Intrinsic DNA damage repair deficiency results in progressive microglia loss and replacement

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Abstract
The DNA excision repair protein Ercc1 is important for nucleotide excision, double strand DNA break, and interstrand DNA crosslink repair. In constitutive Ercc1-knockout mice, microglia display increased phagocytosis, proliferation and an enhanced responsiveness to lipopolysaccharide (LPS)-induced peripheral inflammation. However, the intrinsic effects of Ercc1-deficiency on microglia are unclear. In this study, Ercc1 was specifically deleted from Cx3cr1-expressing cells and changes in microglia morphology and immune responses at different times after deletion were determined. Microglia numbers were reduced with approximately 50% at 2–12 months after Ercc1 deletion. Larger and more ramified microglia were observed following Ercc1 deletion both in vivo and in organotypic hippocampal slice cultures. Ercc1-deficient microglia were progressively lost, and during this period, microglia proliferation was transiently increased. Ercc1-deficient microglia were gradually replaced by nondeficient microglia carrying a functional Ercc1 allele. In contrast to constitutive Ercc1-deficient mice, microglia-specific deletion of Ercc1 did not induce microglia activation or increase their responsiveness to a systemic LPS challenge. Gene expression analysis suggested that Ercc1 deletion in microglia induced a transient aging signature, which was different from a priming or disease-associated microglia gene expression profile.

KEYWORDS
aging, DNA damage repair, Ercc1, microglia, morphometrics

1 INTRODUCTION
Organisms have developed intricate mechanisms to repair different forms of DNA damage, for example, base excision repair (BER), nucleotide excision repair (NER), mismatch repair (MMR), interstrand crosslink repair (ICR), and double-strand break repair (DBR). Repair of DNA damage is important as persistent damage induces cell senescence or cell death (Rodier et al., 2009). Deficiencies in
DNA repair pathways lead to various progeria syndromes both in human and mouse (Schumacher, Garnis, & Hoeijmakers, 2008). In many progeria syndromes, neurologic defects occur, which highlights the vital role of genome stability maintenance for the central nervous system (CNS; Heng, Eggen, Boddeke, & Kooistra, 2017; McKinnon, 2013).

Excision repair cross-complementation group 1 (ERCC1) in complex with XPF is an essential nuclease in the NER, ICR and DBR pathways (Ahmad et al., 2008; Klein Douwel et al., 2014; Li et al., 2019). Besides their role in DNA damage repair, nucleotide excision repair factors like ERCC1 and XPF are recruited to active promoters to facilitate transcription (Le May et al., 2010). Mutations in ERCC1 and XPF cause cerebro-oculo-facioskeletal (COFS) syndrome, Cockayne syndrome (CS), and xeroderma pigmentosum (XP) in humans (Gregg, Robinson, & Niedernhofer, 2011; Kashiyama et al., 2013). Mice carrying a knockout (ko) and a hypomorphic (Δ) allele for Ercc1 (Ercc1Δ/ko) display a range of progeroid changes, including reduced lifespan, loss of the body weight, and various aging-related pathological changes in peripheral organs (Dolle et al., 2011). In addition, constitutive Ercc1-knockout (Ercc1Δ/Δ) mice display premature CNS aging, such as motor abnormalities and cognitive decline, widespread astroglialosis, microgliosis and neuronal degeneration in the brain, and progressive motor neuron loss in the spinal cord (Borgesius et al., 2011; de Waard et al., 2010; Vegh et al., 2012). Microglia in Ercc1Δ/Δ mice exhibit a hypertrophic morphology with thickened primary processes and an increase in soma size (Raj et al., 2014).

Functionally, microglia in Ercc1Δ/Δ mice show increased phagocytosis, proliferation and reactive oxygen species (ROS) production. Notably, microglia in Ercc1Δ/Δ mice displayed increased phagocytic and immune activity and are referred to as disease-associated microglia (DAM) or microglia in neurodegenerative disease (MGN; Butovsky & Weiner, 2018; Keren-Shaul et al., 2017; Krasemann et al., 2017; Mathys et al., 2017). Where constitutive Ercc1 deletion induces microglia priming, the effect of microglia-specific Ercc1-deficiency is unclear. In this study, the Ercc1 gene was deleted from microglia and the effect on microglia density, morphology, survival, proliferation, and responsiveness to LPS-induced inflammation were determined.

2 | MATERIALS AND METHODS

2.1 | Animals

C3H10T1/2 (JAX stock #021160), Ercc1Δ/Δ and Ercc1Δ/Δ mice were crossed to obtain the experimental lines C3H10T1/2, Ercc1Δ/Δ, and C3H10T1/2, Ercc1Δ/Δ mice (Figure S1a). In brief, C3H10T1/2, Ercc1Δ/Δ and C3H10T1/2, Ercc1Δ/Δ mice were generated by crossing Ercc1Δ/Δ and C3H10T1/2, Ercc1Δ/Δ mice (both in a C57BL/6 background). C3H10T1/2, Ercc1Δ/Δ mice were crossed with Ercc1Δ/Δ mice (FVB background), resulting in C3H10T1/2, Ercc1Δ/Δ and C3H10T1/2, Ercc1Δ/Δ mice (Figure S1a). Herein referred to as C3H10T1/2, Ercc1Δ/Δ and C3H10T1/2, Ercc1Δ/Δ mice. The mice were group-housed with 2–4 same-sex littermates per cage under 12-hour light/dark cycle conditions and ad libitum access to food and water. All experiments were performed in the Central Animal Facility (CDP) of the UMCG, with protocol (15360-03-002) approved by the Animal Care and Use Committee of the University of Groningen.

2.2 | Genotyping

Genomic DNA was isolated from ear cuts for genotyping with MyTaq Extract-PCR Kit (Bioline, BIO-21127). Primer information is provided in Table S1. C3H10T1/2 was genotyped by PCR with cre primer pairs (Table S1). Schematic representation of Ercc1Δ, Ercc1Δ, and Ercc1Δ alleles are depicted in Figure S1b. The Ercc1Δ allele consists of a neo cassette insertion interrupting Exon 7, abrogating the essential carboxy-terminal 74 amino acids of Ercc1 (Weeda et al., 1997). In the Ercc1Δ allele, Exons 3–5 are flanked by loxp sites (Doig et al., 2006). After tamoxifen treatment, Exons 3–5 in the Ercc1Δ allele will be deleted by homologous recombination, resulting in a recombined Ercc1Δ allele. In Ercc1Δ/Δ mice, C3H10T1/2, Ercc1Δ/Δ and C3H10T1/2, Ercc1Δ/Δ mouse lines, genotyping of the Ercc1Δ and Ercc1Δ was done by duplex PCR using wt and neo primer pairs (Table S1) as described before (Ahmad et al., 2008). For C3H10T1/2, Ercc1Δ/Δ mice, and C3H10T1/2, Ercc1Δ/Δ mouse lines, a duplex PCR was performed to distinguish Ercc1Δ, Ercc1Δ, and Ercc1Δ alleles using loxp and neo primer pairs (Table S1).

After tamoxifen treatment, the Ercc1Δ alleles in both C3H10T1/2, Ercc1Δ/Δ and C3H10T1/2, Ercc1Δ/Δ mice are recombined (Ercc1Δ) (Figure S1c). This recombination results in full Ercc1Δ deletion in C3H10T1/2, Ercc1Δ/Δ mice, but only partial deletion in C3H10T1/2, Ercc1Δ/Δ mice, since one Ercc1Δ allele is still present (Figure S1c). To confirm the specific deletion of Ercc1 in microglia, all mice were genotyped by genomic PCR on sorted microglia (Figure S1d). Mice of 6–8 weeks of age received tamoxifen to induce Ercc1 gene recombination. At certain time points post tamoxifen treatment, microglia were FACs-sorted and DNA was isolated from these microglia for genotyping. A duplex PCR was performed to distinguish Ercc1Δ, Ercc1Δ, and Ercc1Δ allele using loxp and neo primer pairs (Table S1), and Ercc1Δ was genotyped using rec
primer pair (Table S1). The \(\text{Ercc}^{\text{wt}}\) (~1.000 bp), \(\text{Ercc}^{\text{ko}}\) (~800 bp), and \(\text{Ercc}^{\text{loxP}}\) (~500 bp) products were separated by electrophoresis on 1.2% agarose gels, and the \(\text{Ercc}^{\text{rec}}\) allele generated a ~280 bp PCR product (Figure S1d).

2.3 | Administration of tamoxifen

Mice of 6–8 weeks of age received 2 doses of 500 µl of tamoxifen (20 mg/ml, Sigma-Aldrich, T5648) dissolved in corn oil (Sigma-Aldrich, C8267) via oral gavage with a 48 hr interval as described previously (Parkhurst et al., 2013).

2.4 | LPS treatment

Mice were given an intraperitoneal (i.p.) injection of 1 mg/kg LPS (Sigma-Aldrich, Escherichia coli 011:B4, L4391) dissolved in Dulbecco’s phosphate buffered saline (DPBS, Lonza, BE17512F). Control mice received a respective volume of DPBS. After 3 hr, animals were perfused with saline under deep anesthesia. Brains were placed in Hank’s balanced salt solution (HBSS, Gibco, 14170-088) with 0.6% glucose (Sigma-Aldrich, G8769) and 15 mM HEPES (Lonza, BE17-737E). All the following isolation procedures were performed on ice or at 4°C during centrifugation. Brains were mechanical dissociated using the Potter-Elvehjem tissue homogenizer and centrifuged at 220 g for 10 min. The pellets were resuspended in 25 ml 24% Percoll (GE Healthcare, 17-0891-01) with a 3 ml PBS layer on top, followed by centrifugation for 20 min at 950 g (accelerate 4 and brake 0) to remove myelin. The cell pellets were incubated with CD11b-PE (eBioscience, 12-0112-82), CD45-PE/Cy7 (eBioscience, 25-0451-82), and Ly-6C-APC (Biolegend, 128015) antibodies for 20–30 min on ice. Then the cells were washed once and filtered into FACS tubes. Microglia were FACSort-sorted as DAPI–CD11bhiCD45–Ly6c– events on sorter MoFlo-Astrios or MoFlo-XDP (Beckman Coulter).

Ki67 staining was performed according to the manufacturer’s protocol. In brief, the cell pellets after the Percoll gradient were permeabilized by adding 1 ml cold 70% ethanol drop by drop while vortexing. After 1 h incubation at ~20°C, the cells were twice washed with 1 ml PBS with 10% FBS and incubated with CD11b-PE (eBioscience, 12-0112-82), CD45-FITC (eBioscience, 11-0451-85), Ly-6C-APC/Cy7 (Biolegend, 128025), and Ki67-Alexa Fluor® 647 antibody (Biolegend, 652407) for 30 min. Ki67+ microglia and Ki67− microglia were collected.

2.5 | Microglia isolation and flow cytometry

Microglia were isolated as described in our previous work (Gerrits, Heng, Boddeke, & Eggen, 2020). Mice were perfused with saline or PBS under deep anesthesia. Brains were placed in Hank’s balanced salt solution (HBSS, Gibco, 14170-088) with 0.6% glucose (Sigma-Aldrich, G8769) and 15 mM HEPES (Lonza, BE17-737E). All the following isolation procedures were performed on ice or at 4°C during centrifugation. Brains were mechanical dissociated using the Potter-Elvehjem tissue homogenizer and centrifuged at 220 g for 10 min. The pellets were resuspended in 25 ml 24% Percoll (GE Healthcare, 17-0891-01) with a 3 ml PBS layer on top, followed by centrifugation for 20 min at 950 g (accelerate 4 and brake 0) to remove myelin. The cell pellets were incubated with CD11b-PE (eBioscience, 12-0112-82), CD45-PE/Cy7 (eBioscience, 25-0451-82), and Ly-6C-APC (Biolegend, 128015) antibodies for 20–30 min on ice. Then the cells were washed once and filtered into FACS tubes. Microglia were FACSort-sorted as DAPI–CD11bhiCD45–Ly6c– events on sorter MoFlo-Astrios or MoFlo-XDP (Beckman Coulter).

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2.6 | Genomic DNA and RNA isolation from microglia

The AllPrep DNA/RNA Micro Kit (Qiagen, 80284) was used to extract genomic DNA and total RNA from sorted microglia.

2.7 | Quantification of recombination efficiency in bulk microglia by quantitative real-time PCR (qPCR)

To investigate recombination efficiency, DNA from microglia from Cx3cr1-Ercc1wt/loxP and Cx3cr1-Ercc1wt/loxP mice was analyzed by qPCR. Two primer sets were designed, based on the recombinated \(\text{Ercc}^{\text{rec}}\) allele: \(\text{Ercc}^{\text{rec}}\)1 and \(\text{Ercc}^{\text{rec}}\)2 (Table S1). Without recombination, the primer sets will not give products during the PCR due to the long span between the forward and reverse primer. In addition, a reference primer pair, Ref-Il1, which amplifies a genomic fragment of the Il1 gene was included (Table S1). The PCR reaction mixture contained 5 µl DNA template from microglia samples, 5.5 µl iTaq™ Universal SYBR® Green Supermix (Bio-Rad, 1725125), 0.3 µl ddH2O and 0.2 µl 10 µM primer mix. Each sample was quantified with three technical replicates. The percentage of microglia with a recombinant \(\text{Ercc}^{\text{rec}}\) allele was calculated by the following formula (Livak & Schmittgen, 2001).

\[
\text{Percentage of microglia with } \text{Ercc}^{\text{rec}} \text{ allele} = \frac{\text{microglia } (\text{Ercc}^{\text{rec}}) + \text{microglia } (\text{Ercc}^{\text{loxp}})}{2^{\left(\frac{\text{Ct } \text{Ref-Il1} - \text{Ct } \text{Ref-Il2} - \text{Ct } \text{Ref-Il1}}{2}\right)} - 1}
\]

2.8 | Quantification of recombination efficiency by single-cell qPCR (genomic DNA)

Individual microglia were FACSort-sorted in 384-well PCR plates containing 5 µl ddH2O in each well. The presence of the \(\text{Ercc}^{\text{rec}}\) allele was determined using qPCR primer pair \(\text{Ercc}^{\text{rec}}\text{1-rec1}\). As a positive control, individual microglia were analyzed using \(\text{Ref-Il1}\) primers. As negative controls, individual splenic macrophage (DAPI–CD11b–CD45–Ly6c–) were sorted in 384 well plates and analyzed. 5.5 µl of iTaq™ Universal SYBR® Green Supermix, 0.3 µl ddH2O and 0.2 µl 10 µM primer mix were added to each well. Quantitative PCR reactions were performed using the QuantStudio 7 Real-Time PCR system (Thermo Scientific). In the end, the number of PCR reactions resulting in specific DNA products from \(\text{Ercc}^{\text{rec}}\) (with correct melting curves) were quantified. The percentage of microglia with recombiant \(\text{Ercc}^{\text{rec}}\) allele was calculated by the following formula.

\[
\text{Percentage of microglia with } \text{Ercc}^{\text{rec}} = \frac{\text{number of PCR reactions with products from } \text{Ercc}^{\text{rec}}}{\text{total number of PCR reactions}}
\]
2.9 | cDNA synthesis and qPCR

RNA isolated from microglia was mixed with 1 μl random primers (0.5 μg/μl, Invitrogen, 48190011) and ddH2O to 10 μl. Samples were incubated at 65°C for 15 min and kept on ice. Thereafter, 8 U/μl MuLV reverse transcriptase (Thermo Scientific, EP0442), 0.8 U/μl Ribolock RNase inhibitor (Thermo Scientific, EO0382), 0.5 mM dNTP-mix (Thermo Scientific, R0192) and reverse transcriptase buffer were added and incubated on a thermal cycler at 42°C for 1 hr, at 70°C for 10 min and finally at 4°C. The resulting cDNA was used for qPCR reactions. The PCR reaction contained 5 μl cDNA template from microglia samples, 5.5 μl iTaq™ Universal SYBR® Green Supermix, 0.3 μl ddH2O and 0.2 μl 10 μM primer mix. Each sample was run with three technical replicates. Quantitative PCR reactions were performed using the ABI7900HT Fast Real-Time PCR System (Thermo Scientific), LightCycler® 480 System (Roche) or QuantStudio 7 Real-Time PCR system (Thermo Scientific). To determine relative expression levels, Hprt1 was used as the reference gene. Primer sequences are provided in Table S2.

2.10 | QuantSeq 3’ mRNA-Seqencing and bioinformatic analysis

RNA quantity and quality were analyzed on a Fragment Analyzer (Agilent), only RNA samples with a RIN value >6.5 were used. Sequencing libraries were prepared with the Quant Seq 3’ mRNA-Seq Library Prep Kit FWD (Lexogen, 015.96). Quality control of the raw FASTQ files was performed with FASTQC. Bad quality bases were trimmed with FASTX_trimmer of the FASTX_toolkit (version 0.013). FASTQ files was performed with FASTQC. Bad quality bases were trimmed with FASTX_trimmer of the FASTX_toolkit (version 0.013). Sequences were aligned using default parameters on HiSAT2 version 2.1 to the M. musculus (GRCm38.85) reference template obtained from Ensembl. Quantification of the reads was performed with HTseq-counts (version 0.6.1). Raw count matrices were loaded in R and processed with DESeq2. Genes were identified as differentially expressed with an FDR < 0.05 and fold change >1.5. Normalized values (counts per million) of the differentially expressed genes were used as heatmap input. Gene ontology (GO) term enrichment analysis was performed using Metascape (http://www.metascape.org/).

2.11 | Immunohistochemistry and immunofluorescence

To collect brain tissue for immunostaining, mice were perfused with saline under deep anesthesia. Brains were fixed for 48 hr in 4% paraformaldehyde (PFA) at 4°C. After dehydration in 25% sucrose, the brain samples were embedded with O.C.T. compound (Sakura Finetek, 4583) and stored at −80°C.

For immunohistochemistry, 16 μm sections were prepared by cryo-sectioning. After washing thrice with 1× PBS (identical for all subsequent washing steps), antigen retrieval was performed by pressure cooking in 10 mM sodium citrate, pH 6.0. The sections were washed and incubated in PBS with 1% hydrogen peroxide (H2O2) to block endogenous peroxidases. Again, the sections were washed and blocked for 30 min using 5% normal donkey serum (NDS; Jackson Immuno Research, 017-000-121) in PBS with 0.3% Triton X-100 (PBS*). Afterward, the sections were incubated with the primary rabbit-α-ionized calcium-binding adapter molecule 1 (lba1) antibody (1:1,000; Wako, 01-19741) overnight at 4°C. The following day, the slides were washed and incubated with the biotinylated secondary donkey-α-rabbit IgG antibody (1:400; Jackson Immuno Research, 711-065-152) for 1 hr. After washing, the sections were incubated with ABC solution (VECTASTAIN® ABC Kit, Vector Laboratories, PK-6100) for 30 min. The sections were washed, stained using 0.04% 3,3’-Diaminobenzidine (DAB) and 0.01% H2O2 for 8 min and subsequently dehydrated using a sequence of increasing ethanol concentrations. The slides were air dried for 30 min, mounted with coverslips using DePex (Serva) and stored at room temperature. All the slides were scanned with the NanoZoomer 2.0-HT Digital Pathology system (Hamamatsu Photonics, K.K., Japan) at 40 times magnification.

For immunofluorescence, free-floating brain sections were immunolabeled as described (Sierra et al., 2010). For organotypic hippocampal slice culture, slices were blocked for 1 hr with 5% normal donkey serum and thereafter incubated with a primary antibody against lba1 (1:1,000; Wako, 019-19741) overnight at 4°C. On the next day, after washing thrice with 1× PBS, Alexa Fluor 488 donkey anti-rabbit (1:400; Invitrogen, A21206) secondary antibody was added. After 1 hr of secondary antibody incubation, sections were washed and incubated in Hoechst solution (1 μg/ml, Sigma-Aldrich, 14530) for 5 min. After washing, the slides were mounted with Mowiol mounting medium on glass slides. Image acquisition was performed using a Leica SP8 confocal microscope system (TCS SP8, Leica Microsystems).

2.12 | Microglia density and spatial distribution analysis

Microglia densities in the frontal cortex and in cornu ammonis (CA), and dentate gyrus (DG) were determined by counting all Iba1-positive cells in a specified region of interest (ROI) of known dimensions using the cell counter plugin for the ImageJ software (http://rsb.info.nih.gov/ij/). To assess the spatial distribution of microglia in the frontal cortex, the nearest neighbor distances—that is, the average Euclidian distances between nearest cells—were determined using the NND plugin for ImageJ. 2–3 ROIs were selected per animal per group for the analysis. Three mice per group were used except the 22 months (n = 2).

2.13 | Morphometric analysis of microglia

A pipeline was developed to analyze morphological changes in microglia (Van Weering et al., in prep.). Briefly, single-cell images of
iba1-positive cells were first extracted from the whole slide scans, with at least 20 cells per region per animal. Prior to analysis, the single cell images were preprocessed to cell silhouette images by semi-automated thresholding. Subsequently, the cell silhouettes were converted to cell skeleton images by repeated thinning and pruning of the branch areas. In the cell skeleton, branch endings (end nodes), branch crossings (junctions), and all branch points emanating from the cell soma (start nodes) were tagged to allow node quantification. Both cell silhouette- and cell skeleton images served as input for fully automated morphometric analysis. The outputs of the pipeline included Sholl analysis result and morphometric features per cell. A specified list of morphometric features, as well as a detailed description of the morphometrics pipeline is described elsewhere (van Weering et al., in prep.).

2.14 | Clustering of microglia based on morphometric features

To identify groups of microglia of similar morphology, a nonsupervised clustering approach was applied (described in detail in van Weering et al., in prep.). In brief, after normalization and scaling of all morphometric features, a principal component analysis (PCA) was applied to reduce dimensionality and redundancy in the dataset. Subsequently, a hierarchical clustering (Ward’s method) was performed on the top contributing principal components (PCs) with an eigenvalue >1 (here, PC1–4, Figure S2a), resulting in nine clusters of microglia with distinct morphological properties (Figure S2b). The morphometric properties of each cluster are depicted in Figure S2d.

2.15 | Organotypic hippocampal slice culture

Organotypic hippocampal slice culture (OHSCs) were prepared as described previously (Stoppini et al., 1991) with minor modifications. In brief, brains were rapidly isolated from Cx3cr1-Ercc1wt/loxP and Cx3cr1-Ercc1loxP/loxP mouse pups (p3) after decapitation. Isolated hippocampi from both hemispheres were isolated in ice cold serum-free HBSS supplemented with 0.5% glucose and 15 mM HEPES. Isolated hippocampi were cut into 375 μM thick slices using a tissue chopper (Mcllwain) and were transferred to 0.4 μm culture plate inserts (Mili pore, PICM03050). These culture plate inserts, containing 6 slice cultures each, were placed in 6-well plates containing 1.2 ml of culture medium per well. Culture medium (pH 7.2) consisted of 50% minimum essential medium supplemented with 25% heat-inactivated horse serum (Gibco, 16050-122), 25% basal medium eagle, 2 mM glutamax and 0.65% glucose. The slice cultures were kept at 35°C in a humidified atmosphere (5% CO2). On the first day after preparation, OHSCs were treated with 1 nM 4-hydroxy tamoxifen (Sigma-Aldrich, T176) for 48 hr to induce Ercc1 deletion. OHSCs were kept for up to 3 months and the culture medium was refreshed every 2 days. After fixation with 4% PFA overnight at 4°C, OHSCs were processed for immunofluorescence staining.

2.16 | Quantification and statistical analysis

Statistical significance was determined by either a two-way ANOVA followed by Bonferroni correction or a two-tailed Student’s t-test as indicated in the legends. For the morphometrics data, after hierarchical clustering, a Kruskal-Wallis test followed by a Wilcoxon rank sum test with Bonferroni correction was performed for comparison of morphometric features between microglia clusters. Statistical differences with p values lower than 0.05 were considered significant.

3 | RESULTS

3.1 | Microglia are progressively lost after Ercc1 deletion

Ercc1 is an essential endonuclease component in NER, ICR, and DBR, and microglia will accumulate DNA lesions after Ercc1 deletion. Cell cycle arrest, DNA repair, and apoptosis are the general responses to DNA damage (Norbury & Zhivotovsky, 2004). To obtain conditional Ercc1-deficient mice, Ercc1wt/ko and Cx3cr1wt/creERT2 mice were crossed to generate Cx3cr1wt/creERT2:Ercc1wt/ko mice (Figure S1a). Then, Cx3cr1creERT2/creERT2:Ercc1wt/ko were crossed with Ercc1loxP/loxP mice, resulting in Cx3cr1-Ercc11loxP/loxP and Cx3cr1-Ercc11loxP/loxP mice (Figure S1a). After tamoxifen-induced nuclear translocation of CreER, the Ercc1loxP alleles in both Cx3cr1-Ercc11wt/loxP and Cx3cr1-Ercc11wt/loxP mice microglia were recombined (Ercc11loxP) (Figure S1b,c). This recombination resulted in Ercc1 deficiency in Cx3cr1-Ercc11loxP/loxP mice microglia, but not in Cx3cr1-Ercc11loxP/loxP mice microglia, since one Ercc1loxP allele was still present (Figure S1b,c).

To determine the effect of Ercc1 deletion on microglia, first, the effect on microglia cell density was determined in the frontal cortex (Figure 1a). From 2 months after tamoxifen treatment onward, the density of microglia in the frontal cortex of Cx3cr1-Ercc11loxP/loxP mice was significantly lower than in littermate controls, and this reduction persisted until 12 months after tamoxifen treatment (Figure 1b). A reduction in microglia density after Ercc1 deletion was also observed in other brain regions. In the DG and CA, a significant reduction was observed at 6 and 12 months after tamoxifen treatment (Figure 1b). At 22 months after tamoxifen treatment, microglia density in Cx3cr1-Ercc11loxP/loxP mice was similar to control littermates in all brain regions investigated (Figure 1b). Together with the reduction in microglia density, a significant increase was observed in the nearest neighbor distance between microglia in the frontal cortex of Cx3cr1-Ercc11loxP/loxP mice, suggesting that the observed microglia loss occurred throughout the brain and was not regional (Figure 1c). Similar to our histological data, the number of FACS-sorted microglia from Cx3cr1-Ercc11loxP/loxP mice was significantly lower than from littermate controls. This reduction in the number of isolated microglia persisted from 2 to 12 months after tamoxifen treatment (Figure 1d).

In addition, we generated OHSCs from Cx3cr1-Ercc11loxP/loxP pups and deleted Ercc1 by ex vivo 4-hydroxy-tamoxifen treatment. Similar to our in vivo findings, 3 months after tamoxifen treatment,
FIGURE 1  Legend on next page.
the number of microglia in Cx3cr1-Ercc1<sup>ko/loxP</sup> OHSCs was reduced by approximately 50% compared to control OHSCs, in the DG, CA1, and CA3 regions (Figure S3a,b).

### 3.2 Ercc1-deficient microglia are progressively replaced

After tamoxifen treatment, the Ercc1<sup>loxP</sup> allele recombines into an Ercc1<sup>rec</sup> allele in both Cx3cr1-Ercc1<sup>ko/loxP</sup> and Cx3cr1-Ercc1<sup>wt/loxP</sup> mice (Figure S1b,c). To determine the recombination efficiency, genomic DNA was isolated from FACS-isolated microglia and analyzed by qPCR for the Ercc1<sup>rec</sup> allele. In Cx3cr1-Ercc1<sup>wt/loxP</sup> mice microglia, the percentage of microglia with the Ercc1<sup>rec</sup> allele was high (80–100%) and comparable at all investigated time points after tamoxifen treatment (Figure 2a). The percentage of microglia with an Ercc1<sup>rec</sup> allele in Cx3cr1-Ercc1<sup>ko/loxP</sup> mice initially was same as in littermate controls, but progressively declined over time, to approximately 5% at 12 months after tamoxifen treatment (Figure 2a). Ercc1 recombination efficiency was further determined by single-cell PCR. The percentages of Ercc1<sup>rec</sup> microglia progressively decreased in Cx3cr1-Ercc1<sup>ko/loxP</sup> mice, and very few microglia with an Ercc1<sup>rec</sup> allele were detected at 22 months after tamoxifen treatment (Figure 2b), corroborating our previous observations.

These data indicate that after tamoxifen treatment, Ercc1-deficient microglia (Ercc1<sup>ko/ rec</sup>) were gradually lost in Cx3cr1-Ercc1<sup>ko/loxP</sup> mice and were replaced by Ercc1<sup>ko/loxP</sup> microglia (Figure 2c). The Ercc1<sup>ko/loxP</sup> microglia are likely cells that escaped tamoxifen induced Ercc1 deletion, and still carried a functional Ercc1<sup>rec</sup> allele. At 12 months after tamoxifen treatment, Ercc1-deficient microglia were almost completely replaced by Ercc1<sup>ko/loxP</sup> microglia, but microglia numbers were still reduced by approximately 40–50% (Figures 1 and 2). At 22 months after tamoxifen treatment, Ercc1-deficient microglia were fully replaced and no differences in microglia densities were observed between control and Cx3cr1-Ercc1<sup>ko/loxP</sup> mice (Figures 1 and 2).

### 3.3 Altered microglia morphology in Cx3cr1-Ercc1<sup>ko/loxP</sup> mice

An evident change in microglia morphology was observed in the frontal cortex of Cx3cr1-Ercc1<sup>ko/loxP</sup> mice from 2 to 12 months after tamoxifen treatment when compared to littermate controls (Figure 3a). Some of the microglia became enlarged, with increased soma sizes and branch lengths (Figure 3a). This altered microglia morphology was also observed in other brain regions, including cortex, hippocampus, cerebellum and olfactory bulb (data not shown). Strikingly, at 22 months after tamoxifen treatment, all microglia in the Cx3cr1-Ercc1<sup>ko/loxP</sup> mice displayed a morphology that was comparable to littermate controls (Figure 3a). In OHSCs, a similar change in microglia morphology was observed in Cx3cr1-Ercc1<sup>ko/loxP</sup> mice (Figure S3c). Next, morphological differences in microglia were quantified across groups at different time points after tamoxifen treatment. The generated 23 morphometric features of each microglia cell are provided in Table S3.

To identify subsets of microglia with a similar morphology, we performed hierarchical clustering on principal components. First, a PCA was applied to the morphometric feature dataset. The first four PCs with an eigenvalue >1 were retained for hierarchical clustering (Figure S2a,b), resulting in 9 microglia clusters with distinct morphological properties (Figure 3b). Cell silhouettes representative for each cluster are depicted in Figure 3c. Notably, microglia in cluster I and II were almost exclusively derived from Cx3cr1-Ercc1<sup>ko/loxP</sup> mice, indicating these microglia clusters are Ercc1-deficiency related (Figure 3b). Cluster I and II microglia were characterized by a relatively large soma area, high total branch length values, a large number of end nodes and relatively low cell solidity values (Figures 3c and S2). These findings were corroborated by Sholl analysis with cluster I and II microglia being the largest cells with most extensive ramification patterns compared to other clusters (Figure 3d). Comparisons of all morphometric features between microglia clusters can be found in Table S3. Next, we analyzed the relative distribution of the microglia clusters over the different mouse groups (genotype and time after tamoxifen treatment). In control mice, the relative proportion of cluster II and cluster I microglia remained low or even absent at all timepoints in control animals (Figure 3e). In Cx3cr1-Ercc1<sup>ko/loxP</sup> mice, between 2 and 12 months after tamoxifen treatment, cluster I and II microglia accounted for 45–65% of the total population in the cortex (Figure 3e). At 22 months after tamoxifen treatment, the microglia cluster distribution in Cx3cr1-Ercc1<sup>ko/loxP</sup> and Cx3cr1-Ercc1<sup>wt/loxP</sup> mice was comparable and cluster I and II microglia were almost absent (Figure 3e).

To summarize, upon Ercc1 deletion, CNS microglia numbers were reduced by approximately 50%, which was accompanied by...
the emergence of a microglia subpopulation (Cluster I and II) with relatively large and hyper-ramified cells. This reduction in cell number and changes in microglia morphology persisted until 12 months after tamoxifen treatment. Gradually, microglia numbers and morphology returned to control levels at 22 months after tamoxifen treatment.

3.4 | Increased proliferation compensates for microglia loss after Ercc1 deletion

Under homeostatic conditions, the proliferation rate of microglia is relatively low and the population is maintained by balanced proliferation and apoptosis (Askew et al., 2017). After genetic or
Altered microglia morphology in Cx3cr1–Ercc1<sup>ko/loxP</sup> mice. (a) Representative images of Iba1-stained microglia in the cortex of Cx3cr1–Ercc1<sup>wt/loxP</sup> and Cx3cr1–Ercc1<sup>ko/loxP</sup> mice at different time points after tamoxifen treatment. (b) Hierarchical clustering on principal components resulted in nine cell clusters (I–IX). (c) Representative cells for each microglia cluster. (d) Sholl analysis for microglia clusters I–IX, revealing distinct differences in cell size and branching complexity between clusters. Dots and vertical lines represent means and +/− standard deviations respectively. (e) Distribution analysis of microglia clusters across genotypes at different time points after tamoxifen: wt/loxP: Cx3cr1–Ercc1<sup>wt/loxP</sup>, ko/loxP: Cx3cr1–Ercc1<sup>ko/loxP</sup>
pharmacological depletion of microglia, the remaining microglia rapidly expand and repopulate the CNS (Bruttger et al., 2015; Elmore et al., 2014; Rubino et al., 2018). As Ercc1-deficient microglia were gradually replaced, we determined if microglia proliferation was increased after Ercc1 deletion. The expression levels of Ki67, a gene expressed by proliferating microglia (McDonough et al., 2020), was determined in microglia at early (1 and 2 months) and late (12 months) time points after tamoxifen treatment. Microglia from Cx3cr1-Ercc1ko/loxP mice expressed significantly higher Ki67 levels at 1–2 months after tamoxifen, indicating increased microglia proliferation (Figure 4a). Dividing microglia were observed in the Cx3cr1-Ercc1ko/loxP mouse hippocampus 1.5 months after tamoxifen treatment (Figure 4b, indicated by white arrows). At 12 months after tamoxifen, when almost all Ercc1-deficient microglia were replaced, Ki67 expression levels had

![Image](image-url)
FIGURE 5  Legend on next page.
oscillating microglia from Figure 5a). The expression of these genes was also determined by qPCR in microglia from Cx3cr1-Ercc1ko/loxP mice compared to Cx3cr1-Ercc1WT/loxP controls (Figure 4c,d). Genotyping of Ki67+ microglia population revealed the percentage of microglia with an Ercc1REC allele from both Cx3cr1-Ercc1ko/loxP and Cx3cr1-Ercc1WT/loxP mice was comparable, indicating that Ercc1-deficient microglia (Ercc1ko/loxP) also proliferated (Figure 4e).

In summary, tamoxifen treatment resulted in Ercc1 excision, leading to a progressive loss of Ercc1-deficient microglia in Cx3cr1-Ercc1ko/loxP mice. In parallel, the remaining microglia, including Ercc1-deficient microglia, displayed increased expression of proliferation marker Ki67. At 12 months after tamoxifen treatment, when nearly all Ercc1-deficient microglia (Ercc1ko/loxP) were replaced by Ercc1ko/loxP microglia, Ki67 expression returned to control levels.

### 3.5 | Ercc1-deficient microglia are not immune activated or primed

In Ercc1ko/lox mouse microglia, the expression of genes like Axl, Lgals3, Apoe, and Itpax (Cd11c) is upregulated (Holtman et al., 2015; Figure 5a). The expression of these genes was also determined in Ercc1-deficient microglia from Cx3cr1-Ercc1ko/loxP mice at different time points after tamoxifen treatment. The expression of Axl was significantly higher in Cx3cr1-Ercc1ko/loxP microglia only at 12 months compared to Cx3cr1-Ercc1WT/loxP microglia, but the level of induction was much lower than in Ercc1ko/lox mice (Figure 5a,b). Lgals3, Apoe, and Itpax expression was not significantly induced in microglia from Cx3cr1-Ercc1ko/loxP mice, suggesting microglia were not primed (Figure 5b).

In agreement with our previous findings, basal expression levels of Ccl2 and Tnf were slightly higher in Ercc1ko/lox mice (Figure 5c) and Ercc1ko/lox mice microglia showed an increased responsiveness to LPS compared to controls in terms of Ccl2 and Tnf expression (Raj et al., 2014; Figure 5d). For Cx3cr1-Ercc1ko/loxP microglia, basal expression levels of Ccl2 and Tnf were similar to Cx3cr1-Ercc1WT/loxP microglia, again suggesting Cx3cr1-Ercc1ko/loxP microglia were not primed (Figure 5e). In response to LPS, Cx3cr1-Ercc1ko/loxP microglia only showed a (modest) enhanced Ccl2 expression at 2 and 6 months after tamoxifen treatment (Figure 5f). For Tnf expression, no increase in LPS responsiveness was observed in Cx3cr1-Ercc1ko/loxP microglia (Figure 5f). In summary, microglia-specific deletion of Ercc1 did not induce an immune activated or primed phenotype in microglia.

### 3.6 | Gene expression profiling of microglia after Ercc1 deletion

To delineate the effect of Ercc1 deletion on microglia, we compared the gene expression profiles between Cx3cr1-Ercc1ko/loxP and Cx3cr1-Ercc1WT/loxP microglia, before and at different times after tamoxifen treatment (Figures 6a and 54).

Genes in Cluster 1 were more enriched in Cx3cr1-Ercc1WT/loxP microglia at 12 months than 5 days after tamoxifen treatment (Figure 6b), suggesting changes in expression of these genes are age-related. Cluster 1 genes were associated with GO terms such as brain development, neuronal system, synapse, and morphogenesis (Figure 6c). Some cluster 1 genes, that showed increased expression in microglia isolated from mice 12 months after tamoxifen (from both genotypes), were also transiently increased in Cx3cr1-Ercc1ko/loxP mice microglia at 1 and 2 months after tamoxifen treatment, when most microglia were still Ercc1-deficient (Figures 6b and 2). However, with ongoing microglia replacement, around 6 months after tamoxifen treatment. The expression level of these genes was reduced, suggesting Ercc1 deficiency caused a transient aging phenotype in microglia 1–2 months after tamoxifen treatment (Figure 6b).

When microglia were fully replaced in Cx3cr1-Ercc1ko/loxP mice at 12 months after tamoxifen treatment (Figure 2), the transcriptional profiles of Cx3cr1-Ercc1ko/loxP and Cx3cr1-Ercc1WT/loxP mice microglia were again very similar (Figure 6b).

To get more insight into the transient microglia aging phenotype, several homeostatic microglia signature genes, inflammatory genes expressed by potentially senescent human microglia (Geirsdottir et al., 2019), and DAM genes were investigated (Geirsdottir et al., 2019; Keren-Shaul et al., 2017). Expression levels of homeostatic microglia genes, such as Soll1, Cx3cr1, and Csf1r were unaffected by Ercc1 deletion (Figure S5a). However, at 2 and 6 months after tamoxifen treatment, microglia isolated from Cx3cr1-Ercc1ko/loxP mice displayed a transient aging phenotype with higher expression levels of some senescence-associated genes (Tnf, Ccl3 and Ccl4) and

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**FIGURE 5** | Ercc1-deficient microglia are not immune activated or primed. (a) The expression of Axl, Lgals3, Apoe, and Itpax was determined by qPCR in microglia from Ercc1ko/lox mice and normalized to Hprt1 expression levels. An unpaired two-tailed Student’s t-test was performed for statistical analysis. ***, p < 0.001. (b) Axl, Lgals3, Apoe, and Itpax gene expression was determined by qPCR in microglia from Cx3cr1-Ercc1WT/loxP and Cx3cr1-Ercc1ko/loxP mice at different time points after tamoxifen treatment and normalized to Hprt1 expression levels. A two-way ANOVA followed by a Bonferroni correction for multiple comparisons was performed for statistical analysis. **, p < 0.01. Gene expression levels of Ccl2 and Tnf were determined by qPCR in microglia from Ercc1ko/lox and control mice 3 hr after an i.p. PBS (c) or 1 mg/kg LPS (d) and normalized to Hprt1 expression levels. Each dot represents a mouse. An unpaired two-tailed Student’s t-test was performed for statistical analysis. ***, p < 0.001. Gene expression levels of Ccl2 and Tnf were determined by qPCR in microglia from Cx3cr1-Ercc1WT/loxP and Cx3cr1-Ercc1ko/loxP mice 3 hr after an i.p. PBS (e) or 1 mg/kg LPS (f) and normalized to Hprt1 expression levels. Each dot represents a mouse. A two-way ANOVA followed a Bonferroni correction was performed. **, p < 0.01. ****, p < 0.001
some of the DAM genes (ApoE, Cst7, and Axl) (Figure S5b,c). But this microglia aging phenotype was no longer detected at 12 months after tamoxifen treatment in Cx3cr1-Ercc1<sup>ko/loxP</sup> mice (Figure S5b,c).

The expression of the majority of the genes in Clusters 2 and 3 was increased at 1 and 2 months after Ercc1 deletion, and progressively returned to control levels from 6 to 12 months (Figure 6b).
Some genes in these clusters were associated with mitosis/cell cycle, corroborating the observed increased microglia proliferation at 1 and 2 months after tamoxifen treatment (Figures 6c and 4). In addition, some GO terms of Clusters 2 and 3 genes were related to apoptosis processes such as p53 signaling, release of cytochrome c from mitochondria and intrinsic apoptotic signaling pathway in response to DNA damage (Figure 6c). This supports the excessive loss of Ercc1-deficient microglia between 1 and 6 months after tamoxifen treatment (Figure 1).

Cluster 7 contained genes that were downregulated in Ercc1-deficient microglia at 1, 2-, and 6-months after tamoxifen treatment. GO enrichment indicated that these genes were involved in regulation of protein folding, extracellular exosome assembly, and lysosomal transport (Figure 6c).

To determine if tamoxifen treatment affected gene expression, the transcriptomes of microglia from Cx3cr1-Ercc1$^{+/\Delta ko}$ mice without tamoxifen treatment and Cx3cr1-Ercc1$^{+/\Delta ko}$ mice 5 days after tamoxifen treatment were compared. We only identified 28 differentially expressed genes which were involved in p53 signaling and cell cycle (Figure 6d), indicating a limited influence of tamoxifen treatment on microglia gene expression. No differentially expressed genes were detected between Cx3cr1-Ercc1$^{+/\Delta ko}$ and Cx3cr1-Ercc1$^{+/-\Delta ko}$ mice microglia 5 days after tamoxifen treatment, indicating that 5 days of Ercc1 deficiency was not sufficient to affect microglia gene expression (Figure 6b).

The number of differentially expressed genes, both up- and down-regulated, in microglia at different times after tamoxifen treatment, and associated GO terms are depicted in Figure 6d. The gene lists and enriched GO terms are provided in Table S4.

Microglia-specific deletion of Ercc1 resulted in a gene expression signature distinct from the priming signature of aged and disease-associated microglia we previously reported (Holtman et al., 2015). Only 19 of the 458 genes upregulated at 2 months after tamoxifen treatment in Cx3cr1-Ercc1$^{+/\Delta ko}$ mice overlapped with this priming gene expression module consisting of 295 upregulated genes, suggesting microglia were not primed by intrinsic DNA damage repair deficiency (Figure 6e).

### 4 | DISCUSSION

Here, we analyzed the effect of Ercc1-deficiency on microglia. Deletion of Ercc1 resulted in progressive microglia loss and to compensate for this cell loss, microglia proliferation was transiently increased. Interestingly, the remaining Ercc1-deficient and nondeficient microglia both displayed increased proliferation activity. Gradually, likely due to ongoing loss of Ercc1-deficient microglia, the CNS gradually repopulated with nondeficient (Ercc1$^{+/\Delta ko}$/ko) microglia. Unlike constitutive Ercc1-knockout mice, intrinsic Ercc1 deletion did not induce microglia activation or priming. Microglia-specific deletion of Ercc1 transiently induced an aging-associated gene expression profile, which was different from the gene expression signature of aged and CNS disease-associated microglia (Holtman et al., 2015).

#### 4.1 | Phenotypes of microglia in conditional Cx3cr1-Ercc1$^{+/-\Delta ko}$ mice compared to constitutive Ercc1$^{+/-\Delta ko}$ mice

In constitutive Ercc1$^{+/-\Delta ko}$ mice, Ercc1-deficiency was already present in the zygote, and in all cell types. However, in Cx3cr1-Ercc1$^{+/-\Delta ko}$ mice, Ercc1-deficiency in Cx3cr1-expressing cells was induced by tamoxifen treatment in young adult mice (6-8 weeks of age). In constitutive Ercc1$^{+/-\Delta ko}$ mice, microglia density was increased but in Cx3cr1-Ercc1$^{+/-\Delta ko}$ mice, microglia were lost after Ercc1 deletion. Constitutive Ercc1 deletion resulted in a microglia priming gene expression signature (Holtman et al., 2015). In a previous study, we showed that Camk2$^{creER}$-driven Ercc1 deletion in forebrain neurons also resulted in a microglia phenotype reminiscent of what was observed in Ercc1$^{+/-\Delta ko}$ mice (Raj et al., 2014). In contrast, although microglia from Cx3cr1-Ercc1$^{+/-\Delta ko}$ mice transiently displayed an aging gene expression profile at 2 months after tamoxifen treatment, no clear gene expression signature of priming was detected. Together, these results suggest that microglia priming can be triggered by neuronal genotoxic stress (Raj et al., 2014), but not by microglia-intrinsic genotoxic stress after Ercc1 deletion. Nonetheless, Ercc1 is an essential protein for microglia, as microglia deficient for Ercc1 are progressively replaced by microglia with a functional Ercc1 allele.

#### 4.2 | Turnover of microglia in Cx3cr1-Ercc1$^{+/-\Delta ko}$ mice after tamoxifen treatment

Using genetic labeling and long-term in vivo imaging, Füger et al. reported a median lifetime of mouse neocortical microglia of...
15 months (Füger et al., 2017). In a similar study, the turnover time of mouse microglia was estimated to be 41, 15, and 8 months in the cortex, hippocampus, and olfactory bulb, respectively (Tay et al., 2017). Askew et al. showed that 0.69% of the microglia population is proliferating with an estimated turnover time of 96 days (Askew et al., 2017). Despite these differences in reported turnover times, microglia are relatively long-lived cells in the CNS. Strikingly, after experimental depletion of microglia by either using mice expressing a CD11b-TK transgene (Gowing, Vallières, & Julien, 2006), CSF1R inhibitors (Rice et al., 2017), or the Cx3cr1creERT2;IDTR system, remaining microglia repopulated the CNS very fast (Bruttger et al., 2015; Elmore et al., 2014; Rubin et al., 2018; Varvel et al., 2012). The disadvantage of these approaches is the very fast depletion and repopulation, resulting in (transient) astrogliosis. And the repopulated microglia in the Cx3cr1creERT2;IDTR system have an altered, interferon regulatory factor 7-driven activation phenotype (Rubino et al., 2018; Waisman, Ginchoux, Greter, & Bruttger, 2015).

In the Cx3cr1-Ercc1ko/loxP mice used in this study, Ercc1-deficient microglia were gradually lost, which was associated with a transient increase in proliferation. Similarly, Varol et al. reported a progressive replacement of Dicer-deficient microglia in Cx3cr1creERT2;Dicer ko/loxP mice after tamoxifen treatment. Importantly, they further showed that Dicer-deficiency could induce DNA damage in newborn microglia (Varol et al., 2017). Most likely, Ercc1 deficiency resulted in microglia apoptosis, reflected by the increased expression of apoptosis-related genes in Cx3cr1-Ercc1ko/loxP microglia at 1 and 2 months after tamoxifen treatment (Figure 6bc, Cluster 2 and 3). Importantly, this gradual replacement of microglia did not result in microgliosis.

After pharmacologic depletion by CSF1R inhibitors, the repopulated microglia are derived from the remaining microglia population without contribution from peripheral myeloid cells (Elmore et al., 2014; Huang et al., 2018; Zhan, Sohn, Zhou, Li, & Gan, 2019). For genetic microglia ablation using the Cx3cr1creERT2;IDTR system, Bruttger et al. also showed that microglia exclusively renew from the remaining cells (Bruttger et al., 2015). However, Lund et al. showed that the repopulated microglia originated from remaining CX3CR1/F4/80+Clec12a+ microglia and CX3CR1/F4/80+Clec12a− microglia-like macrophages originated from Ly6C+ monocytes (Lund et al., 2018). These monocyte-derived macrophages can acquire some key features of microglia but still are transcriptionally and functionally distinct from CNS resident microglia even 12 weeks after depletion (Lund et al., 2018). In our study, at 12 months after tamoxifen treatment, the repopulated microglia are transcriptionally similar to microglia from control mice, suggesting that the repopulated microglia in Cx3cr1-Ercc1ko/loxP mice are most likely derived from microglia that escaped Cre recombination.

### 4.4 Spontaneous recombinase activity of CreER

In this study, we used Cx3cr1creERT2 mice to delete Ercc1 in microglia. Microglia deficient for Ercc1 were gradually lost and replaced by Ercc1ko/loxP microglia. The transcriptional changes we determined at different time points after tamoxifen treatment were generated using all microglia isolated from mouse brain. As a consequence, the effects of Ercc1-deficiency on gene expression might be diluted by Ercc1ko/loxP microglia that progressively populate the brain after tamoxifen treatment. In order to separate Ercc1-deficient Ercc1ko/rec microglia and nondeficient Ercc1ko/loxP microglia, we generated a Cx3cr1creERT2;Ercc1ko/loxP;R26CAG-tdTomato mouse line. In these mice, Ercc1ko/rec microglia would also express the tdTomato reporter, where Ercc1ko/loxP microglia do not. This assumes that the excision of the floxed Ercc1 and floxed stop cassette upstream of the tdTomato reporter gene are equally efficient. However, we found 70% of the microglia already expressed the tdTomato reporter prior to tamoxifen treatment (data not shown). This "leaky" Cre activity of the Cx3cr1creERT2 transgene was confirmed by other studies (Chappell-Maor et al., 2020; Fonseca et al., 2017). In addition, for our Cx3cr1creERT2;Ercc1ko/loxP;R26CAG-tdTomato reporter mouse line, we observed that the spontaneous activity of Cre on only expressed the floxed stop cassette upstream of tdTomato, but not the floxed sequence of our target gene Ercc1. This is most likely due to the size of the floxed fragment of our target gene Ercc1, which is around 2.5 kb (Doig et al., 2006), which is longer than the 0.8 kb floxed stop cassette in the tdTomato reporter (Madisen et al., 2010). The different susceptibility of constructs in response to basal CreERT2 activity has also been shown in other studies (Alvarez-Aznar et al., 2020; Van Hove et al., 2020). This tamoxifen-independent Cre activity made it impossible to distinguish between Ercc1-deficient and nondeficient microglia in our Cx3cr1creERT2; Ercc1ko/loxP;R26CAG-tdTomato mice.

In summary, our data indicate that Ercc1 is an essential protein for microglia and its deletion leads to cell death. As a consequence, microglia are gradually replaced by nondeficient Ercc1ko/loxP microglia carrying a functional Ercc1loxP allele. The replacement of Ercc1-deficient microglia by nondeficient microglia is not accompanied by migratory, phagocytic and immune activated (Kettenmann, Hanisch, Noda, & Verkhratsky, 2011). Microglia morphometrics revealed significant alterations in microglia morphology in Cx3cr1-Ercc1ko/loxP mice after tamoxifen treatment. Further analysis revealed this was mainly due to changes in a subset of microglia. Interestingly, this subset of microglia consisted of both Ercc1-deficient and nondeficient microglia, since this subset was detected at both 2 months (when the majority of microglia is Ercc1ko/rec, Figure 2) and 12 months (when the majority of microglia is Ercc1ko/loxP; Figure 2) after tamoxifen treatment. This suggests that, in compensation for the cell loss, both Ercc1-deficient (Ercc1ko/rec) and nondeficient (Ercc1ko/loxP) microglia became larger in size and more extensively ramified, likely in order to surveil a larger parenchymal area. However, potential functional differences between these enlarged cells and homeostatic microglia are still unresolved.

### 4.3 Morphology of microglia after Ercc1 deletion

Microglia are pleomorphic and can adapt to different environments. Ramified microglia are relatively quiescent and surveil the parenchyma, while microglia with a more amoeboid phenotype are more...
extensive immune activation or gliosis in the CNS. These data furthermore indicate that the functionality and gene expression changes observed in constitutive Ercc1Δ/Δ mice are not microglia intrinsic but likely caused by an aging environment, in agreement with earlier postulations (Raj et al., 2014).

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DATA AVAILABILITY STATEMENT
The data that supports the findings of this study are available in the supplementary material of this article

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