

Bis(4-benzhydryl-benzoxazol-2-yl)methane – from a Bulky NacNac Alternative to a Trianion in Alkali Metal Complexes

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Abstract: A novel sterically demanding bis(4-benzhydrylbenzoxazol-2-yl)methane ligand **6** ($^{4-BzhH2}BoxCH_2$) was gained in a straightforward six-step synthesis. Starting from this ligand monomeric [$M(^{4-BzhH2}BoxCH)$] (M = Na (**7**), K (**8**₁)) and dimeric [{ $M(^{4-BzhH2}BoxCH)$ }₂] (M = K (**8**₂), Rb (**9**), Cs (**10**)) alkali metal complexes were synthesised by deprotonation. Abstraction of the potassium ion of **8** by reaction with 18-crown-6 resulted in the solvent separated ion pair [{(THF)₂K@(18crown-6)}{bis(4-benzhydryl-benzoxazol-2-yl)methanide}] (**11**),

Introduction

Over the last decades, monoanionic bidentate N,N'-ligands like β -diketiminates^[1] or amidinates^[2] gained interest due to their relative ease of synthesis and versatility with regards to steric and electronic properties. Based on these ligands a great variety of interesting transition metal,^[1b,3] main group^[4] and lanthanide^[5] compounds were synthesised in the last two decades. For example, in the field of main group chemistry, catalytically active β -diketiminate-derived group 13 metal complexes were synthesised.^[6] Further remarkable results were the synthesis of low-oxidation-state magnesium(I)^[7] and aluminium(I)^[8] complexes, which were facilitated by bulky β diketiminate ligands e.g. DippNacNac (Dipp=2,6-diisopropylphenyl). These findings initiated research into other ligand platforms, which mimic the coordination abilities of the ubiquitous NacNac ligand, in particular the κ^2 -N,N-coordination of the two imine nitrogen atoms to a metal ion to form sixmembered metallaheterocycles. Starting from the bis(2-pyridyl) methane ligand and its isoelectronically N^[9] P^[10] and As^[11] bridged derivatives, the 2-pyridyl moieties were exchanged for

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- Supporting information for this article is available on the WWW under https://doi.org/10.1002/chem.202100616
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including the energetically favoured monoanionic (*E*,*E*)-(^{4-B2hH2}BoxCH) ligand. Further reaction of ^{4-B2hH2}BoxCH₂ with three equivalents KH and two equivalents 18-crown-6 yielded polymeric [{(THF)₂K@(18-crown-6)}{K@(18-crown-6)K(^{4-B2h}BoxCH)}]_n (*n*→∞) (**12**) containing a trianionic ligand. The neutral ligand and herein reported alkali complexes were characterised by single X-ray analyses identifying the latter as a promising precursor for low-valent main group complexes.

benzannulated oxazoline or thiazole sidearms.^[12] Due to this variation of the bridging backbone and therefore connected residues, multiple symmetrical and asymmetrical ligand platforms were developed (Scheme 1).^[13] Although additional chalcogene donor sites (O or S) as well as an optional donating bridging unit (N, P or CH) are available for coordination, in most cases, the oxygen or sulfur do not share the same good Lewisdonor abilities of the ring nitrogen atoms. Hence a sixmembered NacNac-like coordination motif (C_3N_2M , M = cationic metal fragment) is maintained. In recent years the focus has been on symmetrical bis(benzoxazole-2-yl)methanides I to IV in Scheme 1 that can be straightforwardly synthesised by cyclocondensation reaction of two equivalents of corresponding 2aminophenol derivatives and one equivalent of the C3-linker unit ethylbisimidate dihydrochloride derived from malonic acid.^[14] In general, the synthesised bis(benzoxal-2-yl)methanes are utilised as neutral or as monoanionic ligands.^[12a,15] Deproto-



Scheme 1. Symmetrical bis(heterocyclo)amine (A), -phosphane (B), and -methane (C + D) ligands. Bis(benzoxazol-2-yl)methanes C (I, II) are (un) substituted in ortho-position to imine group whereas D is 'Pr (III) or 'Bu (IV) substituted in ortho- and para-position. Asymmetrical and unsubstituted bis (heterocylo)methanes (E).

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nation is accomplished with a rather strong Lewis base e.g. nbutyllithium or potassium hydride.^[16] Furthermore, these alkali metal complexes can be used as precursors in salt elimination reactions for metal complexes e.g. $[X_2AI(I^-)]^{[17]}$ (X = CI, I). Despite of the steric demand provided by ⁱPr- or ^tBu-residues at C4position close to the coordination pocket in III or IV, mimicking the bulkiness provided by NacNac ligands, low-oxidation or low-valent Group 2 or 13 elements could so far not be synthesised. Therefore, we decided to introduce bulkier benzhydryl groups at the ligand scaffold inspired by compounds containing 2,6-dibenzhydrylphenyl residues e.g. [:MH-(L)]^[18] (M = Ge or Sn, (L) =-N(Ar)(SiⁱPr₃), Ar = 4-ⁱPr-C₆H₂Bzh₂) or $[M(ArN)_2CN=C^tBu_2]^{[19]}$ (Ar = 4-^tBu-C₆H₂Bzh₂). In the following paragraph, the six-step synthesis of bis(4-benzhydryl-benzoxazol-2-yl)methane (6, ^{4-BzhH2}BoxCH₂) and its behaviour in alkali metal coordination are discussed.

Results and Discussion

Ligand synthesis and characterisation

To obtain the sterically demanding bis(4-benzhydryl-benzoxazol-2-yl)methane (6, ^{4-BzhH2}BoxCH₂), 2-amino-3-benzhydrylphenol was first synthesised according to the synthetic protocols of Quaranta et al.[20] that were slightly modified (for details of intermediates 1-5 see Supporting Information). Thereafter, two equivalents of 2-amino-3-benzhydrylphenol and one equivalent of ethyl cyanoacetimidate hydrochloride C3-linker unit were reacted by cyclocondensation reaction.^[14,16a,b,21] For this purpose, starting materials were heated in MeOH under reflux (85 °C) for at least 3 d. Ligand 6 and side product 2-(4-benzhydrylbenzoxazol-2-yl) acetamide (6a) precipitated and were separated by filtration. Further work-up (see Supporting Information) and purification by column chromatography on silica gel (THF/ hexane 3:2; R_f (6) = 0.9; R_f (6 a) = 0.48) led to $^{4-BzhH2}BoxCH_2$ (6) in decent yields (YLD(6): 36%; overall YLD: 13%; YLD(6a): 13%). All starting materials and ^{4-BzhH2}BoxCH₂ ligand (6) were analysed by NMR spectroscopy, mass spectrometry, and elemental analysis. Crystals of 6 suitable for single XRD experiments were grown out of a saturated ethyl acetate solution by slow cooling or by vapor diffusion of pentane. It crystallises in the monoclinic space group $P2_1/c$ with one molecule in the asymmetric unit (Figure 1). The bridging carbon atom C8 is coordinated like other symmetrically substituted bisheterocyclo methanes^[12a,21,22] in a distorted tetrahedral fashion with a slightly increased C7-C8-C9 angle of 115.71(14)° (related ligands: $II = 110.79(12)^{\circ}$ to $I = 111.23(9)^{\circ}$.^[12a,21] Furthermore, the two benzoxazole moieties are less twisted 63.25(4)° relative to each other compared to related unsubstituted I = 89.34(13)°[12a] and 4-methyl-substituted $II\!=\!81.14(4)^{\circ [21]}$ which exhibit an almost orthogonal alignment of the heteroaromatic planes.^[12a,21] This is attributed to the bulkier benzhydryl groups. These groups cause wider torsion angles between the C3-linker unit and the nitrogen or oxygen atoms (N1/2-C7-C8-C9=103.5(2)°/ -26.38(9)°; O1/2-C7-C8-C9= $31.47(10)^{\circ}/-71.05(19)^{\circ}$, more similar to 4,6-^tbutyl-substituted III $(twist = 76.50(7)^{\circ},$ N1/N2-C17-C1-C2 = -22.06(9)°/133.20(18)°,



Figure 1. Molecular structure of bis(4-benzhydryl-benzoxazol-2-yl)methane (6, $^{+3chH2}$ BoxCH₂). Anisotropic displacement parameters are depicted at the 50% probability level. All hydrogen atoms are omitted for clarity except for the bridging (methylene) and benzylic ones. Selected bond lengths [Å] and bond angles [°]: N1–C7 1.289(2), N2–C9 1.291(2), C7–C8 1.491(2), C8–C9 1.497(2), C7–C8–C9 115.71(14), N1/2–C7–C8–C9 = 103.5(2)°/–26.38(9)°; O1/2–C7–C8–C9 = 31.47(10)°/–71.05(19)°.

O1/O2-C17-C1-C2 = 23.05(10)°/-46.69(19)°). [16b] Moreover, the solid-state structure of ^{4-BzhH2}BoxCH₂ (6) reveals intermolecular hydrogen bonds, in which one of the acidic methylene hydrogen atom interacts with the imine nitrogen atom (N1...H8'A 2.49 Å) of an adjacent molecule and vice versa. In contrast, single crystal structure determination of ligand II (N1...H8'A 2.39 Å, N1...H8"B 2.63 Å) and III (N1...H8'A 2.58 Å, N1...H8"B 2.64 Å) exhibit intermolecular C–H…N hydrogen bonds of both methylene hydrogen atoms. In addition, an interaction of the less acidic benzylic hydrogen atoms $(pK_a=33)^{[23]}$ in 6 with the carbon atom of the benzoxazol-2-yl (C9) unit of a neighbouring molecule is observed (C9--H16' 2.81 Å). The acidity of the methylene protons correlates with the substitution or rather the electronic properties of related benzoxazole moieties.^[13] UV/Vis spectrophotometric titration experiments^[24] in acetonitrile unveiled a pK_a value of 26.61(6), supporting that $^{4-BzhH2}BoxCH_2$ is a somewhat stronger acid than I (26.89(3)), II (27.59(3)) and III (28.85(3)) but slightly weaker than bis(benzo-thiazol-2-yl) methane (26.14(3)).[13]

Alkali metal complexes of ^{4-BzhH2}BoxCH₂

 ${}^{\scriptscriptstyle 4\text{-}\text{BzhH2}}\text{BoxCH}_2$ (6) was reacted with alkali metal-based deprotonation or reduction agents to better understand its properties and establish an appropriate precursor for salt elimination reactions. All reactions were carried out in toluene at ambient temperature, whereas reaction times and input equivalents of alkali metal (hydrides) were adjusted to the reaction requirements (Scheme 2). In the case of bis(4-benzhydryl-benzoxazol-2-yl) methanide} sodium ([Na(^{4-BzhH2}BoxCH)]) (7), starting material 6 and neat sodium metal were vigorously stirred until all metal was consumed (5 d). After all volatiles had been removed under reduced pressure, 7 was isolated as a white powder in excellent yields (98%). Crystals suitable for single crystal XRD experiments were grown from a saturated toluene solution at -28 °C. Donor-free 7 crystallises in the monoclinic space group $P2_1/n$ with one molecule in the asymmetric unit (Figure 2). The solidstate structure of monomeric [Na(4-BzhH2BoxCH)] (7) exhibits a distorted κ^2 -N,N'-coordination of the sodium ion by the imine ring nitrogen atoms. A negligible dislocation of the cation from

Chem. Eur. J. 2021, 27, 1-9 www.chemeurj.org 2 These are not the final page numbers! Full Paper doi.org/10.1002/chem.202100616



 $M = K(8_2), Rb(9), Cs(10)$

Scheme 2. Synthesis of monomeric complexes 7 to 10 were carried out in toluene and ambient temperature under the following conditions (yiel-d = YLD): a) 1.04 eq. Na, 5 d, YLD: 98%, b) 1.15 eq. KH, 1 d, YLD: 89%, c) 1.10 eq. Rb, 1 d, YLD: 76%, d) 1.10 eq. Cs, 6 h, YLD: 74%.



Figure 2. Molecular structure of {bis(4-benzhydryl-benzoxazol-2-yl) methanide} sodium (7, [Na(^{4-BzhH2}BoxCH)]). Anisotropic displacement parameters are depicted at the 50% probability level. All hydrogen atoms are omitted for clarity except for the bridging (methylene) and benzylic ones.

the C₃N₂-plane of 0.117(2) Å and a minor butterfly folding angle of 4.82(6)° between the two benzoxazol-2-yl moieties is observed (Table 1). Measured nitrogen sodium bond lengths of **7** (Na1–N1 2.3392(16) Å; Na1–N2 2.3456(15) Å) are similar to the Na1–N1 2.358(6) Å bond of the six-membered sodium β -diketiminate complex [(THF)₂NaL]^[25] (L = {N(SiMe₃)C(Ph)}₂CH) but are slightly shorter than neutral five-membered TMEDA based [[{Na{N(SiMe₃)(Dipp)}(TMEDA)}₂] (Na1–N2 2.4726(17) Å; Na1–N3 2.461(2) Å)^[26] or four-membered guadinate complex [[Na{('PrNC (NAr)(HN'Pr))(THF)]₂ (Ar = 2,6-Me₂C₆H₃) (Na1–N1 2.453(1) Å, Na1–N2 2.558(2) Å).^[27] To complete the coordination sphere

around Na⁺ ion π -arene, interactions are formed to one phenyl group of the benzhydryl groups each.^[28] This way the benzyl hydrogen atoms are facing away from the metal. The Na…C_{Ar} distances ranging from 2.8305(18) (C_{ortho}) to 3.0414(18) Å (C_{ipso}) indicate a dihaptic arene coordination.^[29] ¹H NMR spectra of **6** and deprotonated species **7** show a significant deshielding in toluene and simultaneous downfield shift of the methanide bridge from $\delta(-H_2C-) = 3.85$ ppm to $\delta(-HC-) = 5.38$ ppm whereas benzylic protons experience a significant upfield shift from $\delta(-HCPh_2) = 6.41$ ppm to 5.27 ppm. Furthermore, arene protons (Ph and NCOC₆H₃) are slightly upfield shifted in sodium complex **7** compared to ligand **6**. Elemental analysis and mass spectrometry (LIFDI, toluene) confirmed the synthesis of bulk [Na(^{4-BzhH2}BoxCH)] (**7**).

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The reaction of ligand 6 with potassium hydride (Scheme 2) resulted in two different modifications of [K(^{4-BzhH2}BoxCH)]_n with n = 1 (8₁) and 2 (8₂), (YLD: 89%). Colourless crystals were grown from a saturated toluene solution at -30 °C after 1 d. Single crystal XRD experiments revealed a monomeric as well as a dimeric species in the solid state. Monomeric [(toluene)K(4-^{BzhH2}BoxCH)] crystallises in the triclinic space group P1 with one molecule and half a toluene molecule in the asymmetric unit (Figure 3). Compound **8**₁ consists of a distorted κ^2 -*N*,*N*'-coordi-arene interactions to one toluene molecule and one phenyl group of the benzhydryl moieties, respectively. In comparison with the sodium complex 6, a fairly prominent deviation of the cation from the C_3N_2 plane of 1.571(3) Å is present. Due to the larger radius of potassium, longer K–N (K1–N1 2.7768(18) Å; K1–N2 2.6982(18) Å) and a less acute N1–K1–N2 of 70.73(5)° are observed. This is accompanied by an increasing butterfly folding angle to 5.54(8)°. Furthermore, potassium phenyl carbon distances suggest that the cation is η^2 -(K1–C30 3.467(2) Å, K1–C31 3.224(2) Å) and η³-coordinated (K1–C23 3.314(2) Å, K1-C27 3.503(2) Å, K1-C28 3.236(2) Å) by one phenyl residue of each benzhydryl group (A search in the CCDC version 5.41 (Aug. 2020)^[30] for structures with K and phenyl groups excluding η^6 -coordinated ones provided a range of short contacts between K and Cortho of 2.77 to 3.76 Å).[29c,30] In addition, a toluene molecule is capping the cation in a η^3 -fashion (K1–C43 3.344(2) Å, K1–C44 3.098(2) Å, K1–C45 3.152(2) Å) which prevents the formation of a coordination polymer, frequently observed for related potassium complexes based on I-IV^[16a,b,d] or H^{Dipp}NacNac.^[31]

Table 1. Selected bond lengths, distances, and angles of complex 7 to 10.									
	(Z,Z)-(^{4-BzhH2} BoxCH)					(<i>E,E</i>)-(^{4-BzhH2} BoxCH)			
	7	81	8 ₂	9	10		8 ₂	9	10
M–N [Å]	2.3392(16),	2.7768(18),	2.812(2) to	2.924(5) to	3.050(3) to	M—N [Å]	2.775(2),	2.905(5),	3.033(3),
	2.3456(15)	2.6982(18)	3.015(2)	3.063(4)	3.285(3)		2.812(2)	2.948(5)	3.202(3)
N	86.53(5)	70.73(5)	67.57(7),	65.15(13),	62.76(8) to	M…C₃N₂ [Å]	3.170(3) to	3.297(6) to	2.603(10) to
			64.07(6)	63.34(12)	63.52(8)		3.493(3)	3.633(6)	1.252(19)
M…C ₃ N ₂ [Å]	0.117(2)	1.571(3)	1.877(4)	2.036(7),	2.302(5) to	M…HC₂N [Å]	2.18(3),	2.196(9),	2.60(5) to
			2.282(3)	2.303(7)	2.430(5)		0.85(5)	0.699(13)	1.16(8)
Folding	4.82(6)	5.54(8)	23.22(9)	23.63(19)	20.65(14),	Folding	16.75(8)	15.55(14)	17.85(11),
angle [°]					25.24(13)	angle [°]			15.22(10)

Chem. Eur. J. 2021, 27, 1-9

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Figure 3. Molecular structure of monomeric {bis(4-benzhydryl-benzoxazol-2-yl)methanide} potassium ($\mathbf{8}_1$, [K(^{4-B2hH2}BoxCH)]). Anisotropic displacement parameters are depicted at the 50% probability level. All hydrogen atoms are omitted for clarity except for the bridging (methylene) and benzylic ones.

The second species derived from **8** is the dimeric donorbase free $\mathbf{8}_2$ in the monoclinic space group $P2_1/c$. The three toluene lattice molecules in the asymmetric unit do not coordinate to the metal. Isostructural molecular structures were observed for the rubidium (9) (YLD: 76%) and caesium (10) (YLD: 74%) complexes (Figure 4, Table 1). They were synthesised via facile deprotonation of ligand **6** with neat rubidium and



Figure 4. Molecular structure of dimeric {bis(4-benzhydryl-benzoxazol-2-yl) methanide} alkali metal complexes ([{M($^{+BzhH2}BoxCH$)}₂], M = K (**8**₂), Rb (**9**), Cs (**10**)). Anisotropic displacement parameters are depicted at the 50% probability level. All hydrogen atoms are omitted for clarity. Superposition of **8**₂ -**10** showing NacNac-like C₃N₂ unit of (*E*,*E*)- ($^{+BzhH2}BoxCH$) (**A**) as well as twisted (*E*,*E*)-($^{+CzhH2}BoxCH$) (**B**) and their coordinating alkali metal ion, respectively.

caesium metal (Scheme 2). All three compounds show a monoanionic (*E*,*E*)-bis(4-benzhydryl-benzoxazol-2-yl)methanide} moiety coordinating two metal ions in κ^1 , η^2 -fashion (N1–N2–C9; N1–N2–C7) by the NacNac-like C_3N_2 unit (Figure 4A), respectively.^[32] In addition, a single phenyl group of each benzhydryl substituent forms an additional η^{6} - or η^{2} coordination.^[28] A second deprotonated (Z,Z)-isomer of 6 (Figure 4B) bridges both metals in an η^3 -fashion (C49–C48–N3; N4-C50-C49). The coordination sphere of the cations is again satiated by π -arene interactions of the adjacent phenyl groups $(\eta^4 \text{ and } \eta^2)$.^[28] Moreover, close interactions of the potassium ion and a suitable arranged methanide group in dimeric $\mathbf{8}_2$ culminate in rather short potassium hydrogen distances (K1-H49 2.89(3) Å, K2-H49 2.77(3) Å) and an elongated C-H bond to 1.01(3) Å.^[33] The K-H49-C49 angle of 102(2)° for K1 and 127(2)° for K2 also witness these interactions, because they are similar to e.g. [{K(THF)₄}₂Ln{(μ -H)₂BC₈H₁₄}₄] (Ln=Eu, Yb) (K-H 2.64(4) to 3.37(3) Å, K-H-C 117.8 to 120.7°)^[34] or [K(Ar)₂] [M {N(SiMe₃)₂}₃] (M = Mg, Ar = benzene, toluene, p-xylene; M = Zn, Ar = o-xylene) (K-H 2.81 to 3.07 Å, K-H-C 85 to 145°).^[35] The solid-state structures of 9 and 10 also display interactions of metal ions and methanide bridges, but due to the increased residual electron density for the heavier homologues the determination of the freely refined hydrogen atom positions is not that reliable. Rising effective ionic radii,[36] increasing polarizability and softness of the cations are encountered moving down the alkali metal group on account of the shielding effect.^[28a,37]

These features (Table 1) cause rising metal nitrogen bonds for deprotonated (Z,Z)- and (E,E)-(^{4-BzhH2}BoxCH) (8₂: 2.775(2) to 3.015(2) Å; 9: 2.905(5) to 3.063(4) Å; 10: 3.033(3) to 3.285(3) Å), more acute average N–M–N angles (8₂: 65.8°; 9: 64.3°; 10: 62.6°) as well as a growing dislocation of the metal from the C₃N₂ plane (82: 1.877(4) to 2.282(3) Å; 9: 2.036(7) to 2.303(7) Å; 10: 2.302(5) to 2.430(5) Å) as well as from the -HC₂N unit (8₂: 0.85(5) to 2.18(3) Å; 9: 0.699(13) to 2.196(9) Å; 10: 1.16(8) to 2.60(5) Å). Interestingly, the detected butterfly folding angles in both isomers (Z,Z)-(^{4-BzhH2}BoxCH) \approx 23°; (E,E)-(^{4-BzhH2}BoxCH) \approx 16°) are negligibly affected while coordinating the alkali metal ions. Additionally, all alkali metal complexes 82-10 were studied by ¹H and ¹³C{¹H} NMR spectroscopy in [D₈]THF. Recorded ¹H NMR spectra show a distinctive pattern of chemical shifts arising from the C_{2v} symmetry of monoanionic bis(benzoxazol-2-yl) methanide scaffold and the benzylic bound protons, whereas phenylic protons of the benzhydryl groups could not be clearly assigned due to their peak overlap. Deprotonation of 6 results in an accumulation of electron density in the benzene periphery of both benzoxazol-2-yl moieties that again entails a general shielding of corresponding protons and an upfield shift. Concerning the ¹H NMR shifts for the C-H bridging position, a minute but continuous decline along K (4.66 ppm) > Rb (4.65 ppm) > Cs (4.64 ppm) complexes is noticed. The opposite is found for para positioned protons (3-, 13-H) and benzylic protons. These protons exhibit a small but noticeable downfield shift $(\delta(3-, 13-H) = 6.33 \text{ ppm}$ to 6.43 ppm; $\delta(16-, 29-H) =$ 6.03 ppm to 6.17 ppm) which is in tune with the negative charge being restored in the phenylic benzhydryl groups



yielding a growing metal arene interaction. Remaining benzoxazol-2-yl protons in ortho- and meta-position shifts (4-,12- and 5-, 11-H) do not seem to be influenced by varying alkali cations. Although ¹H NMR spectra of **8–10** ([D₈]THF) and sodium complex **7** ([D₈]toluene) were measured in different solvents, benzylic protons (16-, 29-H) display a remarkable change of the average chemical shifts ($\Delta \delta = 0.82$ ppm). The ⁸⁷Rb NMR spectrum of **9** from THF solution shows two singlets. Most probably [{Rb(^{4-BzhH2}BoxCH)}₂] rearranges to give a THF-solvated rubidium cation [Rb(THF)_n] ($\delta = -1.69$ ppm) and a bis(4-benzhydrylbenzoxazol-2-yl)methanide} based anion [Rb(^{4-BzhH2}BoxCH)₂]⁻ ($\delta = -254.69$ ppm). The ¹³³Cs NMR spectrum of **10** displays only one signal at $\delta = -31.12$ ppm. Additionally, elemental analysis and mass spectrometry (LIFDI, THF) confirmed the synthesis of heavier alkali complexes **8–10**.

On the basis of reported (*Z*,*Z*)- and (*E*,*E*)-isomers of complex **8–10** and based on former gained knowledge of sterically less demanding bis(benzoxazol-2-yl)methanide potassium complexes^[16a,b] and their related 18-crown-6 derivatives,^[16c,d,38] we were curious whether the reaction of **8** with 18-crown-6 would prefer the formation of a (*Z*,*Z*), (*E*,*E*) or (*Z*,*E*) isomeric species. Therefore bis(4-benzhydryl-benzoxazol-2-yl)methane was first deprotonated with an excess of potassium hydride in THF at ambient temperature (Scheme 3). After the reaction mixture had been stirred for 1 d, excess of hydride was removed by filtration and



Scheme 3. Reaction of monomeric complexes 8 to solvent separated ion $[(THF)_2K(18-crown-6)]^+[^{4-BzhH2}BoxCH]^-$ (11) via addition of one equivalent 18-crown-6. In solution permanent rearrangement of the (*Z*,*Z*)-, (*Z*,*E*/*E*,*Z*)-, and (*E*,*E*)-(^{4-BzhH2}BoxCH) isomers is ubiquitous.



Figure 5. Molecular structure of {bis(4-benzhydryl-benzoxazol-2-yl) methanide} anion from 11, [(THF)₂K(18-crown-6)]⁺[^{+82hH2}BoxCH]⁻. Anisotropic displacement parameters are depicted at the 50% probability level. All hydrogen atoms are omitted for clarity except for the bridging (methylene) and benzylic ones. Selected bond lengths [Å] and bond angles [°]: N1–C7 1.329(2), N2–C9 1.327(2), C7–C8 1.390(2), C8–C9 1.389(2), C9–C8–C7 131.34(15).

Chem. Eur. J. 2021, 27, 1–9 www.chemeurj.org 5 These are not the final page numbers!

18-crown-6 was added to the solution. Within seconds a strong blue fluorescence ($\lambda_{max} = 454 \text{ nm}$, $\lambda_{ex} = 350 \text{ nm}$, detail see Supporting Information) of the dark red solution becomes entrenched. The reaction mixture was further stirred for 1 d at ambient temperature, volatiles were removed in vacuum and the solvent separated ion pair [(THF)₂K(18-crown-6)]⁺ [⁴⁻ ^{BzhH2}BoxCH]⁻ (11) was isolated as a reddish solid in excellent yields (95%). Crystals suitable for single XRD experiments were grown by vapour diffusion of pentane to a saturated THF solution of 11 at ambient temperature after 3 d. The colourless crystals in the triclinic space group P1 contain one bis(4benzhydryl-benzoxazol-2-yl)methanide anion and two halve (THF)₂K(18-crown-6)} cations in the asymmetric unit (Figure 5). In contrast to [(L)K(18-crown-6)] ($L = I^-$, III⁻)^[16c,d,38] that develop contact ion pairs in the solid, complex 11 was found to form a solvent separated ion pair. Expectedly, the (Z,Z)-(^{4-BzhH2}BoxCH) isomer seems to be preferred over the (Z, E/E, Z)- or (E, E)-(4-BZhH2BoxCH) isomers in the solid-state. In similar manner the formation of a (Z,Z)-isomer accompanied by only an oxygen atom coordination at the metal is found in [(III-)K(18-crown-6)·H₂O].^[16c,38] The molecular structure of $(Z_{z}Z)$ -(^{4-BzhH2}BoxCH) (11) displays a butterfly folding angle of 5.03(9)°, which is almost consistent to the folding angles observed in monomeric species of 8₁ (5.54(8)°) as well as 7 (4.82(6)°). This obtuse-angled benzoxazol-2-zyl moieties and the bond length of C_3N_2 unit indicate a fully conjugated system extended throughout the virtually planar ligand.^[39] Based on these findings, the Gibbs free energy differences $\triangle G$ of the monoanionic (*E*,*E*)-, (*Z*,*E*/*E*,*Z*)or $(E,E)-(^{4-Bzh}BoxCH)$ isomers were calculated by density functional theory at the BP86/def2-SVP level (see details in Supporting Information). Calculated Gibbs free energy differences is for all configurational isomers $\triangle G < 14$ kcal/mol. Hence in solution most likely the permanent rotation around the methanide linker, accompanied by rearrangement of the isomers (Scheme 3), is ubiquitous. Moreover, the synthesis of [{(THF)₂K(18-crown-6)}(^{4-BzhH2}BoxCH)] (11) was confirmed by mass spectrometry (ESI[-],THF) and elemental analysis.

Since Brown or Buncel and Menon ascertained that triphenylmethane is deprotonated in THF by potassium hydride when DMF^[40] or 18-crown-6^[41] is added,^[42] we wanted to examine whether deprotonation of one or both benzylic functions can be accomplished via reaction of one or two equivalents of hydride and crown ether, respectively, to a solution of 8. However, the reactions of different amounts of KH and 18-crown-6 all yielded the same complex [{(THF)₂K(18crown-6)}{K₂(^{4-Bzh}BoxCH)}]_n ($n \rightarrow \infty$) **12** as very sensitive dark purple crystals by storing THF solutions at -30°C. The crystal selection and mounting was challenging due to the dark colour and high sensitivity of compound 12. Thus, a moderate single crystal XRD data set could be collected after many attempts. Furthermore, the synthesis of 12 was improved by using ligand **6** and the exact stoichiometric quantities ($^{4-BzhH2}BoxCH_2:KH:18$ crown-6=1:3:2) (Scheme 4) that are based on the crystal structure. Immediately after adding the starting materials to a solution of 6 in THF, a solid blue fluorescence of the dark red solution was observed. The mixture was stirred at least for 1 d, whereas a red solid precipitated that was separated from the Full Paper doi.org/10.1002/chem.202100616





n [(THF)2K@18-crown-6]+

Scheme 4. Optimised synthesis of polymeric [{(THF)₂K(18-crown-6)}{K(18-crown-6)K($^{4-Bzh}$ BoxCH)}]_n ($n \rightarrow \infty$) (12) by addition of 3.33 equivalents of KH and two equivalents of 18-crown-6.

supernatant solution by decantation. The precipitate was washed with pentane, dried under reduced pressure, and stored at -30 °C for further syntheses or analyses (YLD: 95%). Obtained THF solution was cooled to -30 °C for crystallisation. Complex 12 crystallises in the monoclinic space group P2/n with half a formula unit in the asymmetric unit (Figure 6). The coordination polymer comprises solvent-separated potassium ion (K3) solely coordinated by the oxygen atoms of 18-crown-6 and two THF molecules. This is omitted in Figure 6 for clarity. A second potassium ion (K2) is η^2 -coordinated (C4–K2 3.329(3) Å, C5–K2 3.245(8) Å) to the aryl moieties of two adjacent (E,E)trianions and also coordinated by an equatorial crown ether molecule. The η^2 -coordination provides the link in the polymeric structure. Finally, a third potassium ion in the coordination pocket is surrounded by two nitrogen atoms in κ^2 -fashion (N–K1 2.686(3) Å) and a phenyl ring of each benzhydryl moiety in asymmetrically η^{6} -fashion (C_{Ph}-K1: 2.930(3) Å to 3.291(3) Å), respectively. Determined nitrogen potassium distances and the perfect planarity of the ligand scaffold with the potassium ion



Figure 6. Molecular structure of [{(THF)₂K@(18-crown-6)}{K@(18-crown-6)}K-(^{4-Bzh}BoxCH)}]_n (*n*→∞) (12). Anisotropic displacement parameters are depicted at the 50% probability level. All hydrogen atoms are omitted for clarity except for the bridging (methylene) one. K1–N1 2.686(3), K1–C16 2.930(3), K1–C17 3.084(3), K1–C18 3.255(3), K1–C19 3.291(3), K1–C20 3.154(3), K1–C21 2.993(3), N1–C7 1.324(4), C7–C8 1.393(4), C2–C9 1.478(4), C9–C10 1.459(5), C9–C16 1.426(4), C4–K2 3.329(3) Å, C5–K2 3.245(8) Å, N1–K1–N1A 72.78(11), C1–C2–C3 114.2(3), C6–C5–C4 115.1(4), C15–C10–C11 115.7(3), C17–C16–C21 114.4(3).

in the plane of the trianion are in good agreement with the unsolvated κ^2 -N,N-[K(^{Dipp}NacNac)]^[31] complex (N1–K1 2.710(3) Å, N2-K1 2.652(2) Å). The bond lengths to flanking Cipso-atoms are basically shortened (1.426(4) to 1.478(4) Å) compared to HC_{bzh}-C_{ipso} of complexes 6-11 (1.508(8) Å to 1.538(6) Å) because of the deprotonation of the two benzylic positions (C_{bzh}) (benzhydryl group). The two anionic C9-atoms are essentially in plane with the bound Cipso-atoms (C2, C10, C16) similar to previously studied potassium trityl complexes $[KCPh_3(THF)(PMDTA)]$,^[30] $[KCPh_3(PMDTA)]^{[43]}$ or $[KCPh_3(L)]_n$ (L = diglyme, THF).^[43] Angles around C9 deviate slightly from the typical 120° of sp²-hybridised carbon atoms (for details see Supporting Information), while the steric constraints prevent the phenyl rings from being coplanar leading to a propeller-like arrangement. Within this arrangement, angles φ of the phenyl groups with respect to C2–C10–C16 plane (C3-plane) are $\varphi =$ 32.38(17)° (C10 to C15), and 49.68(10)° (C1 to C6), whereas the smallest angle and shortest bond (C9-C16 1.426(4) Å) involves the phenyl system to which the cation is coordinated. These parameters correspond to an increased π -electron-delocalisation that is correlated with the overlap of the C_3 -plane (C2-C10-C16) and coordinating phenyl ring (C16 to C21) which is dependent on $\cos(\varphi)^2$ function^[30,44] of the corresponding plane angles. Although the phenyl group is twisted ($\varphi =$ 16.4(2)°) the overlap amounts to about 92%, that is a more significant overlap than observed for [KCPh₃(THF)(PMDTA)]^[30] $(\varphi = 17.7^{\circ}, \approx 90\%)$. Further direct analyses of 12 were challenging due to its low solubility in most solvents (toluene, benzene, or THF). To find further evidence for the synthesis of trifold deprotonated anion (4-BzhBoxCH), suspensions of the dark red precipitate (12) were once again protonated (excess \approx 20 eq. H_2O or D_2O) in small scale (NMR experiment). These ¹H and ²D NMR experiments (for details see Supporting Information), as well as mass spectrometry (ESI[-] HRMS, THF), confirmed the previous synthesis of 12.

Conclusion

Within this work, the six-step synthesis and characterisation of a novel bulky bis(benzoxazol-2-yl)methane ligand 6 (4-BzhH2BoxCH2) comprising benzhydryl groups at both C4positions (ortho imine positions) in spatial proximity to the coordination pocket was presented. To get a better knowledge of its properties and find a possible precursor complex for subsequent salt metathesis reactions, 6 was deprotonated with alkali metal bases. Obtained products were analysed by NMR spectroscopy, mass spectrometry as well as single crystal XRD experiments. Crystals grown from toluene solutions unveiled monomeric [$M(^{4-BzhH2}BoxCH)$] (M = Na (7), K (8₁)) and dimeric $[{M(^{4-BzhH2}BoxCH)}_2]$ (M = K (8₂), Rb (9), Cs (10)) species in solid state. Latter alkali metal complexes display distorted κ^2 -N,Ncoordinated (Z,Z)- and an (E,E)-($^{4-BzhH2}$ BoxCH) configurational isomers, which display various polyhaptic metal arene interactions. Furthermore, potassium ion sequestration of 8 by 18crown-6 resulted in [{(THF)₂K(18-crown-6)}(^{4-BzhH2}BoxCH)] (11) a solvent separated ion pair containing energetically favoured

Chem. Eur. J. 2021, 27, 1–9 www.chemeurj.org 6 These are not the final page numbers!



monoanionic (E,E)-(4-BzhH2BoxCH) ligand. Additionally, reaction of ^{4-BzhH2}BoxCH₂ with three equivalents KH and two equivalents 18crown-6 yielded polymeric [{(THF)₂K@(18-crown-6)){K@(18crown-6)K(^{4-Bzh}BoxCH)]_n ($n \rightarrow \infty$) (12) featuring a remarkable trianionic ligand. The single crystal X-ray diffraction experiment of 12 revealed one of the potassium ions to be η^2 -coordinated by the benzoxazol-2-yl scaffold of two adjacent (E,E)-ligands plus a crown ether molecule. A second potassium ion is surrounded by two nitrogen atoms in κ^2 -fashion as well as one phenyl ring of each benzhydryl moiety in a symmetrically η^{6} fashion, respectively. Future work with regard to the mono-(4-BzhH2BoxCH) and trianionic (4-BzhBoxCH) ligand will focus on the coordination of other metal, i.e. those from Group 13 or lanthanides. The benzhydryl groups attached to the benzoxazol-2-yl scaffold seem to exhibit similar qualities to the triphenylmethane, which is able to stabilise carbanionic, radical or carbonium ionic species owing to the extensive π -delocalisation. With this in mind, future research might focus on the synthesis of $^{\rm 4-BzhH2} {\rm BoxCH}_2$ based radical and carbonium ion compounds.

Experimental section

General procedures: All reactions were carried out under an atmosphere of N2 and Ar by Schlenk techniques. Solvents were distilled from Na, K or NaK before used for synthesis. Starting materials were purchased commercially and used without further purification. 1D (1H, 13C, 87Rb, 133Cs) and 2D (1H,13C HSQC, 1H,13C and 1H,15N HMBC) NMR spectroscopic data were recorded on a Bruker Ascend 500 MHz, 400 MHz, and Avance 300 MHz spectrometer and referenced to the deuterated solvent (benzene d_{6} , thf- d_{8} , toluene- d_{8} .^[45] Deuterated benzene and toluene were dried over K (65 $^{\circ}$ C), THF was pre-dried with LiAlH₄ and stored over activated molecular sieve (3 Å) and stored in an argon dry box. Elemental analyses (C, H, N) were performed on a Vario EL3 at the Mikroanalytische Labor, Institut für Anorganische Chemie, University of Göttingen. LIFDI-MS spectra were measured on a Jeol AccuTOF spectrometer and ESI (HR-MS) measurements were performed on Bruker maXis spectrometer. All pK_a measurements in acetonitrile were carried out as in reference [13]. Shock-cooled crystals were selected from a Schlenk flask under argon atmosphere using the X-TEMP2 device.^[46] The data were collected with a Mo- (6, 8₁, 9, 10, 11) or Ag- (7, 8₂, 12) μS microfocus source.^[47] All data were integrated with SAINT,^[48] and a multiscan absorption correction (SADABS^{[49]}) was applied. For structure $\boldsymbol{6}$ a 3λ correction was applied.^[50] The structures were solved by direct methods (SHELXT^[51]) and refined on F² using the full-matrix least-squares methods of SHELXL^[52] within the SHELXLE GUI.^[53] The Crystallographic Information Files (CIF) can be obtained free of charge from the Cambridge Crystallographic Data Centre via www.ccdc.cam.ac.uk/ structures/ using the reference numbers 2031907 (for 6), 2031908 (for 7), 2031909 (for 8₁), 2031910 (for 8₂), 2031911 (for 9), 2031912 (for 10), 2031913 (for 11), 2031914 (for 12).

Acknowledgements

We thank the Danish National Research Foundation (DNRF93) funded Center for Materials Crystallography (CMC) for partial support. The work at UT was supported by the Estonian

Chem. Eur. J. 2021, 27, 1-9 www.chemeurj.org 7 These are not the final page numbers! 77

Research Council grant (PRG690). Open access funding enabled and organized by Projekt DEAL.

Conflict of Interest

The authors declare no conflict of interest.

Keywords: Ligand design · Metalation · Methanides · NacNac · Potassium · s-Block chemistry

- [1] a) L. Bourget-Merle, M. F. Lappert, J. R. Severn, Chem. Rev. 2002, 102, 3031–3066; b) Y.-C. Tsai, Coord. Chem. Rev. 2012, 256, 722–758.
- [2] F. T. Edelmann, Advances in the Coordination Chemistry of Amidinate and Guanidinate Ligands; Elsevier, Burlington, 2008, p. 184–339.
- [3] R. L. Webster, Dalton Trans. 2017, 46, 4483-4498.
- [4] M. Asay, C. Jones, M. Driess, Chem. Rev. 2011, 111, 354–396.
- [5] a) F. T. Edelmann, D. M. M. Freckmann, H. Schumann, Chem. Rev. 2002, 102, 1851–1896; b) H. C. Aspinall, Chem. Rev. 2002, 102, 1807–1850.
- [6] a) P. P. Power, *Nature* 2010, *463*, 171–177; b) Z. Yang, M. Zhong, X. Ma,
 S. De, C. Anusha, P. Parameswaran, H. W. Roesky, *Angew. Chem. Int. Ed.* 2015, *54*, 10225–10229; *Angew. Chem.* 2015, *127*, 10363–10367; c) Z.
 Yang, M. Zhong, X. Ma, K. Nijesh, S. De, P. Parameswaran, H. W. Roesky,
 J. Am. Chem. Soc. 2016, *138*, 2548–2551.
- [7] S. P. Green, C. Jones, A. Stasch, Science 2007, 318, 1754–1757.
- [8] a) C. Cui, H. W. Roesky, H. Hao, H.-G. Schmidt, M. Noltemeyer, Angew. Chem. Int. Ed. 2000, 39, 1815–1817; Angew. Chem. 2000, 112, 4274– 4276.
- [9] a) M. Pfeiffer, F. Baier, T. Stey, D. Leusser, D. Stalke, B. Engels, D. Moigno, W. Kiefer, J. Mol. Model. 2000, 6, 299–311; b) M. Pfeiffer, A. Murso, L. Mahalakshmi, D. Moigno, W. Kiefer, D. Stalke, Eur. J. Inorg. Chem. 2002, 2002, 3222–3234.
- [10] A. Steiner, D. Stalke, Chem. Commun. 1993, 444–446.
- [11] a) H. Gornitzka, D. Stalke, Eur. J. Inorg. Chem. 1998, 3, 311–317; b) A. Steiner, D. Stalke, Organometallics 1995, 14, 2422–2429.
- [12] a) D.-R. Dauer, D. Stalke, *Dalton Trans.* 2014, 43, 14432–14439; b) D.-R. Dauer, M. Flügge, R. Herbst-Irmer, D. Stalke, *Dalton Trans.* 2016, 45, 6136–6148.
- [13] J. Kretsch, I. Koehne, M. Lökov, I. Leito, D. Stalke, Eur. J. Inorg. Chem. 2019, 2019, 3258–3264.
- [14] H. Ben Ammar, J. Le Nôtre, M. Salem, M. T. Kaddachi, P. H. Dixneuf, J. Organomet. Chem. 2002, 662, 63–69.
- [15] a) R. R. Trenholm, D. E. Ryan, Anal. Chim. Acta 1965, 32, 317–322; b) M. T. Ramos, C. Avendano, J. Elguero, F. Florencio, J. Sanz-Aparicio, Inorg. Chim. Acta 1990, 174, 169–173; c) H. A. Elagab, H. G. Alt, Inorg. Chim. Acta 2015, 437, 26–35; d) A. Hijazi, J.-P. Djukic, L. Allouche, A. de Cian, M. Pfeffer, X.-F. Le Goff, L. Ricard, Organometallics 2007, 26, 4180–4196.
- [16] a) I. Koehne, N. Graw, T. Teuteberg, R. Herbst-Irmer, D. Stalke Inorg, *Chem.* 2017, 56, 14968–14978; b) I. Koehne, S. Bachmann, T. Niklas, R. Herbst-Irmer, D. Stalke, *Chem. Eur. J.* 2017, 23, 13141–13149; c) I. Koehne, S. Bachmann, R. Herbst-Irmer, D. Stalke, *Angew. Chem. Int. Ed.* 2017, 56, 15141–15145; *Angew. Chem.* 2017, 129, 15337–15342; d) J. Kretsch, A. Kreyenschmidt, R. Herbst-Irmer, D. Stalke, *Dalton Trans.* 2018, 47, 12606–12612.
- [17] J. Kretsch, R. Herbst-Irmer, D. Stalke, Z. Anorg. Allg. Chem. 2018, 644, 657–660.
- [18] T. J. Hadlington, M. Hermann, G. Frenking, C. Jones, J. Am. Chem. Soc. 2014, 136, 3028–3031.
- [19] A. K. Maity, S. Fortier, L. Griego, A. J. Metta-Magaña, *Inorg. Chem.* 2014, 53, 8155–8164.
- [20] L. Quaranta, O. Corminboeuf, P. Renaud, Org. Lett. 2002, 4, 39-42.
- [21] D.-R. Dauer, M. Flügge, R. Herbst-Irmer, D. Stalke, Dalton Trans. 2016, 45, 6149–6158.
- [22] a) I. Koehne, R. Herbst-Irmer, D. Stalke, *Eur. J. Inorg. Chem.* 2017, 2017, 3322–3326; b) H. Gornitzka, D. Stalke, *Angew. Chem. Int. Ed.* 1994, 33, 693–695; *Angew. Chem.* 1994, 106, 695–698; c) H. Gornitzka, D. Stalke, *Organometallics* 1994, 13, 4398–4405.
- [23] R. Breslow, W. Chu, J. Am. Chem. Soc. 1970, 92, 2165.
- [24] a) I. Kaljurand, A. Kütt, L. Sooväli, T. Rodima, V. Mäemets, I. Leito, I. A. Koppel, J. Org. Chem. 2005, 70, 1019–1028; b) A. Kütt, I. Leito, I.

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Kaljurand, L. Sooväli, V. M. Vlasov, L. M. Yagupolskii, I. A. Koppel, *J. Org. Chem.* **2006**, *71*, 2829–2838; c) A. Kütt, S. Tshepelevitsh, J. Saame, M. Lőkov, I. Kaljurand, S. Selberg, I. Leito, *Eur. J. Org. Chem.* **2021**, *9*, 1407–1419.

- [25] Y. Cheng, P. B. Hitchcock, M. F. Lappert, M. Zhou, Chem. Commun. 2005, 752–754.
- [26] M. Á. Fuentes, A. Zabala, A. R. Kennedy, R. E. Mulvey, Chem. Eur. J. 2016, 22, 14968–14978.
- [27] M. de Tullio, A. Hernán-Gómez, Z. Livingstone, W. Clegg, A. R. Kennedy, R. W. Harrington, A. Antiñolo, A. Martínez, F. Carrillo-Hermosilla, E. Hevia, *Chem. Eur. J.* 2016, 22, 17646–17656.
- [28] a) T. X. Gentner, R. E. Mulvey, Angew. Chem. Int. Ed. 2020, 59, 2–18; Angew. Chem. 2020, 132, 2–20; b) J. M. Gil-Negrete, E. Hevia, Chem. Sci. 2021, 12, 1982–1992.
- [29] a) C. Lambert, P. von Ragué Schleyer, Angew. Chem. Int. Ed. 1994, 33, 1129–1140; Angew. Chem. 1994, 106, 1187–1199; b) G. Rabe, H. W. Roesky, D. Stalke, F. Pauer, G. M. Sheldrick, J. Organomet. Chem. 1991, 403, 11–19; c) U. Pieper, D. Stalke, Organometallics 1993, 12, 1201–1206; d) F. Antolini, P. B. Hitchcock, M. F. Lappert, P. Merle, Chem. Commun. 2000, 1301–1302; e) S.-D. Bai, J.-P. Guo, D.-S. Liu, Dalton Trans. 2006, 2244–2250.
- [30] D. Hoffmann, W. Bauer, P. v R Schleyer, U. Pieper, D. Stalke, Organometallics 1993, 12, 1193–1200.
- [31] A. G. Avent, M. R. Crimmin, M. S. Hill, P. B. Hitchcock, J. Organomet. Chem. 2006, 691, 1242–1250.
- [32] S. Brooker, F. T. Edelmann, T. Kottke, H. W. Roesky, G. M. Sheldrick, D. Stalke, K. H. Whitmire, Chem. Commun. 1991, 325, 144–146.
- [33] a) W. Scherer, V. Herz, A. Brück, C. Hauf, F. Reiner, S. Altmannshofer, D. Leusser, D. Stalke, Angew. Chem. Int. Ed. 2011, 50, 2845–2849; Angew. Chem. 2011, 123, 2897–2902; b) W. Scherer, A. C. Dunbar, J. E. Barquera-Lozada, D. Schmitz, G. Eickerling, D. Kratzert, D. Stalke, A. Lanza, P. Macchi, N. P. M. Casati, J. Ebad-Allah, C. Kuntscher, Angew. Chem. Int. Ed. 2015, 54, 2505–2509; Angew. Chem. 2015, 127, 2535–2539.
- [34] X. Chen, S. Lim, C. E. Plecnik, S. Liu, B. Du, E. A. Meyers, S. G. Shore, *Inorg. Chem.* 2005, 44, 6052–6061.
- [35] a) I. J. Bruno, J. C. Cole, P. R. Edgington, M. Kessler, C. F. Macrae, P. McCabe, J. Pearson, R. Taylor, *Acta Crystallogr. Sect. B* 2002, *58*, 389–397; b) G. C. Forbes, A. R. Kennedy, R. E. Mulvey, B. A. Roberts, R. B. Rowlings, *Organometallics* 2002, *21*, 5115–5121.
- [36] R. Shannon, Acta Crystallogr. 1976, 32, 751-767.

[37] T. L. Brown, H. E. LeMay Jr, B. E. Bursten, C. J. Murphy, P. M. Woodward, *Chemistry*: The central science; Prentice Hall, Upper Saddle River, NJ, 2003, p. 254–259.

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- [38] I. Koehne, S. Bachmann, R. Herbst-Irmer, D. Stalke, Angew. Chem. Int. Ed. 2017, 56, 15141–15145; Angew. Chem. 2017, 129, 15337–15342.
- [39] a) P. Rademacher in Strukturen organischer Moleküle, Physikalische Organische Chemie, Wiley-VCH, Weinheim, 1987, p. 55–88; b) P. Müller, R. Herbst-Irmer, A. L. Spek, T. R. Schneider, M. R. Sawaya in Crytal Structure Refinement A Crystallographer's Guide to SHELXL, Oxford University Press, New York, 2006, p. 1–202.
- [40] C. A. Brown, J. Org. Chem. 1974, 39, 3913-3918.
- [41] E. Buncel, B. Menon, Chem. Commun. 1976, 648-649.
- [42] R. E. Gawley, X. Zhang, Q. Wang in *Encyclopedia of reagents for organic synthesis: potassium hydride*; Wiley, Chichester, **1995**, p. 1–3.
- [43] H. Viebrock, T. Panther, U. Behrens, E. Weiss, J. Organomet. Chem. 1995, 491, 19–25.
- [44] a) R. A. Bartlett, H. V. R. Dias, P. P. Power, J. Organomet. Chem. 1988, 341, 1–9; b) A. C. Benniston, A. Harriman, Chem. Soc. Rev. 2006, 35, 169–179; c) R. M. Williams, Photochem. Photobiol. Sci. 2010, 9, 1018–1026.
- [45] Virtuelles Labor II, "http://www.stalke.chemie.uni-goettingen.de/virtuelles labor/nmr/de.html".
- [46] a) T. Kottke, D. Stalke, J. Appl. Crystallogr. 1993, 26, 615–619; b) D. Stalke, Chem. Soc. Rev. 1998, 27, 171–178.
- [47] T. Schulz, K. Meindl, D. Leusser, D. Stern, J. Graf, C. Michaelsen, M. Ruf, G. M. Sheldrick, D. Stalke, *J. Appl. Crystallogr.* **2009**, *42*, 885–891.
- [48] Bruker AXS Inc, SAINT, Madison, 2016.
 [49] L. Krause, R. Herbst-Irmer, G. M. Sheldrick, D. Stalke, J. Appl. Crystallogr.
- [49] L. Krause, K. Herbst-Irmer, G. M. Sneidrick, D. Staike, J. Appl. Crystallogr. 2015, 48, 3–10.
- [50] L. Krause, R. Herbst-Irmer, D. Stalke, J. Appl. Crystallogr. 2015, 48, 1907– 1913.
- [51] G. M. Sheldrick, Acta Crystallogr. Sect. A 2015, 71, 3-8.
- [52] G. M. Sheldrick, Acta Crystallogr. Sect. C 2015, 71, 3-8.
- [53] C. B. Hübschle, G. M. Sheldrick, B. Dittrich, J. Appl. Crystallogr. 2011, 44, 1281–1284.

Manuscript received: February 17, 2021 Version of record online:

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Bis(4-benzhydryl-benzoxazol-2-yl) methane (^{4-BzhH2}BoxCH₂) provides a versatile platform for monomeric ligands in $[M(^{4-BzhH2}BoxCH)]$ (M = Na, K) and dimeric $[{M(^{4-BzhH2}BoxCH)}_2]$ (M = K, Rb, Cs) alkali metal complexes. Even more remarkable the reaction of ^{4–BzhH2}BoxCH₂ with three equivalents KH and two equivalents 18-crown-6 yields polymeric [{(THF)₂K@(18-crown-6)}{K@(18-crown-6)K($^{4-BzhH2}BoxCH)}]_n$ $(n \rightarrow \infty)$, containing a trianionic ligand. While the monoanions only provide N,N-chelation like NacNac the latter trianion potentially provides C,N,N,Cchelation and should be another promising ligand platform.



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1 – 9

Bis(4-benzhydryl-benzoxazol-2-yl) methane – from a Bulky NacNac Alternative to a Trianion in Alkali Metal Complexes •