



HAL
open science

Variation in clutch size in relation to nest size in birds.

Anders P. Møller, Frank Adriaensen, Alexandr Artemyev, Jerzy Bańbura, Emilio Barba, Clotilde Biard, Jacques Blondel, Zihad Bouslama, Jean-Charles Bouvier, Jordi Camprodon, et al.

► **To cite this version:**

Anders P. Møller, Frank Adriaensen, Alexandr Artemyev, Jerzy Bańbura, Emilio Barba, et al.. Variation in clutch size in relation to nest size in birds.. *Ecology and Evolution*, 2014, 4 (18), pp.3583-3595. 10.1002/ece3.1189 . hal-01092055

HAL Id: hal-01092055

<https://hal.science/hal-01092055v1>

Submitted on 27 May 2020

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.



Distributed under a Creative Commons Attribution 4.0 International License

Variation in clutch size in relation to nest size in birds

Anders P. Møller¹, Frank Adriaensen², Alexandr Artemyev³, Jerzy Bańbura⁴, Emilio Barba⁵, Clotilde Biard⁶, Jacques Blondel⁷, Zihad Bouslama⁸, Jean-Charles Bouvier⁹, Jordi Camprodon¹⁰, Francesco Cecere¹¹, Anne Charmantier¹², Motti Charter^{13,14}, Mariusz Cichoń¹⁵, Camillo Cusimano¹⁶, Dorota Czeszczewik¹⁷, Virginie Demeyrier¹⁸, Blandine Doligez¹⁹, Claire Doutrelant²⁰, Anna Dubiec²¹, Marcel Eens²², Tapio Eeva²³, Bruno Faivre²⁴, Peter N. Ferns²⁵, Jukka T. Forsman²⁶, Eduardo García-Del-Rey²⁷, Aya Goldshtein¹³, Anne E. Goodenough²⁸, Andrew G. Gosler²⁹, Iga Gózdź³⁰, Arnaud Grégoire³¹, Lars Gustafsson³², Ian R. Hartley³³, Philipp Heeb³⁴, Shelley A. Hinsley³⁵, Paul Isenmann³⁶, Staffan Jacob³⁷, Antero Järvinen³⁸, Rimvydas Juškaitis³⁹, Erkki Korpimäki⁴⁰, Indrikis Krams⁴¹, Toni Laaksonen⁴², Bernard Leclercq⁴³, Esa Lehtikoinen⁴⁴, Olli Loukola⁴⁵, Arne Lundberg⁴⁶, Mark C. Mainwaring⁴⁷, Raivo Mänd⁴⁸, Bruno Massa⁴⁹, Tomasz D. Mazgajski⁵⁰, Santiago Merino⁵¹, Cezary Mitrus⁵², Mikko Mönkkönen^{53,54}, Judith Morales-Fernandez⁵⁵, Xavier Morin⁵⁶, Ruedi G. Nager⁵⁷, Jan-Åke Nilsson⁵⁸, Sven G. Nilsson⁵⁹, Ana C. Norte⁶⁰, Markku Orell⁶¹, Philippe Perret⁶², Carla S. Pimentel⁶³, Rianne Pinxten⁶⁴, Ilze Priedniece⁶⁵, Marie-Claude Quidoz⁶⁶, Vladimir Remes⁶⁷, Heinz Richner⁶⁸, Hugo Robles^{69,70}, Seppo Rytkönen⁷¹, Juan Carlos Senar⁷², Janne T. Seppänen⁷³, Luís P. da Silva⁷⁴, Tore Slagsvold⁷⁵, Tapio Solonen⁷⁶, Alberto Sorace⁷⁷, Martyn J. Stenning⁷⁸, János Török⁷⁹, Piotr Tryjanowski⁸⁰, Arie J. van Noordwijk⁸¹, Mikael von Numers⁸², Wiesław Walankiewicz⁸³ & Marcel M. Lambrechts⁸⁴

¹Laboratoire Ecologie, Systematique et Evolution, UMR 8079 CNRS-Université Paris-Sud XI-AgroParisTech, Batiment 362 Université Paris-Sud XI, Orsay Cedex F-91405, France

²Evolutionary Ecology Group, Department of Biology, University of Antwerp, Campus CGB, Antwerp B-2020, Belgium

³Russian Academy of Sciences, Karelian Research Centre, Institute of Biology, Petrozavodsk 185610, Russia

⁴Department of Experimental Zoology & Evolutionary Biology, University of Łódź, Banacha 12/16, Łódź 90-237, Poland

⁵Terrestrial Vertebrates Research Unit "Cavanilles", Institute of Biodiversity and Evolutionary Biology, University of Valencia, C/Catedrático José Beltrán 2, Paterna E-46980, Spain

⁶Laboratoire Ecologie-Evolution, UMR 7625, Equipe Ecophysiologie Evolutive - Evolutionary Ecophysiology Research Group, Université Pierre et Marie Curie – UPMC, 7 quai Saint Bernard, case 237, Paris Cedex 05 F-75252, France

⁷Centre d'Ecologie Fonctionnelle et Evolutive, CEFE UMR 5175, Campus CNRS, 1919 route de Mende, Montpellier Cedex 5 F-34293, France

⁸Research Laboratory "Ecology of Terrestrial and Aquatic Systems", University Badji Mokhtar, Annaba, Algeria

⁹INRA, UR 1115, Plantes et Systèmes de culture Horticoles, Avignon F-84000, France

¹⁰Àrea de Biodiversitat, Grup de Biologia de la Conservació, Centre Tecnològic Forestal de Catalunya, Carretera de St. Llorenç de Morunys, km. 2, Solsona E-25280, Spain

¹¹Strada Bine, 1424, Acquanegra sul Chiese (MM) I-46011, Italy

¹²Centre d'Ecologie Fonctionnelle et Evolutive, CEFE UMR 5175, Campus CNRS, 1919 route de Mende, Montpellier Cedex 5 F-34293, France

¹³University of Haifa, Haifa, Israel

¹⁴Society for the Protection of Nature, University of Lausanne, Lausanne, Switzerland

¹⁵Institution of Environment Science, Jagiellonian University, Krakow, Poland

¹⁶Stazione Inanellamento c/o Dipartimento SEN-FIMIZO, Università di Palermo, Palermo, Italy

¹⁷Department of Zoology, Siedlce University of Natural Sciences and Humanities, Prusa 12, Siedlce PL-08-110, Poland

¹⁸Centre d'Ecologie Fonctionnelle et Evolutive, CEFE UMR 5175, Campus CNRS, 1919 route de Mende, Montpellier Cedex 5 F-34293, France

¹⁹Univ Lyon 1, Department of Biometry & Evolutionary Biology, CNRS UMR 5558, Villeurbanne F-69622, France

²⁰Centre d'Ecologie Fonctionnelle et Evolutive, CEFE UMR 5175, Campus CNRS, 1919 route de Mende, Montpellier Cedex 5 F-34293, France

²¹Museum and Institute of Zoology, Polish Academy of Sciences, Wilcza 64, Warsaw PL-00-679, Poland

²²Campus Drie Eiken, Department of Biology (Ethology), Building C, Antwerp (Wilrijk) B-2610, Belgium

²³Section of Ecology, Department of Biology, University of Turku, Turku FI-20014, Finland

²⁴Université de Bourgogne, UMR CNRS 5561, BioGéoSciences, 6 Boulevard Gabriel, Dijon F-21000, France

²⁵School of Bioscience, Cardiff University, Cardiff CF10 3AX, UK

²⁶Department of Biology, University of Oulu, Oulu FIN-90014, Finland

²⁷Departamento de Ecología, Facultad de Biología, Universidad de La Laguna, La Laguna E-38260, Spain

²⁸Department of Natural and Social Sciences, University of Gloucestershire, Gloucestershire, GL50 4AZ, U.K.

²⁹Department of Zoology, Edward Grey Institute of Field Ornithology & Institute of Human Sciences, South Parks Road, Oxford OX1 3PS, U.K.

³⁰Museum and Institute of Zoology, Polish Academy of Sciences, Wilcza 64, Warszawa PL-00-679, Poland

³¹Centre d'Ecologie Fonctionnelle et Evolutive, CEFE UMR 5175, Campus CNRS, et Université de Montpellier II, 1919 route de Mende, Montpellier Cedex 5 F-34293, France

³²Department of Animal Ecology, Evolutionary Biology Centre, Uppsala University, Uppsala SE-75236, Sweden

³³Lancaster Environment Centre, Lancaster University, Lancaster LA1 4YQ, U.K.

³⁴Laboratoire Évolution & Diversité Biologique, UPS Toulouse III, Bât 4R1, salle 122, 118 route de Narbonne, Toulouse F-31062, France

- ³⁵CEH Wallingford, Maclean Building, Crowmarsh Gifford, Oxfordshire OX10 8BB, U.K.
- ³⁶Centre d'Ecologie Fonctionnelle et Evolutive, CEFE UMR 5175, Campus CNRS, 1919 route de Mende, Montpellier Cedex 5 F-34293, France
- ³⁷Laboratoire Évolution & Diversité Biologique, UPS Toulouse III, Bât 4R1, salle 122, 118 route de Narbonne, Toulouse F-31062, France
- ³⁸University of Helsinki, Kilpisjärvi Biological Station, P.O.Box 17, Helsinki FIN-00014, Finland
- ³⁹Institute of Ecology of Nature Research Centre, Akademijos 2, Vilnius LT-08412, Lithuania
- ⁴⁰Section of Ecology, Department of Biology, University of Turku, Turku FI-20014, Finland
- ⁴¹Institute of Ecology & Earth Sciences, University Tartu, Tartu EE-51014, Estonia
- ⁴²Section of Ecology, Department of Biology, University of Turku, Turku FI-20014, Finland
- ⁴³Crx. St. Pierre, 6 rue Morcueil, Fleurey Sur Ouche F-21410, France
- ⁴⁴Section of Ecology, Department of Biology, University of Turku, Turku FI-20014, Finland
- ⁴⁵Department of Biology, University of Oulu, Oulu FI-90014, Finland
- ⁴⁶Department of Ecology & Evolution, Uppsala University, Uppsala S-75236, Sweden
- ⁴⁷Lancaster Environment Centre, Lancaster University, Lancaster LA1 4YQ, U.K.
- ⁴⁸Department of Zoology, Institute of Ecology and Earth Sciences, University of Tartu, 46 Vanemuise Str., Tartu EE-51014, Estonia
- ⁴⁹Stazione Inanellamento c/o Dipartimento SEN-FIMIZO, Università di Palermo, Palermo, Italy
- ⁵⁰Museum and Institute of Zoology, Polish Academy of Sciences, Wilcza 64, Warszawa PL-00-679, Poland
- ⁵¹Departamento de Ecología Evolutiva Museo Nacional de Ciencias Naturales, Agencia Estatal Consejo Superior de Investigaciones Científicas, CSIC, C/José Gutiérrez Abascal 2, Madrid E-28002, Spain
- ⁵²Department of Zoology, Rzeszów University, Zelwerowicza 4, Rzeszów PL-35-601, Poland
- ⁵³Department of Biological and Environmental Sciences, University of Jyväskylä, POB 35, Jyväskylä FIN-40014, Finland
- ⁵⁴Centre d'Ecologie Fonctionnelle et Evolutive, CNRS, 1919 Route de Mende, Montpellier, France
- ⁵⁵Ecología Evolutiva, Museo Nacional de Ciencias Naturales (CSIC), José Gutiérrez Abascal 2, Madrid E-28006, Spain
- ⁵⁶Centre d'Ecologie Fonctionnelle et Evolutive, Campus CNRS, 1919 route de Mende, Montpellier Cedex 5 F-34293, France
- ⁵⁷Institute of Biodiversity, Animal Health & Comparative Medicine, University of Glasgow, Graham Kerr Building, Glasgow G12 8QQ, U.K.
- ⁵⁸Ecology Building, Animal Ecology, Lund University, Lund SE-223 62, Sweden
- ⁵⁹Department of Biology, Biodiversity, Lund University, Ecology Building, Lund SE-223 62, Sweden
- ⁶⁰Department of Life Sciences, Institute of Marine Research, University of Coimbra, Apartado 3046, Coimbra PT-3001-401, Portugal
- ⁶¹Department of Biology, University of Oulu, P.O. Box 3000, Oulu FIN-90014, Finland
- ⁶²Centre d'Ecologie Fonctionnelle et Evolutive, CEFE UMR 5175, Campus CNRS, 1919 route de Mende, Montpellier Cedex 5 F-34293, France
- ⁶³Centro de Estudos Florestais, Instituto Superior de Agronomia, University of Lisbon, Lisbon 1349-017, Portugal
- ⁶⁴Campus Drie Eiken, Department of Biology (Ethology), Building C, Antwerp (Wilrijk) B-2610, Belgium
- ⁶⁵Latvian Fund for Nature, Dzirnau Street 73-2, Riga LV-1011, Latvia
- ⁶⁶Centre d'Ecologie Fonctionnelle et Evolutive, CEFE UMR 5175, Campus CNRS, 1919 route de Mende, Montpellier Cedex 5 F-34293, France
- ⁶⁷Laboratory of Ornithology, Department of Zoology, Palacky University, Olomouc CZ-77146, Czech Republic
- ⁶⁸Institute of Ecology & Evolution (IEE), University of Bern, Bern CH-3012, Switzerland
- ⁶⁹Faculty of Sciences, Evolutionary Ecology Group (GIBE), University of A Coruña, Campus Zapateira, A Coruña E-15008, Spain
- ⁷⁰Evolutionary Ecology Group (EVECO), Department of Biology, University of Antwerp, Middelheimcampus, Groenenborgerlaan 171, Antwerp B-2020, Belgium
- ⁷¹Department of Biology, University of Oulu, P. O. Box 3000, Oulu FIN-90014, Finland
- ⁷²Unidad Asociada CSIC de Ecología Evolutiva y de la Conducta, Nat-Museu de Ciències Naturals de Barcelona, Barcelona, Spain
- ⁷³Department of Biological and Environmental Science, University of Jyväskylä, Jyväskylä, Finland
- ⁷⁴Department Life Science, IMAR CMA, University Coimbra, Coimbra PT-3004517, Portugal
- ⁷⁵Department of Biology, University of Oslo, Oslo, Norway
- ⁷⁶Luontotutkimus Solonen Oy, Neitsytsaarentie 7b B 147, Helsinki FIN-00960, Finland
- ⁷⁷SROPU, Via R. Crippa 60, Rome, Italy
- ⁷⁸School of Life Sciences, University of Sussex, Brighton, Sussex BN1 9QG, U.K.
- ⁷⁹Behavioral Ecology Group, Department of Systematic Zoology & Ecology, Eötvös Lorand University, Budapest H-1117, Hungary
- ⁸⁰Institute of Zoology, Poznan University of Life Sciences, Wojska Polskiego 71 C, Poznań PL-60-625, Poland
- ⁸¹Netherlands Institute of Ecology (NIOO-KNAW), Doevendaalsesteg, 10, Wageningen NL-6708 BP, the Netherlands
- ⁸²Environmental and Marine Biology, Åbo Akademi University, Artillerigatan 6, Biocity, Åbo FI-20520, Finland
- ⁸³Department of Zoology, Siedlce University of Natural Sciences and Humanities, Prusa 12, Siedlce PL-08-110, Poland
- ⁸⁴Centre d'Ecologie Fonctionnelle et Evolutive, CEFE UMR 5175, Campus CNRS, 1919 route de Mende, Montpellier Cedex 5 F-34293, France

Keywords

Hole nesting, natural holes, nest boxes, reaction norm.

Abstract

Nests are structures built to support and protect eggs and/or offspring from predators, parasites, and adverse weather conditions. Nests are mainly constructed prior to egg laying, meaning that parent birds must make decisions about nest site choice and nest building behavior before the start of egg-laying.

Correspondence

Anders P. Møller
 Laboratoire Ecologie, Systematique et
 Evolution, UMR 8079 CNRS-Université Paris-
 Sud XI-AgroParisTech, Batiment 362
 Université Paris-Sud XI, F-91405 Orsay Cedex,
 France.
 Tel: (+33) 1 69 15 56 88;
 Fax: (+33) 1 69 15 56 96;
 E-mail: anders.moller@u-psud.fr

Funding Information

No funding was provided for this study

Received: 13 May 2014; Revised: 7 July
 2014; Accepted: 15 July 2014

***Ecology and Evolution* 2014; 4(18): 3583–
 3595**

doi: 10.1002/ece3.1189

Parent birds should be selected to choose nest sites and to build optimally sized nests, yet our current understanding of clutch size-nest size relationships is limited to small-scale studies performed over short time periods. Here, we quantified the relationship between clutch size and nest size, using an exhaustive database of 116 slope estimates based on 17,472 nests of 21 species of hole and non-hole-nesting birds. There was a significant, positive relationship between clutch size and the base area of the nest box or the nest, and this relationship did not differ significantly between open nesting and hole-nesting species. The slope of the relationship showed significant intraspecific and interspecific heterogeneity among four species of secondary hole-nesting species, but also among all 116 slope estimates. The estimated relationship between clutch size and nest box base area in study sites with more than a single size of nest box was not significantly different from the relationship using studies with only a single size of nest box. The slope of the relationship between clutch size and nest base area in different species of birds was significantly negatively related to minimum base area, and less so to maximum base area in a given study. These findings are consistent with the hypothesis that bird species have a general reaction norm reflecting the relationship between nest size and clutch size. Further, they suggest that scientists may influence the clutch size decisions of hole-nesting birds through the provisioning of nest boxes of varying sizes.

Introduction

Numerous organisms, including insects, spiders, crustaceans, fish, amphibians, reptiles, birds, and mammals, all construct nests that are used for containing their eggs and/or offspring for a considerable amount of time (Collias and Collias 1984; Taylor *et al.*, 1998; Hansell 2000, 2007). Nest size depends on nest site, and choice of a nest hole as a breeding site may constrain the size of nests. Hence, nest size in hole nesters may depend on the size of the cavity or the nest site. Nests are often larger than the builder(s) themselves and sometimes considerably so, as shown by ants, termites, and some birds such as raptors (Collias and Collias 1984; Hansell 2000, 2007). Nest building is an energetically costly and time-consuming activity that is presumably traded-off against participation in other activities (Collias and Collias 1984; Hansell 2000, 2007; Mainwaring and Hartley 2013).

The parental expenditure on nest building can appear to be paradoxical as nests are often much larger than is required for successful reproduction (Soler *et al.* 1998a,b; and references below). This poses the question as to why animals build nests that are larger than is necessary for raising the offspring (Mainwaring and Hartley 2013).

There are six main, nonmutually exclusive, reasons why large nests may be advantageous. First, the maintenance of a specific nest temperature and humidity (Mertens 1977a,b; Erbeling-Denk and Trillmich 1990; Mainwaring *et al.* 2012). Optimal nest size may provide a microenvironment that is not too cooled or too over-heated for growth and development of temperature regulation

(Mertens 1977a,b; Erbeling-Denk and Trillmich 1990; Mainwaring *et al.* 2012). Larger and better insulated nests built at higher latitudes (Schaefer 1980; Møller 1984; Crossman *et al.* 2011; Mainwaring *et al.* 2012), higher altitudes (Kern and Van Riper 1984), and early during the season when temperatures are cooler at temperate latitudes (Møller 1984; Mainwaring and Hartley 2008) are consistent with thermal benefits of large nests at extreme temperatures. Second, specific types of nest material may provide protection of eggs and nestlings against bacteria and parasites (Wimberger 1984; Mennerat *et al.* 2009; Peralta-Sánchez *et al.* 2010). Third, an optimal nest size will prevent excessive fouling of nests and the associated fitness costs of nestling death by allowing parents to keep the nest clean, as demonstrated experimentally for starlings *Sturnus vulgaris* by Erbeling-Denk and Trillmich (1990). Fourth, sexual selection (Tortosa and Redondo 1992; Soler *et al.* 1998a,b; Møller 2006; Broggi and Senar 2009; Sanz and García-Navas 2011; Tomás *et al.* 2013). Soler *et al.* (1998a) reported for European passerines the relationship between nest size and the number of parent birds involved in nest building, and found that nest size almost doubled when both male and female build nests as compared to nests built by females alone. This was interpreted to have arisen because both parents signal their phenotypic quality via their investment in nest building, despite the fact that clutch size did not differ between species in which only females build nests and species in which both sexes build. This interpretation is supported by intraspecific correlational and experimental studies showing that larger nests are favored because of sexual

selection (Soler *et al.* 1998b, 1999, 2001; de Neve *et al.* 2004; Tomás *et al.* 2013). Fifth, structural support may prevent eggs and nestlings from being lost. Large nests may benefit parent birds through improved insulation and the reduced heat loss from eggs and nestlings. Sixth, large nests may reduce crowding of the offspring and prevent them from falling out of the nest (Slagsvold 1982, 1989; Heenan and Seymour 2011). Thus, reproductive success is positively correlated with nest size (Alabrudzinska *et al.* 2003; Álvarez and Barba 2008), raising the question as to why nest size does not continue to increase.

Nest sizes are restricted by five main selective forces. First, the risk of nest predation and parasitism selects for smaller nest sizes by disproportionately affecting larger nests (Møller 1990; Eeva *et al.* 1994; Soler *et al.* 1999; Antonov 2004). Second, building larger nests means more time spent collecting nesting materials, which are often found on the ground, increasing the risk of predation on nest-building adults (Slagsvold and Dale 1996). Third, large and consequently warm and humid nests do not only benefit offspring, but also pathogens and parasites that inhabit nests, and bacteria multiply at higher rates in warm and humid nest environments compared to ambient temperatures, with potential costs to their avian inhabitants (Stolp 1988). Some nest-building birds have evolved counter-adaptations against exploitation by parasites by using nest material with antimicrobial properties such as feathers and plants with secondary defense substances (Clark and Mason 1985; Gwinner 1997; Mennerat *et al.* 2009; Møller *et al.* 2013) or uropygial secretions that eliminate bacteria from eggs (Soler *et al.* 2012). Fourth, nest size may be limited by individual quality and the construction of larger nests when males participate in nest building may have a signaling function, reflecting the working ability of males for finding specific or scarce nest materials (Møller 2006; Broggi and Senar 2009; Sanz and García-Navas 2011). Fifth, nest size may increase the risk of brood parasitism (Soler *et al.* 1995, 1999), or decrease the probability of rejection of host offspring by brood parasites (Anderson *et al.* 2009; Grim *et al.* 2009). These effects seem minor given that most hole nesters are only rarely parasitized by parasitic cuckoos (Moksnes *et al.* 1991; Moksnes and Røskaft 1995; Davies 2000).

Despite the many and varied factors selecting for or against larger nests, a major factor is likely to be clutch size. Clutch size is often larger when nests are larger (Löhrl 1973, 1980; Karlsson and Nilsson 1977; van Balen 1984; Gustafsson & Nilsson, 1985; Korpimäki 1985; Rendell and Robertson 1993; Wiebe and Swift 2001). This is surprising because nests are generally made well before the first eggs are laid, although parent birds sometimes make adjustments to nests during incubation and even as

late as the nestling period (Møller 1987), posing a potential problem for parents at the time of nest building due to uncertainty about final clutch size. This relationship may arise from both clutch size and nest size being correlated negatively with laying date. This cannot be the entire explanation for experimental studies such as those by Löhrl (1973, 1980), Erbeling-Denk and Trillmich (1990), Rendell and Robertson (1993), and Soler *et al.* (2001), who documented a change in clutch size when nest box size was manipulated after the start of egg laying. These experimental studies showed effects of nest size on clutch size independent of potentially confounding variables. On average, a larger nest after laying is required for raising more offspring. There may be a correlation between the choice of nest site by females and the subsequent clutch size, females building larger nests because such females lay larger clutches, or females laying larger clutches may build larger nests. Several experiments have tried to separate the effects of the size of the nest cavity and the effects of female phenotype on subsequent clutch size by changing nest size through exchanging nest boxes of different sizes after the female had started laying. Exchange experiments performed during egg laying in the great tit resulted in an increase in clutch size when box size increased, but a decrease when a smaller box was provided (Löhrl 1973, 1980; Erbeling-Denk and Trillmich 1990; Rendell and Robertson 1993).

The objective of this study was to investigate the plasticity of clutch size in response to nest size in both hole and open nesting birds. We included all species for which information on nest size and clutch size was available in these analyses because females faced the problem of adjusting clutch size to nest size before hole nesting evolved, and because hole nesting is a derived character in birds (Lack 1968). Therefore, adjustment of clutch size to nest size is likely to have evolved before hole nesting. First, using data in classical experiments on great tits by Löhrl (1973, 1980), we investigated the effects of nest box size on clutch size, when great tits had access to differently sized nest boxes, and when the size of nest boxes was manipulated after laying had started. The latter design prevented differential access of specific females to boxes of a particular size. Second, we examined the plasticity of clutch size to nest size in hole nesters where nest box sizes were varied experimentally within a population. Third, we analyzed the relationship between clutch size and nest size using studies of four secondary hole-nesting species with both single or multiple nest box sizes. We predicted that if hole-nesting birds use a single type of plasticity as a clutch size response to cavity size, then we should find a homogeneous slope linking clutch size to cavity size. Fourth, if different species of hole-nesters respond differently to cavity size, for example, because

populations in different environments encounter different problems in terms of hyperthermia, we should expect heterogeneous relationships between clutch size and cavity size. Finally, we predicted that the slope of the relationship between clutch size and nest size depended on species, latitude and longitude if climatic conditions were underlying variation in slope among studies. We emphasize that the results presented by Møller et al. (2014) only constitute one out of 11 analyses reported here. These previous analyses of 365 populations with more than 79,000 clutches showed that clutch size in the great tit *Parus major*, but not in three other species of hole nesters increased with nest box size, and that nest box size and quality varied nonrandomly among populations (Møller et al. 2014). Here, we broaden this approach to a larger number of species, including open nesting species, by exploring the causality of this relationship and analyzing whether plasticity of clutch size in relation to nest size vary between species and/or regions when correcting for macrogeographic and habitat patterns. Open nesting species differ from hole nesters by being less constrained by nest cavity size as hole nesters. This approach will give insight about the factors governing the relationship between clutch and nest size.

Materials and Methods

Data sets

This study was based on three different data sets: (1) Löhrl's (1973, 1980) data on great tits; (2) a data base on clutch size and nest box size in four secondary hole-nesting birds; and (3) an exhaustive database on the relationship between clutch size and nest size in 21 species of birds for which data on clutch size and nest size were available (Table S1). We defined nest size in hole-nesters as the two-dimensional area of the nest box or nest hole in case the study was based on natural cavities (i.e., nest base area). Similarly, we defined nest size for open nesting species as the horizontal maximum area of the nest including nest cup and nest rim (i.e., nest base area). Nest size refers to this definition in the remainder of the article. We used this measure of nest size because it was available for almost all studies. We could not use total volume of the nest or nest weight as a measure of nest size because only a small number of studies had recorded and/or published such data. Use of our definition of nest size relies on the assumption that nest volume and/or nest weight is positively correlated with nest size as defined here. We were able to explicitly test this assumption for nest size (calculated as $\frac{4}{3} \times \pi \times a^2 \times b$, where a is the radius and b is the height) in five studies of four species, two hole nesters and two open nesters: blue tits

(M. Lambrechts et al., unpubl. data: $F_{1,52} = 176.68$, $r^2 = 0.77$, $P < 0.0001$, slope (SE) = 1.484 (0.112)), great tits (M. Lambrechts et al., unpubl. data: $F_{1,212} = 634.07$, $r^2 = 0.75$, $P < 0.0001$, slope (SE) = 1.280 (0.050)), and great tits (Slagsvold and Amundsen 1992: $F_{1,27} = 207.07$, $r^2 = 0.99$, $P < 0.0001$, slope (SE) = 1.197 (0.083)), the open nesting barn swallows *Hirundo rustica* (A. P. Møller, unpubl. data: $F_{1,1265} = 866.17$, $r^2 = 0.40$, $P < 0.0001$, slope (SE) = 1.202 (0.041)) and rufous bush warblers *Cercotrichas galactotes* (M. Martín-Vivaldi, unpubl. data: $F_{1,48} = 24.28$, $r^2 = 0.34$, $P < 0.0001$, slope (SE) = 0.937 (0.190)) using \log_{10} -transformed data.

We extracted data from Löhrl's (1973, 1980) pioneering studies of great tit clutch size, choice of nest box size and experimental manipulation of nest box size after start of laying. We used these data to estimate the relationship between nest size and clutch size, and in the nest box exchange experiment the relationship between clutch size and the change in nest box size.

We made an exhaustive attempt to obtain information on clutch size, nest size and ecological variables from all studies of four common species of secondary hole nesters in Europe and North Africa, as described in detail elsewhere (Møller et al. 2014). Briefly, we attempted to obtain data on first clutches, or early clutches known to be initiated less than 30 days after the first egg was laid in a given year in a local study plot (cf. Nager and van Noordwijk 1995). In total, we obtained information on 155 study populations of great tits, 121 of blue tits *Cyanistes caeruleus*, 24 of pied flycatchers *Ficedula hypoleuca*, and 65 of collared flycatchers *Ficedula albicollis* for a total of 1479 study years for great tits, 1122 for blue tits, 288 for pied flycatchers, and 592 for collared flycatchers, or in total 79,610 clutches (Møller et al. 2014).

For all studies, we also quantified latitude ($^{\circ}$ N) and longitude ($^{\circ}$ E), main habitat type (deciduous, coniferous, evergreen, or mixed), urbanization (urbanized, or natural/semi-natural habitat), altitude at the center of a study plot, mean study year, nest floor surface as the internal nest base area (in cm^2), and the material used to construct nest boxes (a binary variable classified as either wood or concrete). We defined minimum and maximum size of boxes for all study sites that used two or in one case three nest box sizes by assigning the smallest box size as the minimum size and the largest box size as the maximum size. Further details of how these variables were obtained and quantified can be found in Lambrechts et al. (2010) and Møller et al. (2014).

We made an exhaustive literature search for papers dealing with clutch size and nest size in birds using Web of Science (clutch size, nest size, and birds were used as keywords) combined with a search for other publications in the reference lists of these papers (Lambrechts et al.

2010; Møller *et al.* 2014). If there was missing information on clutch size or nest size in these publications, we attempted to retrieve this information by contacting the authors. The extensive data set on clutch size and nest size in barn swallows was collected by APM by measuring inner and outer diameter and inner depth of all nests in a Danish study site during 1988–2012 (see Møller (2006) for further details).

Ecological characteristics

All studies that allowed an estimation of slope of the relationship between clutch size and base area of the nest were cross-classified with respect to (1) hole nesting, (2) box or natural hole, (3) nest building, (4) body mass and (5) minimum, and (6) maximum size of nests used in a given study to test for an association between the slope of the relationship between clutch size and nest size for each year for all study populations. Species were classified as (1) hole nesters (yes – breeding in cavities, no – not breeding in cavities), (2) nest builders (yes – building or excavating the nest or the cavity, no – not building or excavating the nest or the cavity), as reported in Cramp and Perrins (1977–1994), Dunning (1993) and Poole (2005), and (3) nest box populations (yes, no). All studies with slope estimates of the relationship between clutch size and nest size as defined above and the ecological data are listed in the Table S1.

Statistical analyses

This study presents the results of 11 statistical analyses of the relationship between clutch size and nest size within and among species of birds (we have found no comparable data on any other taxa). This study differs from that by Møller *et al.* (2014) by analyzing an exhaustive compilation of data on both hole and open nesting species and a wider range of factors hypothesized to affect the relationship between clutch size and nest size.

We re-analyzed Löhr's (1973, 1980) data to allow distinction between the effects of original and subsequent nest box size on clutch size following exchange of nest boxes during the laying period. We did so by analyzing clutch size (the response variable) in relation to original and experimental nest box size.

We estimated slopes of the relationship between clutch size and nest base area. These analyses were based on all studies with at least two different sizes of nest boxes (all but one study had two box sizes only), where the analyses were weighted by sample size for the different categories of nest box sizes. Similarly for open nesting species, we estimated slopes by relying on clutch size and base area of nests.

We have previously investigated the relationships between the average size of first clutches and internal base area (cm^2) of artificial cavities after controlling for other factors influencing average clutch size (Møller *et al.* 2014). We used mixed models (GLMM) including (1) plot ID as a random effect to account for differences in sample sizes among plots, and (2) species (great tit, blue tit, pied flycatcher, collared flycatcher) as a random factor that accounted for the difference in level of sampling among species and the following variables as factors: (3) latitude ($32\text{--}65^\circ\text{N}$), (4) longitude ($9^\circ\text{W--}35^\circ\text{E}$), (5) dominant vegetation (deciduous, coniferous, evergreen, mixed), (6) degree of urbanization (urban, rural), (7) altitude (0–1650 m a. s. l.), (8) cavity type (artificial box, tree cavity), (9) material (cavity wall made of wood, cavity wall made of material other than wood, mainly wood-concrete), (10) year (average study year for studies that lasted >1 year), and (11) study period (number of study years). We \log_{10} -transformed altitude to eliminate the skewed distribution adding a constant of 192 m to avoid inclusion of negative values recorded in a few study sites below sea level. We weighted the analyses by sample size to account for the fact that estimated clutch sizes based on large sample sizes will be closer to the actual population mean than those based on small sample sizes.

In the current study, we tested for a difference in slope among 116 populations of hole-nesting and open-nesting bird species using mixed models with slopes weighted by sample size as the response variable and species as a fixed factor (21 species), and latitude and longitude as covariates. Because transformations did not result in normal distributions, we used ranked slopes as a response variable in this analysis. In addition, we used Welch ANOVA for unequal variances weighted by sample size to test for differences in means and variances among species, again relying on ranked slopes. Finally, we also included into our least squares models the minimum and maximum size of nests/boxes used in each study to test for their effects on slope estimates. The justification for this approach was that minimum size would reflect the importance of hyperthermia created in the cramped conditions of small nests, while maximum nest size would reflect problems of hypothermia (Mertens 1977a,b).

When quantifying the effects of base area of nests on clutch size we estimated the change in clutch size for an increase in one standard deviation unit in base area using the linear regression coefficient. In all 11 statistical tests presented in Table 1 we provide the mean and the standard error of the parameter estimates. These were subsequently used to estimate the 95% confidence intervals (CI) for the estimated slopes. If 95% CI of two or more samples did not overlap, they were considered to be

Table 1. Relationship between clutch size and nest base area in different studies and their 95% confidence intervals (CI).

Relationship	Species	F	df	P	Estimate (SE)	95% CI
A. Clutch size and base area in choice test (Löhrl 1973)	<i>Parus major</i>	28.61	1, 49	<0.0001	0.0088 (0.0016)	0.0056, 0.0120
B. Clutch size and base area when box is exchanged during laying (Löhrl 1980)	<i>Parus major</i>	6.48	1, 18	0.020	0.0264 (0.0104)	0.0045, 0.0483
C. Clutch size and change in base area when box is exchanged during laying (Löhrl 1980)	<i>Parus major</i>	6.48	1, 18	0.020	0.0132 (0.0052)	0.0033, 0.0241
D. Clutch size and base area in studies based on two or more nest box sizes	<i>Parus major</i> , <i>Cyanistes caeruleus</i> , <i>Ficedula hypoleuca</i> , <i>Ficedula albicollis</i>	6.50	1, 68	<0.0001	0.0036 (0.0014)	0.0009, 0.0063
E. Clutch size and base area in all studies of four secondary hole nesters	<i>Parus major</i> , <i>Cyanistes caeruleus</i> , <i>Ficedula hypoleuca</i> , <i>Ficedula albicollis</i>	3.37	1, 3371	0.067	-0.0211 (0.0115)	-0.044, 0.0014
F. Clutch size and base area in great tit	<i>Parus major</i>	24.31	1, 1434	<0.0001	0.0071 (0.0014)	0.0044, 0.0098
G. Clutch size and base area in blue tit	<i>Cyanistes caeruleus</i>	2.11	1, 382.2	0.15	-0.0038 (0.0026)	-0.0089, 0.0013
H. Clutch size and base area in pied flycatcher	<i>Ficedula hypoleuca</i>	0.65	1, 22.41	0.43	-0.0016 (0.0020)	-0.0055, 0.0023
I. Clutch size and base area in collared flycatcher	<i>Ficedula albicollis</i>	0.00	1, 32.29	0.99	0.000013 (0.0020)	-0.0039, 0.0039
J. Clutch size and base area in all open-nesting species	Open-nesters	9.90	1, 25	0.0037	0.0033 (0.0010)	0.0013, 0.0054
K. Clutch size and base area in all species	All species	12.51	1, 114	0.0006	0.0036 (0.0010)	0.0056, 0.0063

significantly different. All statistical tests were made with JMP (SAS 2012).

Results

Clutch size and nest size in Löhrl's studies of great tits

First, we investigated whether the relationship between clutch size and nest box base area was significant when great tits could choose between boxes that had base area of either 64 or 314 cm², using data reported by Löhrl (1973). This relationship was indeed significantly positive (Table 1A, slope (SE) = 0.0088 eggs cm⁻² (0.0016)).

Second, Löhrl (1980) provided experimental evidence for great tits adjusting their clutch size directly to the base area of nest boxes by exchanging the nest boxes during the laying period. This justifies that the analyses with clutch size as a response variable and nest base area as a predictor variable were made without inclusion of additional variables. The relationship between clutch size and box base area was statistically significant (Table 1B, slope (SE) = 0.0264 (0.0104)). This relationship and that

reported when great tits were allowed to choose box size were not significantly different, as shown by their overlapping 95% CI (Table 1). These data also allowed a test of whether the relationship between clutch size and change in box base area between the original and the new box was statistically significant. That was indeed the case. The slope was half that of the previous estimate (Table 1C), although the 95% CI of the parameter estimate overlapped with the previous two estimates (Table 1A, B). Mean nest base area in the second dataset for great tits weighted by sample size across all 3447 samples was 118 cm², range 36–400 cm², SD = 123 cm². This implies that an increase by one SD in nest base area was associated with an average increase in clutch size of the great tit by 0.0088 eggs cm⁻² × 123 cm² = 1.08 eggs to 0.0264 eggs cm⁻² × 123 cm² = 3.25 eggs for these two extreme estimates of the relationship.

Clutch size and nest size across species of birds

We obtained an estimate of the clutch size – nest base area relationship in population studies of all four species

of hole-nesting birds (great tits, blue tits, pied flycatchers, and collared flycatchers) with more than one box size since a slope estimate requires at least two box sizes within a study site. The mean slope weighted by sample size was 0.0036 (SE = 0.0014), $N = 69$ populations and four species (Table 1D). This mean slope estimate was not significantly different from those reported by Löhrl (1973, 1980), as shown by the overlapping 95% CI (Table 1).

We then estimated the relationship between clutch size and base area in populations with one or more box sizes for the four secondary hole nesters considered above. This relationship was statistically significant for all species combined (Table 1E). It was also statistically significantly positive for great tit separately (Table 1F), but not for blue tit (Table 1G), collared flycatcher (Table 1H) or pied flycatcher (Table 1I). The interaction between species and nest floor area was highly significant ($F_{3,3436} = 115.12$, $P < 0.0001$), implying that clutch size was related to nest floor area in a species-specific manner. A Welch analysis of variance for unequal variances with ranked slope as the response variable, species as a fixed factor and sample size as a weight revealed a significant effect of means of species ($F_{7,3660.6} = 6.23$, $P < 0.0001$), and a Levene's test showed a significant difference in variance among species ($F_{7,95} = 5.06$, $P < 0.0001$). Thus, a general positive relationship between clutch size and nest size differed significantly among species, but the variances within species also differed significantly. The overall slope estimate was not significantly different from those reported by Löhrl (1973, 1980) for great tits, as shown by the overlapping 95% CI (Table 1).

Open-nesting species also showed an overall positive relationship between clutch size and base area (Table 1J) that overlapped with all the slope estimates reported in Table 1 except for the estimate for clutch size and nest base area in Löhrl's (1973) choice test (Table 1A). We tested whether the relationship between clutch size and base area was statistically significant across all species by inclusion of hole-nesters and open-nesters. The overall slope estimate was statistically significant (Table 1K). It overlapped with most of the slope estimates reported in Table 1 suggesting that even inclusion of open-nesters in an overall estimate across species did not change the conclusion that there was a general positive relationship between clutch size and base area in birds.

We tested whether there was a species-specific relationship between clutch size and nest base area in the data set based on study sites where more than one box size was used and more than one nest size was recorded (Fig. 1). This test was based on all studies for which we had information on the regression coefficient between clutch size and nest base area. Based on these 116 datasets, a Welch

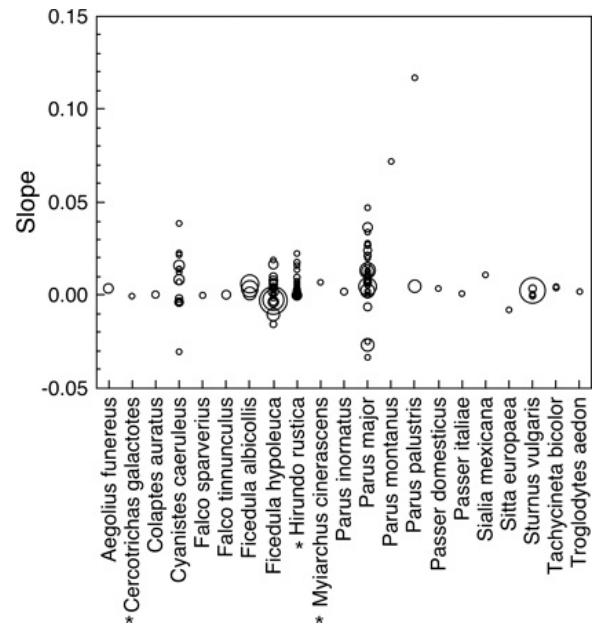


Figure 1. Slopes of the relationship between nest base area and clutch size in different species of birds in each study population. The three open-nesting species are shown with an *. Note that symbol size is related to the number of nests.

analysis of variance for unequal variances with ranked slope as the response variable, species as a fixed factor and sample size as a weight revealed a significant effect of means of species (Fig. 2; $F_{7,3660.6} = 6.23$, $P < 0.0001$), and a Levene's test showed a significant difference in variance among species ($F_{7,95} = 5.06$, $P < 0.0001$). This shows a general positive relationship that differed significantly

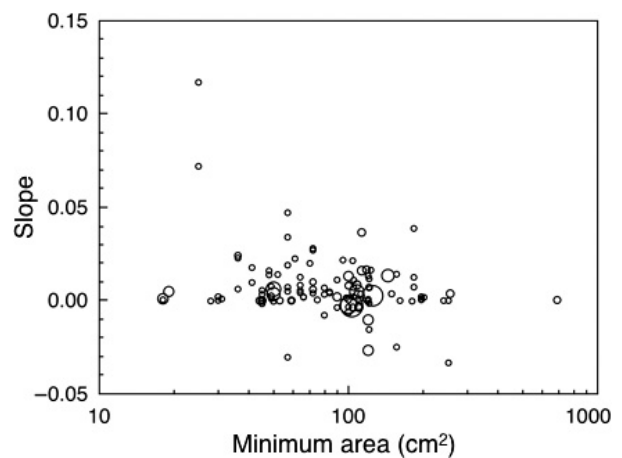


Figure 2. Slopes of the relationship between nest base area and clutch size in relation to the minimum area (cm^2) of nest box bases in different populations and species of birds. Note that symbol size is related to the number of nests.

among species, and the variances within species also differed significantly.

Because ambient temperatures during the breeding season are predictably warmer at lower latitudes, we expected a stronger relationship between clutch size and nest base area in southern populations than in northern populations. However, adding latitude ($F_{1,93} = 1.43$, $P = 0.23$) or longitude ($F_{1,93} = 1.02$, $P = 0.31$) or the squared values to account for nonlinear effects (latitude squared: $F_{1,92} = 0.01$, $P = 0.90$; longitude squared: $F_{1,92} = 0.04$, $P = 0.85$) to the model for the 21 species did not show significant effects. A least squares model with species as a fixed effect showed a significant negative association between slope of the relationship between clutch size and nest size and minimum size (Table 2; Fig. 2) and maximum size of boxes in a given study (Table 2). The effect of species only accounted for 2.7% of the variance and hence was negligible. The effect of minimum size was almost twice as large as the effect of maximum size. All variance inflation factors were smaller than 5 and hence did not pose a problem of collinearity (McClave and Sincich 2003),

Discussion

The main findings of this study of the relationship between clutch size and nest size are that most populations and species of both hole and open nesters show increasing clutch size with increasing nest size, with a general increase that varied within and among species. This relationship did not differ between hole and open nesters.

Clutch size was directly affected by nest size, as shown by Löhrl's (1973, 1980) experiments that manipulated the floor area of nest boxes after clutch initiation and thereby produced significant differences in clutch size. Here, we have extended Löhrl's findings by showing that the

relationship between clutch size and nest size was similar when females could or could not choose a specific box size. Löhrl (1973, 1980) used boxes with base area of 64 or 314 cm², or almost a fivefold difference. However, this difference is similar to natural variation in base area in the great tit (van Balen et al. 1982). This implies that differential access to preferred boxes is not the main cause of bias in estimates of the relationship between clutch size and nest base area. Soler et al. (2001) have shown for magpies *Pica pica* that experimental reduction of the size of the roof reduced clutch size. Given that females responded to a change in the size of the part of the nest that does not hold the eggs and the nestlings, we can conclude that it is nest size and nest building, but not nest cup size that is driving clutch size decisions of the female. Most analyses of the relationship between clutch size and our proxy of nest size that we have reported in Table 1 were statistically significant and slopes were in many cases positive with overlapping 95% confidence intervals among experiments and data sets and for open and hole nesters. Interestingly, the plasticity of the relationship between clutch size and nest size was similar in open and hole-nesting species. Among great tits, there was evidence of one standard deviation in nest base area equaling an increase in clutch size by 1.1–3.3 eggs, which is a considerable change given a modal clutch size of 8.69 eggs in 1453 population-year estimates. We suggest that in natural habitats cavities may differ substantially in size and shape, resulting in the evolution of phenotypic plasticity of clutch size in relation to nest size, although this observation does not explain the difference in reaction norms among hole-nesting species. More recently, competition for nest holes of an optimal size in man-managed forests, where older, larger, and dead trees are regularly removed, may lead to a scarcity of natural cavities for breeding birds (Sandström 1992; Carlson et al. 1998; Sánchez et al. 2007) and may have contributed to the evolution of this extreme reaction norm. We did find evidence of statistical heterogeneity among four secondary hole-nesting species and among more than 17,000 nests belonging to 21 species of birds. Among the four common species of secondary hole-nesting species, great tits showed a significant positive relationship, while blue tits and the two *Ficedula* flycatchers showed no significant relationship (Table 1; Møller et al. 2014). In addition, the 95% CI for great tit did not overlap with the 95% CI for blue tit or the two flycatcher species. Interestingly, intraspecific studies on the relationship between nest size and reproductive success have shown a significant relationship in great tits (Alabrudzinska et al. 2003; Álvarez and Barba 2008), but not in blue tits (Lambrechts et al. 2012). As we stated in the Introduction, the relationship between clutch size and nest size may arise as a consequence of female choice of

Table 2. Least squares model of the relationship between ranked slope relating clutch size to minimum and maximum base area and species (fixed effect). The model had the statistics $F = 2.68$, $df = 22, 93$, $r^2 = 0.24$, $P = 0.0005$.

Variable	Sum of squares	df	F	P	Estimate	Error
Intercept			18.84	<0.0001	264.35	60.94
Minimum base area (cm ²)	847668	1	7.61	0.007	-62.05	22.49
Maximum base area (cm ²)	1242860	1	11.16	0.0012	-37.67	11.28
Species	57274564	20	2.57	0.0012		
Error	10354141	93				

an appropriately sized box in relation to the future clutch size of the female. In fact, nest volume was positively correlated with nest base area as shown in five studies of two species of hole nesters and two species of open nesters. However, it seems unlikely that this relationship is caused by nest site choice as nest cups are usually considerably smaller than the base area of nest boxes, and the relationship between clutch size and nest size did not differ significantly between open and hole nesters.

In addition, the relationship between clutch size and nest size could arise as the reaction norm of individual females to their nest sites. Löhrl's (1980) study of great tits and our re-analysis of his data clearly show that the latter mechanism is at work. Given that the slope of the relationship is stronger for the reaction of the female rather to altered box size than the choice of box size by the female, this implies that the former mechanism may be sufficient for explaining the patterns of phenotypic variation. In addition, an analysis of all studies with two or more nest or box sizes showed significant heterogeneity in slopes among the 21 species of birds including the three open nesting species, but also significant differences in variance within species (Table 2). However, the relationship did not differ significantly between open nesting and hole-nesting species. Given that hole nesting is a derived character, we can conclude that the relationship between nest size and clutch size evolved before hole nesting, justifying the inclusion of open-nesting species in the present study.

Slagsvold (1987) provided the most exhaustive list of hypotheses accounting for the association between clutch size and nest base area within species, using extensive experiments on the pied flycatcher. The seven hypotheses listed by Slagsvold can basically be grouped into hypotheses concerning energetics, nest predation, and sexual selection. Here, we have documented differences in the relationship between clutch size and nest size as reflected by the slope among four species of secondary hole nesters and among 21 species of birds, for which we had data, suggesting that the interspecific differences are real. The phylogenetic signal in these data must be weak at best given that species only accounted for a couple of per cent of the variance. Given that an increase in nest floor area by 100 cm² is equivalent to an additional 0.36 eggs in the 21 species investigated or 1.25 eggs in the four secondary hole nesters, these effects cannot be considered negligible. An increase in floor area of 100 cm² is similar to standard deviations of 5–165 cm² in ten hole-nesting species in natural holes (van Balen *et al.* 1982, Table 14). We found no significant effects of latitude and longitude or a number of other variables predicted to affect this relationship between clutch size and nest size.

The analyses reported here suggest that the reaction norm of the relationship between clutch size and nest size is similar in nest boxes and in natural holes, because the

effect of nest type (nest box or natural hole) did not enter the statistical analyses as a significant predictor. This conclusion is comforting because it suggests that scientists working with nest box breeding birds are studying natural phenomena within a single context (Møller 1989; Lambrechts *et al.* 2010). However, these findings also suggest that comparison of observations among nest box sizes within or among populations should be made with care. We found evidence that in particular the minimum size of nests/boxes used, but less so the maximum size was significantly related to slope estimates, as shown for the analysis of studies with two or more sizes of nests/boxes used. This greater difference for the minimum size of nest boxes used than for the maximum size of nest boxes used suggests that the reaction norm is increasing for the smallest nests/boxes, but less so for the largest nests/boxes for the range of sizes investigated so far. One way forward would be to estimate the relationship between clutch size and nest size experimentally using a range of nest base areas from 25 to 500 cm², while experimentally manipulating clutch size, with replicates from North Africa to Northern Scandinavia. If hyperthermia is the underlying selective factor accounting for the relationship between clutch size and nest size, we should expect stronger negative effects in a small nest box treatment with an enlarged clutch size at low latitudes.

In conclusion, we have shown in this exhaustive analysis of the relationship between clutch size and base area of nests that such a relationship is common across populations and species with differences in effects among populations within species and among species. The findings suggest that scientists by choosing particular nest box sizes affect clutch size and hence potentially reproductive success in a species-specific manner (Møller *et al.* 2014). These findings may be of general importance for nest building taxa of insects, fish and mammals. However, such tests must await future studies of other taxa since no comparable studies are currently available.

Acknowledgments

We thank the hundreds of collaborators and contributors who helped with study plot management, data collection, data management, administration, financial support, and scientific discussion. Listing their names individually would most probably provide a biased picture of all of their contributions. Dr. M. Martín-Vivaldi and Dr. Juan Moreno kindly provided unpublished data.

Conflicts of Interest

None declared.

References

- Alabrudzinska, J., A. Kalinski, R. Slomczynski, J. Wawrzyniak, P. Zielinski, and J. Banbura. 2003. Effects of nest characteristics on breeding success of great tits *Parus major*. *Acta Ornithol.* 38:151–154.
- Álvarez, E., and E. Barba. 2008. Nest quality in relation to adult bird condition and its impact on reproduction in great tits *Parus major*. *Acta Ornithol.* 43:3–9.
- Anderson, M. G., C. Moskát, M. Bán, T. Grim, P. Cassey, and M. E. Hauber. 2009. Egg eviction imposes a recoverable cost of virulence in chicks of a brood parasite. *PLoS ONE* 4: e7725.
- Antonov, A. 2004. Smaller Eastern Olivaceous Warbler *Hippolais pallida elaeica* nests suffer less predation than larger ones. *Acta Ornithol.* 39:87–92.
- van Balen, J. H. 1984. The relationship between nest-box size, occupation and breeding parameters of the great tit *Parus major* and some other hole-nesting species. *Ardea* 72:163–175.
- van Balen, J. H., C. J. H. Booy, J. A. van Franeker, and E. R. Osieck. 1982. Studies on hole-nesting birds in natural nest sites. 1. Availability and occupation of natural nest sites. *Ardea* 70:1–24.
- Broggi, J., and J. C. Senar. 2009. Brighter great tit parents build bigger nests. *Ibis* 155:588–591.
- Carlson, A., U. Sandström, and K. Olsson. 1998. Availability and use of natural tree holes by cavity nesting birds in a Swedish deciduous forest. *Ardea* 86:109–119.
- Clark, L., and J. R. Mason. 1985. Use of nest material as insecticidal and anti-pathogenic agents by the European starling. *Oecologia* 67:169–176.
- Collias, N. E., and E. C. Collias. 1984. Nest building and bird behavior. Princeton Univ. Press, Princeton, NJ.
- Cramp, S., and C. M. Perrins, eds 1977–1994. The birds of the western Palearctic. Vol. 1–9. Oxford Univ. Press, Oxford, U.K.
- Crossman, C. A., V. G. Rohwer, and P. R. Martin. 2011. Variation in the structure of bird nests between northern Manitoba and southeastern Ontario. *PLoS ONE* 6:19086.
- Davies, N. B. 2000. Cuckoos, cowbirds and other cheats. T. & A. D. Poyser, London, U.K.
- Dunning, J. B. Jr. 1993. CRC handbook of avian body masses. CRC Press, Boca Raton, FL.
- Eeva, T., E. Lehtikoinen, and J. Nurmi. 1994. Effects of ectoparasites on breeding success of great tit (*Parus major*) and pied flycatcher (*Ficedula hypoleuca*) in an air pollution gradient. *Can. J. Zool.* 72:624–635.
- Erbeling-Denk, C., and F. Trillmich. 1990. Das Mikroklima im Nistkasten und seine Auswirkungen auf die Nestlinge beim Star. *J. Ornithol.* 131:73–84.
- Grim, T., J. Rutila, P. Cassey, and M. E. Hauber. 2009. The cost of virulence: an experimental study of egg eviction by brood parasitic chicks. *Behav. Ecol.* 20:1138–1146.
- Gustafsson, L., and S. G. Nilsson. 1985. Clutch size and breeding success of pied and collared flycatchers *Ficedula* spp. in nest boxes of different sizes. *Ibis* 127:380–385.
- Gwinner, H. 1997. The function of green plants in nests of European starlings (*Sturnus vulgaris*). *Behaviour* 134:337–351.
- Hansell, M. H. 2000. Bird nests and construction behaviour. Cambridge Univ. Press, Cambridge, U.K.
- Hansell, M. H. 2007. Built by animals. Oxford Univ. Press, Oxford, U.K.
- Heenan, C. B., and R. S. Seymour. 2011. Structural support, not insulation, is the primary driver for avian cup-shaped nest design. *Proc. R. Soc. Lond. B* 278:2924–2929.
- Karlsson, J., and S. G. Nilsson. 1977. The influence of nest-box area on clutch size in some hole-nesting passerines. *Ibis* 119:207–211.
- Kern, M. D., C. Van Riper III. 1984. Altitudinal variations in nests of the Hawaiian honeycreeper *Hemignathus virens virens*. *Condor* 86:443–454.
- Korpimäki, E. 1985. Clutch size and breeding success in relation to nest box size in Tengmalm's Owl *Aegolius funereus*. *Holarctic Ecol.* 8:175–180.
- Lack, D. 1968. Ecological adaptations to breeding in birds. Chapman and Hall, London, U.K.
- Lambrechts, M. M., F. Adriaensen, D. R. Ardia, et al. 2010. The design of artificial nest boxes for the study of secondary hole-nesting birds: a review of methodological inconsistencies and potential biases. *Acta Ornithol.* 45:1–26.
- Lambrechts, M. M., C. Aimé, A. Midamegbe, M.-J. Galan, P. Perret, A. Grégoire, et al. 2012. Nest size and breeding success in first and replacement clutches: an experimental study in blue tits *Cyanistes caeruleus*. *J. Ornithol.* 153:173–179.
- Lima, S. L. 2009. Predators and the breeding bird: behavioural and reproductive flexibility under the risk of predation. *Biol. Rev.* 84:485–513.
- Löhrl, H. 1973. Einfluß der Brutraumfläche auf die Gelegegröße der Kohlmeise (*Parus major*). *J. Ornithol.* 114:339–347.
- Löhrl, H. 1980. Weitere Versuche zur Frage "Brutraum und Gelegegröße" bei der Kohlmeise (*Parus major*). *J. Ornithol.* 121:403–405.
- Mainwaring, M. C., and I. R. Hartley. 2008. Seasonal adjustments in nest cup lining in blue tits *Cyanistes caeruleus*. *Ardea* 96:278–282.
- Mainwaring, M. C., and I. R. Hartley. 2013. The energetic costs of nest building in birds. *Avian Biol. Res.* 6:12–17.
- Mainwaring, M. C., I. R. Hartley, S. Bearhop, K. Brulez, C. R. du Feu, G. Murphy, et al. 2012. Latitudinal variation in blue tit and great tit nest characteristics indicates environmental adjustment. *J. Biogeogr.* 39:1669–1677.
- McClave, J. T., and T. Sincich. 2003. Statistics, 9th ed. Prentice-Hall, Englewood Cliffs, NJ.

- Mennerat, A., P. Mirleau, J. Blondel, P. Perret, M. Lambrecht, and P. Heeb. 2009. Aromatic plants in nests of the blue tit *Cyanistes caeruleus* protect chicks from bacteria. *Oecologia* 161:849–855.
- Mertens, J. A. L. 1977a. Thermal conditions for successful breeding in Great Tits (*Parus major* L.). 1. Relation of growth and development of temperature regulation in nestling great tits. *Oecologia* 28:1–29.
- Mertens, J. A. L. 1977b. Thermal conditions for successful breeding in great tits (*Parus major* L.). 2. Thermal properties of nests and nestboxes and their implications for range of temperature tolerance of great tit broods. *Oecologia* 28:31–56.
- Moksnes, A., and E. Røskaft. 1995. Egg-morphs and host preference in the common cuckoo *Cuculus canorus*: an analysis of cuckoo and host eggs from European museum collections. *J. Zool.* 236:625–648.
- Moksnes, A., E. Røskaft, A. T. Braa, L. Korsnes, H. M. Lampe, and H. C. Pedersen. 1991. Behavioural responses of potential hosts towards artificial cuckoo eggs and dummies. *Behaviour* 116:64–89.
- Møller, A. P. 1984. On the use of feathers in birds' nests: predictions and tests. *Ornis Scand.* 15:38–42.
- Møller, A. P. 1987. Nest lining in relation to the nesting cycle in the swallow *Hirundo rustica*. *Ornis Scand.* 18:148–149.
- Møller, A. P. 1989. Parasites, predators and nest boxes: facts and artefacts in nest box studies of birds? *Oikos* 56:421–423.
- Møller, A. P. 1990. Nest predation selects for small nest size in the blackbird. *Oikos* 57:237–240.
- Møller, A. P. 2006. Rapid change in nest size of a bird related to change in a secondary sexual character. *Behav. Ecol.* 17:108–116.
- Møller, A. P., E. Flensted-Jensen, W. Mardal, and J. J. Soler. 2013. Host-parasite relationship between colonial terns and bacteria is modified by a mutualism with a plant having antibacterial defenses. *Oecologia* 173:169–178.
- Møller, A. P., F. Adriaensen, A. Artemyev, J. Banbura, E. Barba, C. Biard, et al. 2014. Clutch size in European secondary hole-nesting passerines in relation to nest-box floor area, habitat, geographic location and study year. *Methods Ecol. Evol.* 5:353–362.
- Nager, R. G., and A. J. van Noordwijk. 1995. Proximate and ultimate aspects of phenotypic plasticity in timing of great tit breeding in a heterogeneous environment. *Am. Nat.* 146:454–474.
- de Neve, L., J. J. Soler, M. Soler, and T. Perez-Contreras. 2004. Nest size predicts the effect of food supplementation to magpie nestlings on their immunocompetence: an experimental test of nest size indicating parental ability. *Behav. Ecol.* 15:1031–1036.
- Peralta-Sánchez, J. M., A. P. Møller, A. M. Martín-Platero, and J. J. Soler. 2010. Number and colour composition of nest lining feathers predict eggshell bacterial community in barn swallow nests: an experimental study. *Funct. Ecol.* 24:426–433.
- Poole, A., ed. 2005. The Birds of North America Online: <http://bna.birds.cornell.edu.bnaproxy.birds.cornell.edu/BNA/>. Cornell Laboratory of Ornithology, Ithaca, NY.
- Rendell, W. B., and R. J. Robertson. 1993. Cavity size, clutch-size and the breeding ecology of Tree Swallows *Tachycineta bicolor*. *Ibis* 135:305–310.
- Sánchez, S., J. J. Cuervo, and E. Moreno. 2007. Suitable cavities as a scarce resource for both cavity and noncavity nesting birds in managed temperate forests. A case study in the Iberian Peninsula. *Ardeola* 54:261–274.
- Sandström, U. 1992. Cavities in trees: their occurrence, formation and importance for hole-nesting birds in relation to silvicultural practise. Licentiate dissertation, Uppsala Univ., Sweden.
- Sanz, J. J., and V. García-Navas. 2011. Nest ornamentation in blue tits: is feather carrying ability a male status signal? *Behav. Ecol.* 22:240–247.
- SAS. 2012. JMP version 10.0. SAS Institute Inc., Cary, NC.
- Schaefer, V. H. 1980. Geographic variation in the insulative qualities of nests of the northern oriole. *Wilson Bull.* 92:466–474.
- Slagsvold, T. 1982. Clutch size, nest size, and hatching asynchrony in birds: experiments with the fieldfare (*Turdus pilaris*). *Ecology* 63:1389–1399.
- Slagsvold, T. 1987. Nest site preference and clutch size in the Pied Flycatcher *Ficedula hypoleuca*. *Ornis Scand.* 18:189–197.
- Slagsvold, T. 1989. Experiments on clutch size and nest size in passerine birds. *Oecologia* 80:297–302.
- Slagsvold, T., and T. Amundsen. 1992. Do great tits adjust hatching spread, egg size and offspring sex ratio to changes in clutch size? *J. Anim. Ecol.* 61:249–258.
- Slagsvold, T., and S. Dale. 1996. Disappearance of female pied flycatchers in relation to breeding stage and experimentally induced molt. *Ecology* 77:461–471.
- Soler, J. J., M. Soler, J. G. Martínez, and A. P. Møller. 1995. Does the great spotted cuckoo choose magpie hosts according to their parenting ability? *Behav. Ecol. Sociobiol.* 36:201–206.
- Soler, J. J., A. P. Møller, and M. Soler. 1998a. Nest building, sexual selection and parental investment. *Evol. Ecol.* 12:427–441.
- Soler, J. J., J. J. Cuervo, A. P. Møller, and F. de Lope. 1998b. Nest building is a sexually selected behaviour in the barn swallow. *Anim. Behav.* 56:1435–1442.
- Soler, J. J., J. G. Martínez, M. Soler, and A. P. Møller. 1999. Host sexual selection and cuckoo parasitism: an analysis of nest size in sympatric and allopatric magpie *Pica pica* populations parasitised by the great spotted cuckoo *Clamator glandarius*. *Proc. R. Soc. Lond. B* 266:1765–1771.
- Soler, J. J., L. de Neve, J. G. Martínez, and M. Soler. 2001. Nest size affects clutch size and the start of incubation in magpies: an experimental study. *Behav. Ecol.* 12:301–307.

- Soler, J. J., J. M. Peralta-Sánchez, A. M. Martín Platero, M. Martín-Vivaldi, M. Martínez-Bueno, and A. P. Møller. 2012. The evolution of size of the uropygial gland: mutualistic feather mites and uropygial secretion reduce bacterial loads of eggshells and hatching failures of European birds. *J. Evol. Biol.* 25:1779–1791.
- Stolp, H. 1988. *Microbial ecology: organisms, habitats, activities.* Cambridge Univ. Press, Cambridge, U.K.
- Taylor, M. I., G. F. Turner, R. L. Robinson, and J. R. Stauffer. 1998. Sexual selection, parasites and bower height in a bower-building cichlid fish. *Animal Behaviour* 56:379–384.
- Tomás, G., S. Merino, J. Martínez-de la Puente, J. Moreno, J. Morales, and J. Rivero-de-Aguilar. 2013. Nest size and aromatic plants in the nest as sexually selected female traits in blue tits. *Behav. Ecol.* 24:926–934.
- Tortosa, F. S., and T. Redondo. 1992. Frequent copulations despite low sperm competition in white storks (*Ciconia ciconia*). *Behaviour* 121:288–315.
- Wiebe, K. L., and T. L. Swift. 2001. Clutch size relative to tree cavity size in Northern Flickers. *J. Avian Biol.* 32:167–173.
- Wimberger, P. H. 1984. The use of green plant material in bird nests to avoid ectoparasites. *Auk* 101:615–618.

Supporting Information

Additional Supporting Information may be found in the online version of this article:

Table S1. Summary information on studies of the relationship between clutch size and nest size in different species of birds.