Gut bacterial deamination of residual levodopa medication for Parkinson’s disease

Sebastiaan P. van Kessel 1, Hiltje R. de Jong 1, Simon L. Winkel 1, Sander S. van Leeuwen 1,2, Sieger A. Nelemans 3, Hjalmar Permentier 4, Ali Keshavarzian 5 and Sahar El Aidy 1*

Abstract

Background: Parkinson’s disease (PD) is a progressive neurodegenerative disorder characterized by both motor and non-motor symptoms. Gastrointestinal tract dysfunction is one of the non-motor features, where constipation is reported as the most common gastrointestinal symptom. Aromatic bacterial metabolites are attracting considerable attention due to their impact on gut homeostasis and host’s physiology. In particular, *Clostridium sporogenes* is a key contributor to the production of these bioactive metabolites in the human gut.

Results: Here, we show that *C. sporogenes* deaminates levodopa, the main treatment in Parkinson’s disease, and identify the aromatic aminotransferase responsible for the initiation of the deamination pathway. The deaminated metabolite from levodopa, 3-(3,4-dihydroxyphenyl)propionic acid, elicits an inhibitory effect on ileal motility in an ex vivo model. We detected 3-(3,4-dihydroxyphenyl)propionic acid in fecal samples of Parkinson’s disease patients on levodopa medication and found that this metabolite is actively produced by the gut microbiota in those stool samples.

Conclusions: Levodopa is deaminated by the gut bacterium *C. sporogenes* producing a metabolite that inhibits ileal motility ex vivo. Overall, this study underpins the importance of the metabolic pathways of the gut microbiome involved in drug metabolism not only to preserve drug effectiveness, but also to avoid potential side effects of bacterial breakdown products of the unabsorbed residue of medication.

Keywords: Non-motor symptoms, Gastrointestinal motility, *Clostridium sporogenes*, Drug side effects, Bioactive metabolites, Aminotransferase

Background

Gut bacteria produce a wide range of small bioactive molecules from different chemical classes, including aromatic amino acids [1]. Bacterial products from aromatic amino acid degradation have been shown to play a critical role in intestinal barrier function, immune modulation, and gut motility [2–6]. In the lower part of the gastrointestinal (GI) tract, where oxygen is limited, aromatic amino acid degradation by anaerobic bacteria involves reductive or oxidative deamination [7] resulting in production of aromatic metabolites [8–11]. Although the enzymes involved in the deamination pathway of the aromatic amino acids tryptophan, phenylalanine, and tyrosine have been described [11–13], the enzyme involved in the initial transamination step remains unknown.

Recently, small intestinal (SI) microbiota have been implicated in the interference with levodopa drug availability [14, 15]. Early in vivo studies showed that ~ 90%
of levodopa is transported to the circulatory system [16–18], leaving a ~10% unabsorbed fraction of residual levodopa that can act as substrate for other bacterial species associated with the lower, more anaerobic regions of the GI tract [19]. Such bacterial-residual drug interaction might act as bioactive metabolites with an impact on gut homeostasis.

Parkinson’s disease (PD) is often associated with non-motor symptoms especially in the GI tract. GI tract dysfunction such as constipation, drooling, and swallowing disorders occurs frequently in PD patients, especially constipation, which is reported in 80–90% of the PD patients [20]. Importantly, chronic idiopathic constipation is associated with SI motor abnormalities in the esophagus, stomach, jejunum, and ileum [21, 22] and patients with constipation have a longer SI transit time compared to controls [22]. Only recently, SI dysfunction in PD was studied showing that the transit time in the SI was significantly longer in PD patients compared to healthy controls (HC) [23, 24]. Using wireless electromagnetic capsules, the SI transit time was reported to be significantly higher in PD patients (400 min; n = 22) compared to HC (295 min, n = 15) [24].

This study uncovers the aminotransferase responsible for initiating the deamination pathway involved in the transamination of (among others) levodopa and shows that C. sporogenes can effectively deaminate levodopa to 3-(3,4-dihydroxyphenyl)propionic acid through the aromatic amino acid deamination pathway [11]. We show that the deamination product of gut bacterial degradation of the unabsorbed residues of levodopa in fecal samples from PD patients reduces ileal motility ex vivo. Our results highlight the urgency for further research on the effects of bacterial conversion of the unabsorbed residues of medication, which may affect host physiology.

Results

Clostridium sporogenes deaminates levodopa through its deamination pathway

C. sporogenes is able to deaminate proteinogenic aromatic amino acids (PAAAs) through an anaerobic deamination pathway (Fig. 1a) [11–13]. We hypothesized that levodopa, a non-proteinogenic amino acid (NPAA A) and the main treatment in PD, could be deaminated through the same pathway. Together with another NPAAA, 5-hydroxytryptophan (5-HTP, precursor of serotonin, over-the-counter available drug used to treat depression, obesity, insomnia, and chronic headaches [25]), as an analogous control compound derived from tryptophan, we screened for deamination of these compounds in batch cultures of C. sporogenes. Cultures were incubated with 100 μM levodopa or 5-HTP in combination with PAAAs from the growth medium and were followed over a period of 48 h. Analysis of the samples using high-pressure liquid chromatography (HPLC) coupled to an electrochemical detector (ED) revealed that levodopa is completely converted within 24 h to a new metabolite, which was identified by ¹H/¹³C-NMR and LC-MS as 3-(3,4-dihydroxyphenyl)propionic acid, DHPPA (Fig. 1b, c; Additional File 1: Fig. S1A, 1B, 1C). Furthermore, the incubations showed that the PAAAs available from the growth medium did not prevent the deamination of levodopa and that, during the incubation for 48 h, DHPPA remained stable. Similarly, 5-HTP was converted into two new unknown peaks (Additional File 1: Fig. S2A, S2B), albeit to a much lesser extent compared to levodopa. Only the first peak could be detected and assigned by LC-MS as 5-hydroxyindole-3-lactic acid (5-HILA) by its predicted exact mass (Additional File 1: Fig. S2C). The other peak is potentially 5-hydroxyindole-3-propionic acid (5-HIPA), described below.

To further investigate the involvement of the deamination pathway in levodopa and 5-HTP deamination, the enzyme responsible for the dehydratase reaction (encoded by the fldC gene [11–13]) was disrupted using the Clostron mutagenesis system (Additional File 1: Fig. S2D) [26]. The resulting strain C. sporogenes LLLtrB-eryΩfldC (CSΩfldC) was incubated with levodopa, and the PAAAs from the growth medium. Tryptophan and tyrosine were converted to their intermediates ILA (indole-3-lactic acid) and 4-HPLA (3-(4-hydroxyphenyl)lactic acid), respectively, as previously shown [11]. Analogous to tryptophan and tyrosine, levodopa was no longer deaminated to DHPPA but to its intermediate product 3-(3,4-dihydroxyphenyl)lactic acid (DHLA) (Fig. 1d, e). Only a slight production of 4-HPPA (from tyrosine) is observed after 48 h, presumably because of the substitution of FldABC by the similar HadABC proteins from the had-operon in C. sporogenes [11, 12]. HPLC-ED analysis of the 5-HILA production from 5-HTP by the fldC mutant was hampered by the production of coeluting 4-HPLA, the intermediate deamination product produced from tyrosine (described above). However, the analysis revealed that the second unknown peak produced from 5-HTP was no longer produced by CSΩfldC (Additional File 1: Fig. S2E, S2F), demonstrating that 5-HTP conversion is affected and suggesting that the unknown product is 5-HIPA. Overall, the results show that the deamination pathway from C. sporogenes not only is involved in the deamination of PAAAs but also is in the deamination of the NPAAAs, levodopa and 5-HTP.

Identification of the aromatic aminotransferase responsible for initiation of the deamination pathway

The aromatic aminotransferase responsible for the transamination of levodopa and the other (N)PAAAs is crucial for the initiation of the reductive deamination pathway and for the full deamination of the substrates
Fig. 1 (See legend on next page.)
by the dehydrogenases (FldH and AcdA) and dehydratase (FldABC) (Fig. 1a). However, the gene encoding this transaminase remains unidentified. To further investigate this critical step in the pathway, all nine class I/II aminotransferases encoded by \textit{C. sporogenes} were cloned, purified, and screened for their activity on levodopa and the other (N)PAAAs. Screening revealed a single aminotransferase (EDU38870 encoded by CLOSPO_01732) to be involved in their transamination (Fig. 2a). To verify whether other aminotransferases could substitute for the identified aminotransferase in vivo, CLOSPO_01732 was disrupted (resulting in CS\textsuperscript{CLOSPO_01732} (Additional File 1: Fig. S3A) and a targeted metabolomic analysis of all the (N)PAAA metabolites was performed using HPLC-ED (except metabolites from phenylalanine, which were quantified using HPLC-UV). The disruption of \textit{fldC} or CLOSPO_01732 resulted in only a minor reduction of the exponential growth rate in rich broth (doubling time is 55.1 ± 1.2 min and 64.1 ± 1.1 min, respectively, compared to wild type 44.3 ± 1.2 min) all reaching stationary phase within 12h (Additional File 1: Fig. S3B). Comparing the metabolic profiles from wild type \textit{C. sporogenes} (CS\textsuperscript{WT}), CS\textsuperscript{\textit{fldC}}, and CS\textsuperscript{\textit{CLOSPO_01732}} demonstrated that none of the other tested aminotransferases could take over this transaminase reaction effectively, except for the substrate phenylalanine (Fig. 2b and Additional File 1: Table S1). Disrupting CLOSPO_01732 significantly reduced the production of phenyl-3-propionic acid (PPA), 3-(4-hydroxyphenyl)propionic acid (4-HPPA), indole-3-propionic acid (IPA), and 3-(3,4-dihydroxyphenyl)propionic acid (DHPPA) by 16.4%, 79.0%, 97.2%, and 97.7%, respectively, compared to CS\textsuperscript{WT} within 24–48 h (Additional File 1: Table S1). Presumably, the transamination of phenylalanine is substituted by EDU37030 as this substrate by different enzymes (Fig. 2b, Additional File 1: Table S1). Analogous to tryptamine, CS\textsuperscript{CLOSPO_01732} produced significantly more serotonin compared to CS\textsuperscript{\textit{WT}} at 48 h when incubated with 5-HTP (Fig. 2b, Additional File 1: Fig. S3C, Additional File 1: Table S1), though to a much lesser extent (~1% of substrate added) compared to tryptamine. Collectively, the data show that the aromatic aminotransferase (EDU38870) is involved in the initiation of the aromatic amino acid deamination pathway and is crucial for the production of DHPPA, 5-HILA, 5-HIPA, and the previously described metabolites to be circulating in the blood, IPA, and 4-HPPA (Dodd et al. [11]).

3-(3,4-Dihydroxyphenyl)propionic acid elicits an inhibitory effect on ileal muscle contractions ex vivo

Because levodopa is the main treatment of PD patients and is efficiently deaminated to DHPPA within 24 h by the \textit{C. sporogenes} deamination pathway compared to 5-HTP, we further focused on levodopa and its deamination products. DHPPA is a phenolic acid (a molecule in the class polyphenols), and recent findings demonstrated an association between bacterial-derived polyphenol metabolites and gut transit times in humans [27]. Levodopa is mainly absorbed in the proximal small intestine, but significant amounts can reach the distal part of the intestinal tract [17], and these levels increase with age [28]. As levodopa is taken orally, the first intestinal site where anaerobic bacteria such as \textit{C. sporogenes} (\textit{Clostridium} Cluster I) can encounter relevant levels of levodopa is the ileum. Studies on asymptomatic ileostomy subjects established that the core ileal microbiota consists of (facultative) anaerobes including species from \textit{Clostridium} Cluster I [29, 30]. Moreover, the transit time in the SI has been shown to be significantly longer in PD patients compared to healthy controls (with a median increase of 1.75 h in PD patients) [23, 24]. To this end, we tested whether DHPPA (100 μM) could affect the muscle contractility in the ileum. Ileal rings of wild type C57BL/6J mice were suspended in an ex vivo organ bath system to
Identification of the aromatic amino transferase initiating the deamination pathway. In order to identify which aminotransferase is responsible for the initial transaminase reaction, all class I/II aminotransferases were cloned and purified to test the activity against (N)PAAAs. a Transaminase activity (production of glutamate) for all substrates is depicted. EDU38870 (CLOSPO_01732) was involved in all transaminase reactions. EDU37030 showed similar activity as EDU38870, for phenylalanine. Experiment was performed in technical duplicates to screen for candidate genes for mutagenesis in C. sporogenes. b Targeted metabolic quantification of deamination products from CS WT, CS Ω fldC, and CS Ω CLOSPO_01732 reveals that EDU38870 is involved in the transamination of all for all tested (N)PAAAs. All quantified deamination products are normalized to their initial substrate concentration, and the data represents 3 independent biological replicates. Corresponding values are reported, and metabolite concentration differences between WT and Ω fldC or Ω CLOSPO_01732 were tested for significance using Student's t test, in Additional File 1: Table S1. Black squares indicate that quantification was not possible because of a coeluting HPLC-ED peak. As no commercial standards are available for 5-HILA and 5-HIPA, the peaks were quantified assuming a similar ED-detector response as for 5-HTP.
test the effect of DHPPA on muscle contractions. Our initial results indicated that DHPPA displayed an inhibiting effect on natural ileal contractility (Additional File 1: Fig. S4A).

Because acetylcholine is the neurotransmitter constantly produced from the excitatory muscle motor neurons to induce gut smooth muscle contractility [31], we tested whether DHPPA could have an inhibiting effect on acetylcholine-induced contractility in the ileum. The differences in amplitude of the contractions were quantified by measuring the decrease of the observed frequencies after a Fourier transform of 5-min intervals (Fig. 3a). Ileal tissue preparations were tested by initiating an acetylcholinergic twitch by adding 50 μM of acetylcholine (a concentration saturating the muscarinic receptors (Kd = 1.7 ± 0.18 μM [32])). After 5 min, 100 μM DHPPA (a concentration resembling the higher levels detected in fecal samples of PD patients, see

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**Fig. 3** DHPPA inhibits the acetylcholine-induced twitch from mouse ileum. a Experimental setup, where 5 min after adding 50 μM acetylcholine, 100 μM DHPPA is added. The panel below indicates how the amplitude of the frequencies of the observed oscillations (from 5 min bins) is extracted by a Fourier transform of the analog input. b A representative 1-min recording trace before and after the addition of acetylcholine and DHPPA or vehicle (VH) is shown. ACh, acetylcholine; VH, vehicle (0.05% ethanol). c Inhibition of DHPPA on acetylcholine-induced twitch binned in intervals of 5 min shows a decrease in contractility over time (n = 6 biological replicates and experiments were repeated 1–4 times per tissue). Significance was tested using repeated measures (RM) 1-way ANOVA followed by Tukey’s test (*p < 0.0021, ***p < 0.0002, #p < 0.0021). Box represents the median with interquartile range, and whiskers represent the maxima and minima. d Dose response curve of DHPPA on the acetylcholine-induced twitch at the t_{15-20} minute bin (n = 4 biological replicates) with a half maximal inhibitory concentration (IC_{50}) of 20.3 ± 10.6 μM
below) was added and contractions were followed further over a period of 15 min. One-minute traces of the contractility representing one of the experiments are shown before and after addition of acetylcholine, DHPPA, or vehicle (Fig. 3b). A significant decrease in the amplitude (binned in 5-min intervals) of the acetylcholinergic twitch by DHPPA was observed at the 10–15-min (maximal reduction 69%) and 15–20-min interval (maximal reduction 73%) (Fig. 3c). In order to determine the potency of DHPPA, a dose response curve with DHPPA was performed and showed a half maximal inhibitory concentration (IC_{50}) of 20.3 ± 10.6 μM (Fig. 3d). In contrast to DHPPA, incubations with levodopa did not show any significant effect on the acetylcholinergic twitch (Additional File 1: Fig. S4B). Collectively, the data shows that DHPPA can inhibit the acetylcholine-induced muscle contractility of mouse ileum ex vivo.

**Active levodopa deamination pathway in fecal suspensions of patients with Parkinson’s disease**

We hypothesized that if *C. sporogenes* or other bacteria with the deamination pathway (*C. botulinum*, *Peptostreptococcus anaerobius*, or *Clostridium cadyveris* [11]) are present in the GI tract of PD patients, those patients might have considerable amounts of DHPPA in their fecal samples. Because DHPPA can be a product of gut bacterial metabolism of polyphenolic rich foods in the colon such as coffee and fruit (Jenner et al. [33]), fecal samples from 10 PD patients were compared to 10 age-matched HC. Samples were collected in a previous study, and there were no significant differences in macronutrients, dietary fiber, or total calorie intake between groups [34]. Using a catechol extraction targeted for the quantification of DHPPA, we found that the DHPPA concentrations were significantly higher in PD patient’s fecal samples compared to HC (Fig. 4a). Identification of DHPPA was confirmed by LC-MS (Additional File 1: Table S2). The higher amounts (2.2-fold increase) of DHPPA observed in the fecal samples of PD patients are likely to result from levodopa metabolized by the anaerobic bacteria, deaminating levodopa through the FldBC dehydratase (Fig. 1a). In order to investigate the presence and activity of the anaerobic deamination pathway in fecal samples, the dehydratation of the intermediate levodopa metabolite, DHPLA (Fig. 1a), was tested. The levodopa intermediate DHPLA was used as substrate instead of levodopa to prevent an in vitro substrate bias for bacteria that can decarboxylate levodopa to dopamine [14, 15]. Moreover, FldABC is the key protein complex responsible for the production of DHPPA. Screening for the identified transaminase or FldH dehydrogenase upstream of FldABC would not be relevant as many bacterial species harbor these types of enzymes (Additional File 1: Fig. S5). Hence, fecal suspensions (10% w/v) from PD and HC were incubated anaerobically with DHPLA, and samples were collected at 0, 20, and 45 h and were analyzed by HPLC-ED. After 20 h, DHPPA was detected in fecal samples from PD patients, as well as in fecal samples of HC when supplied with the substrate levodopa (Fig. 4b, Additional File 1: Fig. S6A). Moreover, DHPPA was further converted to the downstream dehydroxylated metabolite of DHPPA, 3-(3-hydroxyphenyl)propionic acid (3-HPPA), over time (Fig. 4b, Additional File 1: Fig. S6B). Because DHPPA is further converted to 3-HPPA in vitro, we quantified both the production of DHPPA and/or 3-HPPA in the fecal incubations as measure for the presence of an active deamination pathway. Metabolic profiles of PD or HC samples that produced DHPPA/3-HPPA over time were quantified and merged (Fig. 4c, Additional File 1: Fig. S6B), showing that DHPPA is produced first and is further metabolized to 3-HPPA. The production of DHPPA or 3-HPPA was observed in 50% and 20% of the PD patient’s and HC fecal suspensions, respectively, after 20 h and in 70% and 50% PD patient’s and HC fecal suspensions, respectively, after 45 h (Additional File 1: Fig. S6C). The production of 3-HPPA in vitro is likely to be performed by *Eggerthella lenta*, which has been shown to perform p-dehydroxylations [35]. Indeed, in vitro culturing of *E. lenta* showed p-dehydroxylation of DHPPA (Additional File 1: Fig. S7A [35, 36]). Because DHPPA is further converted to 3-HPPA in vitro, we examined whether 3-HPPA could elicit a similar effect on the acetylcholine-induced contractions in the ileum. Unlike, DHPPA, 3-HPPA did not elicit a significant effect on the acetylcholine-induced twitch (Additional File 1: Fig. S7B [35, 36]). Furthermore, to investigate the genomic abundance levels of bacteria capable of deaminating (N)PAAAs, we analyzed the 16s rDNA sequence data of the fecal samples of patients with Parkinson’s disease [34] that were employed in this study (Additional File 1: Supplementary Results [11–13, 37] and Additional File 1: Fig. S8). A significant positive correlation (r = 0.62, R^2 = 0.38, p = 0.02) was found between bacteria with the deamination pathway and DHPPA/3HPPA production in fecal incubation samples at 20 h (Additional File 1: Fig. S8E). Taken together, the results show that DHPPA can be produced by the microbiota via anaerobic deamination of levodopa. Moreover, our findings indicate that 3-HPPA originates from DHPPA via dehydroxylation potentially by *Eggerthella lenta* and that the aromatic deamination pathway, as measured by the production
Fig. 4 (See legend on next page.)
of DHPPA or 3-HPPA, is active and present in at least 70% of the PD samples.

Discussion
Identifying bacterial pathways and elucidating their potential impact on bacterial drug metabolism are crucial in order, not only, to maximize medication efficacy, but also to recognize and eventually prevent potential side effects that might affect the host’s physiology on an individual basis [14, 36, 38, 39]. Here, we disclosed the reductive anaerobic deamination pathway in C. sporogenes by identifying its initiating enzyme, the aromatic aminotransferase, and expanded the pathway’s relevance by demonstrating its capacity to convert two clinically important NPAAs, levodopa and 5-HTP. We showed that C. sporogenes is able to completely deaminate levodopa to DHPPA and to a much lesser extent 5-HTP (Figs. 1 and 2; Additional File 1: Fig. S1, S2, S3; Additional File 1: Table S1). Disrupting the bacterial transaminase-encoding gene abolished the production of deaminated products, and increased the production of neuromodulators such as tryptamine and, to much lesser extent, serotonin (Fig. 2, Additional File 1: Fig. S3C, and Additional File 1: Table S1). Tryptamine is a natural product produced by C. sporogenes that has been proposed to modulate gut transit time [40]. The application of engineered gut bacteria as a therapeutic strategy to modulate GI motility or host physiology has been also proposed recently in two proof-of-concept studies: by heterologous motility or host physiology has been also proposed re-gut bacteria as a therapeutic strategy to modulate GI microbiota metabolism.

Overall, our results highlight the urgency to unravel potential side effects of this bacterial by-product of the unabsorbed residue of the medication. DHPPA shares similarity with dopamine structure except of the terminal amine group, which is substituted by a carboxyl group in DHPPA. Dopamine and dopamine agonists have been shown to inhibit methacholine (analog of acetylcholine) induced contraction, which is not mediated via dopamine receptors, in guinea pig jejunum in similar concentration ranges to DHPPA (EC50 relaxation by dopamine ~ 290 μM) [41], indicating that DHPPA might act on a similar mechanism. Collectively, although further research is needed to unravel the underlying mechanism, our results show that DHPPA inhibits the acetylcholine-induced muscle contractions in the ileum with implications on intestinal motility, often observed in PD patients.

Conclusions
The present study shows that C. sporogenes can effectively deaminate unabsorbed residues of levodopa in fecal samples from PD patients to 3-(3,4-dihydroxyphenyl)-propionic acid, which reduces ileal motility ex vivo. Overall, our results highlight the urgency to unravel potential effects of gut bacterial processing of (unabsorbed residues of) medication, such as levodopa.

Methods
Growth and incubation of Clostridium sporogenes and Eggerthella lenta
Clostridium sporogenes ATCC15579 was grown in enriched beef broth (EBB) with 2 g/L glucose [14] and 0.1% Tween 80 (EBB/T) anaerobically (10% H2, 10% CO2, 80% N2) in a Don Whitley Scientific DG250 Workstation (LA Biosystems, Waalwijk, The Netherlands) at 37 °C. Eggerthella lenta DSM2243 was grown in
modified DSMZ medium 78 (DSMZ 78: beef extract, 10.0 g/L; casitone, 30.0 g/L; yeast extract, 5.0 g/L; K2HPO4, 5.0 g/L; Tween 80, 0.1%; menadione (vitamin K3), 1 μg/mL; cysteine, 0.5 g/L; hemin, 5 mg/L; L-arginine 0.1–1.5%) anaerobically (1.5% H2, 5% CO2, balance with N2) in a Coy Laboratory Anaerobic Chamber (neolab Miegge GmbH, Heidelberg, Germany) at 37°C in a tube shaker at 500 RPM. Upon use, bacteria were inoculated from ~80°C stored glycerol stocks in the appropriate media and grown for 18–24 h for C. sporogenes and 24–40 h for E. lenta. Overnight turbid cultures were then diluted 1/50 in an appropriate volume EBB/T or CMM for further experiments with 100 μM levodopa (D9628, Sigma), 5-hydroxytryptophan (H9772, Sigma), 50 μM 3-(3,4-dihydroxyphenyl)propionic acid (102601, Sigma), or H2O as control. All experiments were performed in triplicate (3 biological replicates).

Protein production and purification
Transaminase-encoding genes from C. sporogenes (Additional File 1: Table S3) were amplified using Phusion High-fidelity DNA polymerase and primers listed in Additional File 1: Table S3. All amplified genes were cloned in pET15b, except for EDU37032 which was cloned in pET28b (Additional File 1: Table S3). Plasmids were maintained in E. coli DH5α and verified by Sanger sequencing before transformation to E. coli BL21 (DE3). Overnight cultures were diluted 1:50 in fresh LB medium with the appropriate antibiotic and grown to OD600 = 0.7–0.8 shaking at 37°C. Protein translation was induced with 1 mM isopropyl β-D-1-thiogalactopyranoside (IPTG, 1141144-24-9, Megazyme Inc., Wicklow, Ireland), according to the manufacturer’s microplate assay procedure with some modifications. The supplied buffer was substituted for buffer D (described above, to prevent oxidation of the substrates/products). A reaction mix was prepared mixing 50 μL buffer D, 10 μL quenched sample reaction mixture, 20 μL NAD+/iodonitrotetrazolium chloride solution, 5 μL diaphorase solution, and 5 μL glutamate dehydrogenase (GIDH) solution and reconstituted to a final volume of 290 μL with H2O. Absorbance at 492 nm was measured after 10 min of incubation using a microplate reader (Synergy HTX spectrophotometer, BioTek, BioSPX, The Netherlands), and background was subtracted from initial read before addition of GIDH solution.

Transaminase activity test
Purified transaminases were incubated with 1 mM substrate, 2 mM α-ketoglutaric acid, and 0.1 mM PLP (pyridoxal-5-phosphate, P9255, Sigma, The Netherlands) in buffer D with an enzyme concentration of 50 nM for tyrosine, tryptophan, or 5-HTP as substrate and an enzyme concentration of 500 nM for phenylalanine and levodopa as substrate. Enzyme reactions were incubated for 0.5 h at 37°C; the reactions were stopped with 0.7% (v/v) perchloric acid (1:1). Transaminase activity was tested using an L-glutamic acid detection kit (K-GLUT, Megazyme Inc., Wicklow, Ireland), according to the manufacturer’s microplate assay procedure with some modifications. The supplied buffer was substituted for buffer D (described above, to prevent oxidation of the substrates/products). A reaction mix was prepared mixing 50 μL buffer D, 10 μL quenched sample reaction mixture, 20 μL NAD+/iodonitrotetrazolium chloride solution, 5 μL diaphorase solution, and 5 μL glutamate dehydrogenase (GIDH) solution and reconstituted to a final volume of 290 μL with H2O. Absorbance at 492 nm was measured after 10 min of incubation using a microplate reader (Synergy HTX spectrophotometer, BioTek, BioSPX, The Netherlands), and background was subtracted from initial read before addition of GIDH solution.

Targeted mutagenesis
Gene disruptions in Clostridium sporogenes were performed using the ClosTron system [42, 43]. This system facilitates targeted mutagenesis using the group II Ll.LtrB intron of Lactococcus lactis. Introns targeting fbdC (CLOSPO_311) or CLOSPO_1732 (encoding for the transaminase) were designed using the ClosTron intron design tool (http://www.clostron.com) and were ordered in pMTL007C-E2 from ATUM (Newark, CA, USA) resulting in pMTL007C-E2_Cs-fldC-561a and pMT L007C-E2_Cs-CLOSPO_1732-493s, respectively. Plasmids were transferred to C. sporogenes by conjugation as described before [43] using E. coli CA434 (E. coli HB101 (Bio-Rad Laboratories, The Netherlands)) harboring the broad host IncPβ+ conjugational plasmid pRK24 [44] as donor strain. E. coli CA434 harboring pMTL007C-E2_Cs-fldC-561a or pMTL007C-E2_Cs-CLOSPO_1732-493s was verified by Sanger sequencing before transformation.
grown in Luria Broth (LB) with 10 μg/mL tetracycline and 25 μg/mL chloramphenicol (to select for pRK24 and PMTLO07C-E2, respectively). Cell suspensions of 1 mL of overnight culture were washed once with PBS, and the cell pellet was resuspended in 200 μL of C. sporogenes overnight cell suspension. The bacterial mixture was spotted (in drops of 10 μL) on trypticase soy agar (TSA) plates and incubated for 24 h anaerobically at 37 °C. Sequentially, 1 mL of PBS was added to the spotted plates and the donor-recipient mix was scraped of the plate; sequentially, the scraped-off suspension was distributed over TSA plates containing 50 μg/mL neomycin (to prevent growth of E. coli) and 15 μg/mL chloramphenicol to select for C. sporogenes conjugants. Chloramphenicol-resistant colonies of C. sporogenes were re-streaked on TSA plates containing 50 μg/mL neomycin and 2.5 μg/mL erythromycin (to select for intron insertion) for several times. To make sure the plasmids were integrated, colonies were checked and selected for their sensitivity towards chloramphenicol and the genomic DNA was verified using PCR (Additional File 1: Fig. S1F and 2A).

**Fecal samples from patients with Parkinson’s disease and age-matched healthy controls**

Fecal samples from patients diagnosed with PD (n = 10) and age-matched healthy controls (n = 10) were acquired from the Movement Disorder Center at Rush University Medical Center, Chicago, IL, USA, published previously [34]. All study subjects consented to the use of their samples for research. PD was diagnosed according to the UK Brain Bank Criteria as previously described [34]. Study subjects were provided with the supplies and instructions for home feces collection using the BD Gaspak EZ Anaerobe Gas Generating Pouch System with Indicator (Ref 260683; Becton, Dickinson and Company, Sparks, MD) in order to minimize the exposure of the feces to high oxygen ambient atmosphere, which may alter the microflora. Subjects were asked to have a bowel movement within 24 h of their study visit. Subjects kept the sealed anaerobic fecal bag in a cold environment, before bringing the anaerobic fecal bag to the hospital. Fecal samples were then immediately stored at −80 °C until analysis.

**Fecal metabolite incubations from PD patients and HC subjects**

Stool samples were suspended 1:1 (w/v) in EBB/T and incubated anaerobically (10% H2, 10% CO2, 80% N2) in a Don Whitley Scientific DG250 Workstation (LA Biosystems, Waalwijk, The Netherlands) at 37 °C with 100 μM sodium 3-(3,4-dihydroxyphenyl)-DL-lactate (39363, Sigma). Samples were taken at 0, 20, and 45 h and analyzed on HPLC-ED as described below.

**HPLC-ED/UV analysis and sample preparation**

For bacterial cell suspensions, 1 mL of methanol was added to 0.25 mL of cell suspension and stored at −20 °C until further use. For fecal metabolite incubations, 300 μL of methanol was added to 75 μL of fecal suspension and stored at −20 °C until further use. Metabolites from stool samples were extracted by suspending the stool 1:1 (w/v) in water, followed by homogenization by vigorously vortexing while keeping samples as cold as possible. Homogenized suspensions were centrifuged at 3500×g for 20 min at 4 °C, and sequentially, 1.6 mL of methanol was added to 0.4 mL of supernatant. From bacterial, fecal incubation or stool samples, cells and protein precipitates were removed by centrifugation at 20,000×g for 10 min at 4 °C. Supernatant was transferred to a new tube, and the methanol fraction was evaporated in a Savant speed-vacuum dryer (SPD131, Fisher Scientific, Landsmeer, The Netherlands) at 60 °C for 1.5−2 h. The aqueous fraction was reconstituted with 0.7% HClO4 to the appropriate volume. Samples were filtered and injected into the HPLC-ED system (Alliance Separations Module 2695, Waters Chromatography B.V, Etten-Leur, The Netherlands; Dionex ED40 electrochemical detector, Dionex, Sunnyvale, USA, with a glassy carbon working electrode (DC amperometry at 0.8 or 1.0 V, with Ag/AgCl as reference electrode)). Samples were analyzed on a C18 column (Kinetex 5 μM, C18 100 Å, 250 × 4.6 mm, Phenomenex, Utrecht, The Netherlands) using a gradient of water/methanol with 0.1% formic acid (0−10 min, 95−80% H2O; 10−20 min, 80−5% H2O; 20−23 min, 5% H2O; 23−31 min, 95% H2O). Fecal suspension metabolites were injected twice and analyzed at DC amperometry at 0.8 V (for DHPPA) and at 1.0 V (for 3-HPPA). Lowering the voltage makes the detection more selective for more readily oxidizable compounds [45] such as DHPPA, but making 3-HPPA invisible for detection. For the detection of the C. sporogenes metabolites and for peak isolation, another HPLC-ED system was used (Jasco AS2059 plus autosampler, Jasco Benelux, Utrecht, The Netherlands; Knauer K-1001 pump, Separations, H. I. Ambacht, The Netherlands) with the same detector (ED40) and the same gradient as described above. Phenylalanine metabolites were detected by injecting the same samples in an HPLC-UV system (Alliance Separations Module 2695, Waters Chromatography B.V, Etten-Leur, The Netherlands; TSP UV6000LP UV-detector (wavelength, 260 nM) Thermo Scientific, The Netherlands). Samples for peak isolation were separated on a Vydac Semi-preparative C18 column (218TP510, 5 μm, 300 Å, 10 mm × 250 mm, VWR International B.V, Amsterdam, The Netherlands) at 3 mL/min using the same gradient as above. Data recording and analysis were performed using Chromeleon software (version 6.8 SR13).
Significance was tested using a two-sample equal variance (homoscedastic) Student’s *t* test (Microsoft Excel 2019 version 1808).

**Catechol extraction from stool for DHPPA quantification**

Catechols were extracted from PD patients and HC stool samples using activated alumina powder (199966, Sigma) as previously described [14] with a few modifications. A volume of 200 μL 50% stool suspension (described above) was used with 1 mM DHBA (3,4-dihydroxybenzylamine hydrobromide, 858781, Sigma) as an internal standard. Samples were adjusted to pH 8.6 with 800 μL TE buffer (2.5% EDTA; 1.5 M Tris/HCl, pH 8.6), and 5–10 mg of alumina was added. Suspensions were mixed on a roller shaker at room temperature for 20 min and were sequentially centrifuged for 30 s at 20,000×g and washed three times with 1 mL of H 2O by aspiration. Catechols were eluted using 0.7% HClO 4 and filtered before injection into the HPLC-ED-system as described above (DC amperometry at 0.8 V). A standard curve was injected to quantify the concentrations of DHPPA in 50% (w/v) stool samples. Significance was tested using an unpaired nonparametric Mann-Whitney test (GraphPad Prism version 7).

**Organ-bath experiments**

Distal ileal samples were harvested from wild type adult (18–20 weeks) male C57BL/6J mice that were sacrificed for another purpose. Harvested tissue was washed three times with 1 mL of H 2O by aspiration. Catechols were eluted using 0.7% HClO 4 and filtered before injection into the HPLC-ED-system as described above (DC amperometry at 0.8 V). A standard curve was injected to quantify the concentrations of DHPPA in 50% (w/v) stool samples. Significance was tested using a two-sample equal variance (homoscedastic) Student’s *t* test (Microsoft Excel 2019 version 1808).

**HPLC-MS**

HPLC-MS analysis was performed using an Accella1250 HPLC system coupled with the benchtop ESI-MS Orbitrap Exactive (Thermo Fisher Scientific, San Jose, CA, USA) in negative and positive ion mode. Samples were analyzed on a C18 column (Shim Pack Shimadzu XR-ODS 3 × 75 mm) using a gradient of water/acetonitrile with 0.1% formic acid (0–5 min, 98–90% H 2O; 5–10 min, 90–5% H 2O; 10–13 min, 5% H 2O; 13–14 min, 98% H 2O). Data analysis was performed using Qual Browser Thermo Xcalibur software (version 2.2 SP1.48).

HPLC-MS analysis of alumina extraction samples was performed using a Waters Acquity Class-I UPLC (Waters Chromatography B.V., Etten-Leur, The Netherlands) system coupled to a Maxis Plus Q-TOF (Bruker, Billerica, MA, USA) on negative ion mode with post-column addition of 3 μL/min ESI Tune Mix (G1969-85000; Agilent Technologies, Middelburg, The Netherlands) for mass calibration. Samples were analyzed on a C18 column (Shim Pack Shimadzu XR-ODS 3 × 75 mm) using a 2-fold ranging from 8 to 512 μM. Data was recorded and analyzed in BioPac Student Lab 4.1 (Build: February 12, 2015). Frequencies were extracted performing a fast Fourier transform (FFT) on bins of 5-min intervals. The maximum amplitude of all the observed frequencies was extracted, and the average decrease of all frequencies over time was calculated. Significance was tested using repeated measures (RM) 1-way ANOVA followed by Tukey’s test (GraphPad Prism version 7).
gradient of water/acetonitrile with 0.1% formic acid (0–5 min, 98–90% H₂O; 5–10 min, 90–5% H₂O; 10–13 min, 5% H₂O; 13–15 min, 2% H₂O; 15–17 min, 98% H₂O). Data analysis was performed using Bruker Compass Data Analysis (version 4.2 SR1).

Bioinformatics

Phylogenetic trees

Proteins were BLASTed against a local BLAST database constructed from the protein sequences of the NIH Human Microbiome Project (HMP) Roadmap project (PRJNA43021) using BLAST 2.9.0+, NCBI. The top 100 BLASTp hits were aligned in the Constraint-based Multiple Alignment Tool (COBALT, NCBI) and converted to a distance tree using NCBI TreeView (Parameters: Fast Minimum Evolution; Max Seq Difference, 0.85; Distance, Grishin).

Sequence data analysis

The demultiplexed paired-end sequence data from stool and sigmoid colon samples of PD patients and healthy controls from Keshavarzian et al. [34] (bioproject PRJNA268515) were analyzed using Kraken2 (v2.0.9, April 7, 2020), a k-mer taxonomic classification system [46], using the standard Kraken2 database. To further estimate the species abundance, the Kraken2 output was analyzed with Bracken (Bayesian Reestimation of Abundance with Kraken; v2.6.0, April 3, 2020) [47]. The number of mapped reads from bacteria with the fld-gene cluster [11] was extracted from the Bracken results, and the abundance was calculated relative to the total number of mapped bacterial reads.

Supplementary information

Supplementary information accompanies this paper at https://doi.org/10.1186/s12915-020-00876-3.

Additional file 1: Supplementary Results – Metagenomic analysis of deaminating bacteria and E. lenta in PD and HC fecal and mucosal samples. Fig. S1 – NMR and MS confirmation of levodopa product, 3-(3,4-dihydroxyphenyl)propionic acid. Fig. S2 – 5-HT production. Fig. S4 – Initial effect of DHPPA on natural ileal contractility and no effect of levodopa on acetylcholine induced twitch. Fig. S5 – Phylogenetic tree of C. sporogenes FIDH and EDU8870. Fig. S6 – Fecal incubations from healthy age-matched controls. Fig. S7 – 3-HPPA is produced by E. lenta. Fig. S8 – Analysis of 16s RNA metagenomics data of deaminating bacteria and E. lenta in PD and HC fecal and mucosal samples. Table S1 – Values and statistical results corresponding to Fig. 2B. Table S2 – MS confirms that DHPPA is extracted from PD and HC samples using alumina extraction method. Table S3 – Plasmids and primers used in this study.

Additional file 2. All data underlying the main and supplementary figures and tables of the manuscript.

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Authors’ contributions

SP.K. and S.E.A. conceived and designed the study. SP.K., HRJ., SLW., SSL., SAN., HP., and AK. performed the experiments, and SP.K., SSSL., HP., and S.E.A. analyzed the data. SP.K. and S.E.A. wrote the original manuscript that was reviewed by SSSL., SAN., HP., and AK. Funding was acquired by S.E.A. All authors read and approved the final manuscript.

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Availability of data and materials

All data generated or analyzed during this study are included in this published article and its supplementary information files. The metagenomic sequence data collected by Keshavarzian et al. [34] were deposited under bioproject number PRJNA268515.

Ethics approval and consent to participate

Samples from participants were used from Keshavarzian et al. [34] (bioproject PRJNA268515) [46], where all subjects consented to use of their samples for research.

Consent for publication

Not applicable

Competing interests

The authors declare no competing interests.

Author details

1Department of Molecular Immunology and Microbiology, Groningen Biomolecular Sciences and Biotechnology Institute (GBB), University of Groningen, Nijenborgh 7, 9747 AG Groningen, The Netherlands. 2Current Address: Department of Laboratory Medicine, Cluster Human Nutrition & Health, University Medical Center Groningen (UMCG), Hanzeplein 1, 9713 GZ Groningen, The Netherlands. 3Department of Molecular Neurobiology, Groningen Institute for Evolutionary Life Sciences (GELIFES), University of Groningen, Nijenborgh 7, 9747 AG Groningen, The Netherlands. 4Interfaculty Mass Spectrometry Center, University of Groningen, Groningen, The Netherlands. 5Division of Digestive Disease and Nutrition, Section of Gastroenterology, Department of Internal Medicine, Rush University Medical Center, 1725 W. Harrison, Suite 206, Chicago, IL 60612, USA.

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