

Band Depopulation of Graphene Nanoribbons Induced by Chemical Gating with Amino Groups

Jingcheng Li,[†] Pedro Brandimarte,[‡] Manuel Vilas-Varela,[¶] Nestor
Merino-Díez,^{†,‡} César Moreno,[§] Aitor Mugarza,^{§,||} Jaime Sáez Mollejo,[‡] Daniel
Sánchez Portal,^{⊥,‡} Dimas G. de Oteyza,^{⊥,‡,#} Martina Corso,^{⊥,‡} Aran
Garcia-Lekue,^{*,‡,#} Diego Peña,^{*,¶} and Jose Ignacio Pascual ^{*,‡,#}

[†]*CIC nanoGUNE BRTA, Tolosa Hiribidea 76, 20018 Donostia-San Sebastian, Spain*

[‡]*Donostia International Physics Center (DIPC), 20018 Donostia-San Sebastián, Spain*

[¶]*Centro Singular de Investigación en Química Biolóxica e Materiais Moleculares (CiQUS),
and Departamento de Química Orgánica, Universidade de Santiago de Compostela, 15782
Santiago de Compostela, Spain*

[§]*Catalan Institute of Nanoscience and Nanotechnology (ICN2), CSIC and The Barcelona
Institute of Science and Technology, Campus UAB, Bellaterra, 08193 Barcelona Spain*

^{||}*ICREA Institució Catalana de Recerca i Estudis Avançats, 08193 Barcelona, Spain*

[⊥]*Centro de Física de Materiales MPC (CSIC-UPV/EHU), 20018 Donostia-San Sebastián,
Spain*

[#]*Ikerbasque, Basque Foundation for Science, 48013 Bilbao, Spain*

E-mail: wmbgalea@ehu.eus; diego.pena@usc.es; ji.pascual@nanogune.eu

Abstract

The electronic properties of graphene nanoribbons (GNRs) can be precisely tuned by chemical doping. Here we demonstrate that amino (NH_2) functional groups attached at the edges of chiral GNRs (chGNRs) can efficiently gate the chGNRs and lead to the valence band (VB) depopulation on a metallic surface. The NH_2 -doped chGNRs are grown by on-surface synthesis on Au(111) using functionalized bianthracene precursors. Scanning tunneling spectroscopy resolves that the NH_2 groups significantly up-shift the bands of chGNRs, causing the Fermi level crossing of the VB onset of chGNRs. Through density functional theory simulations we confirm that the hole-doping behavior is due to an upward shift of the bands induced by the edge NH_2 groups.

Keywords

scanning tunneling microscope, density functional theory, chiral graphene nanoribbons, doping, amino, chemical gating, band depopulation

Graphene nanoribbons (GNRs) have emerged as a promising material for nanoelectronics,¹ since they combine many of the extraordinary properties of the parent graphene^{2,3} with a high tunability of their electronic band structure.⁴ Contrary to graphene, GNRs are frequently semiconducting materials, thus offering excellent perspectives for their utilisation as electronic components such as diodes⁵ or transistors.^{1,3,6,7} A fascinating aspect of narrow GNRs is that their electronic properties can be tuned through the precise control of their atomic structure, which can be achieved thanks to the recent development of bottom-up on-surface synthesis (OSS) strategies.^{8,9} These strategies rely on the careful design of suitable molecular precursors with specific shape and chemical composition to steer a step-wise reaction on a metal surface, leading to extended and atomically precise GNRs. An ample library of precursors and reaction pathways has been constructed in the last years, incorporating successful examples of precise control over the GNR's width,^{10–17} orientation and edge topology,^{18–25} which resulted in large variations of their electronic structure. Furthermore, OSS strategies allow to easily combine several precursors on a surface for fabricating complex structures such as atomically sharp graphene heterojunctions,^{26,27} quantum dots,^{28,29} or hybrid systems composed of metal-organic molecules embedded in GNRs,^{30–32} demonstrating

their potential role for GNRs based nanoelectronics.

Since most GNRs are semiconducting, a promising method for tuning their band structure is the electrostatic gating effect induced by doping.³³ Chemical doping of GNRs has been achieved through modification of the molecular precursors to either incorporate substitutional heteroatoms in the carbon backbone^{26,28,34–41} or by adding functional groups at the edges.⁴² However, the band structure of the substitutionally-doped systems could not always be simply related to the pristine one (CB);^{28,35,36,38,39} the embedded heteroatoms have a profound impact on the bands' symmetries and character.⁴³ Instead, chemical functionalization of the GNR edges promisingly behaved as an efficient method to fine-tune the electronic structure of pristine GNRs. In the most simple scenarios, the effect of attached chemical groups can be described as an effective electrostatic gating of the native band structure.^{26,40,42} The magnitude of edge chemical gating is small, to date barely restricted to sub-gap down-shifts of the band structure.

Here, we report on the very efficient chemical gating of amino(NH₂) functional groups attached at the edges of narrow chiral GNRs with a sequence of 3 zigzag and 1 armchair sites ((3,1) chGNRs). The electron donating character of NH₂ is inherited by the GNR, resulting in an effective GNR charging by hole injection and valence band depopulation on a Au(111) surface. Experimentally, NH₂ end groups were substituted at the edges of pristine chGNRs through bottom-up synthesis with modified precursors. Our results satisfactorily demonstrate that the electropositive affinity of the dopant is transmitted to the chGNR carbon backbone,⁴² which on a Au(111) surface becomes strongly hole doped. Through a combination of scanning tunnelling microscopy (STM) and spectroscopy (STS), X-ray photoelectron spectroscopy (XPS) measurements and density functional theory (DFT) calculations we show that the NH₂ groups induce a substantial up-shift of the GNR bands, and cause their VB to cross the Fermi level. The effective depopulation of the VB depends on the number of NH₂ groups per unit cell surviving the reaction, as well as on their relative alignment.

Results/Discussion

Synthesis of the molecular precursor: To functionalize (3,1) chGNRs with NH₂ groups at the edge (named as NH₂-chGNRs), we prepared the molecular precursor 2,2'-dibromo-[9,9'-bianthracene]-10,10-diamine (**3**, Fig. 1) in two steps from 2,2'-dibromo-9,9'-bianthracene (**1**). First, selective double nitration of compound **1** led to the synthesis of compound **2**,

which was isolated with 64% yield. Then, Fe-promoted reduction of bianthracene **2** in acetic acid afforded the GNR precursor **3** with 98% yield (see Supporting Information for details).

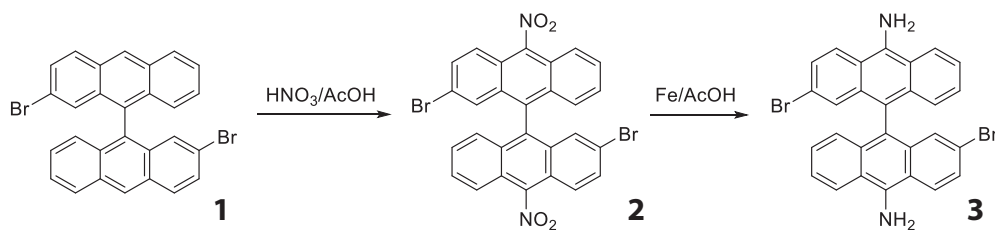


Figure 1: Synthetic route to obtain 2,2'-dibromo-[9,9'-bianthracene]-10,10-diamine (**3**).

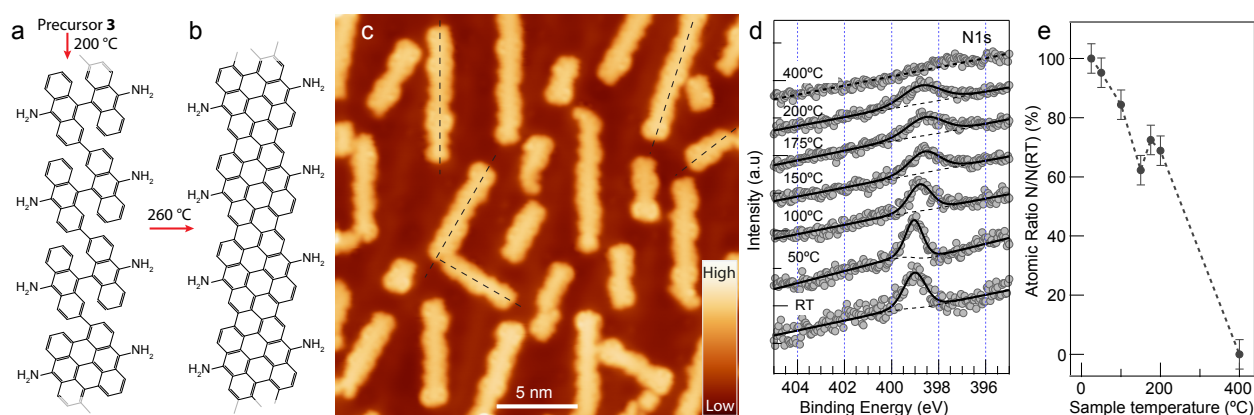


Figure 2: Synthesis of NH₂-chGNRs. (a),(b) Reaction scheme of the polymerization and cyclodehydrogenation of molecular precursors respectively. For simplicity, only one chirality of NH₂-chGNRs is shown here. (c) STM overview image of NH₂-GNRs grown on Au(111), taken at $V_s = 0.2$ V, $I_t = 20$ pA. The dashed lines show the preferred orientations of NH₂-chGNRs on the substrate. Same color scale is used for all the STM images. (d,e) Characterization of NH₂ groups at different annealing temperatures by N1s peaks in XPS measurements.

The molecular precursor **3** was sublimated from a Knudsen cell at 212 °C onto a clean Au(111) substrate kept at room temperature. The precovered sample was gradually annealed first to 200 °C to induce the polymerization of compound **3** through Ullmann-like coupling reactions (Fig. 2(a)). Subsequently, the sample temperature was further raised to 260 °C to trigger the cyclodehydrogenation step causing the planarization of the polymers into NH₂-chGNRs (Fig. 2(b)). Fig. 2(c) shows a STM overview image of the resulting NH₂-chGNRs on a Au(111) surface. The doped chGNRs, as the pristine ones, display preferential orientations on the substrate, but notably shorter than pristine chGNRs,^{21,24} reaching lengths of up to 15 nm. We speculate that the attachment of NH₂ groups lowers the diffusion of precursors and

thus hinders their polymerization processes⁴⁴. The edges of the resulting NH₂-chGNRs are quite inhomogeneous, and their width is larger than pristine chGNRs, suggesting the presence of a certain amount of NH₂ groups attached to the edges (see Supporting Information).

To prove the survival of NH₂ groups during the on-surface synthesis (OSS) process, we characterized the thermal evolution during the OSS with X-ray photoelectron spectroscopy (XPS) measurements (Fig. 2(d,e)). We find a gradual decrease of the N-1s XPS peak intensity with increasing annealing temperatures, which completely vanishes at 400°C. Since the cyclodehydrogenation step completing the synthesis of chGNRs occurs at temperatures slightly above 200°C,^{21,24} the XPS results suggest that about 60% of the NH₂ groups survive the OSS.

Fig. 3(a) shows the constant current STM image of a NH₂-chGNR which shows irregular edges. The superimposed chemical structure of the NH₂-chGNR indicates the loss of some NH₂ groups on the edges. To further prove this, we acquired high resolution STM image of the same GNR with a CO-terminated tip,^{45,46} as shown in Fig. 3(b). The high resolution image resolves the backbone structure of the ribbon as composed of zig-zag edged segments,²⁴ and also shows distinctive triangular features at many of the zigzag edges (indicated by arrows in Fig. 3(b)), which we attribute to surviving NH₂ groups as shown in Fig. 3(a). The carbon atoms with missing NH₂ groups are then passivated by hydrogen atoms during the on-surface reactions⁴⁷. Most of the chGNR segments maintain at least one of the NH₂ edge groups attached at one side, and some appear with both of them intact. From such high resolution images on more than 80 GNRs, we extract that about 75% of the NH₂ groups survived the on-surface reaction. This value is even larger than the one obtained from the XPS results, most probably due to the different annealing processes (see methods). This is a remarkable doping value when compared with the chemical doping surviving the reaction for cyano(CN) functionalization of 7-AGNRs.⁴² We note that edge functionalization in such chiral ribbons is so efficient due to the lower temperatures required for their complete synthesis, in contrast with *e.g.* 7-AGNRs, which need more than 300°C for their OSS.

Electronic structure of doped chGNRs: NH₂ groups have an electron donating character. To study their effect on the electronic properties of the chiral nanoribbons, we recorded differential conductance plots (dI/dV) directly over singly and doubly NH₂-doped segments, and compare them with measurements on pristine chGNRs. As shown in Fig. 3(c), the characteristic 0.7 eV energy gap of pristine chGNRs²⁴ is severely modified on the NH₂-

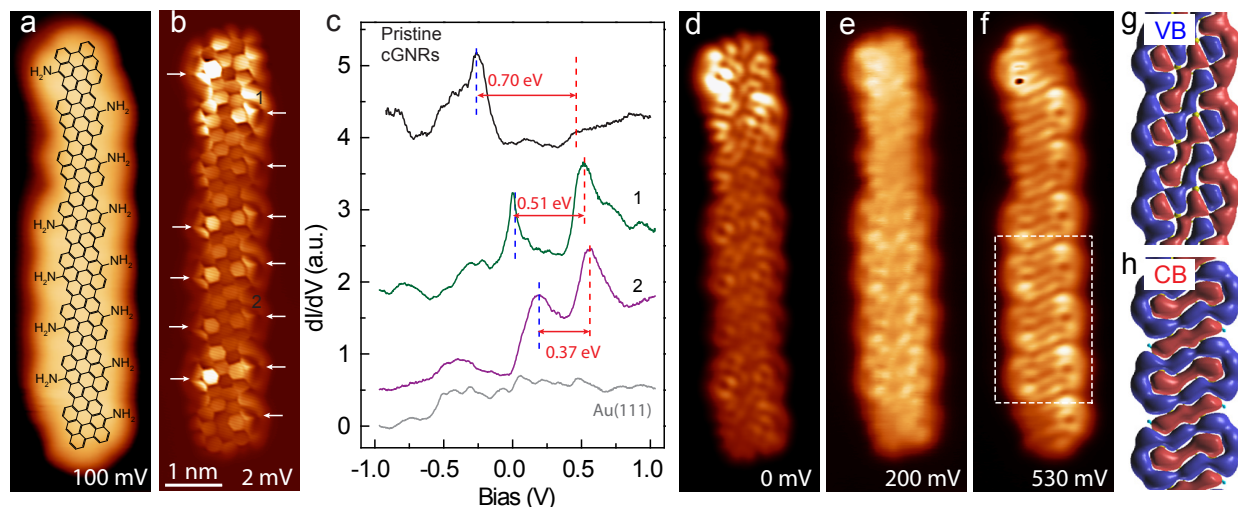


Figure 3: Characterization of the atomic and electronic structures of NH_2 -chGNRs. (a) Constant current image of a single NH_2 -chGNRs with its chemical structure superimposed on top. (b) Constant height current image ($V_s = 2$ mV) of the same NH_2 -chGNRs in (a) taken with CO-terminated tip. (c) dI/dV spectra taken on the locations as indicated with numbers in (b). For comparison, dI/dV spectra taken on pristine chGNRs (black curve on top) are also shown here. Blue and red dashed lines indicate the energy locations of VB and CB, respectively. The values of the band gaps are noted in the figure. (Open-feedback parameters: $V_s = 1$ V, $I_t = 1$ nA; modulation voltage $V_{\text{rms}} = 10$ mV). (d-f) Constant height dI/dV maps of the ribbon in (a) taken at different bias values as indicated in the figures respectively (modulation voltage $V_{\text{rms}} = 10$ mV). All STM images share the same scale bar indicated in (b). (g),(h) DFT simulations of the wave function corresponding to states at the onset of VB and CB (at the Γ point) of NH_2 -chGNRs with all NH_2 groups attached. Red and blue colors represent isosurfaces of positive and negative wave function amplitudes, for an isovalue of $0.015 \text{ \AA}^{-3/2}$.

substituted segments, which exhibit instead two pronounced peaks, indicated by red and blue dashed lines in Fig. 3(c). Over the singly doped segment, one of the peaks is pinned at the Fermi level (zero bias), while the other appears at 500 mV above. On the doubly-doped units, both peaks appear slightly shifted upwards, and their separation noticeably decreases.

To elucidate the origin of the peaks, we acquired dI/dV maps at bias values corresponding to the peak energies, which unveil the spatial distribution of these states (Fig. 3(d-f)). dI/dV maps at 0 mV and 200 mV (the bias values of the lower peaks in each case) resemble the VB shape of pristine chGNRs at the singly- and doubly-doped segments, respectively. In contrast, the dI/dV map at 530 mV (in coincidence with the higher-bias peak in both cases) reproduces a wave-front pattern that is characteristic of the CB of pristine chGNRs.²⁴ Furthermore, these dI/dV maps show very good agreement with the density functional theory

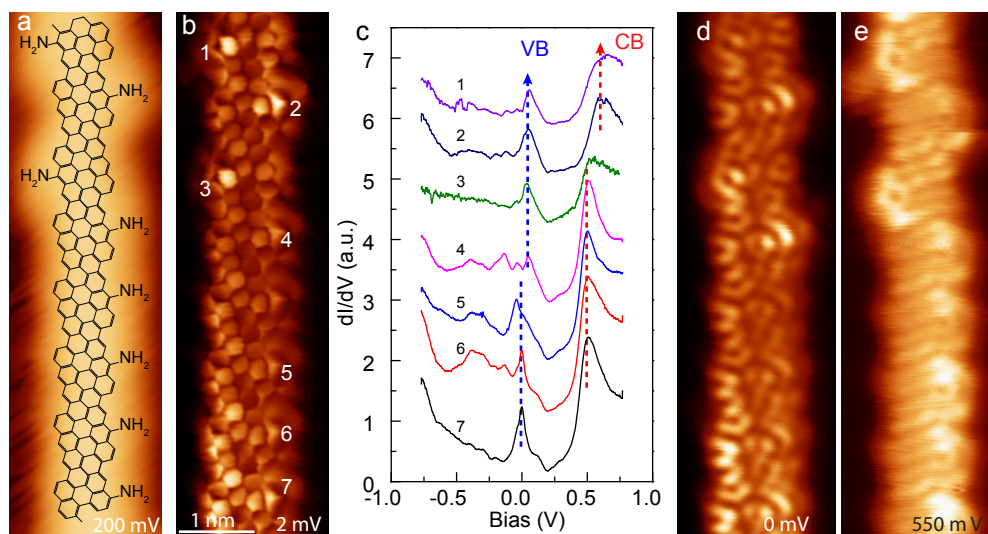


Figure 4: Electronic structures of NH_2 -chGNR with half NH_2 groups lost. (a) Constant current image of a single NH_2 -chGNRs with its chemical structure superimposed on top. (b) Constant height current image ($V_s = 2$ mV) of the same NH_2 -chGNRs in (a) taken with CO-terminated tip. (c) dI/dV spectra taken on the locations indicated with numbers in (b). Blue and red dashed lines indicate the energy onsets of VB and CB, respectively. (Open-feedback parameters: $V_s = 1$ V, $I_t = 1$ nA, modulation voltage $V_{\text{rms}} = 10$ mV). (d-e) Constant height dI/dV maps of the ribbon in (a) taken at different bias values as indicated in the figures respectively (modulation voltage $V_{\text{rms}} = 10$ mV). All STM images share the same scale bar indicated in (b).

simulations of wave function amplitude of valence and conduction bands along a fully NH_2 -doped ribbon (Fig. 3(g,h)). Based on these observations, we ascribe the two dI/dV peaks to the VB and CB onsets of NH_2 -chGNRs, respectively.

Compared to pristine chGNRs, both VB and CB onsets of the NH_2 -doped segments appear shifted upwards, and the VB-CB energy gap is significantly reduced (as labelled in Fig. 3(c)). This upwards shift of the frontier bands can be attributed to a significant hole-doping introduced by the edge NH_2 groups, whose effect increases with the number of groups surviving in each segment. Particularly striking is the alignment of the VB onset with the Fermi-level for the singly-doped segments, and its further shift to ~ 0.18 eV when a second NH_2 is present, unveiling a substantial band depopulation by edge-doping, and underpinning of chGNRs on Au(111).

Intriguingly, the number of NH_2 -dopants per unit is not the only parameter tuning the bands' onset, but also their relative arrangement. This was concluded from the inspection of ribbons with only one edge group per unit. In order to induce the more diluted function-

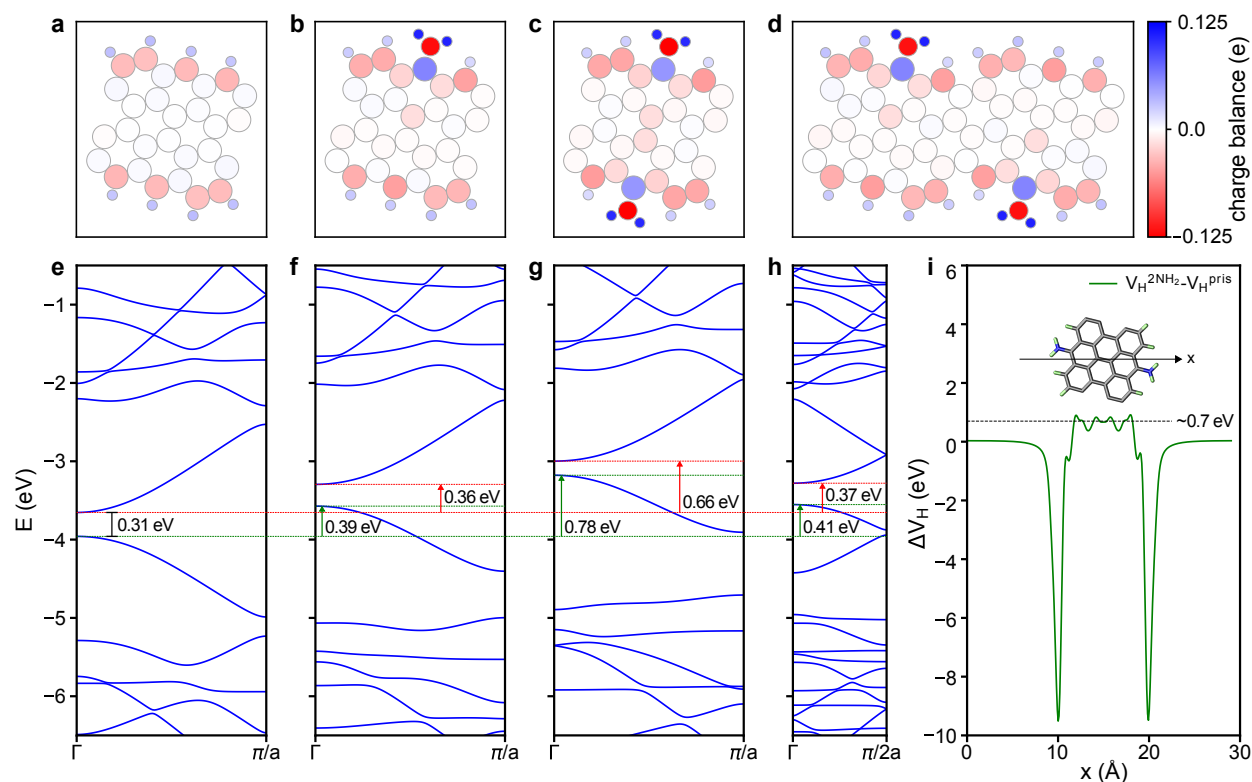


Figure 5: Simulated electronic structure and charges. Charge distribution calculated with a Hirshfeld population analysis for: (a) pristine chGNR, (b) chGNR with NH₂ groups on one side (1a-NH₂-chGNR), (c) NH₂ groups on both sides (2-NH₂-chGNR), and (d) on alternating sides (1b-NH₂-chGNR). Corresponding band structures obtained for (e) pristine chGNR, (f) 1a-NH₂-chGNR, (g) 2-NH₂-chGNR and (h) 1b-NH₂-chGNR. The shift in energy of the top of the VB and the bottom of the CB as compared to the pristine chGNR are marked in green and in red, respectively. (i) Electrostatic potential difference between the 2-NH₂-chGNR and a pristine chGNR, taken at the plane of the ribbon and averaged along the periodic direction (y).

alization observed in Fig. 4, we annealed the sample to a slightly higher temperature during the OSS process (*e.g.* to 280 °C for the data shown in Fig. 4). The result was a reduction of the edge functionalization to about 50%, with a nice preference to maintain one group per unit cell rather than two or none. We found that NH₂ groups could either line-up along the same edge (*e.g.* as between segments 4 and 7 in Fig. 4(a,b)), or alternate at opposite sides (as between 1 and 4 in Fig. 4(a,b)), what had an apparent impact in their corresponding band alignment. While the dI/dV spectra (Fig. 4(c)) and maps (Fig. 4(d) and (e)) appear in every case similar to the ones shown in Fig. 3, both VB and CB peaks in regions with alternating NH₂ groups (spectra 1 to 4 in Fig. 4(c)) appear shifted upwards by ~ 50 meV with respect to the regions where NH₂ groups line up in the same side (spectra 5 to 7).

Theoretical analysis: Further insight into the doping effects of NH_2 groups is obtained by DFT calculations of free-standing nanoribbons (see Methods for details on the simulations). In particular, we studied the band structure of infinite (3,1) chGNRs with the edge structures shown in Fig. 5(a-d), namely, the three experimentally observed scenarios: NH_2 groups only on one side (1a- NH_2 -chGNR), on both sides (2- NH_2 -chGNR), and in alternating positions (1b- NH_2 -chGNR), as well as the pristine chGNRs for comparison. The corresponding band structures (Fig. 5(e-h)) show that, in all three cases, the presence of NH_2 groups at the edge induces an upward shift of the bands with respect to the pristine case. For the singly-doped cases, the VB band shifts about 0.4 eV. When both edges are doped (2- NH_2 -chGNR), the upwards shift of the VB is almost doubled (0.78 eV), showing that it scales with the number of dopants.

The band structure also reproduces the clear reduction of the VB-CB energy gap with doping seen in the experiment. In particular, we find that the band gap of the singly-doped GNR decrease $\approx 10\%$ with respect to the pristine case, while this reduction amounts to $\approx 40\%$ for the doubly-doped species. A similar band gap closure with doping in GNRs was attributed to the extension of the π -conjugated electron cloud outside the graphene backbone with the addition of the functional groups.⁴² On the other hand, we note that the calculated shifts of the frontier bands are larger than the measured ones, and the gap reduction obtained numerically is smaller than in experiments. These discrepancies might be related to screening effects of the metallic substrate, which are not included in our calculations. Yet, our DFT results are in very good qualitative agreement with the experimental measurements.

To explain the origin of the band shifts, we studied the charge redistribution induced by the NH_2 edge groups in the chGNRs. As seen from the Hirshfeld plots in Fig. 5(a-d), the NH_2 groups redistribute their charges by the covalent bonding to the edge and induce a small electron depletion at the connecting C and accumulation right at the middle of the ribbon. Interestingly, the lower electron density at the edge coincides with the higher current of these rings in the chGNR backbone (Fig. 3(a) and Fig. 4(a)). We speculate that the higher current signal over the edge phenyl rings bonding to NH_2 and lower current over the neighbour rings at the center are related to the calculated electron depletion and accumulation, respectively, and to their influence on the effective tunneling.

As a result of the charge redistribution, a strong edge dipole of ~ 3.6 D per unit cell pointing outwards from the ribbon backbone emerges around the NH_2 group. Considering

the edge dipole per unit cell of the pristine chGNR (~ 2.1 D), and since both CH and NH₂ dipoles are pointing in the same direction, we can estimate that each NH₂ group contributes with a dipole of about 2.0 D. This edge dipole increases the internal electrostatic potential, and consequently, shifts their occupied bands upwards. For example, for the 2-NH₂-chGNR case (Fig. 5(i)), we find that the ~ 0.7 eV higher electrostatic potential is the main responsible of the ~ 0.7 eV up-shift of its VB. The result is that the NH₂ groups endow their electron-donating character to the ribbons. On the Au(111) surface, this leads to a substantial hole-doping from the metal, reaching the point where the VB is partially depopulated. We note that this scenario differs from the previously reported case of armchair GNRs doped with CN groups. CN induced a dipole pointing opposite, *i.e.* towards the ribbon's backbone, leading to a total dipole smaller in magnitude and opposite in sign to the one presented here. This caused a small band down shift, contrary to the situation found here, but the semiconducting character remained unchanged.⁴²

Conclusions

Our results demonstrate a chemical-gating strategy of graphene nanoribbons, by which incorporating electron donating chemical groups to the edge of GNRs causes a significant hole-doping on a Au(111) surface. As found previously with CN groups,⁴² the GNR inherits the electron affinity of the attached chemical group. Favored by the lower reaction temperatures for the synthesis of chiral GNRs, we found that a large fraction of NH₂ groups survived the on-surface reaction, and caused the depopulation of its VB, signaling its charging. Based on DFT simulations, we explained the enhanced electron-donating character of the doped ribbons as arising from a substantial charge redistribution induced by the dopant. From these results, we foresee that combination of edge terminations with different electron affinity character can be a promising route to engineer hybrid GNRs with active electronic properties.

Methods/Experimental

The experiments were performed on a home-made low temperature (5 K) STM under ultra-high vacuum (UHV) conditions. The Au(111) substrate was cleaned in UHV by repeated

cycles of Ne^+ ion sputtering and subsequent annealing to 460 °C. The molecular precursor was sublimated at 210 °C from a Knudsen cell onto the clean Au(111) substrate kept at room temperature. Then the sample was first annealed at 200 °C for 15 minutes and then annealed at 260 °C for 5 minutes. A tungsten tip functionalized with a CO molecule was used for high resolution images. The high resolution images were acquired in constant height mode, at very small voltages, and junction resistances of typically 20 M Ω . The dI/dV signal was recorded using a lock-in amplifier with a bias modulation of $V_{\text{rms}} = 10$ mV at 760 Hz. XPS measurements were carried out using a Specs PHOIBOS 150 hemispherical energy analyzer using a monochromatic X-ray source (Al $K\alpha$ line with an energy of 1486.6 eV and 400 W) and energy referenced to the Fermi level. Surface cleanliness was confirmed with x-ray photoelectron spectroscopy (XPS) prior molecules deposition. Organic molecules were sublimated in UHV from a home-made Knudsen-cell type evaporator with an alumina crucible, with the sample kept at room temperature. Then the sample was step-wisely annealed to 400 °C. The sample was kept at each temperature step in Fig. 2d for 10 minutes for the XPS measurements.

Computational Procedures

The optimized geometry and electronic structure of free-standing pristine and NH_2 doped ch-GNRS were calculated using density functional theory, as implemented in the SIESTA code.⁴⁸ The nanoribbons were relaxed until forces on all atoms were < 0.01 eV/Å, and the dispersion interactions were taken into account by the nonlocal optB88-vdW functional.⁴⁹ The basis set consisted of double- ζ plus polarization (DZP) orbitals for all species, with an energy shift parameter of 0.01 Ry. A $1 \times 1 \times 100$ Monkhorst-Pack mesh was used for the k-point sampling of the Brillouin zone, where the 100 k-points are taken along the direction of the ribbon. A cutoff of 300 Ry was used for the real-space grid integrations. The atomic population analysis was performed using the Hirshfeld scheme for partitioning the electron density.⁵⁰ The dipole moments were calculated integrating the valence electron density at each point times the distance of that point to the ribbon backbone (central axis). The corresponding neutralizing contribution due to the nuclei is also added. More details can be find in Carbonell-Sanromà *et al.*⁴².

Supporting Information

The Supporting Information includes the detailed methods for the synthesis of molecular precursor; the characterization of molecular precursor by NMR spectra and the comparison of line profiles between NH₂-chGNRs and pristine chGNRs.

Acknowledgement

We acknowledge financial support from: i) AEI/FEDER-EU through grants no. MAT2016-78293-C6, FIS2017-83780-P (AEI/FEDER,EU), the Maria de Maeztu unit of excellence MDM-2016-0618, and the Severo Ochoa program (ICN2) SEV-2017-0706; ii) the European Research Council (grant agreement no. 635919); iii) the Xunta de Galicia (Centro singular de investigación de Galicia, accreditation 20162019, ED431G/09); iv) the EU project SPRING (863098); v) the European Regional Development Fund (ERDF) under the program Interreg V-A España-Francia-Andorra (Contract No. EFA 194/16 TNI), vi) the CERCA Program/Generalitat de Catalunya, and vii) the Gobierno Vasco-UPV/EHU (project IT1246-19).

References

1. Llinas, J. P.; Fairbrother, A.; Borin Barin, G.; Shi, W.; Lee, K.; Wu, S.; Yong Choi, B.; Braganza, R.; Lear, J.; Kau, N.; Choi, W.; Chen, C.; Pedramrazi, Z.; Dumsclaff, T.; Narita, A.; Feng, X.; Müllen, K.; Fischer, F.; Zettl, A.; Ruffieux, P. *et al.* Short-channel Field-effect Transistors with 9-atom and 13-atom Wide Graphene Nanoribbons. *Nat. Commun.* **2017**, *8*, 633.
2. Han, W.; Kawakami, R. K.; Gmitra, M.; Fabian, J. Graphene Spintronics. *Nat. Nanotechnol.* **2014**, *9*, 794–807.
3. Moreno, C.; Vilas-Varela, M.; Kretz, B.; Garcia-Lekue, A.; Costache, M. V.; Paradinas, M.; Panighel, M.; Ceballos, G.; Valenzuela, S. O.; Peña, D.; Mugarza, A. Bottom-up Synthesis of Multifunctional Nanoporous Graphene. *Science* **2018**, *360*, 199–203.

4. Yazyev, O. V. A Guide to the Design of Electronic Properties of Graphene Nanoribbons. *Acc. Chem. Res* **2013**, *46*, 2319–2328.
5. Kargar, A.; Lee, C. Graphene Nanoribbon Schottky Diodes Using Asymmetric Contacts. *2009 9th IEEE Conference on Nanotechnology (IEEE-NANO)* **2009**, *8*, 243–245.
6. Schwierz, F. Graphene Transistors. *Nat. Nanotechnol* **2010**, *5*, 487–496.
7. Bennett, P. B.; Pedramrazi, Z.; Madani, A.; Chen, Y. C.; De Oteyza, D. G.; Chen, C.; Fischer, F. R.; Crommie, M. F.; Bokor, J. Bottom-up Graphene Nanoribbon Field-effect Transistors. *Appl. Phys. Lett* **2013**, *103*, 1–5.
8. Talirz, L.; Ruffieux, P.; Fasel, R. On-Surface Synthesis of Atomically Precise Graphene Nanoribbons. *Adv. Mater.* **2016**, *28*, 6222–6231.
9. Clair, S.; De Oteyza, D. G. Controlling a Chemical Coupling Reaction on a Surface: Tools and Strategies for On-Surface Synthesis. *Chem. Rev* **2019**, *119*, 4717–4776.
10. Cai, J.; Ruffieux, P.; Jaafar, R.; Bieri, M.; Braun, T.; Blankenburg, S.; Muoth, M.; Seitsonen, A. P.; Saleh, M.; Feng, X.; Mullen, K.; Fasel, R. Atomically Precise Bottom-up Fabrication of Graphene Nanoribbons. *Nature* **2010**, *466*, 470–473.
11. Chen, Y.-C.; De Oteyza, D. G.; Pedramrazi, Z.; Chen, C.; Fischer, F. R.; Crommie, M. F. Tuning the Band Gap of Graphene Nanoribbons Synthesized from Molecular Precursors. *ACS Nano* **2013**, *7*, 6123–6128.
12. Abdurakhmanova, N.; Amsharov, N.; Stepanow, S.; Jansen, M.; Kern, K.; Amsharov, K. Synthesis of Wide Atomically Precise Graphene Nanoribbons from Para-oligophenylene Based Molecular Precursor. *Carbon* **2014**, *77*, 1187 – 1190.
13. Basagni, A.; Sedona, F.; Pignedoli, C. A.; Cattelan, M.; Nicolas, L.; Casarin, M.; Sami, M. Molecules Oligomers Nanowires Graphene Nanoribbons: A Bottom-Up Step-wise On-Surface Covalent Synthesis Preserving Long-Range Order. *J. Am. Chem. Soc* **2015**, *137*, 1802–1808.
14. Kimouche, A.; Ervasti, M. M.; Drost, R.; Halonen, S.; Harju, A.; Joensuu, P. M.; Sainio, J.; Liljeroth, P. Ultra-narrow Metallic Armchair Graphene Nanoribbons. *Nat. Commun* **2015**, *6*, 10177.

15. Zhang, H.; Lin, H.; Sun, K.; Chen, L.; Zagranyarski, Y.; Aghdassi, N.; Duhm, S.; Li, Q.; Zhong, D.; Li, Y.; Mllen, K.; Fuchs, H.; Chi, L. On-Surface Synthesis of Rylene-Type Graphene Nanoribbons. *J. Am. Chem. Soc* **2015**, *137*, 4022–4025.
16. Talirz, L.; Söde, H.; Dumsclaff, T.; Wang, S.; Sanchez-Valencia, J. R.; Liu, J.; Shinde, P.; Pignedoli, C. A.; Liang, L.; Meunier, V.; Plumb, N. C.; Shi, M.; Feng, X.; Narita, A.; Müllen, K.; Fasel, R.; Ruffieux, P. On-Surface Synthesis and Characterization of 9-Atom Wide Armchair Graphene Nanoribbons. *ACS Nano* **2017**, *11*, 1380–1388.
17. Merino-Díez, N.; Garcia-Lekue, A.; Carbonell-Sanromà, E.; Li, J.; Corso, M.; Colazzo, L.; Sedona, F.; Sánchez-Portal, D.; Pascual, J. I.; De Oteyza, D. G. Width-Dependent Band Gap in Armchair Graphene Nanoribbons Reveals Fermi Level Pinning on Au(111). *ACS Nano* **2017**, *11*, 11661–11668.
18. Han, P.; Akagi, K.; Federici Canova, F.; Mutoh, H.; Shiraki, S.; Iwaya, K.; Weiss, P. S.; Asao, N.; Hitosugi, T. Bottom-Up Graphene-Nanoribbon Fabrication Reveals Chiral Edges and Enantioselectivity. *ACS Nano* **2014**, *8*, 9181–9187.
19. Liu, J.; Li, B.-W.; Tan, Y.-Z.; Giannakopoulos, A.; Sanchez-Sanchez, C.; Beljonne, D.; Ruffieux, P.; Fasel, R.; Feng, X.; Mllen, K. Toward Cove-Edged Low Band Gap Graphene Nanoribbons. *J. Am. Chem. Soc* **2015**, *137*, 6097–6103.
20. Ruffieux, P.; Wang, S.; Yang, B.; Sánchez-Sánchez, C.; Liu, J.; Dienel, T.; Talirz, L.; Shinde, P.; Pignedoli, C. A.; Passerone, D.; Dumsclaff, T.; Feng, X.; Müllen, K.; Fasel, R. On-surface Synthesis of Graphene Nanoribbons with Zigzag Edge Topology. *Nature* **2016**, *531*, 489–492.
21. De Oteyza, D. G.; García-Lekue, A.; Vilas-Varela, M.; Merino-Díez, N.; Carbonell-Sanromà, E.; Corso, M.; Vasseur, G.; Rogero, C.; Guitián, E.; Pascual, J. I.; Ortega, J. E.; Wakayama, Y.; Peña, D. Substrate-Independent Growth of Atomically Precise Chiral Graphene Nanoribbons. *ACS Nano* **2016**, *10*, 9000–9008.
22. Rizzo, D. J.; Veber, G.; Cao, T.; Bronner, C.; Chen, T.; Zhao, F.; Rodriguez, H.; Louie, S. G.; Crommie, M. F.; Fischer, F. R. Topological Band Engineering of Graphene Nanoribbons. *Nature* **2018**, *560*, 204–208.

23. Gröning, O.; Wang, S.; Yao, X.; Pignedoli, C. A.; Borin Barin, G.; Daniels, C.; Cupo, A.; Meunier, V.; Feng, X.; Narita, A.; Müllen, K.; Ruffieux, P.; Fasel, R. Engineering of Robust Topological Quantum Phases in Graphene Nanoribbons. *Nature* **2018**, *560*, 209–213.
24. Merino-Díez, N.; Li, J.; Garcia-Lekue, A.; Vasseur, G.; Vilas-Varela, M.; Carbonell-Sanromà, E.; Corso, M.; Ortega, J. E.; Peña, D.; Pascual, J. I.; de Oteyza, D. G. Unraveling the Electronic Structure of Narrow Atomically Precise Chiral Graphene Nanoribbons. *The J. Phys. Chem. Lett* **2018**, *9*, 25–30.
25. Moreno, C.; Panighel, M.; Vilas-Varela, M.; Sauthier, G.; Tenorio, M.; Ceballos, G.; Peña, D.; Mugarza, A. Critical Role of Phenyl Substitution and Catalytic Substrate in the Surface-Assisted Polymerization of Dibromobianthracene Derivatives. *Chem. Mater* **2019**, *31*, 331–341.
26. Cai, J.; Pignedoli, C. A.; Talirz, L.; Ruffieux, P.; Söde, H.; Liang, L.; Meunier, V.; Berger, R.; Li, R.; Feng, X.; Müllen, K.; Fasel, R. Graphene Nanoribbon Heterojunctions. *Nat. Nanotechnol* **2014**, *9*, 896–900.
27. Chen, Y.-C.; Cao, T.; Chen, C.; Pedramrazi, Z.; Haberer, D.; de OteyzaDimas G.; Fischer, F. R.; Louie, S. G.; Crommie, M. F. Molecular Bandgap Engineering of Bottom-up Synthesized Graphene Nanoribbon Heterojunctions. *Nat. Nanotechnol* **2015**, *10*, 156–160.
28. Carbonell-Sanromà, E.; Brandimarte, P.; Balog, R.; Corso, M.; Kawai, S.; Garcia-Lekue, A.; Saito, S.; Yamaguchi, S.; Meyer, E.; Sánchez-Portal, D.; Pascual, J. I. Quantum Dots Embedded in Graphene Nanoribbons by Chemical Substitution. *Nano Lett.* **2017**, *17*, 50–56.
29. Wang, S.; Kharche, N.; Costa Giro, E.; Feng, X.; Mllen, K.; Meunier, V.; Fasel, R.; Ruffieux, P. Quantum Dots in Graphene Nanoribbons. *Nano Lett.* **2017**, *17*, 4277–4283.
30. Li, J.; Merino-Díez, N.; Carbonell-Sanromà, E.; Vilas-Varela, M.; De Oteyza, D. G.; Peña, D.; Corso, M.; Pascual, J. I. Survival of Spin State in Magnetic Porphyrins Contacted by Graphene Nanoribbons. *Sci. Adv* **2018**, *4*, eaaq0582.

- 1
2
3
4
5
6
7
8
9
10
11
12
13
14
15
16
17
18
19
20
21
22
23
24
25
26
27
28
29
30
31
32
33
34
35
36
37
38
39
40
41
42
43
44
45
46
47
48
49
50
51
52
53
54
55
56
57
58
59
60
31. Su, X.; Xue, Z.; Li, G.; Yu, P. Edge State Engineering of Graphene Nanoribbons. *Nano Lett.* **2018**, *18*, 5744–5751.
32. Li, J.; Sanz, S.; Corso, M.; Choi, D. J.; Peña, D.; Frederiksen, T.; Pascual, J. I. Single Spin Localization and Manipulation in Graphene Open-shell Nanostructures. *Nat. Commun.* **2019**, *10*, 200.
33. Krull, C.; Robles, R.; Mugarza, A.; Gambardella, P. Site- and Orbital-dependent Charge Donation and Spin Manipulation in Electron-doped Metal Phthalocyanines. *Nat. Mater.* **2013**, *12*, 337–343.
34. Bronner, C.; Stremlau, S.; Gille, M.; Brauße, F.; Haase, A.; Hecht, S.; Tegeder, P. Aligning the Band Gap of Graphene Nanoribbons by Monomer Doping. *Angew. Chem* **2013**, *52*, 4422–4425.
35. Kawai, S.; Saito, S.; Osumi, S.; Yamaguchi, S.; Foster, A. S.; Spijker, P.; Meyer, E. Atomically Controlled Substitutional Boron-doping of Graphene Nanoribbons. *Nat. Commun* **2015**, *6*, 8098.
36. Cloke, R. R.; Marangoni, T.; Nguyen, G. D.; Joshi, T.; Rizzo, D. J.; Bronner, C.; Cao, T.; Louie, S. G.; Crommie, M. F.; Fischer, F. R. Site-Specific Substitutional Boron Doping of Semiconducting Armchair Graphene Nanoribbons. *J. Am. Chem. Soc* **2015**, *137*, 8872–8875.
37. Nguyen, G. D.; Toma, F. M.; Cao, T.; Pedramrazi, Z.; Chen, C.; Rizzo, D. J.; Joshi, T.; Bronner, C.; Chen, Y. C.; Favaro, M.; Louie, S. G.; Fischer, F. R.; Crommie, M. F. Bottom-Up Synthesis of $N = 13$ Sulfur-Doped Graphene Nanoribbons. *J. Phys. Chem. C* **2016**, *120*, 2684–2687.
38. Durr, R. A.; Haberer, D.; Lee, Y.-L.; Blackwell, R.; Kalayjian, A. M.; Marangoni, T.; Ihm, J.; Louie, S. G.; Fischer, F. R. Orbitally Matched Edge-Doping in Graphene Nanoribbons. *J. Am. Chem. Soc* **2018**, *140*, 807–813.
39. Carbonell-Sanromà, E.; Garcia-Lekue, A.; Corso, M.; Vasseur, G.; Brandimarte, P.; Lobo-Checa, J.; de Oteyza, D. G.; Li, J.; Kawai, S.; Saito, S.; Yamaguchi, S.; Ortega, J. E.; Sánchez-Portal, D.; Pascual, J. I. Electronic Properties of Substitutionally

- Boron-Doped Graphene Nanoribbons on a Au(111) Surface. *J. Phys. Chem. C* **2018**, *122*, 16092–16099.
40. Wang, X. Y.; Urgel, J. I.; Barin, G. B.; Eimre, K.; Di Giovannantonio, M.; Milani, A.; Tommasini, M.; Pignedoli, C. A.; Ruffieux, P.; Feng, X.; Fasel, R.; Müllen, K.; Narita, A. Bottom-Up Synthesis of Heteroatom-Doped Chiral Graphene Nanoribbons. *J. Am. Chem. Soc* **2018**, *140*, 9104–9107.
41. Kawai, S.; Nakatsuka, S.; Hatakeyama, T.; Pawlak, R.; Meier, T.; Tracey, J.; Meyer, E.; Foster, A. S. Multiple Heteroatom Substitution to Graphene Nanoribbon. *Sci. Adv.* **2018**, *4*, 1–8.
42. Carbonell-Sanromà, E.; Hieulle, J.; Vilas-Varela, M.; Brandimarte, P.; Iraola, M.; Baragán, A.; Li, J.; Abadia, M.; Corso, M.; Sánchez-Portal, D.; Peña, D.; Pascual, J. I. Doping of Graphene Nanoribbons via Functional Group Edge Modification. *ACS Nano* **2017**, *11*, 7355–7361.
43. Cao, T.; Zhao, F.; Louie, S. G. Topological Phases in Graphene Nanoribbons: Junction States, Spin Centers, and Quantum Spin Chains. *Phys. Rev. Lett.* **2017**, *119*, 076401.
44. Bronner, C.; Marangoni, T.; Rizzo, D. J.; Durr, R. A.; Jørgensen, J. H.; Fischer, F. R.; Crommie, M. F. Iodine versus Bromine Functionalization for Bottom-Up Graphene Nanoribbon Growth: Role of Diffusion. *J. Phys. Chem. C* **2017**, *121*, 18490–18495.
45. Gross, L.; Mohn, F.; Moll, N.; Liljeroth, P.; Meyer, G. The Chemical Structure of a Molecule Resolved by Atomic Force Microscopy. *Science* **2009**, *325*, 1110–1114.
46. Kichin, G.; Weiss, C.; Wagner, C.; Tautz, F. S.; Temirov, R. Single Molecule and Single Atom Sensors for Atomic Resolution Imaging of Chemically Complex Surfaces. *J. Am. Chem. Soc* **2011**, *133*, 16847–16851.
47. Talirz, L.; Söde, H.; Cai, J.; Ruffieux, P.; Blankenburg, S.; Jafaar, R.; Berger, R.; Feng, X.; Müllen, K.; Passerone, D.; Fasel, R.; Pignedoli, C. A. Termini of Bottom-Up Fabricated Graphene Nanoribbons. *J. Am. Chem. Soc.* **2013**, *135*, 2060–2063.
48. Soler, J. M.; Artacho, E.; Gale, J. D.; García, A.; Junquera, J.; Ordejón, P.; Sánchez-Portal, D. The SIESTA Method for ab Initio order-N Materials Simulation. *J. Phys. Condens. Matter* **2002**, *14*, 2745–2779.

- 1
2
3 49. Klimeš, J.; Bowler, D. R.; Michaelides, A. Chemical Accuracy for The Van der Waals
4 Density functional. *J. Phys. Condens. Matter* **2009**, *22*, 022201.
5
6
7 50. Hirshfeld, F. Bonded-Atom Fragments for Describing Molecular Charge Densities. *Theor.*
8 *Chim. Acta* **1977**, *44*, 129–138.
9
10
11
12
13
14
15
16
17
18
19
20
21
22
23
24
25
26
27
28
29
30
31
32
33
34
35
36
37
38
39
40
41
42
43
44
45
46
47
48
49
50
51
52
53
54
55
56
57
58
59
60

Graphical TOC Entry

