



Cite this: *Nanoscale*, 2023, **15**, 17621

One-year post-exposure assessment of ^{14}C -few-layer graphene biodistribution in mice: single *versus* repeated intratracheal administration†

Antoine Sallustrau,^a Mathilde Keck,^a Peggy Barbe,^b Dominique Georgin,^a Nathalie Fresneau,^{c,a} Stephane Campidelli,^c Baptiste Pibaleau,^d Mathieu Pinault,^d Martine Mayne-L'Hermite,^d Christine Granotier-Beckers,^{e,f} Michel Schlegel,^g Viviana Jehová González,^h Ester Vázquez,^h Denis Servent^b and Frédéric Taran^a

Research on graphene-based nanomaterials has experienced exponential growth in the last few decades, driven by their unique properties and their future potential impact on our everyday life. With the increasing production and commercialization of these materials, there is significant interest in understanding their fate *in vivo*. Herein, we investigated the distribution of ^{14}C -few-layer graphene (^{14}C -FLG) flakes (lat. dim. ~ 500 nm) in mice over a period of one year. Furthermore, we compared the effects of repeated low-dose and acute high-dose exposure by tracheal administration. The results showed that most of the radioactivity was found in the lungs in both cases, with longer elimination times in the case of acute high-dose administration. In order to gain deeper insights into the distribution pattern, we conducted *ex vivo* investigations using μ -autoradiography on tissue sections, revealing the heterogeneous distribution of the material following administration. For the first time, μ -autoradiography was used to conduct a comprehensive investigation into the distribution and potential presence of FLG within lung cells isolated from the exposed lungs. The presence of radioactivity in lung cells strongly suggests internalization of the ^{14}C -FLG particles. Overall these results show the long-term accumulation of the material in the lungs over one year, regardless of the administration protocol, and the higher biopersistence of FLG in the case of an acute exposure. These findings highlight the importance of the exposure scenario in the context of intratracheal administration, which is of interest in the evaluation of the potential health risks of graphene-based nanomaterials.

Received 28th July 2023,
 Accepted 4th October 2023
 DOI: 10.1039/d3nr03711h
rsc.li/nanoscale

Introduction

Graphene based nanomaterials (GBN) have emerged as highly promising nanomaterials across a wide range of research fields.¹ Due to their distinctive and unique properties such as chemical inertness, exceptional mechanical strength, high thermal conductivity, and excellent transmittance, GBN have shown great potential in diverse domains such as energy, electronics, photonics, sensors and biomedical applications.² GBN can now be produced in large quantities, which has led to a significant boost in the development of graphene-based technologies,³ with market estimations expected to reach more than £1100 million by 2025.⁴ Given their increasing exploitation and the potential for both occupational and public exposure, the impact of GBN on health has become a significant concern.⁵ A recent example is the use of graphene-coated face masks produced and distributed all over the world to

^aUniversité Paris Saclay, CEA, INRAE, Département Médicaments et Technologies pour la Santé (DMTS), SCBM, 91191 Gif-sur-Yvette, France.

E-mail: antoine.sallustrau@cea.fr

^bUniversité Paris Saclay, CEA, INRAE, Département Médicaments et Technologies pour la Santé (DMTS), SIMos, 91191 Gif-sur-Yvette, France

^cUniversité Paris-Saclay, CEA, CNRS, NIMBE, LICSEN, 91191 Gif-sur-Yvette, France

^dUniversité Paris-Saclay, CEA, CNRS, NIMBE, LEDNA, 91191 Gif-sur-Yvette, France

^eUniversité Paris Cité, Inserm, CEA, Stabilité Génétique Cellules Souches et Radiations/iRCM, 92265 Fontenay-aux-Roses, France

^fUniversité Paris-Saclay, Inserm, CEA, Stabilité Génétique Cellules Souches et Radiations/iRCM, 92265 Fontenay-aux-Roses, France

^gUniversité Paris Saclay, CEA, Service de Recherche en Matériaux et Procédés Avancés, 91191 Gif-sur-Yvette, France

^hInstituto Regional de Investigación Científica Aplicada (IRICA), Universidad de Castilla-La Mancha, 13071 Ciudad Real, Spain

†Electronic supplementary information (ESI) available. See DOI: <https://doi.org/10.1039/d3nr03711h>

‡These authors contributed equally to this work.



prevent COVID-19 transmission, which were withdrawn in some countries owing to their potential toxic effects on the respiratory tract.^{6–8} In this context, the assessment of their fate *in vivo* remains of great interest.

Since graphene discovery by Novoselov *et al.*,⁹ many graphene related materials have been developed and *in vivo* studies have been conducted to assess their absorption, distribution, and excretion, mainly driven by their physicochemical properties (lateral dimensions, thickness, and carbon-to-oxygen ratio), their functionalization and the route of administration.¹⁰ However, when assessing the potential risks of GBN on human health, the respiratory system is considered as one of the most concerning routes of exposure.¹¹ Following intranasal or tracheal instillation, previous studies have shown that GBN primarily accumulate in the lungs. Over time, signs of clearance can occur as GBN translocate to the blood, either directly from the lungs or through intestinal adsorption. Notably, in the case of tracheal administration, large amounts of the material could be detected in the stomach and intestines, potentially attributed to mechanisms such as mucociliary clearance, swallowing, and redistribution of inhaled GBN to the gastrointestinal tract.^{14,15} Furthermore, intranasal exposure investigations revealed that graphene oxide could be found in the brain and remain there for over 30 days, showing signs of biodegradation over time.^{12,13}

Overall, when considering respiratory exposure, there is evidence that various GBN are able to cross physiological barriers, reaching distant secondary organs and showing in some cases severe lung toxicity.¹⁶ Noteworthy, most of these studies consider single-dose administration. Long-term *in vivo* studies exploring repeated dose administration are still scarce despite their high relevance when considering occupational exposure.^{17,18} In 2015, Shin *et al.* exposed rats to commercial graphene nanoplatelets (GNP) (lateral dimensions: ~500 nm, thickness: 8 nm) through 5 day repeated exposure (6 h per day), using a nose-only inhalation system.¹⁹ The repeated exposure to GNP did not change neither the body weight nor the organ weight of the rats during the recovery period of 28 days. No signs of lung damage were noticed at any time point. In a second study, rats were exposed to GNP (lat. dim.: sizes up to 2 µm, thickness: 20–30 layers) for 6 h per day, 5 days per week, for 4 weeks.²⁰ The animals were sacrificed at 1, 28, and 90 days after exposure. Inhaled materials were found in macrophages, with translocation also to lung lymph nodes, but without inducing any toxicity regardless of the dose applied. More recently, the same group studied repeated exposure of graphene oxide (GO) using the same nose-only inhalation exposure system over 5 days for 6 hours per day.²¹ Exposure impact was assessed at 1, 3 and 21 days by haematological and biochemical analyses and no significant toxic effects of graphene oxide were found. Recently, a reproductive toxicity study was carried out in male mice after the intratracheal instillation of commercial GO once a week for 7 weeks. The increased neutrophil number in the bronchoalveolar lavage fluid suggested pulmonary inflammation.²² In addition to their scarcity, these investigations used histopathological examination to provide

distribution and toxicity evaluation. Undoubtedly, *in vivo* qualitative and quantitative biodistribution analysis of GBN represents a big challenge due to their carbon and oxygen composition, two elements also present as the major part of organic matter.

Radiolabelling has been used in numerous studies to investigate the biodistribution of GBN using tags such as ¹¹¹In and ^{99m}Tc or ¹⁸F, ⁸⁹Zr, and ⁶⁸Ga in tandem with radio imaging techniques.^{23–28} These highly sensitive detection and quantification techniques allow the monitoring of very low concentrations of materials and the study of realistic environmental doses *in vivo* but require surface chemical modification that potentially alters the physicochemical properties of the material or may cause a loss of the tag during *in vivo* circulation. Direct incorporation of carbon isotopes (¹³C, ¹⁴C) into the backbone of carbon-based nanomaterials has proven to be a method of choice to overcome these limitations. Carbon-13 was recently used to structurally label FLG by González *et al.*²⁹ in order to trace and quantify its distribution inside the different cellular compartments of HepG2 cells. ¹³C-labeling is indeed mostly used to study metabolic pathways or for the preparation of internal standards. Nevertheless, in the context of *in vivo* biodistribution investigations, the utilization of ¹⁴C radiolabelling, coupled with exceptionally sensitive detection and quantification techniques, is a reference as it allows following extremely low material concentrations (ranging from ng L⁻¹ to µg L⁻¹) and facilitates the examination of realistic environmental doses. Moreover, carbon-14 is considered in the pharmaceutical industry as one of the gold standards for absorption, distribution, metabolism, and excretion (ADME) studies due to its long half-life (5730 years), ideal for long-term distribution investigations. Our group recently used ¹⁴C-labeled GO to develop a tandem imaging technique combining digital autoradiography (β-imager) with mass spectrometry imaging (MSI), hence providing high resolution and quantitative assessment of its distribution *in vivo*.³⁰ In 2013, a graphitized material was labelled with ¹⁴C to assess and measure its uptake and excretion in *Daphnia magna*.³¹ A year later, Czarny *et al.* conducted a study where carbon nanotubes (CNTs) were labelled with ¹⁴C, revealing their translocation across the air-blood barrier to distant organs.³² Recently, Lu *et al.*³³ utilized ¹⁴C to study the fate of smaller size few layer graphene (FLG) (SLG, lat. dim.: 20–40 nm) and larger size FLG (LLG, lat. dim.: 330–630 nm) in mice over a one-year period after intravenous administration. Their findings demonstrated that liver cells were able to degrade larger ¹⁴C-FLG (LLG) into ¹⁴C-CO₂. In addition to the typical quantification of radioactivity in collected organs, the researchers captured and quantified the exhaled ¹⁴C-CO₂ after intravenous injection in mice.

Considering the need for long-term *in vivo* studies exploring repeated dose administration of GBN, we propose here the assessment of the distribution of FLG over one year, from the whole animal to the cellular level, using ¹⁴C radiolabelling to track the material and compare repeated low dose exposure against single acute dose exposure after intratracheal administration in mice (Fig. 1).



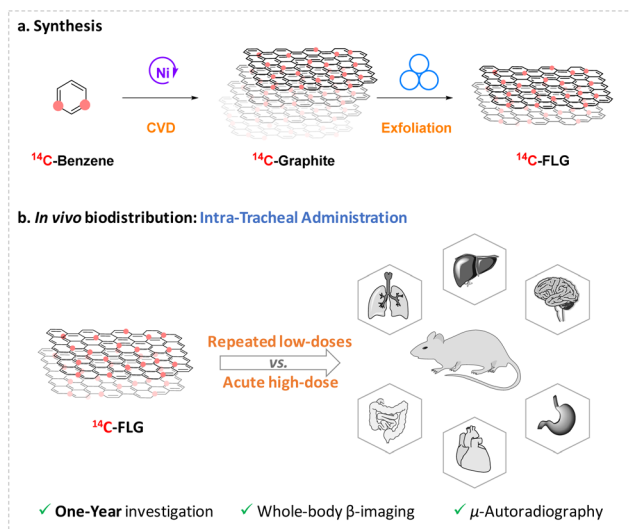


Fig. 1 (A) Strategy for the synthesis of ^{14}C -few-layer graphene (^{14}C -FLG). (B) *In vivo* biodistribution investigation of ^{14}C -FLG in mice over 1 year after intra-tracheal administration. Red circles correspond to the incorporation of ^{14}C into the structure.

Experimental

Chemicals

Organic chemicals and solvents (Sigma-Aldrich) were used without further purification. ^{14}C -benzene (319 mg, 18.5 GBq, 4.77 GBq per mmol, 58 MBq per mg) was purchased from Quotient BioResearch Radiochemicals Ltd with radiochemical and chemical purity > 99.9%. Scintillation fluid (Gold Star, Meridian) was purchased from PerkinElmer, USA. The de-ionized water used in the experiment was prepared using a Millipore Milli-Q ultra-purification system (resistance > 18.2 M Ω cm). Porous nickel foams (95%, 5 cm $\times$ 1 cm $\times$ 1 mm) for the growth of graphite were purchased from Metal Material Co., Ltd.

Materials and methods

Graphite and FLG were analysed by transmission electron microscopy (TEM) or scanning electron microscopy (SEM) (Philips CM 30; CEA Saclay, DEN-LM2E, France, and LVEM5 low-voltage transmission electron microscope, CORDOUAN Technologies, Pessac, France) at an accelerating voltage of 5 kV. For the TEM studies in high resolution mode (HRTEM), we used a Titan ETEM 300 microscope (from FEI/Thermo Fisher Scientific) working at 80 kV and equipped with an image aberration corrector and a K2 *in situ* direct detection camera from Gatan. Raman spectra of ^{12}C -FLG were recorded on an InVia Renishaw micro-spectrometer equipped with a 532 nm point-based laser. In all cases the power density was kept below 1 mW μm^{-2} to avoid laser heating effects. Raman samples were measured in the solid state under ambient conditions. The resulting spectra (after at least 30–40 random

locations on each sample) were fitted with Lorentzian-shaped bands in their D, G and 2D peaks to ascertain band positions, widths and intensities. μRaman spectra of ^{14}C -FLG were recorded on a LabRam HR800 spectrometer with an Olympus microscope and a 50 $\times$ long-range objective. Thermogravimetric analyses (TGA) were performed with a TGA Q50 (TA Instruments) at 10 $^{\circ}\text{C min}^{-1}$ under nitrogen flow, from 100 $^{\circ}\text{C}$ to 800 $^{\circ}\text{C}$. The chemical composition of the non-radiolabelled graphite was evaluated by XPS (X-ray induced photoelectron spectroscopy) analysis using a Kratos Analytical Axis Ultra DLD spectrometer with monochromatic Al KR X-ray radiation ($h\nu = 1486.6$ eV). Milling treatments were carried out in a Retsch PM100 planetary mill using a 50 mL stainless steel milling ball and 10 stainless steel beads. UV-vis spectra were recorded using a Cary 5000 UV-vis-NIR spectrophotometer with 1 cm quartz cuvettes. The measurements of ^{12}C -graphite and ^{12}C -FLG were performed at 271 nm and 660 nm for 12.5 and 48 hours, respectively, at various time intervals in the dual beam mode and using baseline correction. Graphite and FLG concentrations were determined from the optical absorption coefficient at the maximum absorbance, using the formula $A = \alpha \times l \times c$, where l (m) represents the light path length, c (g L^{-1}) corresponds to the concentration, and α ($\text{L g}^{-1} \text{m}^{-1}$) corresponds to the absorption coefficient. The absorption coefficient α was determined to be 21.9 $\text{L g}^{-1} \text{m}^{-1}$ for graphite at 271 nm and 690 $\text{L g}^{-1} \text{m}^{-1}$ for FLG at 660 nm. The relationship between optical absorbance divided by cell length and concentration confirmed Lambert–Beer behavior.

Few-layer graphene synthesis

Few-layer graphene was prepared by exfoliation of graphite obtained by a chemical vapour deposition process (CVD). First developed and optimized on carbon-12, this strategy was employed to synthesize ^{14}C -FLG.

^{12}C -graphite preparation. Graphite was produced by a CVD process specifically developed for the use of benzene as a carbon source. The standard experimental setup involved an aerosol generator, a quartz reactor placed in a furnace, and traps for the exhaust gases and was appropriately modified to be used with small amounts of radioactive liquid carbon precursors. A volume of 3 mL of ^{12}C -benzene was placed in the reservoir and injected for two minutes in a pre-heated oven (1000 $^{\circ}\text{C}$) using H_2 as a vehicle gas (flow: 55 sccm). Five nickel foams of 250 mg each (5 cm $\times$ 1 cm $\times$ 1 mm) were previously placed in the middle of the isothermal zone of the oven as a support for the growth of graphite flakes. At the end of the injection, the oven was cooled down over 10 min (100 $^{\circ}\text{C min}^{-1}$) to come back to room temperature and the nickel foams were collected and then placed in a 3 M HCl solution heated at 100 $^{\circ}\text{C}$ for 5 hours. After complete dissolution of the nickel, the graphite was filtered on a PTFE filter and rinsed with water, ethanol and Et_2O . 25 mg of graphite was obtained.

^{12}C -FLG preparation. A stainless steel jar (50 mL) containing 10 stainless steel balls (1 cm in diameter each) was successively charged with 5 mg of the previously prepared graphite and 300 mg of D-glucose in a ventilated fume hood. The jar



was then closed with an inox clip to avoid spillage during transport and placed in a planetary mill (PM100). The clip was removed and the reaction mixture was stirred for 2 hours at 250 rpm with a spinning rotation change every 3 min with a stop of 3 s each time. In the meantime, a dry dialysis membrane (35 cm, 3.5 kDa) was hydrated in Milli-Q water for 30 min. After the end of the milling step, the reactor was closed again with the inox clip and transported back to the fume hood. 20–25 mL of Milli-Q water was added to the open reactor to disperse graphene flakes. The jar containing the graphene suspension was sonicated for 5 min to recover all the graphene from the walls of the reactor. The reactor was partially dipped into the sonication bath. The graphene suspension was transferred to the dialysis membrane with a 10 mL syringe. The cap and balls were also carefully rinsed. The filled dialysis membrane was placed in a Erlenmeyer flask containing 500 mL of Milli-Q water and heated up to 70 °C for 1.5 h. Every 1.5 h, 9 consecutive times, water was removed, replaced with new Milli-Q water and sonicated for 15 s before heating it up again. The dispersion was transferred to a clear glass flask and decanted for 24 h. The supernatant was recovered and lyophilised, leading to the desired ^{14}C -FLG.

^{14}C -graphite and ^{14}C -FLG preparation. Starting from 0.3 mL of ^{14}C -benzene (18.5 GBq, 4.77 GBq mmol $^{-1}$, 58 MBq mg $^{-1}$) diluted with 2.7 mL of ^{12}C -benzene, an identical protocol was followed, providing 25 mg of ^{14}C -graphite (135 MBq, 5.4 MBq mg $^{-1}$). From 5 mg of this ^{14}C -graphite, an aqueous suspension of ^{14}C -FLG (18.12 MBq, 607 kBq mL $^{-1}$, 5.4 MBq mg $^{-1}$) was obtained before being lyophilised.

^{14}C -FLG dispersion preparation. The suspensions used for *in vivo* administrations were prepared by the dispersion of the lyophilised ^{14}C -FLG in Milli-Q water followed by a 5 min sonication treatment.

In vivo experiments

Animals. Female Balb/cJ mice aged 6 weeks (from Janvier Laboratory, St Berthevin, France) were housed in groups of 4–8 animals in polysulfone cages under standard conditions: room temperature (20 ± 2 °C), hygrometry (55 ± 10%), light/dark cycle (12 h/12 h), air replacement (15–20 volumes per hour), drinking water and a pelleted maintenance diet (LASQDiet® Rod16, Genobios, France) *ad libitum*. All procedures were validated by the local animal ethics committee (CETEA DSV – comité n° 44) and the authorization APAFIS#32015-2021061612169288 was obtained from the Ministry of National Education, Higher Education and Research (France).

Tracheal administration. One-hundred and five mice were randomized in three groups corresponding to the administration of vehicle ($n = 25$ mice) or an aqueous suspension of ^{14}C -FLG either at a unique high dose of 13 µg (74 kBq, $n = 25 + 15$ mice) or 0.65 mg per kg b.w. for the acute administration or at a low dose of 3.4 µg (18.5 kBq, $n = 25 + 15$ mice) or 0.17 mg per kg b.w. that was repeated once a week for 4 weeks for the chronic administration. The vehicle or suspension of ^{14}C -FLG was administered by the tracheal route using an endotracheal

intubation kit (Kent Scientific, Torrington, CT, USA) with a final volume of 20 µL per mouse.

Sample preparation. Mice were sacrificed at 1, 30, 90, 180, and 360 days after the administration by cervical dislocation or, for whole-body sections, with an intraperitoneal injection of Exagon® (pentobarbital, Gedeon Richter, Austria).

Blood and plasma. Before sacrifice, blood sampling *via* the retro-orbital route was performed under anesthesia (2% of iso-flurane) on five distinct mice of each group at each time of sacrifice. The plasma was separated by centrifugation of the blood in a heparin tube at 1500g for 15 min.

Urine and faeces. The urine and faeces of eight mice administered with ^{14}C -FLG were collected for 30 days following administration thanks to individual metabolic cages (Techniplast, France). The urine and faeces were stored at –20 °C before analysis.

Organs. At 1, 30, 90, 180, and 360 days after the administration of ^{14}C -FLG, five mice of both groups were sacrificed by perfusion with 0.9% NaCl (10 mL per mouse, 200 mL h $^{-1}$) for collection of organs (brain, thymus, heart, trachea, lungs, liver, spleen, stomach, stomach bowl, small intestine, small intestine content, large intestine, large intestine content, and kidneys). The organs were immediately frozen at –80 °C by immersion in a mixture of dry ice and isopentane to prevent redistribution of the ^{14}C -FLG. The organs were stored at –20 °C before analysis.

Whole-body and tissue sections. At 1, 30, 90, 180, and 360 days after the administration of ^{14}C -FLG, three mice of both groups were sacrificed for whole-body and tissue sections. The carcass or the organs were frozen and sectioned at 30 µm and 20 µm thickness, respectively, with a CM3050 S cryostat (Leica Biosystems, Wetzlar, Germany) in which the chamber temperature was adjusted to –18 °C and the object temperature to –20 °C. The cryotome was carefully cleaned with a water/ethanol solution between each sample to remove any traces of radioactivity and to avoid contamination between different mice. The whole-body or tissue sections were placed on glass slides and were kept at room temperature for 1 day in the presence of silica gel to ensure complete drying for radio imaging.

Primary culture cells. At 1 and 30 days after the administration of ^{14}C -FLG, three mice of both groups were sacrificed by perfusion with 0.9% NaCl (10 mL per mouse, 200 mL h $^{-1}$) for lung collection. Almost whole lungs were frozen and sectioned at 20 µm thickness, as described above. A small part of the lungs was cut in small pieces and digested at 37 °C for 15 min with 0.25% trypsin (Gibco, Thermo Fisher Scientific, Massachusetts, USA). The digested organs were filtered on a 100 µm cell strainer (Corning, Dutscher, Issy-les-Moulineaux, France). Then, 100 µL of cells were diluted in 900 µL of RPMI medium plus 10% fetal bovine serum and 1% penicillin–streptomycin (Gibco, Thermo Fisher Scientific, Massachusetts, USA). Cells were plated on LabTek chambers (Thermo Fisher Scientific, Massachusetts, USA) and incubated overnight at 37 °C with 5% CO $_2$. The plated cells from lungs were immersed for 5 min in 4% formalin for fixation.



Analysis of clinical chemistry, elimination and organ distribution

Clinical chemistry. The collected plasma samples were freshly analyzed on Piccolo Xpress® (from Abaxis Europe GmbH, Griesheim, Germany) with an AmLyte 13 disk to determine the following parameters: glucose, blood urea nitrogen, bilirubin, albumin, alanine aminotransferase, aspartate aminotransferase, creatine kinase, sodium, potassium, and calcium.

Elimination via urine and faeces. Whole urine was quantified using a liquid scintillation analyzer Tri-Carb 4910 TR (PerkinElmer, Villebon-sur-Yvette, France). Due to the strong background induced by faeces, several samples of 30 mg of faeces were quantified using a liquid scintillation analyzer Tri-Carb 4910 TR. Then, the amount of radioactivity in all faeces was extrapolated from the data of 30 mg count relative to the total weight of faeces harvested.

Organ distribution. Organs were shredded with Precellys® (Ozyme, Saint-Cyr-l'École, France) and radioactivity contained in the whole organs was quantified using a liquid scintillation analyzer Tri-Carb 4910 TR (PerkinElmer, Villebon-sur-Yvette, France).

Autoradiography from the whole body to cells

β-Imager. A high-performance autoradiography imager (β-imager 2000, Biospace Lab, Paris, France) allowing real-time radioactive imaging through direct β-particle counting and absolute radioactivity quantification (detection threshold of 0.01 cpm per mm² for ¹⁴C) was used to measure radioactivity in dried whole-body sections. The time of acquisition on the β-Imager was between 1 and 4 h for whole-body sections.

μ-Autoradiography

¹⁴C-FLG was detected in tissue sections and cells by μ-autoradiography following the detailed protocol described in the ESI.† Briefly, slides were dipped in a liquid radiosensitive emulsion (NTB, Carestream), and after drying under a humidified atmosphere, the slides were exposed for 4 days at 4 °C. The slides were then developed in Kodak D-19 (Kodak) and then stained with Mayer's hemalum (Merck).

Results and discussion

With the aim of assessing the long-term effects of repeated graphene exposure and comparing them to those of single exposure, we conducted an investigation into the distribution of ¹⁴C-labelled few-layer graphene (¹⁴C-FLG) in mice. Our approach adopted a comprehensive “top to bottom” strategy, analysing the distribution from the whole animal down to the cellular level. To produce ¹⁴C-FLG, we developed a two-step methodology. Initially, ¹⁴C-labelled graphite was obtained through a chemical vapor deposition (CVD) process. Subsequently, a mechano-chemical exfoliation step was employed to achieve the desired ¹⁴C-FLG. The methodology,

previously established with carbon-12 materials, was effectively applied to synthesize ¹⁴C-FLG.

Preparation of ¹⁴C-graphite

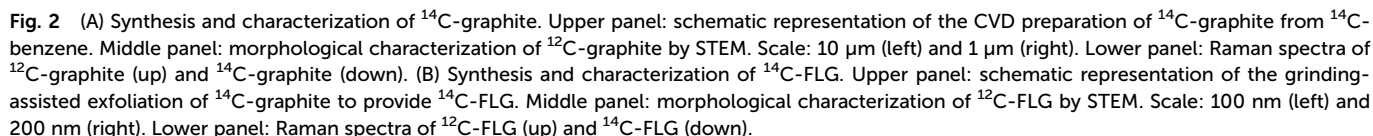
¹⁴C-graphite synthesis proceeded first with the injection of diluted ¹⁴C-benzene (3 mL) in the CVD system. It was pushed using argon pressure (2 bars) from the tank and then injected into the rest of the device (Fig. S1†) using an injection head connected to a computer-assisted control system. This system adjusts the injection frequency and the valve opening time to control the amount of injected benzene. Once injected, the carbon source was carried by a flow of hydrogen regulated by a flow meter and then heated in the vaporiser. At this stage, the fine droplets of ¹⁴C-benzene are transformed into gas, which was sent into the quartz tube and heated to 1000 °C by means of a tube furnace with a 12 cm isothermal zone. Based on the work of Chen *et al.*,³⁴ five nickel foams, playing the role of a support for the growth of graphite, were placed in the middle of this isothermal zone. Out of the oven, the gas passes through a water cooler and a bubbler filled with acetone, which isolates the device from ambient air and allows the gas flow to be monitored. Finally, a stream of air was directed at the walls of the quartz tube to control the cooling rate of the furnace (up to 100 °C min⁻¹). After cooling down to room temperature, the nickel foams were collected and placed in a 3 M HCl solution heated at 100 °C for 5 hours. After complete dissolution of the nickel, the ¹⁴C-graphite was filtered, rinsed and dried, providing 25 mg of ¹⁴C-graphite (135 MBq, 5.4 MBq mg⁻¹).

Physicochemical characterization of ¹²C-graphite and ¹⁴C-graphite was performed to confirm the consistency of the production method and the quality of the engineered materials. Electron microscopy (TEM, SEM, STEM) analysis of ¹²C-graphite performed directly after CVD showed flakes of graphite covering the entire structure of the nickel foam (Fig. S2†). After dissolution and removal of the nickel, the recovered ¹²C-graphite appeared as micrometre scale particles (>1 μm) (Fig. 2A and S3†).

All graphite samples analysed by Raman spectroscopy presented the characteristic bands of graphene-based materials: D, G and 2D. The D band stands for the presence of defects in the structure since it is related to Csp³ carbons and the G band corresponds to Csp² carbons and the stretching of the C–C bond in graphitic materials. Then, the I_D/I_G ratio can be used to evaluate the defects in the structure and I_{2D}/I_G can be related to the thickness of the material. The 2D band can also be used to assess and determine the number of layers using the full width at half-maximum (FWHM)³⁵ of the band, *e.g.*, a narrow 2D band indicates a low number of layers.³⁶

Just after the chemical vapour deposition, ¹²C-graphite deposited on nickel presented a very similar Raman spectrum (Fig. S4†) to ¹²C-graphite without nickel. Interestingly in the latter case, the 2D band is slightly shifted and the intensity of the D peak decreased, which indicated that the structural quality of the graphene was preserved and the disorder effects associated with the nickel have disappeared. Graphite on





XPS analyses were performed on ^{12}C -graphite only, to avoid any radioactive contamination of the equipment. The deconvolution C 1s spectrum of ^{12}C -graphite exhibits one major asymmetric C 1s peak at 284 eV and minor ones at 286 and

Nanoscale, 2023, 15, 17621-17632 | 17627

from the animal down to the cellular level. The mice used were outbred Balb/cJ mice, which are the most practical and commonly used species for pre-clinical toxicity studies. The protocol was conducted in female mice (nulliparous and non-pregnant), which are generally considered more sensitive than male mice in toxicity studies.

Biodistribution and elimination

Two groups of female Balb/cJ mice were exposed to ^{14}C -FLG by tracheal administration to either a unique high dose of 13 μg (74 kBq) or a low dose of 3.4 μg (18.5 kBq) that was repeated once a week for 4 weeks. The distribution of ^{14}C -FLG in the mice was subsequently monitored for one year after the end of the administration period, corresponding to the completion of the fourth and final repeated administration. The mice were sacrificed at day 1, 30, 90, 180, and 360 post-exposure. At each of these times, radioactivity was imaged on whole-body sections using a β -imager and quantified using scintillation counting in the following organs: brain, thymus, heart, trachea, lungs, liver, spleen, stomach, stomach bowl, small intestine, small intestine content, large intestine, large intestine content, and kidneys. Radioactivity was also quantified in urine and faeces to follow the elimination of FLG.

Whole-body β -imaging demonstrated that radioactivity remained predominantly concentrated in the lungs, even after 360 days, irrespective of the administration protocol (Fig. 3A, S21 and S22†). Upon further quantification of all collected and shredded organs, it was confirmed that the lungs contained the majority of radioactivity, accounting for $72 \pm 3\%$ after acute administration and $69 \pm 5\%$ after repeated administration. This trend was consistent when considering the total organs, with the lungs comprising $80 \pm 8\%$ after acute administration and $71 \pm 5\%$ after repeated administration (Fig. 3B). Lung distribution of large nanoparticles (size $> 1 \mu\text{m}$) and FLG had also been demonstrated after intravenous administration,³³ showing that these nanoparticles could be trapped in the capillary beds of lungs. FLG exhibited reduced distribution to other organs, such as the trachea, stomach, and large intestine, with these organs containing less than 5% of the administered dose only one day after the administration. In agreement with previous results,^{14,15} tracheal instillation of FLG leads to mucociliary clearance and to the redistribution of inhaled FLG to the gastrointestinal tract. However, conversely to several other reports, no radioactivity was detected in the brain, even at low levels (Fig. S22 and S23†), which suggests that FLG flakes of around 500 nm administered *via* the tracheal route are unlikely to pass through the blood–brain barrier.

Interestingly, when examining the amount of radioactivity remaining after a year, we observed distinct differences in the biopersistence of FLG following acute *versus* repeated administration. For instance, a significant decrease in the radioactivity content was observed in the lungs only after 180 days following acute exposure, with $25 \pm 5\%$ of the administered dose persisting in all organs after 360 days (Fig. 3B, left panel and Fig. S22†). Surprisingly, the content of FLG in the lungs

decreased much faster after repeated exposure, by 57, 72, and 92%, at 30, 90, and 180 days post-exposure, respectively (Fig. 3B, right panel and Fig. S23†). During the six remaining months, the radioactivity found in the lungs after repeated exposure stabilised to $8 \pm 3\%$, indicating a bioaccumulation of the graphene in that organ. Dose and frequency of FLG exposure play a crucial role in influencing the biodistribution of FLG in the lungs.

No radioactivity was found in the kidneys at any time point (Fig. S22 and S23†), and only 0.01–0.10% of the injected dose was recovered in urine (Table S3†), suggesting no clearance of the ^{14}C -FLG *via* glomerular filtration. While no significant radioactivity was detected in urine, most of the elimination was through faeces, peaking at 18 ± 9 and $23 \pm 0.4\%$ at 1 day after acute and repeated administration, respectively (Fig. 3C, left panel). Then, 38 ± 3 and $41 \pm 1\%$ of radioactivity was eliminated through faeces in the next 30 days after acute and repeated administration, respectively.

Tissue and cellular localization

Following the investigation of FLG distribution at both the whole-body and organ levels, we further explored the sub-organ distribution of ^{14}C -FLG on tissue slices by μ -autoradiography. This technique is used to provide a subcellular assessment of the localisation of radiolabelled compounds.^{44,45} Silver grains (black spots on images) emerge as a result of the interaction between the emitted β -particles from the radioelement (in this case, ^{14}C) and a radiosensitive emulsion containing silver bromide nanocrystals.⁴⁶

To visualise ^{14}C -FLG at the cellular level, we first optimized the autoradiography conditions on HeLa cells (Fig. S25†). We then applied them on tissue sections from exposed mice. Lung and liver slices were analysed 30 and 90 days after ^{14}C -FLG administration, and acute exposure and repeated tracheal exposure were compared (Fig. 4A). To optimize the resolution of the μ -autoradiography images, different times of exposure to the radiosensitive nuclear emulsion were investigated (1, 4 and 15 days). After 4 days of exposure of lung and liver slices (from mice 30 days after administration) to the emulsion, μ -autoradiography images showed a heterogeneous distribution of radioactivity in the lungs with larger spots of silver grains in the central region of the lungs and smaller ones in the peripheral region (black dots in Fig. 4A, S26 and S28†). These observations may be attributed to FLG tendency to aggregate at the entrance of the lungs shortly after administration. In line with the quantification of radioactivity in organs, no radioactivity was observed in liver sections as indicated by the absence of black dots in Fig. S27 and S29† for both exposure modes. Additionally, we examined lung slices from mice after 90 days of both acute and repeated tracheal administrations of ^{14}C -FLG. Over time, we observed fewer, smaller, and more dispersed aggregates in the lung slices. However, lung slices from mice after 90 days were not further investigated due to the extended exposure time to the emulsion, which exceeded the period needed to yield a significant silver grain signal (greater than 15 days).



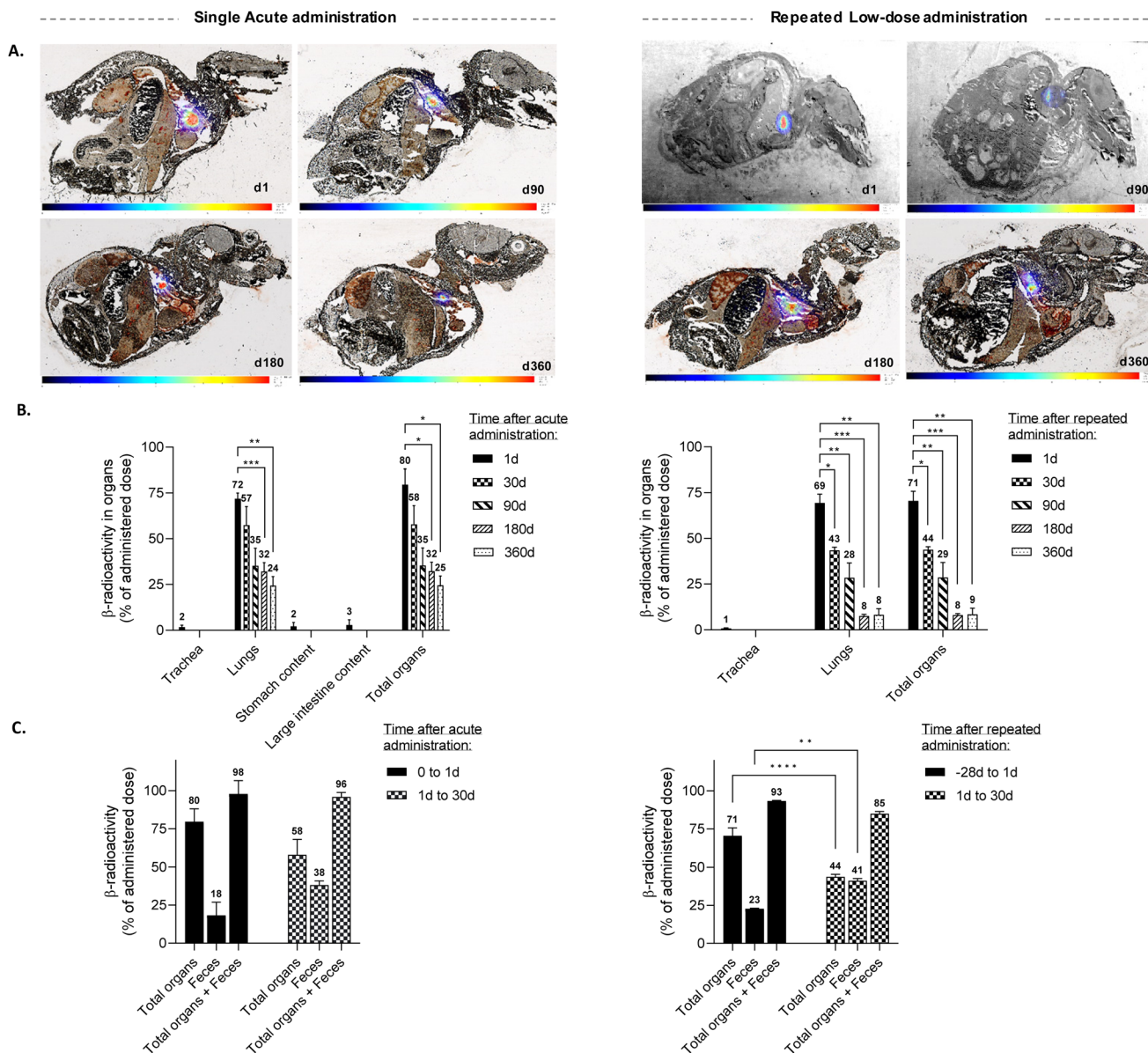


Fig. 3 (A) Whole-body slices (30 μ m thick) of female Balb/cJ mice observed by β -imaging after ^{14}C -FLG single (left panel) or repeated (right panel) tracheal administration over 360 days. A slice is representative of 6 slices per mouse of $n = 3$ mice per time and per group. (B) Mean $\pm$ SEM of the percentage of organ β -radioactivity administered dose over time following ^{14}C -FLG single (left panel) or repeated (right panel) tracheal administration. Organs with $<1\%$ of administered dose are not presented. $n = 5$ female Balb/cJ mice per time and per group. Mixed-effect model followed by Dunnett's multiple comparison test; * $p < 0.05$, ** $p < 0.01$, *** $p < 0.001$ compared to 1 day for each organ. (C) Mean $\pm$ SEM of the percentage of β -radioactivity administered dose contained in faeces and total organs over 30 days following ^{14}C -FLG single (left panel) or repeated (right panel) tracheal administration. $n = 8$ mice per time and per group.

To obtain a more in-depth understanding of the material fate, we decided to examine primary culture cells from the lungs and determine if we could detect radioactivity within these cells using μ -autoradiography. As the lungs are the primary targeted organs, primary culture cells were derived from lung tissue samples collected 30 days after administration. Lung tissues were mechanically dissociated, enzymatically digested, and then subjected to centrifugation to isolate lung cells, which were subsequently cultured. Autoradiography

revealed that primary lung cells consistently exhibited radioactivity and that ^{14}C -FLG is mainly located in their cytoplasm (Fig. 4B, S30 and S31†), suggesting the internalization of the nanomaterials into the cells, as it seems unlikely that the material would have remained solely on the cell surface during the primary culture cell mechanical and enzymatic treatments.

Noteworthy, the studied lung cells associated with radioactivity exhibit a morphological aspect more indicative of alveolar macrophages rather than epithelial cells. However, to



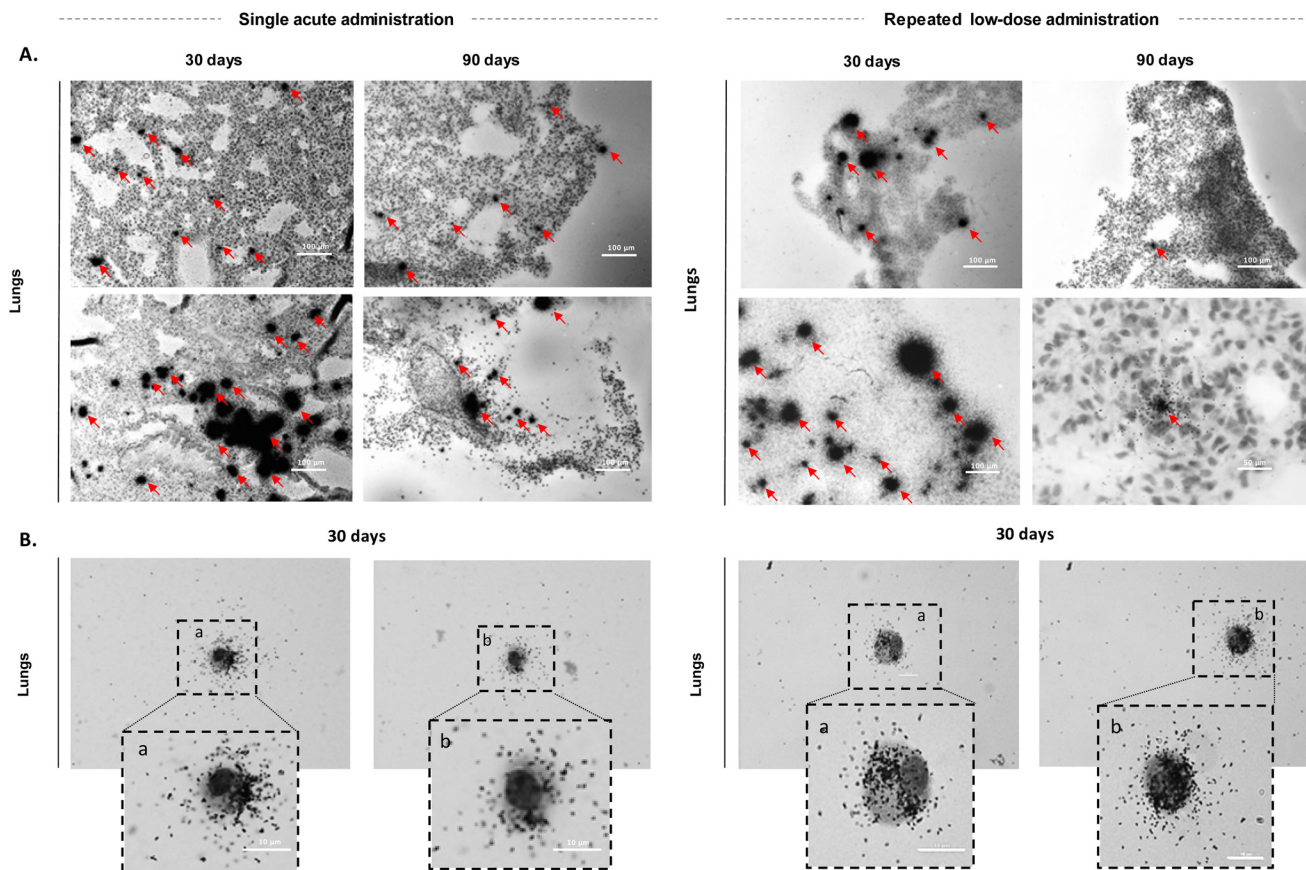


Fig. 4 (A) μ -Autoradiography images of lung tissue sections of exposed organs at 30 and 90 days after ^{14}C -FLG single (left panel) or repeated (right panel) tracheal administration. The upper images focus on the peripheral region of the lung and the lower images focus on the central region. The red arrows correspond to the presence of radioactivity. Scale: 100 μm . (B) μ -Autoradiography analysis of primary culture cells of the lungs after ^{14}C -FLG single (left panel) or repeated (right panel) tracheal administration. Scale: 10 μm .

confirm this suggestion, it would be essential to achieve accurate identification using specific cellular markers. Interestingly, a very recent comparative study of the impact of several GBN *in vivo* reported evidence of FLG (lat. dim. ~ 200 nm) degradation through macrophages and neutrophils in the lungs, and suggested that there may be a size threshold (<1 μm) for more efficient degradation of GBMs by macrophages and faster elimination from the lungs.⁴⁷

Clinical chemistry

The toxicological effect of ^{14}C -FLG exposure was also investigated in the plasma from mice at different times after acute and repeated administration. Hepatic, renal, pancreatic, metabolic, and muscle parameters, as well as electrolytes, have been quantified over one year.

Regardless of the administration protocol, no significant signs of perturbations of plasma chemistry were detected over time (Fig. S32 and S33†). There was a non-significant increase in alanine and aspartate aminotransferases during the first month, which decreased over time for repeated administration but not for acute administration. Albumin showed a slight increase after one year of acute administration. Significant

variations in creatine kinase were observed, attributed to sample haemolysis. However, overall, electrolytes remained within standard thresholds, suggesting the absence of systemic toxicity over time under these conditions, regardless of the frequency of administration. However, local lung toxicity, where FLG is mostly accumulated, cannot be excluded. This should be confirmed using specific biomarkers for lung toxicity. Further studies should be performed to explore this hypothesis.

Conclusions

With the extended production and commercialization of graphene related materials, the assessment of their fate *in vivo* remains a main public health priority, particularly when considering the effect of repeated low-dose exposure over a long period of time.

We showed that FLG was mainly present in the lungs of mice when an acute high dose or repeated low doses of ^{14}C -FLG were administered, in agreement with previous reports. We have not been able to detect even low doses of radioactivity in the brain, indicating that such a material



(500 nm and 6 layers thickness) is unlikely to pass through the blood–brain-barrier. The biopersistence of FLG differs between acute high-dose and repeated low-dose administrations. Actually, the content of FLG in the lungs showed a more rapid decrease after repeated low-dose exposure, indicating reduced biopersistence compared to that in acute administration. Furthermore, we showed the long-term accumulation of the material in the lungs over one year, regardless of the administration protocol, which was not correlated to any significant perturbations of the plasma chemistry under these conditions but without excluding potential lung toxicity. Our findings also described the elimination of graphene nanomaterials of around 500 nm through faeces, but not *via* glomerular filtration. Finally, ^{14}C -FLG materials were found to be present in lung cells and more particularly within the cytoplasm, suggesting a possible internalisation of the nanomaterials.

Although the physico-chemical characteristics of the material (size, thickness, carbon/oxygen ratio) play a dominant role, it is clear that the *in vivo* fate of FLG after intratracheal administration shows similar biodistribution trends to GO with a clear and major accumulation in the lungs that decreases over time. However, in contrast to GO, FLG is demonstrated to be eliminated less rapidly in the long term. Both have been found to be internalized in lung cells, most probably by macrophages which have been shown to participate actively in the degradation of the materials.^{33,47}

In summary, this work provides an initial glimpse into the long-term impact of repeated FLG exposure through the respiratory tract and shows that radiolabelling and autoradiography are powerful techniques to assess its distribution from the animal to the cell level. Further and more comprehensive analyses would be interesting to gain a deeper understanding of the degradation mechanisms of FLG *in vivo*.

Author Contributions

A. S. and M. K. contributed equally to this work. The manuscript was written through contributions of all authors. All authors have given approval to the final version of the manuscript.

Ethical statement

All animal procedures were performed in accordance with Care and Use of Laboratory Animals of the Directive 2010/63/EU of the European Parliament, which has been approved by the Ministry of Agriculture, France. The project was approved by the French Ethics Committee CEEA (*Comité d'Ethique en Expérimentation Animale*) which is responsible for overseeing the provisions for the care and well-being of animals used for research and educational purposes at CEA-Saclay and serves the public by ensuring compliance with all legal and ethical standards regarding the use of vertebrate animals in research and teaching at CEA-Saclay.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

This work was supported by the Commissariat à l'Energie Atomique et aux Energies Alternatives. This project has also received funding of the GrapheneCore3 project from the European Union's Horizon 2020 Research and Innovation Program under grant agreement no. 881603. This study forms part of the Advanced Materials programme and was supported by MCIN with funding from the European Union NextGenerationEU (PRTR-C17.I1) and by the Junta de Comunidades de Castilla-La Mancha. The authors thank G. Patriarche for the support and help with the transmission electron microscopy.

References

- 1 A. C. Ferrari, F. Bonaccorso, V. Fal'ko, K. S. Novoselov, S. Roche, P. Bøggild, S. Borini, F. H. L. Koppens, V. Palermo, N. Pugno, J. A. Garrido, R. Sordan, A. Bianco, L. Ballerini, M. Prato, E. Lidorikis, J. Kivioja, C. Marinelli, T. Ryhänen, A. Morpurgo, *et al.*, *Nanoscale*, 2015, 7, 4598.
- 2 G. Reina, J. M. Gonzalez-Dominguez, A. Criado, E. Vázquez, A. Bianco and M. Prato, *Chem. Soc. Rev.*, 2017, 46, 4400.
- 3 T. Reiss, K. Hjelt and A. C. Ferrari, *Nat. Nanotechnol.*, 2019, 14, 907.
- 4 Global Graphene Market Size and Forecast To 2025. BBC Research. (2018) <https://www.bccresearch.com/partners/verified-market-research/global-graphene-market-size-and-forecast.html>.
- 5 A. B. Seabra, A. J. Paula, R. de Lima, O. L. Alves and N. Duran, *Chem. Res. Toxicol.*, 2014, 27, 159.
- 6 G. Pullangott, U. Kannan, S. Gayathri, D. V. Kiran and S. M. Maliyekkal, *RSC Adv.*, 2021, 11, 6544.
- 7 C. M. White *Are graphene-coated face masks a COVID-19 miracle – or another health risk?*, 2021. <https://the-conversation.com/are-graphene-coated-face-masks-a-covid-19-miracle-or-another-health-risk-159422>.
- 8 K. Pal, G. Z. Kyzas, S. Kralj and F. Gomes de Souza, *J. Mol. Struct.*, 2021, 1233, 130100.
- 9 K. S. Novoselov, A. K. Geim, S. V. Morozov, D. Jiang, Y. Zhang, S. V. Dubonos, I. V. Grigorieva and A. A. Firsov, *Science*, 2004, 306, 666.
- 10 B. Fadeel, C. Bussy, S. Merino, E. Vázquez, E. Flahaut, F. Mouchet, L. Evariste, L. Gauthier, A. J. Koivisto, U. Vogel, C. Martín, L. G. Delogu, T. Buerki-Thurnherr, P. Wick, D. Beloin-Saint-Pierre, R. Hischier, M. Pelin, F. Candotto Carniel, M. Tretiach, F. Cesca, F. Benfenati, D. Scaini, L. Ballerini, K. Kostarelos, M. Prato and A. Bianco, *ACS Nano*, 2018, 12, 10582.



- 11 A. Schinwald, F. A. Murphy, A. Jones, W. MacNee and K. Donaldson, *ACS Nano*, 2012, **6**, 736.
- 12 L. Newman, A. F. Rodrigues, D. A. Jasim, I. A. Vacchi, C. Ménard-Moyon, A. Bianco, C. Bussy and K. Kostarelos, *Cell Rep. Phys. Sci.*, 2020, **1**, 100176.
- 13 A. F. Rodrigues, L. Newman, D. A. Jasim, S. P. Mukherjee, J. Wang, I. A. Vacchi, C. Ménard-Moyon, A. Bianco, B. Fadeel, K. Kostarelos and C. Bussy, *Adv. Sci.*, 2020, **7**, 1903200.
- 14 B. Li, J. Yang, Q. Huang, Y. Zhang, C. Peng, Y. Zhang, Y. He, J. Shi, W. Li, J. Hu and C. Fan, *NPG Asia Mater.*, 2013, **5**, e44.
- 15 L. Mao, M. Hu, B. Pan, Y. Xie and E. J. Petersen, *Part. Fibre Toxicol.*, 2015, **13**, 7.
- 16 M. Ema, M. Gamo and K. Honda, *Regul. Toxicol. Pharmacol.*, 2017, **85**, 7.
- 17 A. Spinazzè, A. Cattaneo, D. Campagnolo, V. Bollati, P. A. Bertazzi and D. M. Cavallo, *Aerosol Sci. Technol.*, 2016, **50**, 812.
- 18 J. H. Lee, J. H. Han, J. H. Kim, B. Kim, D. Bello, J. K. Kim, G. H. Lee, E. K. Sohn, K. Lee, K. Ahn, E. M. Faustman and I. J. Yu, *Inhalation Toxicol.*, 2016, **28**, 281.
- 19 J. H. Shin, S. G. Han, J. K. Kim, B. W. Kim, J. H. Hwang, J. S. Lee, J. H. Lee, J. E. Baek, T. G. Kim, K. S. Kim, H. S. Lee, N. W. Song, K. Ahn and I. J. Yu, *Nanotoxicology*, 2015, **9**, 1023.
- 20 J. K. Kim, J. H. Shin, J. S. Lee, J. H. Hwang, J. H. Lee, J. E. Baek, T. G. Kim, B. W. Kim, J. S. Kim, G. H. Lee, K. Ahn, S. G. Han, D. Bello and I. J. Yu, *Nanotoxicology*, 2016, **10**, 891.
- 21 Y. H. Kim, M. S. Jo, J. K. Kim, J. H. Shin, J. E. Baek, H. S. Park, H. J. An, J. S. Lee, B. W. Kim, H. P. Kim, K. H. Ahn, K. Jeon, S. M. Oh, J. H. Lee, T. Workman, E. M. Faustman and I. J. Yu, *Nanotoxicology*, 2018, **12**, 224.
- 22 A. Skovmand, A. Jacobsen Lauvås, P. Christensen, U. Vogel, K. S. Hougaard and S. Goericke-Pesch, *Part. Fibre Toxicol.*, 2018, **15**, 10.
- 23 D. A. Jasim, C. Menard-Moyon, D. Begin, A. Bianco and K. Kostarelos, *Chem. Sci.*, 2015, **6**, 3952.
- 24 D. A. Jasim, S. Murphy, L. Newman, A. Mironov, E. Prestat, J. McCaffrey, C. Ménard-Moyon, A. F. Rodrigues, A. Bianco, S. Haigh, R. Lennon and K. Kostarelos, *ACS Nano*, 2016, **10**, 10753.
- 25 A. Bianco, *Angew. Chem., Int. Ed.*, 2013, **52**, 4986.
- 26 B. Cornelissen, S. Able, V. Kersemans, P. A. Waghorn, S. Myhra, K. Jurkshat, A. Crossley and K. A. Vallis, *Biomaterials*, 2013, **34**, 1146.
- 27 D. A. Jasim, H. Boutin, M. Fairclough, C. Ménard-Moyon, C. Prenant, A. Bianco and K. Kostarelos, *Appl. Mater. Today*, 2016, **4**, 24.
- 28 H. Hong, K. Yang, Y. Zhang, J. W. Engle, L. Feng, Y. Yang, T. R. Nayak, S. Goel, J. Bean, C. P. Theuer, T. E. Barnhart, Z. Liu and W. Cai, *ACS Nano*, 2012, **6**, 2361.
- 29 V. J. González, J. Frontiñan-Rubio, M. V. Gomez, T. Montini, M. Durán-Prado, P. Fornasiero, M. Prato and E. Vázquez, *ACS Nano*, 2023, **17**, 606.
- 30 H. Cazier, C. Malgorn, D. Georgin, N. Fresneau, F. Beau, K. Kostarelos, C. Bussy, S. Campidelli, M. Pinault, M. Mayne-L'Hermite, F. Taran, C. Junot, F. Fenaille, A. Sallustrau and B. Colsch, *Nanoscale*, 2023, **15**, 5510.
- 31 X. Guo, S. Dong, E. J. Petersen, S. Gao, Q. Huang and L. Mao, *Environ. Sci. Technol.*, 2013, **47**, 12524.
- 32 B. Czarny, D. Georgin, F. Berthon, G. Plastow, M. Pinault, G. Patriarche, A. Thuleau, M. Mayne-L'Hermite, F. Taran and V. Dive, *ACS Nano*, 2014, **8**, 5715.
- 33 K. Lu, S. Dong, T. Xia and L. Mao, *ACS Nano*, 2021, **15**, 396.
- 34 Z. Chen, W. Ren, L. Gao, B. Liu, S. Pei and H. M. Cheng, *Nat. Mater.*, 2011, **10**, 424.
- 35 K. R. Paton, E. Varrla, C. Backes, R. J. Smith, U. Khan, A. O'Neill, C. Boland, M. Lotya, O. M. Istrate, P. King, T. Higgins, S. Barwich, P. May, P. Puczkarski, I. Ahmed, M. Moebius, H. Pettersson, E. Long, J. Coelho, S. E. O'Brien, *et al.*, *Nat. Mater.*, 2014, **13**, 624.
- 36 Y. Hao, Y. Wang, L. Wang, Z. Ni, Z. Wang, R. Wang, C. K. Koo, Z. Shen and J. T. Thong, *Small*, 2010, **6**, 195.
- 37 S. Wang, S. Suzuki and H. Hibino, *Nanoscale*, 2014, **6**, 13838.
- 38 C. Zhang, Q. Li, B. Tian, Z. Huang, W. Lin, H. Li, D. He, Y. Zhou and W. Cai, *Mech. Astron.*, 2014, **57**, 1817.
- 39 Y. Anno, K. Takei, S. Akita and T. Arie, *Phys. Status Solidi RRL*, 2014, **8**, 692.
- 40 M. Kalbáč, L. Kavan, M. Zukalová and L. Dunsch, *Small*, 2007, **3**, 1746.
- 41 J. Ren and S. Licht, *Sci. Rep.*, 2016, **6**, 27760.
- 42 V. J. González, A. M. Rodríguez, V. León, J. Frontiñan-Rubio, J. L. G. Fierro, M. Durán-Prado, A. B. Munoz-García, M. Pavone and E. Vázquez, *Green Chem.*, 2018, **20**, 3581.
- 43 V. J. González, A. M. Rodríguez, I. Payo and E. Vázquez, *Nanoscale Horiz.*, 2020, **5**, 331.
- 44 C. Granotier, G. Pennarun, L. Riou, F. Hoffschir, L. R. Gauthier, A. De Cian, D. Gomez, E. Mandine, J.-F. Riou, J.-L. Mergny, P. Mailliet, B. Dutrillaux and F. D. Boussin, *Nucleic Acids Res.*, 2005, **33**, 4182.
- 45 S. Mokrani, C. Granotier-Beckers, O. Etienne, T. Kortulewski, C. Grisolia, J.-P. de Villartay and F. D. Boussin, *DNA Repair*, 2020, **88**, 102801.
- 46 E. G. Solon, A. Schweitzer, M. Stoeckli and B. Prideaux, *AAPS J.*, 2010, **12**, 1.
- 47 T. Loret, L. A. Visani de Luna, M. A. Lucherelli, A. Fordham, N. Lozano, A. Bianco, K. Kostarelos and C. Bussy, *Small*, 2023, 2301201.

