Vigilance of mustached tamarins in single-species and mixed-species groups—the influence of group composition

Mojca Stojan-Dolar · Eckhard W. Heymann

Abstract Species that participate in mixed-species groups (MSG) may have complementary roles in antipredator strategies. We studied vigilance in mustached tamarins (Saguinus mystax), small arboreal primates that form stable mixed-species groups with saddleback tamarins (Saguinus fuscicollis), in order to examine how the direction of vigilance changes with different species group compositions and whether the division of labor between the two species can be confirmed. We did so by comparing quantitative and qualitative differences in vigilance behavior between same individuals in and out of association (case A); MSG and single-species groups of the same total group size from two different populations (case B); and MSG of the same group size but with a different ratio of conspecifics to heterospecifics (case C). We predicted that individuals would increase downward scanning when heterospecifics are absent or their percentage is low, but total vigilance would increase only in case A due to the group size effect. However, mustached tamarins increased total vigilance due to horizontal scanning in cases A and C, and the predictions were confirmed only in small-sized groups in case B. Thus, we found indications that associating tamarin species in MSG might complement each other in the direction of vigilance, but the division of labor alone does not satisfactorily explain all the findings. There appear to be other mechanisms at work that define how direction of vigilance changes with group size and species composition. Complementarity of species probably occurs due to species vertical stratification rather than differences in the direction of vigilance.

Keywords Vigilance · Mixed-species groups · Saguinus mystax · Antipredator behavior · Species complementarity · Primates

Introduction

Mixed-species groups (MSG) can be observed in a number of vertebrate taxa including fish (Ehrlich and Ehrlich 1973; Semeniuk and Dill 2006), birds (Morse 1977; Thompson and Barnard 1983; Metcalfe 1984; Munn 1986) and mammals (Struhsaker 1981; Cords 1990; Fitzgibbon 1990; Shelden et al. 1995; Herzing and Johnson 1997; Heymann and Buchanan-Smith 2000). Benefits of forming such associations are considered in terms of predator detection and avoidance, increased foraging efficiency, and reproductive advantages (reviewed in Stensland et al. 2003). However, antipredator benefits are most often considered as the ultimate cause leading to the evolution of MSG (Thompson and Barnard 1983; Munn 1986; Fitzgibbon 1990; Noë and Bshary 1997; Semeniuk and Dill 2006), and in primates strong predation pressure may even be one of the preconditions for MSG formation (Struhsaker 1981; Gautier-Hion et al. 1983; Terborgh 1990). Larger single-species groups (SSG) are supposed to be safer due to both the detection (Pulliam 1973) and dilution effects (Hamilton 1971). Individuals in MSG have the same antipredator advantages...
as animals forming SSG, while avoiding increased intraspecific competition for food and mates, which poses a limitation on how large SSG can be (Terborgh 1990; Caine 1993; Hardie and Buchanan-Smith 1997).

Besides contributing to group size, heterospecífics might also have a complementary role in antipredator behavior. Associated species may occupy different microhabitats or show physiological differences in perception and could therefore differ in the type of stimuli they are more likely to detect. In primates, species mainly residing in higher forest strata are often more likely to detect and to warn against aerial predators, while species mainly residing in lower strata are more likely to detect and to warn against terrestrial predators (Gautier-Hion et al. 1983; Peres 1993). The presence of one species can change the direction of vigilance and habitat use of another (Bshary and Noé 1997; Porter 2001; McGraw and Bshary 2002; Wolters and Zuberbühler 2003).

Mustached tamarins (Saguinus mystax) are cooperatively breeding arboreal New World primates that provide an interesting model for a study of vigilance. They have a highly cooperative social system with low levels of agonistic within-group interactions and lack an apparent dominance hierarchy (Heymann 1990b, 1996). Due to their small body size, they are a subject to strong predation pressure, mostly by large raptores but also by snakes and terrestrial predators such as ocelot and tayra (Galef et al. 1976; Terborgh 1983; Heymann 1990a; Peres 1993; Oversluijs Vasquez and Heymann 2001; Shahuano Tello et al. 2002). They form stable mixed-species groups with saddle-back tamarins (Saguinus fuscicollis), and shared vigilance is proposed to be one of the major reasons for the formation of such groups (Heymann and Buchanan-Smith 2000). Peres (1993) suggested that the two species, which show vertical stratification in the forest, might additionally enhance predator detection by functional division of labor in antipredator vigilance. He found that mustached tamarins that occupy higher levels of the forest were usually the first to emit alarm calls to aerial predators, and lower-living saddleback tamarins were more likely to warn against terrestrial predators. Similarly, in captive MSG of tamarins the higher-ranging species was quiker to discover objects at higher levels, and the lower-ranging species was quicker to discover objects on the floor (Hardie and Buchanan-Smith 2000). The aim of the present study is to examine whether this division of roles is reflected also in the direction of vigilance. The results in the literature are inconsistent: captive red-bellied tamarins (Saguinus labiatus) and wild mustached tamarins were found to spend more time looking upwards or sideways than lower-living saddleback tamarins, and saddlebacks spent more time looking downwards than the higher-living species (Peres 1993; Buchanan-Smith and Hardie 1997). While this supports the idea of the division of labor, Smith et al. (2004) and Kirchhof (2003) came to opposing results in wild mustached-saddleback MSG. When focusing on the direction of vigilance in mustached tamarins only, results from different studies are confusing already due to differences in how direction of vigilance was recorded: Peres (1993) reports more looks sideways or up than downwards, Kirchhof (2003) slightly more looks upwards than downwards, and Smith et al. (2004) more side-sweeps than look-ups. In a previous study, we already demonstrated the group size effect and the importance of proximity of conspecifics as well as heterospecifics on vigilance of mustached tamarins living in associations with saddle-back tamarins (Stojan-Dolar and Heymann 2009), which indicates that the presence of heterospecifics contributes to safety in tamarin MSG. Here, we examined how the direction of vigilance changes with different species group compositions and whether the division of labor between the two species can be confirmed. We did so by comparing quantitative and qualitative differences in vigilance behavior between MSG and SSG. In the wild, this kind of comparison is very difficult because tamarin MSG are so stable that they can rarely be observed out of association (Buchanan-Smith and Hardie 1997). We solved this problem by including an isolated population of mustached tamarins that lives on an island where no other primates are present and forms SSG of comparable size to that of MSG in the wild. The following comparisons were made (Fig. 1): MSG and SSG within the same wild population (same individuals in and out of association) (comparison A); MSG and SSG of the same total group size from two different populations (comparison B), and MSG of the same group size but with a different ratio of conspecifics to heterospecifics (comparison C). If larger groups are safer and the lower-living saddleback tamarins are more likely to detect terrestrial threats, the following predictions can be made: (1) Individuals in groups that usually associate with saddleback tamarins will increase their total vigilance when out of association and especially increase the percentage of time spent scanning downwards. (2) Individuals in SSG will show similar levels of vigilance as individuals in MSG of the same size but would spend more time scanning downwards. As for the third comparison, it is possible that a mere difference in the percentage of heterospecifics does not cause a response in individual vigilance behavior, but if it does individuals in the group with less heterospecifics would be expected to (3) spend more time looking downwards, whereas total vigilance levels in groups of the same size should not change.

Methods

Study sites and subjects

Wild mustached tamarins were observed at two study sites in the northeastern lowlands of Peruvian Amazonia. The
first site, Estación Biológica Quebrada Blanco (EBQB), is located about 70 km southeast of Iquitos (4°21′S, 73°09′W) and is covered by primary tierra firme forest (see Heymann 1995 for details). Mixed-species groups of tamarins have been continuously followed for at least 6 years before the onset of this study. Three groups of mustached tamarins were observed for 136 days between February 2006 and March 2007, covering both rainy and dry season. During the rainy season, each group was observed for 5–6 days every month, and during the dry season each group was observed for 5–6 days every second month. The mean group size (±SE) was 6.6±0.2 (range, 5–9), including 2.9±0.1 adult males (range, 2–5) and 1.6±0.1 adult females (range, 1–2). They formed stable mixed-species groups with 5.1±0.3 saddleback tamarins.

The second location, Padre Isla (PI), is an approximately 8 km² island on the Amazon River near Iquitos (3°44′S, 73°14′W) (described in Heymann 1990b). No primates originally lived on the island, but in 1977, 1978, and 1980, 87 wild-trapped S. mystax were released there by the Peruvian Primatological Project (Ique2000). The island is covered by seasonally flooded (January–May) secondary forest and fruit and yuca plantations. Observations were only possible during the dry season. There are human settlements on the island, so this population is used to human presence, and groups have occasionally been followed by researchers (e.g., Heymann 1990b; Garber et al. 1993; Garber and Pruett 1995; Tomow et al. 2006). However, in order to minimize the potential problems resulting from comparing populations with different degrees of habituation, we followed the monkeys for 2 months before starting the observations. After this additional habituation period, the monkeys did not pay more attention to observers than those at EBQB. Two groups of mustached tamarins were observed for 63 days between August 2006 and January 2007. Each group was observed for approximately 10 days every second month. The mean group size (±SE) was 10.75±0.68 (range, 8–14), including 2.42±0.47 adult males (range, 1–5) and 2.75±0.13 adult females (range, 2–3). Since no other primates are present, mustached tamarins on PI live in single-species groups (SSG). Animals at both sites were individually identified by differences in pelage patterns, body size, and shape and pigmentation of reproductive organs (Löttker et al. 2004). None of the observed females was in estrus.

Observation methods

Vigilance was defined as observable head movements in stationary animals where scanning is directed beyond arm’s reach (Treves 1998). Looking at their own body, body of a grooming partner, objects close to their own body, or objects held in hands was not considered as vigilance. Looking beyond arm’s reach was excluded in two cases, since it is reasonable to assume that in these situations animals cannot effectively perceive the environment: (a) staring without head movements in an unusual head position (when head is turned upside-down and eyes are lower than the mouth) while being groomed, and (b) when an animal is scratching its head in such a way that the head shakes.

Continuous focal sampling (Martin and Bateson 1990) was used to record vigilance status (vigilant, not vigilant), the direction of scanning (up, down, horizontal), and individual activity. Scanning upwards was defined as looking towards the sky at an angel of at least 45°, scanning downwards was defined as looking towards the ground at an angle of at least 45°, and scanning in between was considered as horizontal. With every observation, data on the height of the focal individual was also collected, and a height class was ascribed to every observation (0, ground;
1, 0–3 m; 2, 3–6 m; 3, 6–9 m; 4, 9–12 m; 5, 12+ m). We chose the focal animal based on which individuals were visible in order to maximize the amount of data collected. When a choice was possible, we chose the individual with least observations, unless it had been the focal animal during the previous 30 min. The activities were defined as follows: resting—sitting or laying without being engaged in any activity that requires visual attention; feeding—ingesting food or searching for prey using one or both hands to investigate leaves, bark, holes etc.; allogrooming—going through the fur of another individual with the hands; autogrooming—going through own fur with the hands. In order to evaluate the suitability and reliability of the definition of vigilance, a reliability test was performed prior to data collection.

Observation conditions allowed for 30–120 s focal observations. Data were entered directly into a hand-held computer using The Observer Mobile® software. The two species were considered as a MSG when at least one individual from one species was less than 20 m apart from one individual from the other species (Struhsaker 1981). In order to ensure independence of observations, the same individual was never resampled within a 30-min time period.

Data analysis

For every individual, the proportion of time spent vigilant was taken as a measure of vigilance. Only individuals that had at least ten observations under specified conditions were considered in the analyses. Percentages of time spent vigilant were averaged for each individual, using only observations from groups of the specified composition. These individual mean values were used as data points in further analyses. In all cases, we used nonparametric statistical tests. The following comparisons were made:

A. Comparison of vigilance behavior of individuals at EBQB in and out of association. Since tamarins in the wild are very rarely found in SSG, we only had enough data to compare vigilance in nine individuals from different groups.

B. Comparison between MSG at EBQB and SSG at PI of the same total group size. The size of all the study groups was varying in time. There was a sufficient amount of data to include two pairs of MSG and SSG. The pair of smaller groups had a total group size of eight, and the pair of larger groups had a total group size of 13. For this analysis, it was possible to divide comparisons by activity: vigilance in larger groups was compared during resting and during feeding, whereas for smaller groups the data only allowed a comparison during resting.

C. Comparison between two MSG of the same size (14 individuals) but different species composition (43% and 64% of mustached tamarins). The amount of data under these specific conditions did not allow for splitting by activity.

To compare heights of different groups, mean of the height class values was calculated for every individual under the specified conditions.

Results

At EBQB (comparison A), individuals increased their overall vigilance in the absence of heterospecifics (Wilcoxon exact: \( T=1, N=9, p=0.020 \); Fig. 2) due to an increase in horizontal scanning (\( T=1, N=9, p=0.008 \)). There was no difference in looking upwards (\( T=4, N=9, p=1.000 \)) or looking downwards (\( T=4, N=9, p=0.652 \)).

When comparing groups of the same size at different sites (comparison B), we first compared the rate of alarm calls, which did not differ between the two sites (0.33±0.23 per 10 h of observation per individual for EBQB (mean ± SD)
and 0.33±0.11 for PI; Mann–Whitney U: n1=27; n2=6; Z=−0.420; p=0.674). The group compositions of our study groups enabled us to compare two pairs of groups: smaller groups with a total of eight individuals and larger groups with a total of 13 individuals. However, in both cases, data at EBQB were collected during the rainy season, while observations on PI were only possible during the dry season. Therefore, we first checked whether season affected vigilance at EBQB to make sure that observations collected at different times of the year are comparable. We found that vigilance during feeding was higher in the rainy season (total vigilance: T=2, N=14, p=0.019; up: T=5, N=14, p=0.140; horizontal: T=4, N=14, p=0.084; down: T=6, N=14, p=0.331; Wilcoxon exact), but vigilance during resting was not affected by season (total vigilance: T=7, N=17, p=0.906; up: T=8, N=17, p=0.795; horizontal: T=7, N=17, p=0.356; down: T=6, N=17, p=0.309; Wilcoxon exact). Possible reasons for this difference are beyond the scope of this study, but in order to avoid confounding effects we conducted comparison B only with data on vigilance during resting. The only significant difference was found in smaller groups (eight individuals) where individuals in SSG spent more time looking downwards than in MSG (Fig. 3, Table 1). Larger SSG at PI and MSG at EBQB (13

Fig. 3 Differences in the proportion of a time spent looking upwards; b time spent looking downwards; c time spent looking horizontally, and d total time spent vigilant between MSG at EBQB and SSG at PI controlled for the effect of the group size. Data points represent individual mean values. Open circles individuals in groups of total size 8 during resting; black circles individuals in groups of total size 13 during resting. ** p=0.01 level of significance
individuals) did not differ in any component of their vigilance behavior.

In all cases, the monkeys spent two to five times more time looking horizontally than upwards or downwards (Table 2). They also tended to look more downwards than upwards during resting; this difference was significant for the larger MSG group at EBQB and for both groups at PI (Table 2).

In MSG of the same size but different species composition (comparison C), individuals in the group with a higher percentage of conspecifics were more vigilant (Mann–Whitney U: n1=5; n2=6; Z=−2.373; p=0.046) due to differences in horizontal scanning (Z=−2.556; p=0.032, Fig. 4). There were no differences in looking upwards (Z=−0.365; p=1.000) or downwards (Z=−0.183; p=0.886).

Individuals in SSG that were temporarily out of association (comparison A) were found higher in the forest than when they were in MSG (Wilcoxon: T=0, N=5, p=0.008). The same was true for individuals in SSG at PI compared to MSG at EBQB (comparison B) for both larger (Mann–Whitney U: n1=5; n2=8; Z=−2.635; p=0.008) and smaller (n1=5; n2=8; Z=−2.781; p=0.005) pair of groups. Individuals in MSG with different percentages of conspecifics (comparison C) did not differ in heights (Mann–Whitney U: n1=5; n2=6; Z=−1.551; p=0.121).

### Discussion

We found weak support for complementarity of the direction of vigilance between the two associating tamarin species. Only the comparison between SSG and MSG of the same size supported our prediction (comparison B, prediction 2), and that was true only for individuals in smaller groups during resting where individuals in the population that never associated with other species looked downwards more than individuals in MSG (Table 2, Fig. 3). In larger groups, there were no differences between MSG and SSG of the same size, which could imply that heterospecifics play a more important role in smaller groups, but data on more groups would be needed for firmer conclusions.

The idea of the division of labor was further supported by the analysis of directions of vigilance within groups (Table 2). Mustached tamarins in all groups tended to look downwards more than upwards, but while the difference at EBQB was of only a few percent and

### Table 1

Results of Mann–Whitney U exact test for differences between MSG at EBQB and SSG at PI

| Direction of vigilance | Group size | N\(\text{(EBQB)}\) | N\(\text{(PI)}\) | U   | Z     | p    |
|------------------------|------------|------------------|----------------|-----|-------|------|
| Up                     | 8          | 5                | 8              | 14.0| −0.878| 0.435|
|                        | 13         | 5                | 8              | 12.0| −1.171| 0.284|
| Down                   | 8          | 5                | 8              | 0.0 | −2.928| 0.002***|
|                        | 13         | 5                | 8              | 11.0| −1.317| 0.222|
| Horizontal             | 8          | 5                | 8              | 8.0 | −1.757| 0.093|
|                        | 13         | 5                | 8              | 18.0| −0.293| 0.833|
| Total                  | 8          | 5                | 8              | 20.0| 0.000 | 0.999|
|                        | 13         | 5                | 8              | 9.0 | −1.610| 0.127|

Differences between \(N\text{(PI)}\) and total group size emerge because not all the individuals fulfilled the conditions to be included in the analysis (see “Methods”)

**\(p=0.01\) level of significance

***\(p=0.001\) level of significance

### Table 2

Mean, median, minimum, and maximum percentage of time individuals in groups of different group composition spent looking in the specified direction

| Group       | Horizontal | Up     | Down   | Wilcoxon |
|-------------|------------|--------|--------|----------|
|             | Mean | Median | Min | Max | Mean | Median | Min | Max | Mean | Median | Min | Max | N | T | p   |
| SSG EBQB    | 51.9 | 53.1   | 38.2 | 64.4 | 11.6 | 11.4   | 7.3  | 18.5 | 13.7 | 11.3   | 6.0  | 22.3 | 9 | 15 | 0.374 |
| MSG EBQB    | 43.9 | 41.9   | 34.0 | 52.5 | 11.2 | 11.6   | 9.6  | 12.5 | 13.4 | 13.5   | 10.4 | 17.6 | 9 | 1  | 0.011* |
| 8a rest EBQB| 56.2 | 59.0   | 46.3 | 62.3 | 15.3 | 13.8   | 13.1 | 18.7 | 18.5 | 19.5   | 15.4 | 20.3 | 5 | 2  | 0.138 |
| 13 rest EBQB| 55.8 | 57.1   | 50.8 | 57.7 | 14.2 | 13.8   | 11.6 | 17.6 | 16.6 | 16.4   | 14.6 | 18.7 | 5 | 0  | 0.043* |
| 8 rest PI   | 50.3 | 49.7   | 44.4 | 55.3 | 13.6 | 13.4   | 9.6  | 18.2 | 26.6 | 25.7   | 21.0 | 32.0 | 8 | 0  | 0.012* |
| 13 rest PI  | 56.5 | 57.5   | 47.0 | 62.7 | 12.2 | 12.5   | 8.3  | 16.0 | 20.7 | 20.9   | 14.3 | 26.7 | 8 | 1  | 0.017* |

Wilcoxon exact test was used to test for the difference between upward and downward scanning

*The numbers in this column indicate the group size

*p=0.05 level of significance
could hardly have an ecological significance, it was about
twofold at PI. It seems that tamarins in the population that
has no contact with other primate species compensate for
the absence of heterospecifics by increased scanning
downwards.

However, comparisons of vigilance behavior in cases A
and C gave results that were less straightforward. Individu-
als that were temporarily out of association (case A)
increased their total vigilance as predicted by the group size
effect, but that was due to horizontal, not downward,
scanning. A similar behavioral response was observed
when comparing individuals in the group with a higher
percentage of conspecifics with those in the group with a
smaller percentage of conspecifics (case C), although we
predicted no difference in total vigilance levels in groups of
the same size. We offer three explanations for why
horizontal scanning increases in these two cases. First,
non-antipredator functions such as food searching and
especially social vigilance are probably much more included
in horizontal than in upward or downward scanning
because conspecifics are usually found at about the same
height as the focal individual. Having other functions in
addition to antipredator detection is probably also the
reason why horizontal scanning takes up much more of
animals’ time than looking upwards or downwards (Table 2).
In a group with more conspecifics (comparison C), there is
probably an increased need for social vigilance. Our
previous study suggests that, while the major function of
vigilance is predator-related, the most probable reason for
monitoring conspecific group members in mustached
tamarins is maintaining group cohesion (Stojan-Dolar and
Heymann 2009). This certainly becomes more challenging
with an increased number of conspecifics in the group. Yet,
this interpretation does not explain why horizontal scanning
increases when mustached tamarins are temporarily found
out of association (comparison A). Other functions of social
vigilance such as avoiding food stealing or aggression that
are usually mentioned in studies on other species (e.g.,
Jones 1998; Steenbeek et al. 1999; Hirsch 2002) have not
been found in the relatively non-aggressive society of these
cooperatively breeding primates.

Second, due to the arboreal nature of neotropical
felines callitrichids might increase horizontal or even
upward scanning as a response to terrestrial predators
(Barros et al. 2008). Neotropical felines and tayras can
climb trees occasionally (Emmons 1987; Wang 2002), and
the absence of the lower-living species certainly increases
the probability that a predator climbing up a tree will
remain unnoticed. This may be why mustached tamarins
in both temporary and permanent SSG were found at
greater heights than those in MSG (even though the forest
at PI was actually lower than at EBQB), which gives an
impression that this is a general reaction to the absence
of heterospecifics that is not conditioned with the type of
predators. The height of saddleback tamarins, on the other
hand, is not affected by the presence or absence of
congeners (Buchanan-Smith 1999), which may mean that
they are more specialized to their specific niche and/or
that there are antipredator benefits of living in lower forest
strata.

And third, since mustached tamarins are found at
higher levels in the forest and raptors are ambush
predators that often perch within canopies (Rettig 1978;
Robinson 1994), it might not be necessary to elevate the
gaze for more than 45° in order to detect them—especially
since monkeys even further increase their height when in
SSG. Horizontal scanning might therefore also have an
important role in detection of aerial predators, and it has
been understood as such also in some other vigilance
studies (Peres 1993; Bshary and Noë 1997). Increased
vigilance against raptors in a group of the same total size
but with fewer heterospecifics (case C) or in the absence
of heterospecifics (case A) can imply that lower-ranging
heterospecifics have a complementary role that cannot be
compensated by adding additional conspecifics, not only
in the detection of terrestrial but also aerial predators. This
might occur because the lower-ranging saddlebacks see
the upper levels of the forest and the sky from a different
angle—a fact that could be important in a habitat with
dense vegetation. But it can also emerge simply because
the benefits of contributing vigilance in such a group are
greater than in a group with more heterospecifics. In this
case, contributing vigilance can be seen as a case of a
component tragedy of the commons, where vigilance is a

![Fig. 4 Proportion of time spent vigilant for individuals from two MSG of the same total size but with different percentage of conspecifics. Data points represent individual mean values. **p=0.05 level of significance.](image-url)
social good (Rankin et al. 2007). When the percentage of conspecifics is high, it is more likely that a predator would take a conspecific than when the percentage of conspecifics is low. Here, it is important to note that tamarins live in a highly cooperative society where reproductive success depends heavily on help from conspecifics (Caine 1993; Garber 1997) and where the degree of relatedness within the group is relatively high (Huck et al. 2005). Hence, if a predator kills a conspecific, this can have direct consequences for the fitness of other individuals—they lose a helper who is also quite likely to be a close relative. The associating species therefore both contribute vigilance to improve group safety, but the ratio between costs and benefits of contributing depends on the percentage of conspecifics in MSG. When the benefits/percentage of conspecifics are high, they tend to contribute more; when the benefits/percentage of conspecifics are low, they tend to free-ride more. However, vigilance can never drop under a certain level because individuals also increase their personal safety by being watchful.

In addition to that, individuals that are temporarily out of association (case A) are probably more vigilant already due to the group size effect. However, it remains unusual that upward scanning that certainly also serves detecting birds of prey did not increase.

A general observation based on our results is that the two populations did not respond in the same way to changes in group size and species composition. Tamarins from EBQB reacted by increasing total vigilance due to increased horizontal scanning, whereas the comparison with same-sized groups from PI revealed either no differences in vigilance or the expected changes in downward scanning (Fig. 1). Summarizing the above-described mechanisms, we may explain the observed results as follows: In case A, individuals that are temporarily out of association increased their vigilance due to the group size effect, the absence of the low-ranging species, and/or increased benefits of contributing vigilance. In case B, individuals in the smaller groups from PI increased their vigilance against terrestrial predators due to the permanent absence of the low-ranging species. And in case C, individuals in the group with a higher percentage of conspecifics increased their horizontal scanning due to an increased need to monitor conspecifics in order to maintain group cohesion and/or increased benefits of contributing vigilance.

It is difficult to speculate about the reasons for these mechanisms to act only under certain circumstances. However, there are two major differences between the populations that could give some clue as to why this occurs. First, the forest on PI is less dense and lower than at EBQB, and as a consequence it is possible that there are different species of birds of prey present that might use different attack tactics. By recording alarm calls, we measured perceived predation risk, but we were not able to identify species that actually prey upon tamarins. Birds or conspecifics within the canopy are easier to detect in conditions of lower vegetation density, and therefore, benefits of increasing horizontal vigilance in order to monitor conspecifics or to detect aerial predators are not the same as at EBQB. Furthermore, there are no felines or tyras on the island, which means that horizontal scanning is unlikely to be intended to detect terrestrial predators. However, terrestrial threat is not absent as monkeys perceive dogs as potential predators. And second, tamarins at PI do not have any experience with other primate species, while animals at EBQB have had an opportunity to learn to adjust their vigilance according to the group species composition. Being only temporarily out of association or having a lower percentage of heterospecifics in MSG might elicit different reactions than living permanently in SSG and not having any experience with other primates.

Comparison with results of other studies on primate MSG that are analogous to our comparison A also implies that there is no uniform explanation for changes of vigilance patterns under different MSG/SSG conditions (Table 3). Individuals in different combinations of African cercopithecids have been observed to increase, decrease, or not to change their total vigilance at all when they are temporarily found in SSG, which appears to depend on which species they associate with (Cords 1990; Chapman and Chapman 1996; Bshary and Noë 1997; Treves 1999). With regard to callitrichids, a study on captive red-bellied (S. labiatus) and saddleback tamarins revealed that higherranging red-bellied tamarins increased “scanning” when they were housed without saddlebacks (Buchanan-Smith and Hardie 1997; Hardie and Buchanan-Smith 1997). Since the definition of “scanning” in that study included also looking sideways, these results might be consistent with our findings.

Mixed results are not uncommon in vigilance literature. The effects of sex, age, distance to cover, and also group size on vigilance vary greatly across different species and studies (reviewed in Caro 2005; Beauchamp 2008). Inconsistent results from studies on different species could occur due to differences in predation pressure between sites and species, differences in group size of different species, but also differences in methodology employed by different researchers. In order to facilitate comparison between species and populations, future studies should differentiate between all three possible directions of vigilance and take into account that vigilance can have a social as well as an antipredator component that might show different patterns (Klose et al. 2009). Different studies...
also operate with different measures of vigilance (percentage of time vigilant, frequency/duration of vigilance bouts) that appear to give different results (e.g., Trouilloud et al. 2004; Beauchamp 2008).

To sum up, we found indications that associating tamarin species in MSG might complement each other in the direction of vigilance, but the division of labor alone does not satisfactorily explain all the findings. There appear to be other mechanisms at work that define how direction of vigilance changes with group size and species composition. However, this does not directly contradict the notion that heterospecifics have a complementary role in antipredator strategies of MSG. The two species notice different predators already because they live in different forest strata (Gautier-Hion et al. 1983; Peres 1993) and might additionally complement each other by seeing the same predators from different angles; therefore, additional adjustments in the direction of vigilance may not always be necessary. In this case, division of labor between species is still very important, it just does not occur due to differences in the direction of vigilance but rather due to vertical stratification per se.

### Table 3
An overview of primate studies comparing vigilance behavior in and out of association (analog to comparison A in this study)

| Study                        | Species                        | Forest stratum | Changes of vigilance patterns when temporarily in SSG |
|------------------------------|--------------------------------|----------------|------------------------------------------------------|
| Cords 1990[^a]               | Blue monkeys (Cercopithecus mitis) | Higer          | When feeding on plants, frequency of look-ups increases |
|                              | Redtail monkeys (Cercopithecus ascanius) | Lower          | When feeding on plants, frequency of look-ups increases |
| Bshary and Noë 1997          | Diana monkeys (Cercopithecus Diana) | Higher         | No change in vigilance                               |
|                              | Red colobus (Colobus badius)     | Lower          | When foraging, frequency of looking down and looking sideways increases |
| Treves 1999[^a]              | Red colobus (Colobus badius)     | Higher         | No change in vigilance                               |
|                              | Redtail monkeys (Cercopithecus ascanius) | Lower          | No change in vigilance                               |
| Chapman and Chapman 1996[^a] | Black-and white colobus (Colobus guereza) | Highest       | No change in vigilance                               |
|                              | Red colobus (Colobus badius)     | In between     | Depending on which species they associate with, frequency of look-ups increases, decreases, or does not change |
|                              | Mangabeys (Cerocebus albigena)    | In between     | No change in vigilance                               |
|                              | Blue monkeys (Cercopithecus mitis) | In between     | Depending on which species they associate with, frequency of look-ups increases, decreases, or does not change |
|                              | Redtail monkeys (Cercopithecus ascanius) | Lowest         | Depending on which species they associate with, frequency of look-ups increases, decreases, or does not change |
| Stanford 1998[^a]            | Red colobus (Colobus badius)     | Highest        | Frequency of look-ups increases in both associations |
|                              | Blue monkeys (Cercopithecus mitis) | In between     | Not measured                                         |
|                              | Redtail monkeys (Cercopithecus ascanius) | Lowest         | Not measured                                         |
| Wolters and Zuberbühler 2003 | Diana monkeys (Cercopithecus Diana) | Higher         | Percentage of time spent vigilant increases           |
|                              | Campbell’s monkeys (Cercopithecus campbellii) | Lower         | Percentage of time spent vigilant increases           |
| Buchanan-Smith and Hardie 1997| Captive red-bellied tamarins (Saguinus labiatus) | Higher         | Percentage of time spent scanning[^b] increases |
|                              | Captive saddleback tamarins (Saguinus fuscicolli) | Lower         | Percentage of time spent looking up and scanning[^b] increases |
| This study                   | Mustached tamarins (Saguinus mystax) | Higher         | Percentage of time spent looking sideways increases |
|                              | Saddleback tamarins (Saguinus fuscicolli) | Lower          | Not measured                                         |

[^a]: Approximate forest strata based on Gebo and Chapman (1995).

[^b]: Scanning in this study is defined as “moving the head from side to side and/or up and down in a continuous flow of movement, excluding fixated stares or scanning the floor area.” Look-ups are defined as “deliberate large single head movement upwards or a stare in an upward direction where the head is usually angled 45° or more and not moving side to side”
Interestingly, increasing the height in the forest appears to be a more uniform reaction to the absence of heterospecifs than changes in vigilance patterns. Data on more groups and populations and on responses of lower-ranging saddleback tamarins would be needed to gain clearer insight into mechanisms and ecological relevance of complementarity of antipredator behavior in species participating in MSG.

Acknowledgments We would like to thank the following people and organizations: Instituto Veterinario de Investigaciones Tropicales y de Altur (IVITA) for the permission to work on Padre Isla; our field assistants Camilo Flores Amsafu, Ney Shahuano Tello, and Manuel Shahuano Tello for their invaluable help in the forest; Carlos Ique for helpful practical advice; and Markus Port and other members of the Department for Behavioural Ecology and Sociobiology of the German Primate Centre for many fruitful discussions. The manuscript benefited greatly from helpful advice of two anonymous reviewers. Financial support was partially provided by Deutscher Akademischer Austausch Dienst (DAAD). Field work was authorized by the Instituto Nacional de Recursos Naturales (authorization no. 018-2006-INRENA-JFF-DCB) and complied with Peruvian laws.

Open Access This article is distributed under the terms of the Creative Commons Attribution Noncommercial License which permits any noncommercial use, distribution, and reproduction in any medium, provided the original author(s) and source are credited.

References

Barros M, Alencar C, Silva MAD, Tomaz C (2008) Changes in experimental conditions alter anti-predator vigilance and sequence predictability in captive marmosets. Behav Process 77:351–356
Beauchamp G (2008) What is the magnitude of the group-size effect on vigilance? Behav Ecol 19:1361–1368
Bshary R, Noé R (1997) Red colobus and Diana monkeys provide mutual protection against predators. Anim Behav 54:1461–1474
Buchanan-Smith HM (1999) Tamarin polyspecific associations: forest utilization and stability of mixed-species groups. Primates 40:233–247
Buchanan-Smith HM, Hardie SM (1997) Tamarin mixed-species groups: an evaluation of a combined captive and field approach. Folia Primatol 68:272–286
Caine NG (1993) Flexibility and co-operation as unifying themes in Saguinus social organization and behaviour: the role of predation pressures. In: Rylands AB (ed) Marmosets and tamarins: systematics, behaviour, and ecology. Oxford University Press, Oxford, pp 200–219
Caro T (2005) Antipredator defenses in birds and mammals. University of Chicago Press, Chicago
Chapman CA, Chapman LJ (1996) Mixed-species primate groups in the Kibale forest: ecological constraints on association. Int J Primatol 17:31–50
Cords M (1990) Vigilance and mixed-species association of some East African forest monkeys. Behav Ecol Sociobiol 26:297–300
Ehrlich PR, Ehrlich AH (1973) Coevolution—heterotypic schooling in Caribbean reef fishes. Am Nat 107:157–160
Emmons LH (1987) Comparative feeding ecology of felids in a neotropical rainforest. Behav Ecol Sociobiol 20:271–283
Fitzgibbon CD (1990) Mixed-species grouping in Thomson and Grant gazelles—the antipredator benefits. Anim Behav 39:1116–1126
Galef BG, Mittermeier RA, Bailey RC (1976) Predation by tayra (Eira barbara). J Mammal 57:760–761
Garber PA (1997) One for all and breeding for one: cooperation and competition as a tamarin reproductive strategy. Evol Anthropol 5 (6):187–199
Garber PA, Pruett JD (1995) Positional behavior in moustached tamarin monkeys—effects of habitat on locomotor variability and locomotor stability. J Hum Evol 28:411–426
Garber PA, Pruett JD, Isaacson J (1993) Patterns of range use, range defense, and intergroup spacing in moustached tamarin monkeys (Saguinus mystax). Primates 34:11–25
Gautier-Hion A, Quiris R, Gautier JP (1983) Monospecific vs. polyspecific life—a comparative study of foraging and antipredatory tactics in a community of Cercopithecus monkeys. Behav Ecol Sociobiol 12:325–335
Gebo DL, Chapman CA (1995) Positional behavior in 5 sympatric Old-World monkeys. Am J Phys Anthropol 97:49–76
Hamilton WD (1971) Geometry for the selfish herd. J Theor Biol 31:295–311
Hardie SM, Buchanan-Smith HM (1997) Vigilance in single- and mixed-species groups of tamarins (Saguinus labiatus and Saguinus fuscicollis). Int J Primatol 18:217–234
Hardie SM, Buchanan-Smith HM (2000) Responses of captive single- and mixed-species groups of Saguinus to novel nonthreatening objects. Int J Primatol 21:629–648
Herzing D, Johnson C (1997) Interspecific interactions between Atlantic spotted dolphins (Stenella frontalis) and bottlenose dolphins (Tursiops truncatus) Bahamas, 1985–1995. Aqua Mamm 23(2):85–99
Heymann EW (1990a) Reactions of wild tamarins, Saguinus mystax and Saguinus fuscicollis to avian predators. Int J Primatol 11:327–337
Heymann EW (1990b) Social behaviour and infant carrying in a group of moustached tamarins, Saguinus mystax (Primates: Platyrhini: Callitrichidae), on Padre Isla, Peruvian Amazonia. Primates 31:183–196
Heymann EW (1995) Sleeping habits of tamarins, Saguinus mystax and Saguinus fuscicollis (Mammalia; Primates; Callitrichidae), in north-eastern Peru. J Zool 237:211–226
Heymann EW (1996) Social behavior of wild moustached tamarins, Saguinus mystax, at the Estacion Biologica Quebrada Blanco, Peruvian Amazonia. Am J Primatol 38:101–113
Heymann EW, Buchanan-Smith HM (2000) The behavioural ecology of mixed-species troops of callitrichine primates. Biol Rev 75:169–190
Hill RA, Dunbar RIM (1998) An evaluation of the roles of predation rate and predation risk as selective pressures on primate grouping behaviour. Behaviour 135:411–430
Hirsch BT (2002) Social monitoring and vigilance behaviour in brown capuchin monkeys (Cebus apella). Behav Ecol Sociobiol 52:458–464
Huck M, Löttker P, Bohle UR, Heymann EW (2005) Paternity and kinship patterns in polyandrous moustached tamarins (Saguinus mystax). Am J Phys Anthropol 127:449–464
Ique C (2000) Manejo semi-extensivo de Saguinus mystax en Padre Isla, río Amazonas. In: Howard F, García Podestá M (eds) La Primatologia en el Perú 2 Master Graf Editores, Lima, pp 131–143
Jones ME (1998) The function of vigilance in sympatric marsupial carnivores: the eastern quoll and the Tasmanian devil. Anim Behav 56:1279–1284
Kirchhof J (2003) Raubfeindvermeidung bei Tamarinen im Amazonas-Regenwald von Peru: Alarmrufsysteme und Vigilanz in polyspezifischen Assoziationen. PhD thesis. Freie Universität Berlin, Berlin
Klose S, Welbergen J, Goldizen A, Kalko E (2009) Spatio-temporal vigilance architecture of an Australian flying-fox colony. Behav Ecol Sociobiol 63:371–380
Löttker P, Huck M, Heymann EW (2004) Demographic parameters and events in wild moustached tamarins (Saguinus mystax). Am J Primatol 64:425–449
Martin P, Bateson P (1990) Measuring behaviour—an introductory guide. Cambridge University Press, Cambridge
McGraw WS, Bshary R (2002) Association of terrestrial mangabeys (Cercocebus atys) with arboreal monkeys: experimental evidence for the effects of reduced ground predator pressure on habitat use. Int J Primatol 23:311–325
Metcalfe NB (1984) The effects of mixed-species flocking on the vigilance of shorebirds: who do they trust? Anim Behav 32:986–993
Morse DH (1977) Feeding behavior and predator avoidance in heterospecific groups. Bioscience 27:332–339
Munn CA (1986) Birds that cry wolf. Nature 319:143
Noë R, Bshary R (1997) The formation of red colobus-diana monkey associations under predation pressure from chimpanzees. Proc R Soc B-Biol Sci 264:253–259
Oversluijs Vasquez MR, Heymann EW (2001) Crested eagle (Morphnus guianensis) predation on infant tamarins (Saguinus mystax and Saguinus fuscicollis, Callitrichinae). Folia Primatol 72:301–303
Peres CA (1993) Anti-predation benefits in a mixed-species group of Amazonian tamarins. Folia Primatol 61:61–76
Porter LM (2001) Benefits of polyspecific associations for the goeldi's monkey (Callimico goeldii). Am J Primatol 54:143–158
Pulliam HR (1973) On advantages of flocking. J Theor Biol 38:419–422
Rankin DJ, Bargum K, Kokko H (2007) The tragedy of the commons in evolutionary biology. Trends Ecol Evol 22:643–651
Rettig NL. (1978) Breeding behavior of the harpy eagle (Harpia harpyja). Auk 95:629–643
Robinson SK (1994) Habitat selection and foraging ecology of raptors in Amazonian Peru. Biotropica 26:443–458
Semeniuk CAD, Dill LM (2006) Anti-predator benefits of mixed-species groups of cowtail stingrays (Pastinachus sephen) and whiprays (Himantura uarnak) at rest. Ethology 112:33–43
Shahuano Tello N, Huck M, Heymann EW (2002) Boa constrictor attack and successful group defence in moustached tamarins, Saguinus mystax. Folia Primatol 73:146–148
Shelden KEW, Baldridge A, Withrow DE (1995) Observations of rissos dolphins (Grampus griseus) with gray whales (Eschrichtius robustus). Mar Mamm Sci 11:231–240
Smith AC, Kelez S, Buchanan-Smith HM (2004) Factors affecting vigilance within wild-mixed-species troops of saddleback (Saguinus fuscicollis) and moustached tamarins (S. mystax). Behav Ecol Sociobiol 56:18–25
Stanford C (1998) Chimpanzee and red colobus: the ecology of predator and prey. Harvard University Press, Cambridge
Steenbeek R, Piek R, van Buul M, van Hooff JARAM (1999) Vigilance in wild Thomas's langurs (Presbytis thomasi): the importance of infanticide risk. Behav Ecol Sociobiol 45:137–150
Stensland E, Angerbjorn A, Berggren P (2003) Mixed species groups in mammals. Mamm Rev 33:205–223
Stoian-Dolar M, Heymann EW (2009) Vigilance in a cooperatively breeding primate. Int J Primatol (in press)
Struhsaker TT (1981) Polyspecific associations among tropical rainforest primates. Zeitschrift Für Tierpsychologie 57:268–304
Terborgh J (1983) Five new world primates: a study in comparative ecology. Princeton University Press, Princeton
Terborgh J (1990) Mixed flocks and polyspecific associations: costs and benefits of mixed groups to birds and monkeys. Am J Primatol 21:87–100
Thompson DBA, Barnard CJ (1983) Anti-predator responses in mixed-species associations of lapwings, golden plovers and black-headed gulls. Anim Behav 31:585–593
Tornow MA, Ford SM, Garber PA, Sauerbrunn ED (2006) Dentition and benefits of mixed groups to birds and monkeys. Am J Primatol 21:87–100
Thompson DBA, Barnard CJ (1983) Anti-predator responses in mixed-species associations of lapwings, golden plovers and black-headed gulls. Anim Behav 31:585–593
Tornow MA, Ford SM, Garber PA, Sauerbrunn ED (2006) Dentition and benefits of mixed groups to birds and monkeys. Am J Primatol 21:87–100
Trevor A. (1998) The influence of group size and neighbors on vigilance in two species of arboreal monkeys. Behaviour 135:453–481
Trevor A. (1999) Has predation shaped the social systems of arboreal primates? Int J Primatol 20:35–67
Trouwillo W, Delisle A, Kramer DL (2004) Head raising during foraging and pausing during intermittent locomotion as components of antipredator vigilance in chipmunks. Anim Behav 67:789–797
Wang E (2002) Diets of ocelots (Leopardus pardalis), margays (L. wiedii), and oncillas (L. tigrinus) in the Atlantic rainforest in southeast Brazil. Stud Neotrop Fauna E 37:207–212
Wolters S, Zuberbühler K (2003) Mixed-species associations of Dane and Campbell's monkeys: the costs and benefits of a forest phenomenon. Behaviour 140:371–385