

## Spring migration of birds in relation to North Atlantic Oscillation

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**A b s t r a c t.** Long-term spring phenological instants of 57 migratory bird species, i.e. arrival in summer visitors and departure in winter visitors, were recorded in South Moravia (Czech Republic) from 1952 through 2001 and evaluated for annual correspondence with the North Atlantic Oscillation (NAO) weather system. The migration instants occurred significantly earlier following positive winter/spring NAO index values (causing periods warmer than normal in Europe) in a number of short-distance migrants with a European winter range (e.g., *Alauda arvensis*, *Columba palumbus*, *Corvus frugilegus*, *Motacilla alba*, *Phoenicurus ochruros*, *Phylloscopus collybita*, *Serinus serinus*, *Sturnus vulgaris*, *Vanellus vanellus*), whereas they did not correlate with NAO in most long-distance migrants having a sub-Saharan winter range (e.g., *Acrocephalus* spp., *Anthus trivialis*, *Apus apus*, *Cuculus canorus*, *Delichon urbica*, *Ficedula albicollis*, *Hippolais icterina*, *Hirundo rustica*, *Jynx torquilla*, *Lanius collurio*, *Locustella* spp., *Muscicapa striata*, *Oriolus oriolus*, *Phylloscopus sibilatrix*, *Riparia riparia*, *Streptopelia turtur*, *Sylvia* spp.). The winter/spring (especially February and March) NAO conditions thus affect the migration timing of short-distance migrants that winter in western or southern Europe, and could explain their earlier than normal arrival that had been observed in Europe since the 1980s.

**Key words:** phenology, migration, weather, climate, temperature, NAO

### Introduction

Recent climate warming is often regarded as the cause of the advanced spring arrival and egg-laying dates of migratory birds that has been recorded in Europe and North America since the 1980's (e.g., Forchhammer et al. 1998, Winkler et al. 1999, Both & Visser 2001, Sokolov 2001, Zalakevicius & Zalakeviciute 2001). An approach to test this hypothesis could be to evaluate whether the bird arrival data correlate with some global measure of the weather/climate change. It has been demonstrated that El Niño/Southern Oscillation (ENSO) system of ocean and atmosphere circulation in the tropical Pacific affects the biota in the Americas, Australia, southern Asia and eastern Africa either directly, through extreme precipitation or temperature, or indirectly, through ecosystem changes (Jacobs et al. 1994, Karl et al. 1995, Silllett et al. 2000, Nott et al. 2002, Stenseth et al. 2002). Another major mode of the world climate variability is the North Atlantic Oscillation (NAO) that profoundly affects the weather in continental Europe and eastern North America (Wallace & Gutzler 1981, Barnston & Livezey 1987, Hurrell 1995, Yoo & D'Odorico 2002). The NAO system can be expressed quantitatively using simple monthly, seasonal or annual indices that make a correlation analysis possible and relatively easy. A number of papers have reported a correspondence between the NAO fluctuation and plant phenology (Post & Stenseth 1999, Chmielewski & Rotzer 2001, Post et al. 2001), cephalopod migration (Simms et al. 2001), marine fish migration (Alheit & Hagen 1997, Reid et al. 2001),

mammalian population dynamics (Post & Stenseth 1999, Tkadlec 2000, Post & Forchhammer 2002, Stenseth et al. 2002) or the avian breeding phenology and productivity (Forchhammer et al. 1998, Przybylo et al. 2000, Saether et al. 2000, Both & Visser 2001, Sokolov 2001, Møller 2002, Nott et al. 2002, Sanz 2002). However, the effect of NAO on the variability of bird migration instants in Europe has not been studied until very recently (Forchhammer et al. 2002, Jonzén et al. 2002, Tryjanowski et al. 2002, Huppopp & Huppopp 2003).

## Material and Methods

### Study area

Observations of birds were carried out in the Břeclav area (48°40'–48°50'N, 16°30'–17°00'E) of South Moravia, Czech Republic. The relief is a mostly flat to slightly undulating lowland (150 to 200 m a.s.l.), formed largely by accumulations of Pleistocene loess and riverine sand deposits on Pliocene sedimentaries. The climate is relatively dry and warm (annual precipitation 550 mm; annual mean air temperature 9.5 °C (January -1.8 °C, July 19.1 °C). There are agroecosystems (mostly arable fields), forest ecosystems (floodplain forests, subxerophilic mixed deciduous oak stands), fishponds and big water reservoirs (Nové Mlýny) in the wider area. Local community is composed of about 150 species of breeding terrestrial and water birds (Hájek 1992, Štastný et al. 1996, Hubálek 1997, Zuna-Kratky et al. 2000), and many migrants use the area for stopovers.

### Bird phenology data

Longitudinal records of migratory birds were documented from 1969 through 2001. Spring phenological instants recorded for each species involved the first occurrence (spring arrival) in summer visitors, and the last occurrence (spring departure) in winter visitors. In addition to the author's observations, data of Hájek (1992) were used for the years 1952 to 1969. Total length of the period thus covered 50 years (1952–2001), and 57 common bird species with a distinct pattern of spring phenology and a sufficient number of annual records were selected for the correlation and regression analysis.

### North Atlantic Oscillation data

Monthly (January; February; March; April) and seasonal (December to March, DJFM; January to March, JFM; February to April, FMA) NAO indices for particular years 1952 through 2001 were extracted from the Internet at <http://www.cgd.ucar.edu/~jhurrell/nao.html> (Hurrell 1995: NAO<sub>H</sub>) and <http://www.cpc.ncep.noaa.gov/data/teledoc/nao.html> (Barnston & Livezey 1987: NAO<sub>BL</sub>; Wallace & Gutzler 1981: NAO<sub>EA</sub>, the East Atlantic Pattern). The indices are based on normalized sea-level pressure differences between two points or areas: (1) Ponta Delgada (Azores) and Stykkisholmur/Reykjavik (Iceland) in the eastern sector of North Atlantic (NAO<sub>H</sub>); (2) the central latitudes (35°N–40°N) and Greenland (NAO<sub>BL</sub>, reflecting thus spatially a more broad-scale atmospheric pressure pattern in the western North Atlantic sector); and (3) NAO<sub>EA</sub> covers the eastern North Atlantic sector. The positive NAO index generally shows that the atmospheric pressure over the subtropical part of the North Atlantic is higher than normal while that over

the northern sector of the North Atlantic is lower than normal; this increased pressure difference between the two sectors results in more and stronger storms crossing the Atlantic Ocean and, in turn, causing warm and wet weather (especially in winter) in northern and central Europe. The negative NAO index reflects an opposite pattern of height and pressure anomalies over these sectors; this reduced pressure gradient results in fewer and weaker storms crossing the Ocean, bringing cold air to northern (and central) Europe and moist, often cold air into the Mediterranean (Hurrell 1995).

## Statistical analysis

Calendar data of phenological instants were transformed into sequential numbers (e.g., 1 for 1<sup>st</sup> January, 32 for 1<sup>st</sup> February, 60 for 1<sup>st</sup> March, 91 for 1<sup>st</sup> April, 121 for 1<sup>st</sup> May 1; in leap-years, the sequential numbers were corrected by adding 1, starting from 1<sup>st</sup> March). Linear correlation and regression models were then used to examine relationships between avian spring phenological instants of particular bird species and the three NAO indices (NAO<sub>H</sub>, NAO<sub>BL</sub>, NAO<sub>EA</sub>); Pearson's correlation coefficient values were calculated and statistical test done for all comparisons using SOLO 4.0 (BMDP Statistical Software, Los Angeles, CA).

## Results

The late-winter and early-spring NAO weather system conditions affected significantly the spring arrival (in summer visitors) or departure (in winter visitors) instants of some bird species (Tables 1 and 2, Fig. 1). The species significantly inversely correlated with both the 'JFM' and 'FMA' seasonal NAO indices (i.e., the phenological instants occurred earlier following the January-February-March and February-March-April seasonal air pressure difference in the North Atlantic higher than normal) were *Alauda arvensis*, *Anas crecca*, *Aythya ferina*, *Bucephala clangula*, *Columba palumbus*, *Corvus frugilegus*, *Gallinago gallinago*, *Mergus merganser*, *Motacilla alba*, *Phoenicurus ochruros*, *Phylloscopus collybita*, *P. trochilus*, *Pyrrhula pyrrhula*, *Regulus regulus*, *Remiz pendulinus*, *Saxicola torquata*, *Serinus serinus*, *Sterna hirundo*, *Sturnus vulgaris*, *Sylvia atricapilla*, *Tringa glareola*, *Turdus philomelos* and *Vanellus vanellus*. In some species, there was a single statistically significant seasonal (either 'JFM' or 'FMA') NAO index (*Acrocephalus scirpaceus*, *Anas querquedula*, *Charadrius dubius*, *Hirundo rustica*, *Luscinia megarhynchos*, *Phylloscopus sibilatrix*, *Streptopelia turtur*, *Sylvia communis*, *Tringa nebularia*, *T. totanus*), whereas the remaining species did not correlate with NAO at  $p < 0.05$ : *Acrocephalus arundinaceus*, *A. palustris*, *A. schoenobaenus*, *Actitis hypoleucos*, *Anthus trivialis*, *Apus apus*, *Cuculus canorus*, *Delichon urbica*, *Ficedula albicollis*, *Hippolais icterina*, *Jynx torquilla*, *Lanius collurio*, *Locustella fluviatilis*, *L. luscinoides*, *L. naevia*, *Motacilla flava*, *Muscicapa striata*, *Nycticorax nycticorax*, *Oriolus oriolus*, *Riparia riparia*, *Sylvia borin*, *S. curruca*, *S. nisoria* and *Upupa epops*.

The winter NAO<sub>H</sub> index (DJFM data, not given in Table 1) correlated significantly ( $p < 0.01$ ) with phenological instants of *Columba palumbus* (-0.45), *Mergus merganser* (-0.68), *Phylloscopus collybita* (-0.49), *Sturnus vulgaris* (-0.44), *Tringa nebularia* (-0.51) and *Vanellus vanellus* (-0.40). According to monthly NAO indices (data not given in Table 1), the number of avian species significantly ( $p < 0.05$ ) correlated in their phenology with NAO<sub>H</sub>/NAO<sub>EA</sub> during January, February, March and April were 8/8, 12/12, 9/13 and 3/3, respectively. Therefore, the NAO conditions especially in February and March have

**Table 1.** Spring phenology instants of 57 migratory bird species in South Moravia, 1952–2001, with the number of years (n), the mean date of arrival or departure (mm/dd), principal wintering area (W), and Pearson's correlation coefficient value (*r*) between the avian phenology data and the mean seasonal indices (JFM, January to March; FMA, February to April) of NAO<sub>H</sub> (H), NAO<sub>BL</sub> (BL) and NAO<sub>EA</sub> (EA); *r* values in bold are significant (*p*<0.05). The bird species are arranged according to the mean arrival/departure date.

Bird species	n	Mean date	W	JFM			FMA		
				H	BL	EA	H	BL	EA
<i>Alauda arvensis</i> <sup>A</sup>	49	02/28	E	-0.16	-0.09	<b>-0.47</b>	<b>-0.35</b>	-0.19	<b>-0.31</b>
<i>Sturnus vulgaris</i> <sup>A</sup>	50	03/01	E	<b>-0.51</b>	<b>-0.44</b>	<b>-0.43</b>	<b>-0.53</b>	<b>-0.45</b>	<b>-0.28</b>
<i>Mergus merganser</i> <sup>D</sup>	18	03/05	E	<b>-0.61</b>	<b>-0.60</b>	<b>-0.47</b>	<b>-0.59</b>	<b>-0.54</b>	-0.33
<i>Turdus philomelos</i> <sup>A</sup>	39	03/08	E	+0.06	+0.25	<b>-0.33</b>	-0.06	+0.15	<b>-0.44</b>
<i>Corvus frugilegus</i> <sup>D1/2</sup>	35	03/09	E	<b>-0.52</b>	<b>-0.41</b>	<b>-0.62</b>	<b>-0.55</b>	<b>-0.38</b>	<b>-0.54</b>
<i>Motacilla alba</i> <sup>A</sup>	47	03/10	E	-0.11	-0.08	<b>-0.42</b>	<b>-0.31</b>	-0.22	<b>-0.45</b>
<i>Vanellus vanellus</i> <sup>A</sup>	47	03/11	E	<b>-0.34</b>	-0.28	<b>-0.39</b>	<b>-0.42</b>	-0.27	<b>-0.29</b>
<i>Aythya ferina</i> <sup>A</sup>	24	03/12	E	<b>-0.46</b>	<b>-0.45</b>	<b>-0.50</b>	<b>-0.42</b>	-0.40	<b>-0.42</b>
<i>Anas crecca</i> <sup>A</sup>	23	03/15	E	<b>-0.42</b>	<b>-0.51</b>	<b>-0.52</b>	<b>-0.42</b>	<b>-0.51</b>	-0.34
<i>Pyrrhula pyrrhula</i> <sup>D</sup>	27	03/19	E	-0.14	-0.15	<b>-0.39</b>	-0.18	-0.27	<b>-0.52</b>
<i>Bucephala clangula</i> <sup>D</sup>	30	03/20	E	-0.10	-0.20	<b>-0.52</b>	-0.26	-0.21	<b>-0.57</b>
<i>Columba palumbus</i> <sup>A</sup>	38	03/23	E	<b>-0.55</b>	<b>-0.55</b>	<b>-0.38</b>	<b>-0.58</b>	<b>-0.56</b>	-0.19
<i>Phylloscopus collybita</i> <sup>A</sup>	36	03/23	E	<b>-0.60</b>	<b>-0.52</b>	<b>-0.40</b>	<b>-0.54</b>	<b>-0.44</b>	-0.20
<i>Phoenicurus ochrurus</i> <sup>A</sup>	38	03/24	E	<b>-0.38</b>	<b>-0.33</b>	<b>-0.54</b>	<b>-0.34</b>	<b>-0.32</b>	<b>-0.42</b>
<i>Saxicola torquata</i> <sup>A</sup>	35	03/24	E	<b>-0.46</b>	<b>-0.40</b>	-0.02	<b>-0.47</b>	<b>-0.46</b>	+0.14
<i>Regulus regulus</i> <sup>D</sup>	23	03/26	E	<b>-0.44</b>	<b>-0.55</b>	-0.26	<b>-0.49</b>	<b>-0.62</b>	-0.21
<i>Anas querquedula</i> <sup>A</sup>	29	03/26	A	-0.28	-0.30	<b>-0.36</b>	-0.19	-0.22	-0.23
<i>Gallinago gallinago</i> <sup>A</sup>	26	03/26	E	<b>-0.56</b>	<b>-0.48</b>	-0.29	<b>-0.43</b>	-0.30	+0.04
<i>Serinus serinus</i> <sup>A</sup>	38	03/30	E	<b>-0.44</b>	<b>-0.36</b>	-0.23	<b>-0.34</b>	-0.29	-0.06
<i>Remiz pendulinus</i> <sup>A</sup>	37	03/30	E	<b>-0.38</b>	-0.26	<b>-0.58</b>	<b>-0.49</b>	<b>-0.39</b>	<b>-0.41</b>
<i>Tringa totanus</i> <sup>A</sup>	27	04/01	E	<b>-0.46</b>	<b>-0.39</b>	-0.33	<b>-0.24</b>	-0.27	-0.13
<i>Sylvia atricapilla</i> <sup>A</sup>	33	04/03	E	-0.12	-0.15	<b>-0.53</b>	-0.10	-0.24	<b>-0.56</b>
<i>Phylloscopus trochilus</i> <sup>A</sup>	37	04/06	A	-0.18	-0.19	<b>-0.44</b>	-0.20	-0.25	<b>-0.42</b>
<i>Hirundo rustica</i> <sup>A</sup>	46	04/09	A	-0.17	-0.26	-0.03	-0.20	<b>-0.34</b>	-0.01
<i>Charadrius dubius</i> <sup>A</sup>	33	04/10	E	<b>-0.48</b>	<b>-0.38</b>	-0.07	-0.15	-0.25	+0.11
<i>Anthus trivialis</i> <sup>A</sup>	45	04/12	A	+0.18	+0.08	+0.13	+0.11	+0.07	-0.11
<i>Motacilla flava</i> <sup>A</sup>	35	04/12	A	-0.30	-0.15	-0.12	-0.19	-0.24	+0.13
<i>Jynx torquilla</i> <sup>A</sup>	42	04/13	A	-0.08	-0.16	-0.12	-0.15	-0.07	-0.29
<i>Nycticorax nycticorax</i> <sup>A</sup>	18	04/13	A	-0.34	-0.23	-0.22	-0.21	-0.21	-0.15
<i>Upupa epops</i> <sup>A</sup>	38	04/14	A	-0.21	-0.12	-0.31	-0.13	-0.14	-0.27
<i>Sylvia curruca</i> <sup>A</sup>	44	04/15	A	-0.12	-0.03	-0.14	-0.05	+0.09	-0.03
<i>Ficedula albicollis</i> <sup>A</sup>	45	04/16	A	+0.10	+0.17	-0.03	+0.15	+0.13	-0.08
<i>Acrocephalus schoenobaenus</i> <sup>A</sup>	32	04/19	A	-0.26	-0.13	-0.07	-0.17	-0.25	+0.03
<i>Actitis hypoleucos</i> <sup>A</sup>	41	04/20	A	-0.00	-0.02	-0.13	-0.08	-0.17	-0.26
<i>Delichon urbica</i> <sup>A</sup>	40	04/21	A	+0.24	+0.23	+0.06	+0.29	+0.13	+0.17
<i>Phylloscopus sibilatrix</i> <sup>A</sup>	30	04/21	A	+0.00	-0.06	<b>-0.36</b>	+0.14	+0.01	<b>-0.41</b>
<i>Cuculus canorus</i> <sup>A</sup>	50	04/22	A	-0.13	-0.21	-0.14	-0.09	-0.23	-0.10
<i>Luscinia megarhynchos</i> <sup>A</sup>	39	04/23	A	<b>-0.36</b>	-0.31	+0.02	-0.26	-0.23	+0.24
<i>Tringa nebularia</i> <sup>A</sup>	29	04/23	E	<b>-0.45</b>	<b>-0.42</b>	<b>-0.56</b>	-0.27	-0.30	-0.23
<i>Locustella luscinioides</i> <sup>A</sup>	28	04/24	A	-0.16	-0.23	-0.06	-0.30	-0.37	-0.14
<i>Streptopelia turtur</i> <sup>A</sup>	34	04/24	A	-0.22	-0.25	<b>-0.39</b>	-0.08	-0.18	-0.23
<i>Sylvia communis</i> <sup>A</sup>	44	04/24	A	<b>-0.33</b>	<b>-0.33</b>	-0.12	-0.26	-0.21	-0.04
<i>Riparia riparia</i> <sup>A</sup>	23	04/25	A	-0.13	+0.13	-0.28	+0.08	+0.12	-0.35
<i>Tringa glareola</i> <sup>A</sup>	31	04/26	E	<b>-0.39</b>	-0.33	-0.21	<b>-0.45</b>	-0.33	+0.03
<i>Sterna hirundo</i> <sup>A</sup>	29	04/27	E	<b>-0.43</b>	<b>-0.52</b>	-0.01	<b>-0.53</b>	<b>-0.36</b>	+0.01

**Table 1** – continued.

<i>Sylvia borin</i> <sup>^</sup>	37	04/29	A	-0.26	-0.29	-0.11	-0.19	-0.20	-0.04
<i>Apus apus</i> <sup>^</sup>	37	04/29	A	-0.17	-0.20	-0.25	-0.12	-0.01	-0.27
<i>Acrocephalus arundinaceus</i> <sup>^</sup>	31	04/30	A	+0.24	+0.20	-0.02	+0.10	+0.03	-0.12
<i>Acrocephalus scirpaceus</i> <sup>^</sup>	22	05/02	A	-0.34	-0.44	-0.20	<b>-0.45</b>	<b>-0.45</b>	-0.06
<i>Locustella naevia</i> <sup>^</sup>	26	05/04	A	-0.12	-0.21	-0.03	+0.03	-0.04	+0.01
<i>Oriolus oriolus</i> <sup>^</sup>	42	05/05	A	+0.12	+0.15	-0.08	+0.13	+0.17	-0.15
<i>Hippolais icterina</i> <sup>^</sup>	44	05/07	A	-0.04	-0.07	+0.15	+0.07	+0.08	+0.14
<i>Locustella fluviatilis</i> <sup>^</sup>	29	05/07	A	+0.23	+0.26	-0.03	+0.16	+0.25	-0.13
<i>Muscicapa striata</i> <sup>^</sup>	33	05/07	A	+0.08	-0.07	-0.03	+0.06	-0.02	-0.12
<i>Sylvia nisoria</i> <sup>^</sup>	41	05/07	A	-0.05	+0.05	-0.12	-0.04	-0.02	-0.10
<i>Lanius collurio</i> <sup>^</sup>	31	05/09	A	+0.01	-0.16	-0.06	-0.15	-0.22	-0.10
<i>Acrocephalus palustris</i> <sup>^</sup>	24	05/10	A	+0.05	-0.10	-0.03	-0.38	-0.38	-0.24

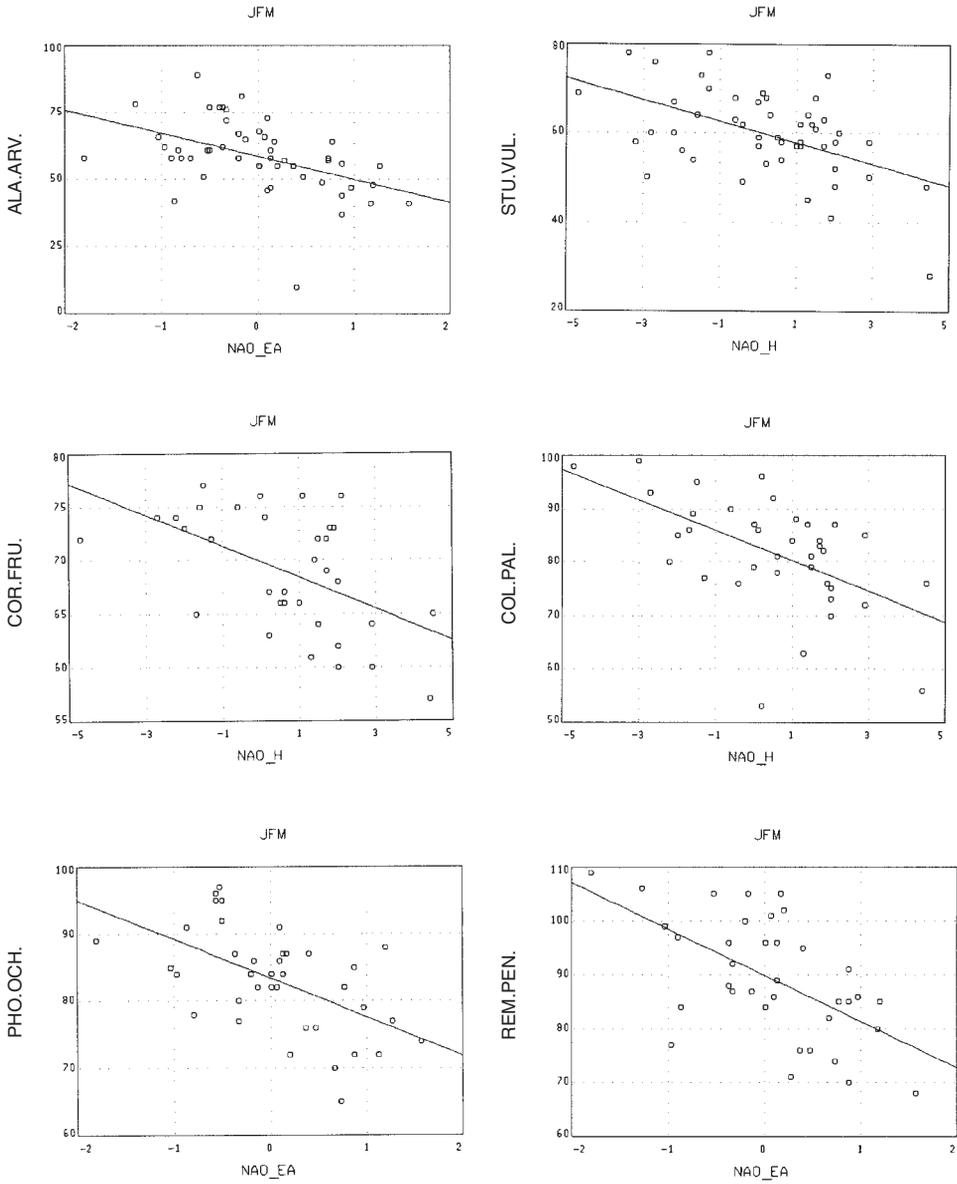
<sup>^</sup> arrival of the first bird; <sup>D</sup> departure (the last observed bird in the spring); <sup>D1/2</sup> departure of about half of the wintering population. Principal wintering area (W): E, southern, western or central Europe, Mediterranean; A, sub-Saharan Africa (or southwestern Asia).

**Table 2.** Regression ( $y = a + b.x$ ) of 18 selected bird species on seasonal NAO index (see Fig. 1).

Bird species	W	NAO index		Regression slope ( <i>b</i> )	Intercept ( <i>a</i> )	R <sup>2</sup>
		Type	Season			
<i>Alauda arvensis</i>	E	EA	JFM	-8.5757***	58.71	0.216
<i>Sturnus vulgaris</i>	E	H	JFM	-2.4273***	60.26	0.261
<i>Corvus frugilegus</i>	E	H	JFM	-1.4594**	69.87	0.275
<i>Columba palumbus</i>	E	H	JFM	-2.8513***	83.05	0.308
<i>Phoenicurus ochruros</i>	E	EA	JFM	-5.7958***	83.44	0.306
<i>Remiz pendulinus</i>	E	EA	JFM	-8.5900***	89.97	0.338
<i>Phylloscopus collybita</i>	E	H	JFM	-2.2751***	84.23	0.356
<i>Phylloscopus sibilatrix</i>	A	H	JFM	+0.0008	111.31	0.000
<i>Sylvia atricapilla</i>	E	EA	JFM	-4.9794**	93.40	0.282
<i>Sylvia curruca</i>	A	EA	JFM	-1.1621	105.36	0.018
<i>Charadrius dubius</i>	E	H	JFM	-2.4511**	100.26	0.231
<i>Actitis hypoleucos</i>	A	H	JFM	-0.0652	110.08	0.000
<i>Hirundo rustica</i>	A	EA	JFM	-0.7587	99.47	0.006
<i>Delichon urbica</i>	A	EA	JFM	+0.2758	111.03	0.001
<i>Ficedula albicollis</i>	A	H	JFM	+0.2707	105.87	0.011
<i>Muscicapa striata</i>	A	H	JFM	+0.1748	127.38	0.006
<i>Hippolais icterina</i>	A	H	FMA	+0.2002	127.19	0.004
<i>Cuculus canorus</i>	A	H	FMA	-0.2552	112.19	0.009

W, principal wintering area. NAO index type: EA, NAO<sub>EA</sub>; H, NAO<sub>H</sub> ). Other abbreviations as for Table 1. \*\*, significant at  $p < 0.01$ ; \*\*\*, significant at  $p < 0.001$ .

a profound effect on many early-spring migrants that usually winter in western or southern Europe (Mediterranean). On the other hand, NAO does not seem to affect significantly the timing of long-distance migrants arriving in Central Europe later, having wintering grounds largely in sub-Saharan Africa. The seasonal (JFM) NAO<sub>H</sub> index explained on the average 18% (range 1–37%) variability in the phenological instants of the 25 short-distance migrants, whereas only 4% (range, 0–13%) variability in the 32 long-distance migrants; the difference between the two categories of migratory birds is highly significant ( $p < 0.001$ ) as found with either t-test or nonparametric Mann-Whitney two-sample test.



**Fig. 1.** Scatter plot diagrams of relations between seasonal (JFM or FMA) NAO indices ( $NAO_H$  or  $NAO_{EA}$ ) and spring migration instants in 18 selected bird species (acronyms) in South Moravia, 1952–2001. Y-axis shows calendar data of the phenological instants transformed into sequential numbers. For the regression analysis, see Table 2.

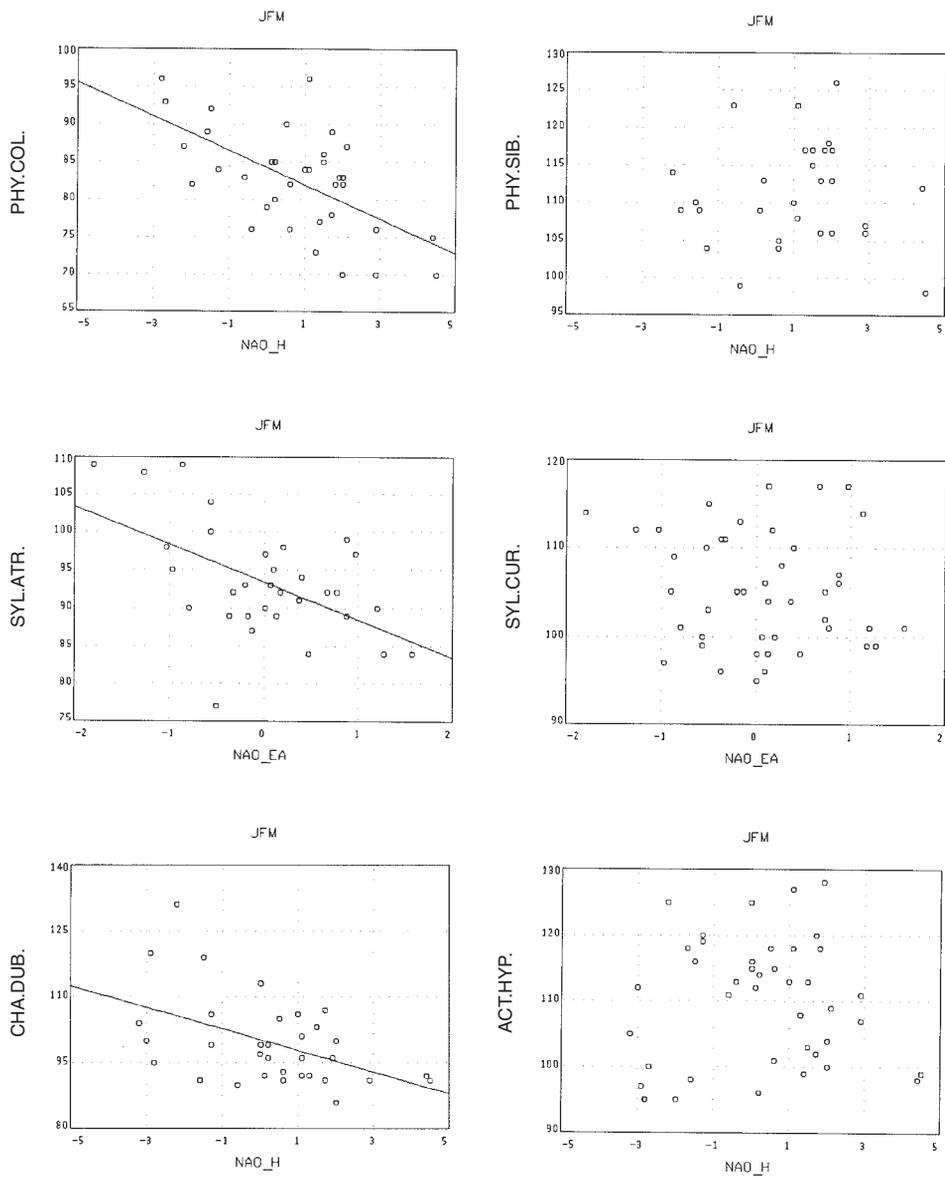


Fig. 1 – continued.

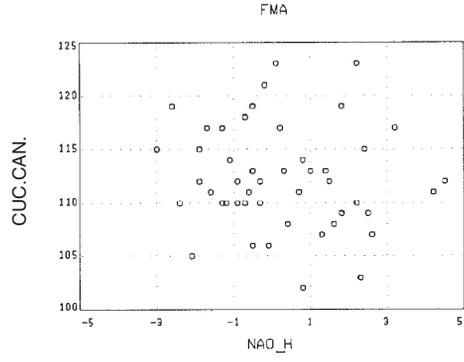
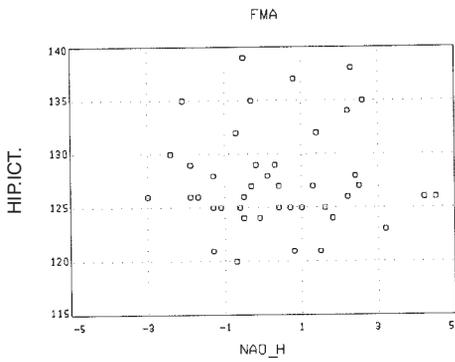
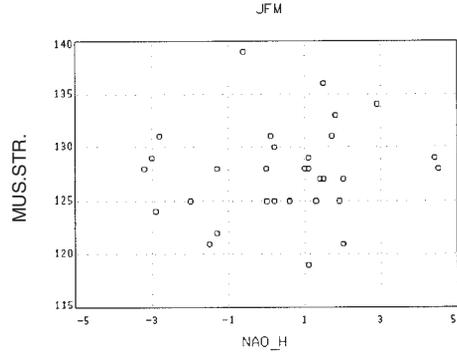
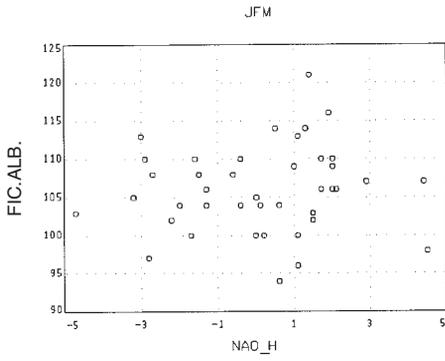
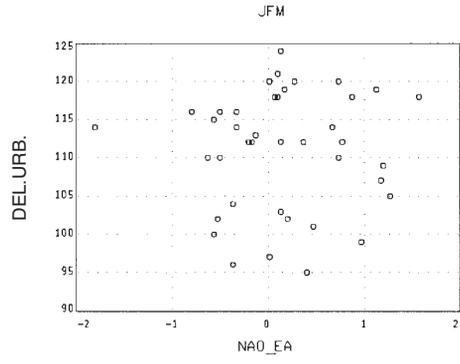
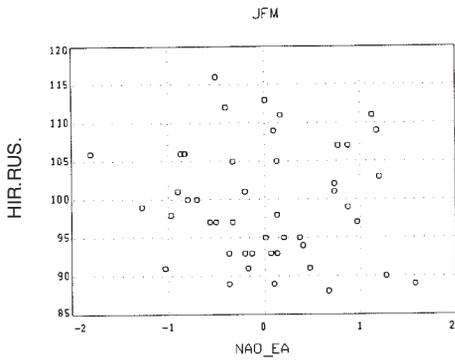


Fig. 1 – continued.

## Discussion

Tryjanowski & Sparks (2001) suggested that the relationship between arrival date and population size should be considered in particular bird species: an increasing population of *Lanius collurio* showed an advanced arrival in western Poland during 1983–2000. In the present study, usually common and abundant avian species were taken into account whose population sizes did not change considerably with time. The exceptions with the population steadily low (l), on decline (d) or increasing (i) were *Acrocephalus* spp. (d), *Anas* spp. (d), *Ficedula albicollis* (i), *Gallinago gallinago* (d), *Lanius collurio* (i), *Motacilla flava* (d), *Nycticorax nycticorax* (l), *Sylvia nisoria* (l), *Tringa nebularia* (l), *T. totanus* (d), *Upupa epops* (d) and *Vanellus vanellus* (d) (Hájek 1992, Štátný et al. 1996, Hubálek 1997, etc.). However, the results yielded in these species do not seem to contradict the general conclusion that long-distance migrants are not affected significantly with the NAO activity, in contrast to short-distance migratory species.

The positive NAO index values mean generally a stronger than usual subtropical high pressure center and a deeper than usual subarctic atmospheric low. The increased pressure difference causes intense atmospheric mass circulation resulting in more winter storms crossing the Atlantic Ocean on a more northerly track, a milder winter and the whole year in Europe, and specifically a warmer (but drier) than normal weather over central Europe (Hurrell 1995). For the Czech Republic, a significant correlation was found between the winter (DJFM) NAO<sub>H</sub> index and local winter air temperature ( $r = 0.78$ ), but that between this index and winter precipitation ( $r = -0.30$ ) was insignificant (Tkadlec 2000). The milder temperatures associated with the positive NAO phase lead to higher invertebrate loads in spring which might benefit short-distance migrants more than long-distance migratory species (Nott et al. 2002).

Three indices that characterize NAO were used in this study. Although NAO<sub>H</sub> and NAO<sub>BL</sub> correlated well ( $r$  value for JFM season was 0.87), NAO<sub>BL</sub> seemed to be less sensitive in the phenology of European migratory birds (in terms of the number of significant correlations with avian species) than NAO<sub>H</sub>, probably because it describes weather conditions in the western sector of the North Atlantic and may thus be more relevant for the North American continent than for Europe. The eastern Atlantic measures NAO<sub>H</sub> and NAO<sub>EA</sub> were comparable in their phenological sensitivity although they did not correlate well ( $r$  value for JFM season was very low, 0.17): in contrast to NAO<sub>H</sub>, NAO<sub>EA</sub> revealed a marked correspondence between NAO activity and phenological instants in *Alauda arvensis*, *Bucephala clangula*, *Motacilla alba*, *Pyrrhula pyrrhula*, *Sylvia atricapilla* and *Turdus philomelos*, whereas NAO<sub>H</sub>, contrary to NAO<sub>EA</sub>, yielded significant correlations with phenological data of *Charadrius dubius*, *Gallinago gallinago*, *Larus ridibundus*, *Regulus regulus*, *Saxicola torquata*, *Serinus serinus*, *Sterna hirundo*, *Tringa totanus* and *T. glareola*. At present, it is difficult to evaluate which of the two indices is better for the use in the European phenology; they might be used as complementary measures.

In an earlier study, a cluster analysis of temporal spring migration patterns of birds in Moravia between 1881 and 1960 revealed several groups (called ‘migrans’) of co-related avian species: the first (‘Mediterranean’) migran consisted almost exclusively of short-distance migrants wintering in southern or western Europe (*Alauda arvensis*, *Sturnus vulgaris*, *Fringilla coelebs*, *Columba palumbus*, *Motacilla alba*, *Vanellus vanellus*, *Turdus philomelos*, *Erithacus rubecula*, *Scolopax rusticola*), whereas the remaining migrans had been called ‘African’ in that they involved long-distance migrants *Hirundo rustica*, *Jynx*

*torquilla*, *Delichon urbica*, *Cuculus canorus*, *Luscinia megarhynchos*, *Streptopelia turtur*, *Apus apus*, *Oriolus oriolus* and *Coturnix coturnix*, i.e. species having their winter quarters in sub-Saharan Africa (H u b á l e k 1985). In the present study, the seasonal winter/spring NAO index did not correlate significantly with the arrival of a vast majority of long-distance migratory bird species wintering in tropical and southern Africa (only 2 of 32 those species tested revealed significant both 'JFM' and 'FMA' seasonal measures). Timing of their departure from the winter grounds must therefore be based on principles other than the weather system fluctuation at northern latitudes. However, significant inverse relationship was found between the arrival of nearly all (22) short-distance migrants wintering in western and southern Europe (25 species tested) and the seasonal winter/spring NAO index, indicating that a higher than normal air pressure difference over the North Atlantic during the winter/spring (especially in February and March) determines an earlier than normal arrival of these birds in Central Europe. This result is not in accord with a very recently published study of F o r c h h a m m e r et al. (2002), who found no significant difference in the effect of NAO on spring arrival between long-distance and short-distance migrants (but only three species of each group were tested) breeding in Norway, although it corresponds well with the results of N o t t et al. (2002).

Positive values of the winter NAO index prevailed in 11 out of 15 years between 1987 and 2001 (H u r r e l l 1995), as did the bird phenological data in South Moravia during the same period: earlier than normal early-spring bird instants occurred in 12 out of 15 years, whereas those markedly later than normal in one year only (H u b á l e k 1997, H u d e c et al. 1999, unpublished data for 1998–2001). The present study revealed that the NAO weather system affects spring phenological instants in a number of bird species and that this effect could explain the earlier than normal arrival of common migratory species (e.g., *Alauda arvensis*, *Sturnus vulgaris*, *Vanellus vanellus*, *Columba palumbus*, *Motacilla alba*, *Phoenicurus ochruros*, *Phylloscopus collybita* and *Serinus serinus*) that has been observed in Central Europe since the 1980s. When we accept that the NAO system is an indicator (or even the proximate causative mechanism) of the climate change, then the earlier than normal spring arrival of migratory birds should be considered as both a result of, and evidence for, the climate warming in the last two decades.

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#### L I T E R A T U R E

- ALHEIT J. & HAGEN E. 1997: Long-term climate forcing of European herring and sardine populations. *Fish. Oceanogr.* 6: 130–139.
- BARNSTON A. G. & LIVEZEY R. E. 1987: Classification, seasonality and persistence of low-frequency atmospheric circulation patterns. *Monthly Weath. Rev.* 115: 1083–1126.
- BOTH C. & VISSER M. E. 2001: Adjustment to climate change is constrained by arrival date in a long-distance migrant bird. *Nature* 411: 296–298.
- CHMIELEWSKI F. M. & ROTZER T. 2001: Response of tree phenology to climate change across Europe. *Agricul. Forest Meteorol.* 108: 101–112.

- FORCHHAMMER M. C., POST E. & STENSETH N. C. 1998: Breeding phenology and climate. *Nature* 391: 29–30.
- FORCHHAMMER M. C., POST E. & STENSETH N. C. 2002: North Atlantic Oscillation timing of long- and short-distance migration. *J. Anim. Ecol.* 71: 1002–1014.
- HÁJEK V. 1992: [Bird fauna of South Moravia, 1952–1991]. *Manuscript*, 324 pp. (in Czech).
- HUBÁLEK Z. 1985: Cluster analysis of the spring migration of birds in Moravia (Czechoslovakia), 1881–1960. *Věstník Českosl. Spol. Zool.* 49: 81–86.
- HUBÁLEK Z. 1997: Trends of bird populations in a managed lowland riverine ecosystem. *Folia Zool.* 46: 289–302.
- HUDEK K., HUBÁLEK Z. & VAČKAŘ J. 1999: [Spring arrival of birds in the South Moravian region in 1997]. *Zprávy MOS*, 57: 117–128 (in Czech, with English summary).
- HÜPPOO O. & HÜPPOO K. 2003: North Atlantic Oscillation and timing of spring migration in birds. *Proc. Roy. Soc. Lond., B* 270: 233–240.
- HURRELL J. W. 1995: Decadal trends in the North Atlantic Oscillation: regional temperatures and precipitation. *Science*, 269: 676–679.
- JACOBS G. A., HURLBURT H. E., KINDLE J. C., METZGER E. J., MITCHELL J. L., TEAGUE W. E. & WALLCRAFT A. J. 1994: Decade-scale trans-Pacific propagation and warming effects of an El Niño anomaly. *Nature* 370: 360–363.
- JONZÉN N., HEDENSTRÖM A., HJORT C., LINDSTRÖM Å., LUNDBERG P. & ANDERSSON A. 2002: Climate patterns and the stochastic dynamics of migratory birds. *Oikos* 97: 329–336.
- KARL D. M., LETELIER R., HEBEL D., TUPAS L., DORE J., CHRISTIAN J. & WINN C. 1995: Ecosystem changes in the North Pacific subtropical gyre attributed to the 1991–92 El Niño. *Nature* 373: 230–234.
- MØLLER A. P. 2002: North Atlantic Oscillation (NAO) effects of climate on the relative importance of first and second clutches in a migratory passerine bird. *J. Anim. Ecol.* 71: 201–210.
- NOTT M. P., DESANTE D. F., SIEGEL R. B. & PYLE P. 2002: Influences of the El Niño/Southern Oscillation and the North Atlantic Oscillation on avian productivity in forests of the Pacific Northwest of North America. *Global Ecol. Biogeogr.* 11: 333–342.
- POST E. & FORCHHAMMER M. C. 2002: Synchronization of animal population dynamics by large-scale climate. *Nature* 420: 168–171.
- POST E., FORCHHAMMER M. C., STENSETH N. C. & CALLAGHAN T.V. 2001: The timing of life-history events in a changing climate. *Proc. Roy. Soc. Lond., B* 268: 15–23.
- POST E. & STENSETH N. C. 1999: Climatic variability, plant phenology, and northern ungulates. *Ecology* 80: 1322–1339.
- PRZYBYLO R., SHELDON B. C. & MERILA J. 2000: Climatic effects on breeding and morphology: evidence for phenotypic plasticity. *J. Anim. Ecol.* 69: 395–403.
- REID P. C., BORGESS M. D. & SVENDSEN E. 2001: A regime shift in the North Sea circa 1988 linked to changes in the North Sea horse mackerel fishery. *Fish. Res.* 50: 163–171.
- SAETHER B. E., TUFTO J., ENGEN S., JERSTAD K., ROSTAD O. W. & SKATAN J. E. 2000: Population dynamical consequences of climate change for a small temperate songbird. *Science* 287: 854–856.
- SANZ J. J. 2002: Climate change and breeding parameters of great and blue tits throughout the western Palearctic. *Global Change Biol.* 8: 409–422.
- SILLETT T. S., HOLMES R. T. & SHERRY T. W. 2000: Impacts of a global climate cycle on population dynamics of a migratory songbird. *Science* 288: 2040–2042.
- SIMS D. W., GENNER M. J., SOUTHWARD A. J. & HAWKINS S. J. 2001: Timing of squid migration reflects North Atlantic climate variability. *Proc. Roy. Soc. Lond., B* 268: 3607–3611.
- SOKOLOV L. V. 2001: Spring ambient temperature as an important factor controlling timing of arrival, breeding, post-fledging dispersal and breeding success of Pied Flycatcher *Ficedula hypoleuca* in Eastern Baltic. *Avian Ecol. Behav.* 5: 79–104.
- ŠTASTNÝ K., BEJČEK V. & HUDEK K. 1996: [Atlas of Breeding Birds in the Czech Republic, 1985–1989]. *H&H, Jinočany* (in Czech).
- STENSETH N. C., MYSTERUD A., OTTERSEN G., HURRELL J. W., CHAN K.-S. & LIMA M. 2002: Ecological effects of climate fluctuations. *Science* 297: 1292–1296.
- TKADLEC E. 2000: Brown hare population dynamics in the Prostějov district. *Přírodovědné Studie Muzea Prostějovska* 3: 133–149 (in Czech with English summary).
- TRYJANOWSKI P., KUZNIAK S. & SPARKS T. 2002: Earlier arrival of some farmland migrants in western Poland. *Ibis* 144: 62–68.

- TRYJANOWSKI P. & SPARKS T. H. 2001: Is the detection of the first arrival date of migrating birds influenced by population size? A case study of the Red-Backed Shrike *Lanius collurio*. *Internat. J. Biometeorol.* 45: 217–219.
- WALLACE J. M. & GUTZLER D. S. 1981: Teleconnections in the geopotential height field during the Northern Hemisphere winter. *Monthly Weath. Rev.* 109: 784–812.
- WINKLER D. W., DUNN P. O. & McCULLOCH C. E. 2002: Predicting the effects of climate change on avian life-history traits. *Proc. Nat. Acad. Sci.* 99: 13595–13599.
- YOO J. C. & D'ODORICO P. 2002: Trends and fluctuations in the dates of ice break-up of lakes and rivers in Northern Europe: the effect of the North Atlantic Oscillation. *J. Hydrol.* 268: 100–112.
- ZALAKEVICIUS M. & ZALAKEVICIUTE R. 2001: Global climate change impact on birds: a review of research in Lithuania. *Folia Zool.* 50: 1–17.
- ZUNA-KRATKY T., KALIVODOVÁ E., KÜRTHY A., HORAL D. & HORÁK P. 2000: Die Vögel der March-Thaya-Auen im österreichisch-slowakisch-tschechischen Raum. *Distelverein, Deutsch Wagram*.