. This project was financially supported by the Swiss National Science Foundation (SNSF).

## **References**

1. Bergmüller R., Johnstone R.A., Russell A.F., Bshary R. 2007 Integrating cooperative breeding into theoretical concepts of cooperation. *Behav. Processes* **76**(2), 61-72. (doi:10.1016/j.beproc.2007.07.001).

2. Lehmann L., Keller L. 2006 The evolution of cooperation and altruism - a general framework and a classification of models. *J. Evol. Biol.* **19**(5), 1365-1376. (doi:10.1111/j.1420-9101.2006.01119.x).

3. Hamilton W.D. 1964 The genetical evolution of social behaviour I. *J. Theor. Biol.* **7**(1), 1-&. (doi:10.1016/0022-5193(64)90038-4).

4. Hamilton W.D. 1964 The genetical evolution of social behaviour II. *J. Theor. Biol.* **7**(1), 17-&. (doi:10.1016/0022-5193(64)90039-6).

5. Woolfenden G.E. 1975 Florida scrub jay helpers at nest. *Auk* **92**(1), 1-15.

6. Kokko H., Johnstone R.A., Clutton-Brock T.H. 2001 The evolution of cooperative breeding through group augmentation. *Proc. R. Soc. B* **268**(1463), 187-196. (doi:10.1098/rspb.2000.1349).

7. Brown J.L., Brown E.R. 1984 Parental facilitation: Parent-offspring relations in communally breeding birds. *Behav. Ecol. Sociobiol.* **14**(3), 203-209. (doi:10.1007/bf00299620).

8. Komdeur J. 1996 Influence of helping and breeding experience on reproductive performance in the Seychelles warbler: a translocation experiment. *Behav. Ecol.* **7**(3), 326-333.

9. Zahavi A. 1974 Communal nesting by Arabian babbler: a case of individual selection. *Ibis* **116**(1), 84-87. (doi:10.1111/j.1474-919X.1974.tb00225.x).

10. Gaston A.J. 1978 The Evolution of group territorial behavior and cooperative breeding. *Am. Nat.* **112**(988), 1091-1100. (doi:10.1086/283348).

11. Kokko H., Johnstone R.A., Wright J. 2002 The evolution of parental and alloparental effort in cooperatively breeding groups: when should helpers pay to stay? *Behav. Ecol.* **13**(3), 291-300. (doi:10.1093/beheco/13.3.291).

12. Hamilton I.M., Taborsky M. 2005 Unrelated helpers will not fully compensate for costs imposed on breeders when they pay to stay. *Proc. R. Soc. B* **272**(1561), 445-454. (doi:10.1098/rspb.2004.2961).

13. Mulder R.A., Langmore N.E. 1993 Dominant males punish helpers for temporary defection in superb fairy-wrens. *Anim. Behav.* **45**(4), 830-833. (doi:10.1006/anbe.1993.1100).

14. Zöttl M., Fischer S., Taborsky M. 2013 Partial brood care compensation by female breeders in response to experimental manipulation of alloparental care. *Anim. Behav.* **85**(6), 1471-1478. (doi:10.1016/j.anbehav.2013.03.045).

15. Bergmüller R., Taborsky M. 2005 Experimental manipulation of helping in a cooperative breeder: helpers ‘pay to stay’ by pre-emptive appeasement. *Anim. Behav.* **69**(1), 19-28. (doi:10.1016/j.anbehav.2004.05.009).

16. Mitchell J.S. 2003 Social correlates of reproductive success in false clown anemonefish: subordinate group members do not pay-to-stay. *Evol. Ecol. Res.* **5**(1), 89-104.

17. Balshine-Earn S., Neat F.C., Reid H., Taborsky M. 1998 Paying to stay or paying to breed? Field evidence for direct benefits of helping behavior in a cooperatively breeding fish. *Behav. Ecol.* **9**(5), 432-438. (doi:10.1093/beheco/9.5.432).

18. Taborsky M. 1985 Breeder-helper conflict in a cichlid fish with broodcare helpers: an experimental analysis. *Behaviour* **95**, 45-75. (doi:10.1163/156853985x00046).

19. Bergmüller R., Heg D., Taborsky M. 2005 Helpers in a cooperatively breeding cichlid stay and pay or disperse and breed, depending on ecological constraints. *Proc. R. Soc. B* **272**(1560), 325-331. (doi:10.1098/rspb.2004.2960).

20. Bruintjes R., Taborsky M. 2008 Helpers in a cooperative breeder pay a high price to stay: effects of demand, helper size and sex. *Anim. Behav.* **75**(6), 1843-1850. (doi:10.1016/j.anbehav.2007.12.004).

21. Heg D., Taborsky M. 2010 Helper response to experimentally manipulated predation risk in the cooperatively breeding cichlid *Neolamprologus pulcher*. *Plos ONE* **5**(5), e10784. (doi:10.1371/journal.pone.0010784).

22. MacLeod K.J., Nielsen J.F., Clutton-Brock T.H. 2013 Factors predicting the frequency, likelihood and duration of allonursing in the cooperatively breeding meerkat. *Anim. Behav.* **86**(5), 1059-1067. (doi:10.1016/j.anbehav.2013.09.012).

23. Taborsky M. 1984 Broodcare helpers in the cichlid fish *Lamprologus brichardi*: their costs and benefits. *Anim. Behav.* **32**(NOV), 1236-1252. (doi:10.1016/s0003-3472(84)80241-9).

24. Zöttl M., Heg D., Chervet N., Taborsky M. 2013 Kinship reduces alloparental care in cooperative cichlids where helpers pay-to-stay. *Nat. Commun.* **4**, 1341. (doi:10.1038/ncomms2344).

25. Stiver K.A., Dierkes P., Taborsky M., Gibbs H.L., Balshine S. 2005 Relatedness and helping in fish: examining the theoretical predictions. *Proc. R. Soc. B* **272**(1572), 1593-1599. (doi:10.1098/rspb.2005.3123).

26. Zöttl M., Frommen J.G., Taborsky M. 2013 Group size adjustment to ecological demand in a cooperative breeder. *Proc. R. Soc. B* **280**(1756), 20122772. (doi:10.1098/rspb.2012.2772).

27. Zöttl M., Chapuis L., Freiburghaus M., Taborsky M. 2013 Strategic reduction of help before dispersal in a cooperative breeder. *Biol. Lett.* **9**(1), 20120878. (doi:10.1098/rsbl.2012.0878).

28. Koenig W.D., Walters E.L. 2011 Age-related provisioning behaviour in the cooperatively breeding acorn woodpecker: testing the skills and the pay-to-stay hypotheses. *Anim. Behav.* **82**(3), 437-444. (doi:10.1016/j.anbehav.2011.05.028).

29. Reeve H.K. 1992 Queen activation of lazy workers in colonies of the eusocial naked mole-rat. *Nature* **358**(6382), 147-149. (doi:10.1038/358147a0).

30. Cant M.A. 2011 The role of threats in animal cooperation. *Proc. R. Soc. B* **278**(1703), 170-178. (doi:10.1098/rspb.2010.1241).

31. Raihani N.J., Thornton A., Bshary R. 2012 Punishment and cooperation in nature. *Trends Ecol. Evol.* **27**(5), 288-295. (doi:10.1016/j.tree.2011.12.004).

32. Wong M., Balshine S. 2011 The evolution of cooperative breeding in the African cichlid fish, *Neolamprologus pulcher*. *Biol. Rev.* **86**(2), 511-530. (doi:10.1111/j.1469-185X.2010.00158.x).

33. Taborsky M., Limberger D. 1981 Helpers in Fish. *Behav. Ecol. Sociobiol.* **8**(2), 143-145. (doi:10.1007/bf00300826).

34. Dierkes P., Heg D., Taborsky M., Skubic E., Achmann R. 2005 Genetic relatedness in groups is sex-specific and declines with age of helpers in a cooperatively breeding cichlid. *Ecol. Let.* **8**(9), 968-975. (doi:10.1111/j.1461-0248.2005.00801.x).

35. Heg D., Brouwer L., Bachar Z., Taborsky M. 2005 Large group size yields group stability in the cooperatively breeding cichlid *Neolamprologus pulcher*. *Behaviour* **142**, 1615-1641. (doi:10.1163/156853905774831891).

36. Balshine S., Leach B., Neat F., Reid H., Taborsky M., Werner N. 2001 Correlates of group size in a cooperatively breeding cichlid fish (*Neolamprologus pulcher*). *Behav. Ecol. Sociobiol.* **50**(2), 134-140. (doi:10.1007/s002650100343).

37. Cant M.A., Hodge S.J., Bell M.B.V., Gilchrist J.S., Nichols H.J. 2010 Reproductive control via eviction (but not the threat of eviction) in banded mongooses. *Proc. R. Soc. B* **277**(1691), 2219-2226. (doi:10.1098/rspb.2009.2097).

38. Clutton-Brock T.H., Brotherton P.N., Smith R., McIlrath G.M., Kansky R., Gaynor D., O'Riain M.J., Skinner J.D. 1998 Infanticide and expulsion of females in a cooperative mammal. *Proc. R. Soc. B* **265**(1412), 2291-2295. (doi:10.1098/rspb.1998.0573).

39. Mulder R.A. 1995 Natal and breeding dispersal in a cooperative, extra-group-mating bird. *J. Avian Biol.* **26**(3), 234-240. (doi:10.2307/3677324).

40. Johnstone R.A., Cant M.A. 1999 Reproductive skew and the threat of eviction: a new perspective. *Proc. R. Soc. B* **266**(1416), 275-279.

41. Bruintjes R., Bonfils D., Heg D., Taborsky M. 2011 Paternity of subordinates raises cooperative effort in cichlids. *Plos ONE* **6**(10). (doi:10.1371/journal.pone.0025673).

42. Heg D., Hamilton I.M. 2008 Tug-of-war over reproduction in a cooperatively breeding cichlid. *Behav. Ecol. Sociobiol.* **62**(8), 1249-1257. (doi:10.1007/s00265-008-0553-0).

43. Heg D., Jutzeler E., Bonfils D., Mitchell J.S. 2008 Group composition affects male reproductive partitioning in a cooperatively breeding cichlid. *Mol. Ecol.* **17**(19), 4359-4370. (doi:10.1111/j.1365-294X.2008.03920.x).

44. Heg D., Bergmüller R., Bonfils D., Otti O., Bachar Z., Burri R., Heckel G., Taborsky M. 2006 Cichlids do not adjust reproductive skew to the availability of independent breeding options. *Behav. Ecol.* **17**(3), 419-429. (doi:10.1093/beheco/arj056).

45. Dierkes P., Taborsky M., Kohler U. 1999 Reproductive parasitism of broodcare helpers in a cooperatively breeding fish. *Behav. Ecol.* **10**(5), 510-515. (doi:10.1093/beheco/10.5.510).

46. Doerr E.D., Doerr V.A.J. 2007 Positive effects of helpers on reproductive success in the brown treecreeper and the general importance of future benefits. *J. Anim. Ecol.* **76**(5), 966-976. (doi:10.1111/j.1365-2656.2007.01280.x).

47. Clutton-Brock T.H., Hodge S.J., Flower T.P. 2008 Group size and the suppression of subordinate reproduction in Kalahari meerkats. *Anim. Behav.* **76**(3), 689-700. (doi:10.1016/j.anbehav.2008.03.015).

48. Field J., Cant M.A. 2009 Social stability and helping in small animal societies. *Phil. Trans. R. Soc.* **364**(1533), 3181-3189. (doi:10.1098/rstb.2009.0110).

49. Duftner N., Sefc K.M., Koblmüller S., Salzburger W., Taborsky M., Sturmbauer C. 2007 Parallel evolution of facial stripe patterns in the *Neolamprologus brichardi/pulcher* species complex endemic to Lake Tanganyika. *Mol. Phylogenet. Evol.* **45**(2), 706-715. (doi:10.1016/j.ympev.2007.08.001).

50. Limberger D. 1983 Pairs and harems in a cichlid fish, *Lamprologus brichardi*. *J. Comp. Eth.* **62**(2), 115-144.

51. Heg D., Bachar Z., Brouwer L., Taborsky M. 2004 Predation risk is an ecological constraint for helper dispersal in a cooperatively breeding cichlid. *Proc. R. Soc. B* **271**(1555), 2367-2374. (doi:10.1098/rspb.2004.2855).

52. Bruintjes R., Taborsky M. 2011 Size-dependent task specialization in a cooperative cichlid in response to experimental variation of demand. *Anim. Behav.* **81**(2), 387-394. (doi:10.1016/j.anbehav.2010.10.004).

53. Bruintjes R., Hekman R., Taborsky M. 2010 Experimental global food reduction raises resource acquisition costs of brood care helpers and reduces their helping effort. *Funct. Ecol.* **24**(5), 1054-1063. (doi:10.1111/j.1365-2435.2010.01715.x).

54. Taborsky M. 1982 Brutpflegehelfer beim Cichliden *Lamprologus brichardi*, Poll (1974): Eine Kosten/Nutzen Analyse [Dissertation], University of Vienna.

55. Hamilton I.M., Heg D., Bender N. 2005 Size differences within a dominance hierarchy influence conflict and help in a cooperatively breeding cichlid. *Behaviour* **142**, 1591-1613. (doi:10.1163/156853905774831846).

56. Reddon A.R., O'Connor C.M., Marsh-Rollo S.E., Balshine S. 2012 Effects of isotocin on social responses in a cooperatively breeding fish. *Anim. Behav.* **84**(4), 753-760. (doi:10.1016/j.anbehav.2012.07.021).

57. Bolker B.M., Brooks M.E., Clark C.J., Geange S.W., Poulsen J.R., Stevens M.H., White J.S. 2009 Generalized linear mixed models: a practical guide for ecology and evolution. *Trends Ecol. Evol.* **24**(3), 127-135. (doi:10.1016/j.tree.2008.10.008).

58. Engqvist L. 2005 The mistreatment of covariate interaction terms in linear model analyses of behavioural and evolutionary ecology studies. *Anim Beh* **70**(4), 967-971. (doi:10.1016/j.anbehav.2005.01.016).

59. R Core Development Team. 2013 *R: A language and environment for statistical computing*.Vienna, Austia

60. Bates D., Maechler M., Bolker B. 2013 *lme4: Linear mixed-effects models using S4 classes. R package version 0.999999-2.*

61. Fischer S., Zöttl M., Groenewoud F., Taborsky B. 2014 Group-size dependent punishment of idle subordinates in a cooperative breeder where helpers pay to stay. *Dryad Digital Respository* (doi:10.5061/dryad.s3720)

62. McDonald P.G., Kazem A.J.N., Wright J. 2009 Cooperative provisioning dynamics: fathers and unrelated helpers show similar responses to manipulations of begging. *Anim. Behav.* **77**(2), 369-376. (doi:10.1016/j.anbehav.2008.10.009).

63. Bell M.B.V. 2007 Cooperative begging in banded mongoose pups. *Curr. Biol.* **17**(8), 717-721. (doi:10.1016/j.cub.2007.03.015).

64. Wright J., Dingemanse N.J. 1999 Parents and helpers compensate for experimental changes in the provisioning effort of others in the Arabian babbler. *Anim. Behav.* **58**, 345-350. (doi:10.1006/anbe.1999.1152).

65. Cant M.A., Field J. 2005 Helping effort in a dominance hierarchy. *Behav. Ecol.* **16**(4), 708-715. (doi:10.1093/beheco/ari051).

66. Heinsohn R., Legge S. 1999 The cost of helping. *Trends Ecol. Evol.* **14**(2), 53-57. (doi:10.1016/s0169-5347(98)01545-6).

67. Hert E. 1985 Individual recognition of helpers by the breeders in the cichlid fish *Lamprologus brichardi* (Poll, 1974). *J. Comp. Eth.* **68**(4), 313-325.

68. Balshine-Earn S., Lotem A. 1998 Individual recognition in a cooperatively breeding cichlid: evidence from video playback experiments. *Behaviour* **135**, 369-386.

69. Carpenter J.P. 2007 Punishing free-riders: how group size affects mutual monitoring and the provision of public goods. *Game. Econ. Behav.* **60**(1), 31-51. (doi:10.1016/j.geb.2006.08.011).

70. Boyd R., Richerson P.J. 1988 The evolution of reciprocity in sizable groups. *J. Theor. Biol.* **132**(3), 337-356. (doi:[http://dx.doi.org/10.1016/S0022-5193(88)80219-4)](http://dx.doi.org/10.1016/S0022-5193%2888%2980219-4%29).

71. Santema P., Clutton-Brock T.H. 2012 Dominant female meerkats do not use aggression to elevate work rates of helpers in response to increased brood demand. *Anim. Behav.* **83**(3), 827-832. (doi:10.1016/j.anbehav.2011.12.032).

72. Riehl C. 2013 Evolutionary routes to non-kin cooperative breeding in birds. *Proc. R. Soc. B* **280**(1772), 7. (doi:10.1098/rspb.2013.2245).

73. Baglione V., Canestrari D., Chiarati E., Vera R., Marcos J.M. 2010 Lazy group members are substitute helpers in carrion crows. *Proc. R. Soc. B* **277**(1698), 3275-3282. (doi:10.1098/rspb.2010.0745).

74. Clutton-Brock T.H., Brotherton P.N., O'Riain M.J., Griffin A.S., Gaynor D., Sharpe L., Kansky R., Manser M.B., McIlrath G.M. 2000 Individual contributions to babysitting in a cooperative mongoose, *Suricata suricatta*. *Proc. R. Soc. B* **267**(1440), 301-305. (doi:10.1098/rspb.2000.1000).

75. Wright J., McDonald P.G., te Marvelde L., Kazem A.J., Bishop C.M. 2010 Helping effort increases with relatedness in bell miners, but 'unrelated' helpers of both sexes still provide substantial care. *Proc. R. Soc. B* **277**(1680), 437-445. (doi:10.1098/rspb.2009.1360).

76. Clutton-Brock T.H. 2009 Cooperation between non-kin in animal societies. *Nature* **462**(7269), 51-57. (doi:10.1038/nature08366).

**Table and Figure captions**

**Table 1:** Comparison of the (a) the absolute frequency of defence behaviours, (b) submissive behaviour and (c) received aggression of focal helpers in the prevented help and control treatments. Reference categories for estimates of factor 'Treatment': control treatment; of factor 'Group size': large groups. *N*=21 groups; *P*-values <0.05 are highlighted in bold, and 0.05<*P*<0.1 are italicised.

**Table 2**: Comparisons of (a) the total group defence and (b) the defence of all other group members against conspecifics and predators in the prevented help and control treatments. Reference categories for estimates of factor 'Treatment': control treatment; of factor 'Group size': large groups; and of factor 'Class': breeders. *N*=21 groups; *P*-values <0.05 are highlighted in bold.

**Table 3**:Comparison of the per capita received aggression of focal helpers in the prevented help treatment. Reference categories for estimates of factors 'Larger helper' and 'Similar helpers': breeder; and of factor 'Group size': large groups. N=20 groups; *P*-values <0.05 are highlighted in bold, and 0.05<*P*<0.1 are italicised.

**Fig. 1:** Comparisons between social behaviours per 10 minutes of observations in the control treatment and prevented-help treatment (medians and interquartile ranges). (a) Defence behaviour of focal helpers, (b) submission of focal helpers; (c) aggression received by focal helpers from all other group members. Triangles represent small groups and circles large groups.

**Fig. 2**: Influence of group size on defence behaviours per 10 minutes of observation in the control and the prevented help treatment [(a)-(c)] and eviction rates in the removal treatment (d). (a) Total group defence; (b) defence behaviour of other helpers; (c) defence behaviour of breeders; (d) comparison of the number of accepted and evicted focal helpers between small and large groups in the helper removal treatment. (a)-(c) Triangles represent small groups and circles large groups; medians and interquartile ranges are shown; in (d) dark grey bars represent accepted focal helpers and light grey bars evicted focal helpers.

**Fig. 3**: (a) Comparison of the probability of the per capita received aggression from breeders, larger and similar-sized helpers between small and large groups in the prevented help treatment (medians and interquartile ranges). Triangles represent small groups and circles large groups. (b) Comparison of the number of accepted and evicted focal helpers between the prevented help and the helper removal treatments; dark grey bars represent accepted focal helpers and light grey bars evicted focal helpers.

**Table 1**

|  |  |  |  |
| --- | --- | --- | --- |
| **Factors** | **Estimate ± SE** | **z-value** | ***P*-value** |
| **(a) Focal defence** |
| Treatment | -0.116 ± 0.307 | -0.38 | 0.71 |
| Group size | -0.977 ± 0.611 | -1.60 | 0.11 |
| SL | 0.094 ± 0.053 | 1.76 | *0.078* |
| Treat × Group size | 1.372 ± 0.525 | 2.61 | **0.009** |
| **(b) Submissive behaviour** |
| Treatment | 3.197 ± 0.729 | 4.37 | **<0.0001** |
| Group size | 1.633 ± 0.930 | 1.76 | *0.079* |
| SL | -0.102 ± 0.068 | -1.51 | 0.13 |
| Treat x Group size | -2.731 ± 0.854 | -3.2 | **0.001** |
| **(c) Received aggression** |
| Treatment | 3.47 ± 0.521 | 6.66 | **<0.0001** |
| Group size | -1.535 ± 0.539 | -2.85 | **0.004** |
| SL | 0.066 ± 0.065 | 1.01 | 0.31 |

**Table 2**

|  |  |  |  |
| --- | --- | --- | --- |
| **Factors** | **Estimate ± SE** | **z-value** | ***P*-value** |
| **(a) Total defence** |
| Treatment | -0.621 ± 0.14 | -4.44 | **<0.0001** |
| Group size | -0.8 ± 0.234 | -3.84 | **0.0001** |
| Treat × Group size | 0.625 ± 0.238 | 2.62 | **0.009** |
| **(b) Defence of other group members** |
| Treatment | 1.18 ± 0.573 | 2.06 | **0.04** |
| Group size | 1.0 ± 0.573 | 3.49 | **0.0005** |
| Class | 3.385 ± 0.51 | 6.64 | **0.0001** |
| Treat x Group size | -1.53 ± 0.65 | -2.36 | **0.02** |
| Treat x Class | -2.166 ± 0.5 | -3.61 | **0.0003** |
| Group size x Class | -3.847 ± 0.598 | -6.44 | **<0.0001** |
| Treat x Group size x Class | 1.734 ± 0.786 | 2.21 | **0.027** |

**Table 3**

|  |  |  |  |
| --- | --- | --- | --- |
| **Factors** | **Estimate ± SE** | **z-value** | ***P*-value** |
| **Per capita received aggression** |
| Larger helper | -0.421 ± 0.906 | -0.47 | 0.64 |
| Similar helper | 1.433 ± 1.022 | 1.40 | 0.16 |
| Group size | 0.879 ± 0.957 | 0.92 | 0.36 |
| Larger helper x Group size | -2.718 ± 1.571 | -1.730 | *0.084* |
| Similar helper x Group size | -4.572 ± 1.641 | -2.79 | **0.005** |