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RESEARCH PAPER



Freeze-dried fecal samples are biologically active after long-lasting storage and suited to fecal microbiota transplantation in a preclinical murine model of *Clostridioides difficile* infection

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ABSTRACT

Fecal microbiota transplantation is now recommended for treating recurrent forms of *Clostridioides difficile* infection. Recent studies have reported protocols using capsules of either frozen or freeze-dried stool allowing oral administration in in- and out-patient settings. However, a central question remains the viability, engraftment, and efficacy of the microbiome over time during storage life. This study shows that both the freeze-drying and freezing procedures for fecal samples allowed preserving viability, short-chain fatty acids concentration, and anti-*Clostridioides difficile* properties of microbiota without significant alteration after storage for 12 months. Fecal transplantation with freeze-dried microbiota allowed engraftment of microbiota leading to clearance of *Clostridioides difficile* infection in a preclinical murine model with a survival rate of 70% versus 53–60% in mice treated with frozen inocula, and 20% in the untreated group. Moreover, the freeze-dried powder can be used to fill oral hard capsules using a very low amount (0.5%) of glidant excipient, allowing oral formulation. Altogether, this study showed that freeze-dried inocula can be used for the treatment of *Clostridioides difficile* infection with long-lasting stability of the fecal microbiota. This formulation facilitates biobanking and allows the use of hard capsules, an essential step to simplify patient access to treatment.

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Introduction

Clostridioides difficile (CD, formerly *Clostridium difficile*), are one of the most common health care-associated infections, representing a major burden in terms of mortality, morbidity, and cost.^{1,2} The increasing prevalence, severity, and mortality related to CD infections in the last decade have led to a growing interest in new therapeutics based on the manipulation of the microbiota. Many concordant studies, including randomized controlled trials, have shown that fecal microbiota transplantation (FMT) is highly effective for treating recurrent forms of CD infection relative to conventional antibiotic treatment, with cure rates reaching 85% to 95%,^{3–6} and a recent report suggests it may be life-saving in

severe infections.⁷ Based on these trials, FMT is now strongly recommended by the European and US infectious disease societies.^{8,9} Beneficial effects of FMT have also been described in case series and small prospective trials for several other conditions associated with gastrointestinal microbial dysbiosis, including inflammatory bowel diseases,¹⁰ metabolic disorders,¹¹ neuropsychiatric conditions,¹² and the eradication of multi-resistant bacteria, i.e. extended-spectrum β -lactamase and/or carbapenem-resistant *Enterobacteriaceae*.¹³ However, no evidence-based recommendations have yet emerged for such conditions, except for clinical research.

The initial methods for FMT proposed the use of fresh material re-administered through retention enema, colonoscopy, or naso-duodenal delivery

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within a maximum period of 6 h between donor stool emission and recipient administration. Now, frozen fecal material is widely used and several randomized controlled trials have confirmed that the use of frozen-and-thawed material is non-inferior (*versus* fresh) in terms of cure rates for recurrent CD infection.^{4,6} A few studies have reported protocols using encapsulated frozen stool allowing oral administration in in- and out-patient settings with high cure rates in patients with recurrent infections.^{14–16} Pharmaceutical grade glycerol with final concentration ranging from 10% (G10) to 80% (G80) is currently used as microbial cryoprotectant for the preparation of fecal microbiota transplants.^{14–17} Costello *et al.* tested bacterial viability after 6 months of storage at -80°C using culture on plate media and showed that the microbiome remained largely unchanged when stored in G10.¹⁸ Recently, protocols using freeze-dried (FD) encapsulated fecal microbiota have been reported with similar cure rates.^{19–21} These new formulations for FMT represent important advances toward improving access to this therapy. However, apart from the health screening, no standard microbial and biological characterization of fecal transplant does exist and current practice remains mainly empirical. Minimal guidelines exist for the preparation and preservation of fecal transplant^{17,22} but a central question remains the consequence of long-term storage. In a model of CD infection, Jiang *et al.* showed that mice treated with frozen or FD samples stored for ≤ 7 -months were protected from post-CD challenge diarrhea.²³ Thus, the freeze-drying procedure seems to be adapted to FMT, but we lack scientific rationale establishing the biological activity over time of fecal samples after freeze-drying.

In the present study, we followed over 12 months of storage, the bacterial viability, metabolite production, engraftment in a recipient germ-free model, and CD inhibition of new FD *versus* classical frozen (G10 and G80) fecal transplants. The anti-CD activity was evaluated, both *in vitro* and in a murine preclinical model of CD infection. We show that the freeze-drying procedure preserves the biological activity of fecal samples after storage for 12 months at 4°C . This formulation is also compatible with the filling of hard oral capsules allowing oral formulation, and represents an easier solution for biobanking.

Results

Microbiota viability in freeze-dried and frozen fecal samples

We compared the viability of representative cultivable microbes from six human stool samples (donors #S1 to #S6) either fresh or preserved for a twelve-month period by either freeze-drying (samples FD, patent application WO2017103225A1) or freezing in 10% or 80% glycerol (samples G10 and G80). Total number of cultivable bacteria was 10.9 ± 0.6 log CFU/g of feces in fresh stool samples. No significant decrease in either total culturable bacteria or Extremely Oxygen-Sensitive bacteria (EOS) was observed until 12 months of storage regardless the procedure of preservation (Figure 1(a) and Table S1 supplementary data). Concerning anaerobic genera, there was no effect on the viability of *Bifidobacterium* spp. for up to 12 months, regardless of the procedure. In contrast, there was significantly less *Bacteroides* immediately after processing in the FD and G80 samples than in fresh stools ($p < .001$) and a subsequent decrease in the FD samples over time ($p < .01$ at M6 and M12) (Figure 1(a), Table S1 supplementary data). Concerning facultative aero-anaerobic genera, there was less *Enterobacteriaceae* after M3 in the FD samples ($p = .04$ at M3 and $p < .01$ at M6 and M12) and less *Enterococcus* spp. after 6 months of storage ($p < .01$ at M6 and $p = .04$ at M12) than in fresh stools. No other significant difference was observed regardless of the procedure (Figure 1(b), Table S1 supplementary data). No variation in *Lactobacilli* was observed over time whatever the procedure. Cultivable *Clostridia* (i.e. cluster I–II) were at very low levels within the microbiota whatever the sample, making it impossible to count. CD was only detected in one out of the six samples. Finally, the different profiles observed within the six donors at inclusion were preserved during storage whatever the process of preservation. No further significant modification was observed in two samples until 18 months of storage (Table S1, supplementary data).

Short-chain fatty acids concentrations in fecal samples after 12 months of storage

Total short-chain fatty acids (SCFA) concentrations ranged from 114.8 ± 31 μmol per g of feces in FD

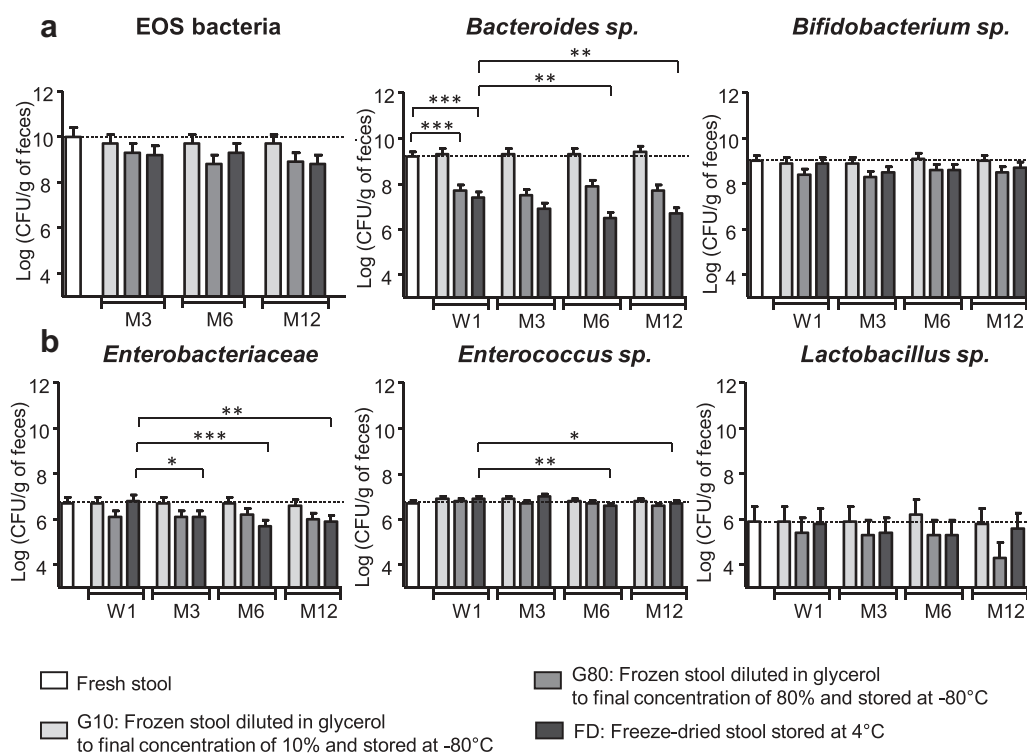


Figure 1. Microbiota viability of freeze-dried and frozen human fecal samples over 12 months of storage. (a) Dominant microbiota: EOS bacteria ($n = 6$), *Bacteroides sp.* ($n = 6$), and *Bifidobacterium sp.* ($n = 6$) (b) Subdominant microbiota: Enterobacteriaceae ($n = 6$), *Enterococcus sp.* ($n = 5$), and *Lactobacillus sp.* ($n = 3$). Time study: Week 1, W1; Month 3,6 and 12: M3, M6, M12. Results are expressed as the mean \pm SEM. Statistical significance: * <0.05 , ** <0.01 , *** <0.001 .

samples to 52.7 ± 30.5 μmol per g of feces in samples frozen without any cryoprotectant at M12. SCFA concentrations were significantly higher in stool samples preserved in G10, G80, and FD than in the corresponding samples stored without cryoprotectant ($p < .001$). This was related to higher concentrations of acetate in the G10 ($p = .028$), G80 ($p = .049$), and FD ($p < .001$) samples (Figure 2(a), Table S2 supplementary data).

In vitro anti-Clostridioides difficile activity in freeze-dried and frozen samples after 12 months of storage

A comparison of the effect of the freeze-drying and freezing procedures on anti-CD activity was carried out after 12 months of storage in comparison to pure bacterial culture used as controls. *Escherichia coli*, *Staphylococcus aureus*, and *Klebsiella pneumoniae* reference strains (negative controls) did not inhibit the growth of CD, whereas both *Lactobacillus spp.* and *Bifidobacterium infantis* reference strains were able to inhibit the growth of CD with an antagonistic

effect significantly higher for *Lactobacillus* (inhibition ratio normalized to lactic acid 500 mM: $161 \pm 3\%$ versus $129 \pm 3\%$, $p < .001$). All fecal samples were antagonistic toward CD R20291 (027 type strain), with results similar for the three conditions and results similar to those observed with the *Bifidobacterium* reference strain. The inhibition ratio normalized to lactic acid (500 mM) was $131 \pm 6\%$ for FD samples; $141 \pm 6\%$ for G10 samples and $138 \pm 6\%$ for G80 samples (Figure 2(b)).

Colonization of germ-free mice with rehydrated freeze-dried microbiota

We studied the ability of rehydrated FD human fecal microbiota, to colonize the digestive tract of germ-free mice. Two donors were randomly selected (#S4-FD and #S5-FD). After storage for 12 months at 4°C, the rehydrated FD microbiota was inoculated by one oral gavage in recipient germ-free mice ($n = 4$ per stool sample) and fecal samples were collected for microbial analysis at day 1 (D1), 4 (D4), 11 (D11), 18 (D18) and 28 (D28)

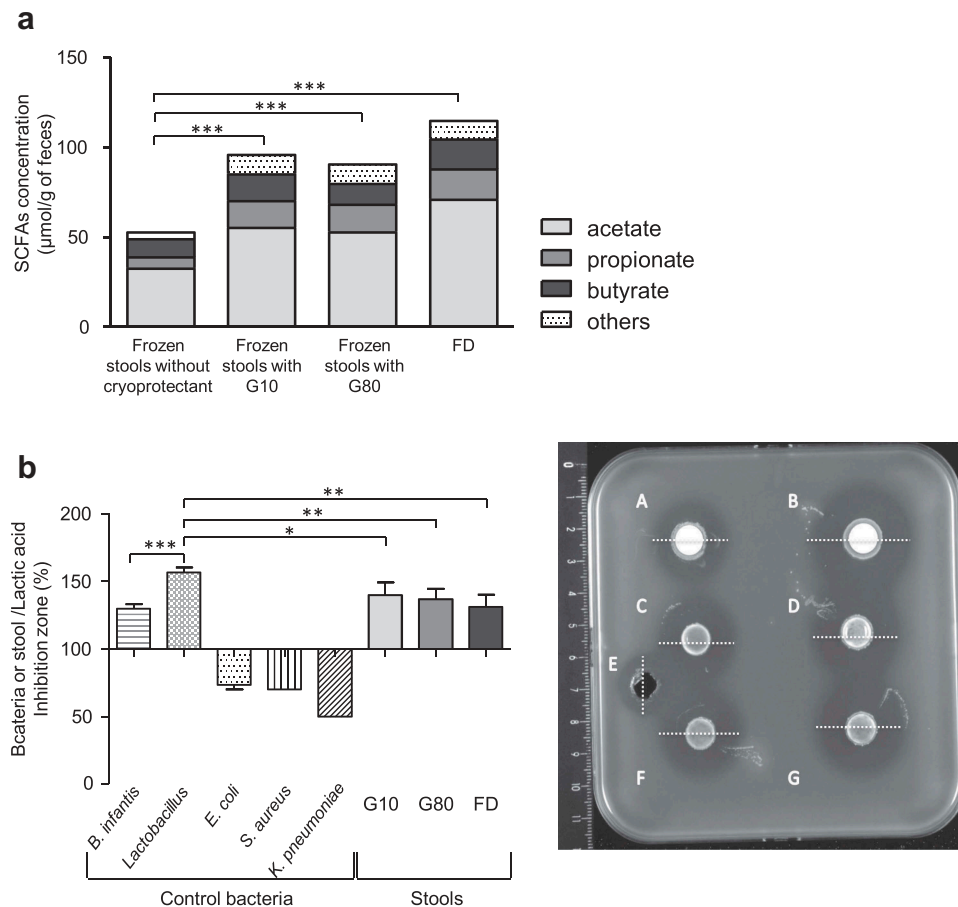


Figure 2. Fecal metabolic activity of frozen and freeze-dried human fecal microbiota after 12 months of storage. (a) Amount of Short-Chain Fatty Acids at M12. Results are expressed as the mean \pm SEM. (b) Anti-CD activity of stool samples at M12. Left: results are given as the ratio between inhibition diameters of pure bacteria or stool samples and chemical control, and expressed as the mean \pm SEM. Right: representative image of an anti-CD agar spot test: A: *Bifidobacterium infantis longum* CUETM 89-215, B: *Lactobacillus* spp., C: stool frozen in G10, D: stool frozen in G80, E: chemical control (lactic acid, 500 mM), F: stool frozen in G10, G: stool lyophilized FD. White lines represent inhibition diameter. Statistical significance: * <0.05 , ** <0.01 , *** <0.001 .

after gavage (Figure 3(a)). Engraftment of fecal microbiota was rapid and stable over time for the two stool samples (Figure 3(b)). The total bacterial load and colonization by major groups of commensals like the *Clostridium coccoides* cluster were high as soon as D1. Maximal colonization by *Clostridium leptum* and *Bacteroides* groups was achieved by D4, reaching 10^9 and 10^{10} CFU/g of feces, respectively. Colonization by EOS bacteria, such as *Faecalibacterium prausnitzii* and *Akkermansia muciniphila* was also high as soon as D1, without any significant evolution over the survey period (Figure 3(b)). Engraftment of the microbiota led to the morphological remodeling of epithelium

with a significant increase in crypt depth for #S5-FD inoculated mice compared to that of germ-free mice: $188.3 \mu\text{m}$ [(25-75% percentile: 181.5–194.4) versus $166 \mu\text{m}$ (25-75% percentile: 124–174), $p < .05$] (Figure 3(b)). A similar trend was observed for #S4-FD [$172.4 \mu\text{m}$, 25-75% percentile: 145–186.6], but the difference relative to germ-free mice did not reach significance. There was no significant difference in the percentage of Ki67-positive cells between groups: germ-free mice: 20.1% (25-75%: 16.5–23.0); #S4-FD: 21.8% (25-75%: 20.7–29.0), and #S5-FD: 18.5% (25-75%: 17–19.8) (Figure 3(c)). The 2 FD fecal samples were thus revivable after 12 months of storage and

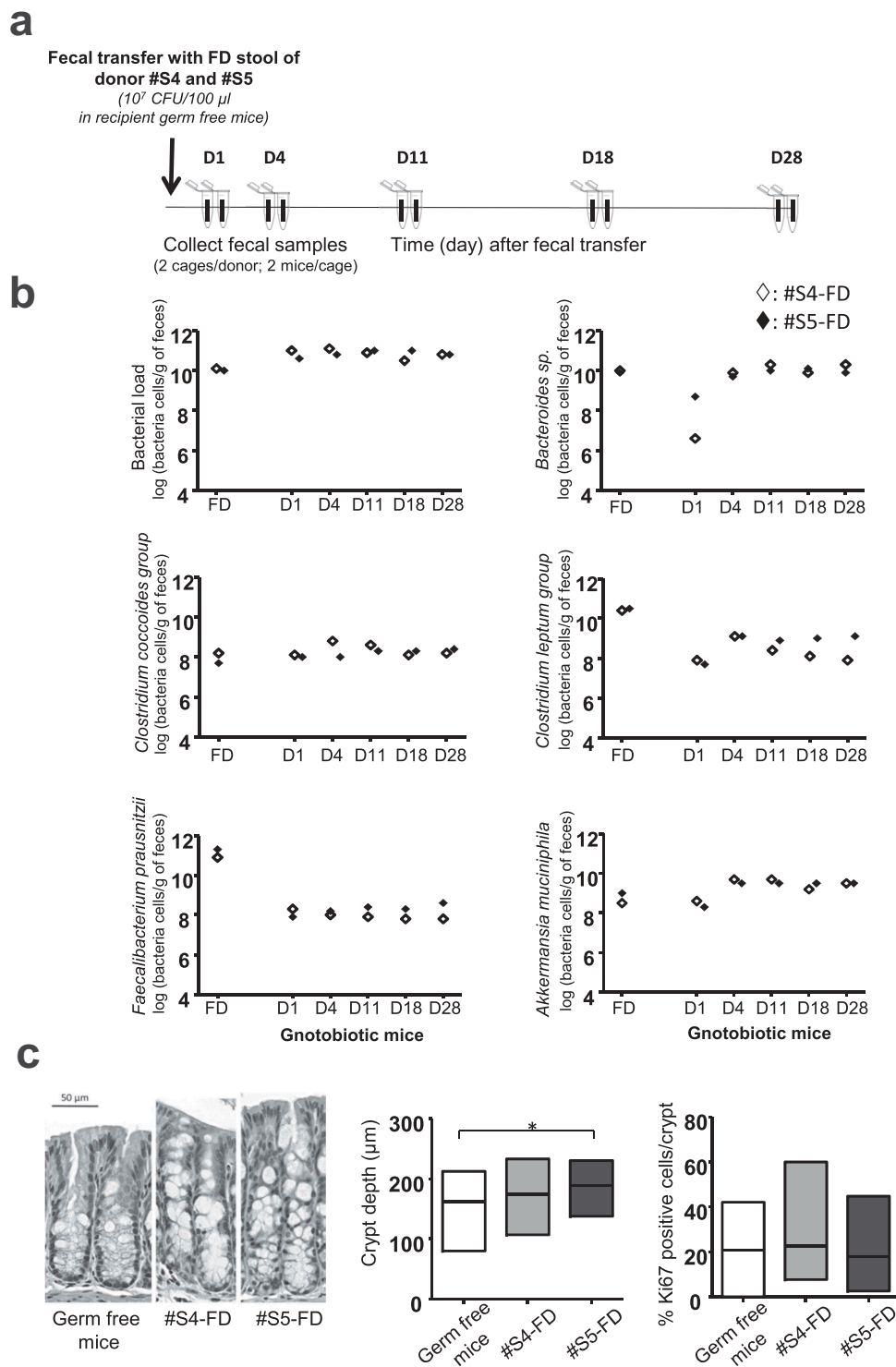


Figure 3. Ability of freeze-dried human fecal microbiota to colonize the digestive tract of germ-free mice. (a) Scheme of the experiment. (b) Follow-up of main bacterial groups per gram of feces after colonization with rehydrated FD microbiota [; #S4-FD (n = 4); #S5-FD (n = 4), feces sampled per cage]. Results are expressed as the mean for each of the 2 FD samples. (c) Representative image of a colon section from germ-free mice colonized with rehydrated FD microbiota (D30). Scale bar, 50 μ m; crypt depth in the colon of germ free and mice colonized with rehydrated FD microbiota (30 crypts per mouse were measured) and Ki67-positive cells expressed as the percentage of total cell number in colon crypts from germ-free mice colonized with rehydrated FD microbiota (60 crypts per mouse were analyzed). Statistical significance: $* < 0.05$.

were able to colonize the digestive tract of recipient germ-free mice without hyper-proliferation of the intestinal epithelium.

Therapeutic efficacy of freeze-dried and frozen human fecal microbiota in *Clostridioides difficile* infected mice

We then evaluated the protective effect of FMT performed with stools from two donors randomly

selected (#S1 and #S6) either frozen in G10 (FMT-G10 group, n = 10 per stool sample) or G80 (FMT-G80 group, n = 10 per stool sample) or FD (FMT-FD group, n = 10 per stool sample), in adult C57/BL6 mice orally challenged with $8 \cdot 10^3$ CFU of the vegetative form of CD R20291 strain 027 after antibiotic pretreatment (Figure 4(a)). Four uninfected mice were used as negative controls (CD- group). CD challenge led to 80% of the mice becoming moribund by day 2 or 3 in untreated animals (CD+ group,

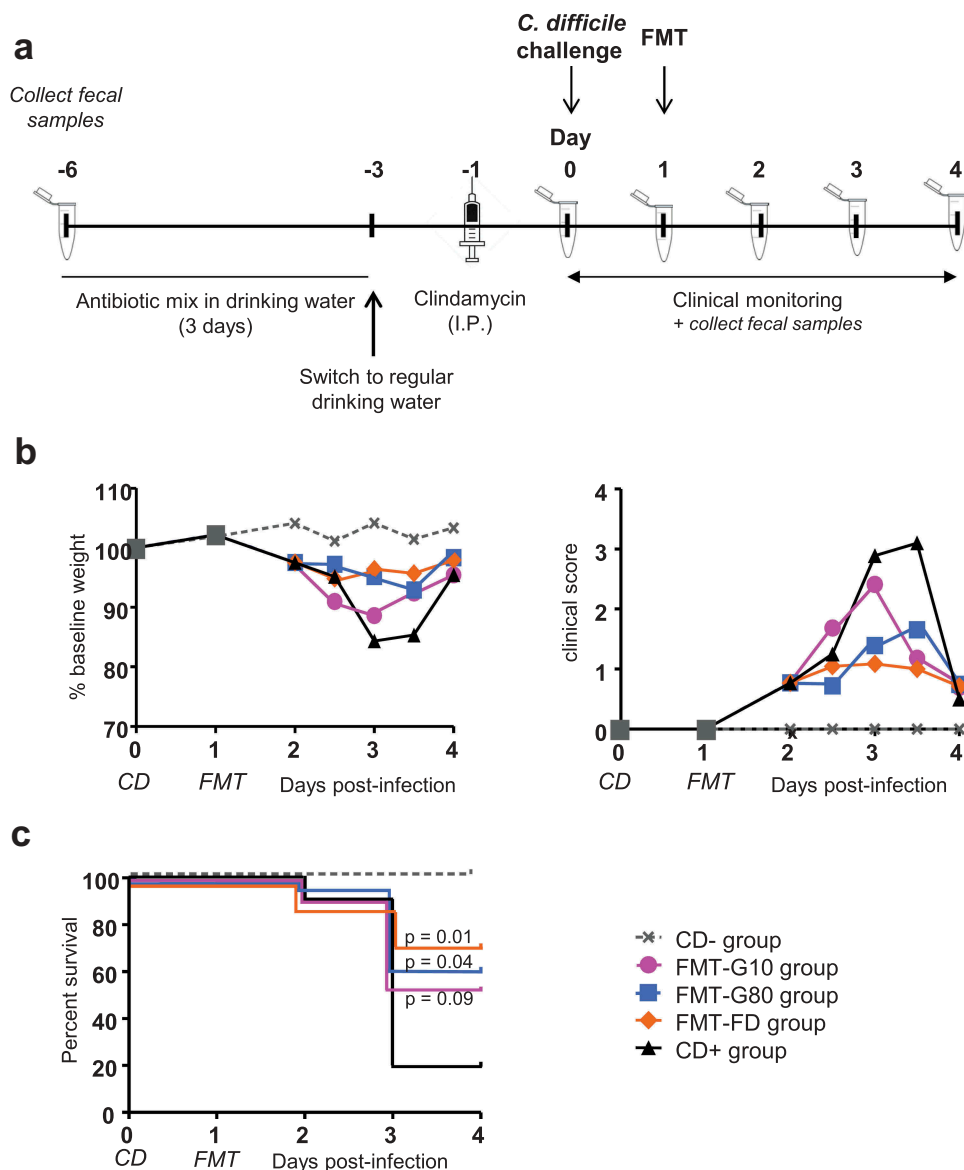


Figure 4. Evolution of clinical parameters after frozen and freeze-dried FMT in *Clostridioides difficile*-infected mice. (a) Experimental design of the CD-infected-FMT mouse model. Two stools samples were evaluated #S1 and #S6. (b) Monitoring of body weight and a clinical score of all survival mice (up to the day of death). Daily measurements were made starting from Day 0 (day of infection). Grey squares show the total number of mice before randomization into the various FMT groups. Results are expressed as the mean (c). Survival curves of CD-infected mice receiving FMT with FD or frozen microbiota. CD-: control group receiving PBS (n = 4), CD+: CD-infected mice (n = 10); FMT-G10: CD-infected mice receiving FMT with frozen microbiota diluted in G10; FMT-G80: CD-infected mice receiving FMT with frozen microbiota diluted in G80 and FMT-FD: CD-infected mice receiving FMT with freeze-dried microbiota. For each CD group receiving FMT, n = 20, i.e. 10 mice per stool sample.

Table 1. Clinical score performed daily on animals after *Clostridioides difficile* infection. The score includes the variables “weight” and “behavior,” with a score from 0 to 3. The final score was obtained by adding the two scores. Mice with a total score > 3 were euthanized within 2 h.

score	Behaviour	Weight
0	Normal	No weight loss
1	Decrease in grooming	Weight loss < 10%
2	Matted fur and/or slitted eyes	10% ≤ weight loss
	Decrease in mobility	≤20%
3	Prostration associated with matted fur and slitted eyes	Weight loss > 20%

n = 10), leading to euthanasia. CD+ surviving mice displayed a major loss in body weight compared to day 0 ($-16.0 \pm 2.4\%$) and a high clinical score (3.0 ± 0.4) at day 3 post-infection (Figure 4(b), Table 1).

The survival rate of mice at day 4 post-infection was significantly higher in those receiving FMT, reaching 70%, 53%, and 60% for the FMT-FD, FMT-G10 and FMT-G80 group, respectively ($p = .01$, 0.09 and 0.04 compared to the CD+ group, Figure 4(c)). This was associated with a reduced loss in body weight in the FMT-FD and FMT-80 groups ($-4.5 \pm 1.7\%$, $p < .001$, and $-5.1 \pm 1.7\%$, $p = .002$, compared to CD+ mice) (Figure 4(b), Table 2). The clinical score and cecal load were also significantly reduced for the FMT-FD and FMT-G80 groups ($p < .01$) (Figures 4(b) and 5(a), Table 2).

At the time of necropsy, colonic myeloperoxidase (MPO) was measured as a marker of inflammation.

Median MPO levels reached 2465 ng/g of colon tissue in CD+ untreated mice. In contrast, animals that received FMT, performed either with rehydrated FD or frozen and thawed stools showed significantly lower colonic inflammation, with median MPO levels at least five times lower (<500 ng/g, $p < .01$) (Figure 5(a), Table 2). Histopathological analyses were performed at day 4 post-infection for mice receiving FMT and for CD-group, and at the time of necropsy for mice from the CD+ group. Six randomly selected animals were analyzed per group except for the CD-group (n = 4). There were histopathological changes in the cecal and colonic tissues of all CD-infected mice, but of varying severity, depending on the type of treatment (Figure 5(b)). We observed the maximal histopathological score in CD+ mice: 7.5 (25-75% percentile: 5.5–9) in the cecum and 7 (25-75% percentile: 5.75–7.25) in the colon. Treatment with FD-FMT was the most efficient to reduce histological damages ($p < .01$) (Figure 5(c), Table 2).

Finally, the infection was associated with a marked decrease in α -diversity of microbiota compared to CD-mice, as shown by the OTU number and Shannon indices (Figure 6(a,c)). FMT performed at day 1 post-infection restored the α -diversity only for #S1 ($p < .05$), regardless of the procedure of conservation of fecal samples (Figure 6(a)), whereas there was no major impact on β -diversity (Figure 6(b)). However, microbiota composition was clearly modified following FMT.

Table 2. Main characteristics of the effect of FMT in the preclinical mouse model of *Clostridioides difficile* infection. Clinical data were measured at day 3 post-infection and biological results were measured at the time of necropsy. Clinical data are given as mean \pm SEM and biological and histological data as median, 25-75% percentile. FMT-FD: CD-infected mice receiving FMT with freeze-dried microbiota; FMT-G10: CD-infected mice receiving FMT with frozen microbiota diluted in G10; FMT-G80: CD-infected mice receiving FMT with frozen microbiota diluted in G80. For each CD group treated with FMT, n = 20, i.e. 10 mice per stool sample. CD-infected mice (n = 10). All results are compared to those of untreated CD+ mice.

	FMT-FD	FMT-G10	FMT-G80	CD-infected mice
Body weight loss vs day 0	$-4.5 \pm 1.7\%$ ($p < .001$)	$-11.8\% \pm 1.8$ (NS)	-5.1 ± 1.7 ($p = .002$)	-16.0 ± 3.0
Clinical score	1.2 ± 0.3 ($p < .01$)	2.5 ± 0.3 (NS)	1.4 ± 0.3 ($p < .01$)	3.0 ± 0.4
CD load (log CFU/g of cecal content)	4.9 (3.3–6.3) ($p = .007$)	5.8 (4.8–6.7) ($p = .06$)	4.9 (3.6–6.4) ($p = .005$)	6.8 (6.3–9.9)
MPO (ng/g of colonic tissue)	300 (209.3–1054) ($p = .0005$)	384 ng (286.2–1529) ($p = .002$)	446 (290.3–890.8) ($p = .0002$)	2465 (1512–8928)
Histopathological score	1.5 0–2.75 ($p = .003$)	4.5 3–5.75 ($p = .03$)	4.5 2.75–6.75 4.5 2–7.5	7.5 5.5–9 7 5.75–7.25
● Cecum				
● Colon	1.5 0.75–4.5 ($p = .005$)	3 1.75–4.25 ($p = .004$)		

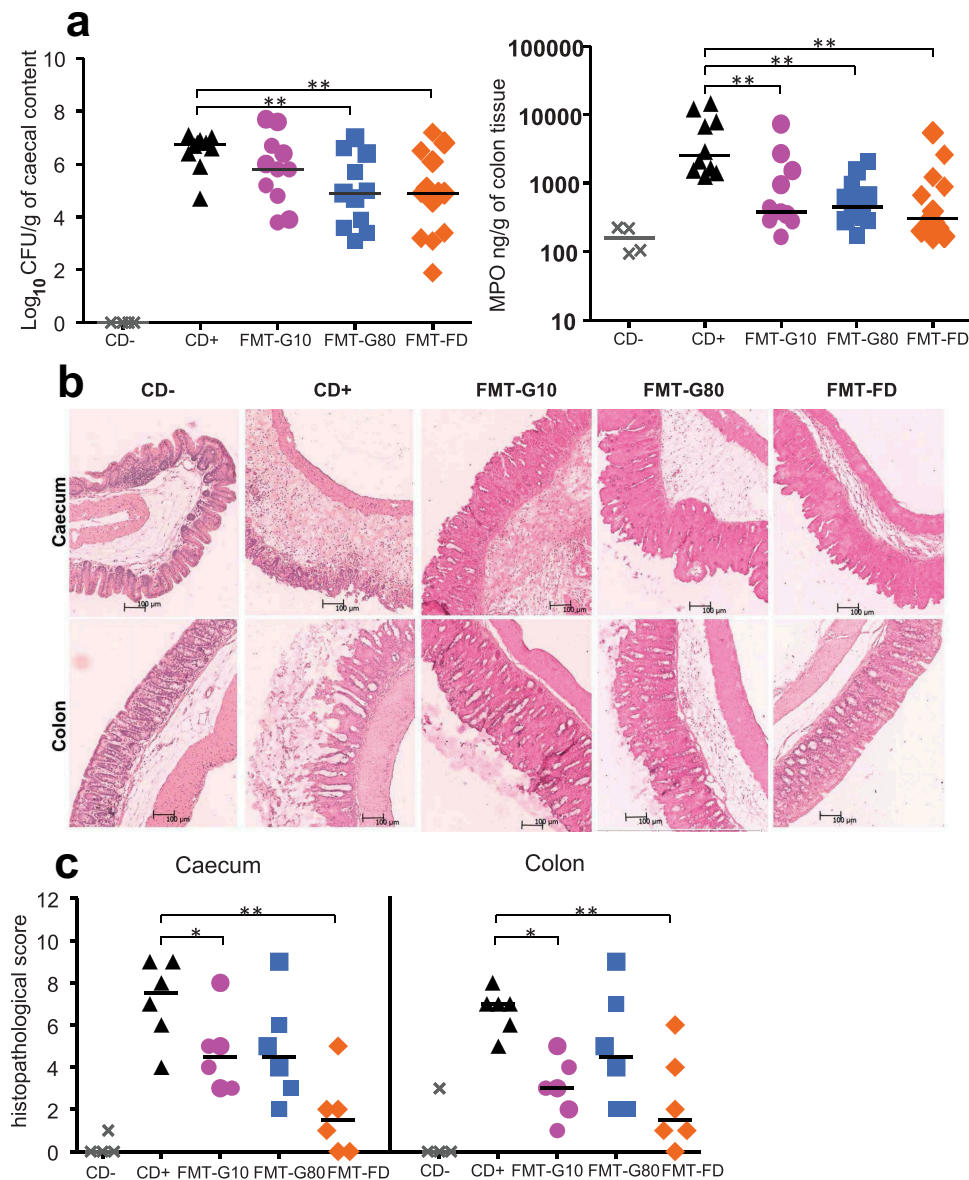


Figure 5. Evolution of the biological parameters of *Clostridioides difficile*-infected mice following frozen or freeze-dried FMT (a) Level of CD in the caecal content and level of MPO in colonic tissue at the time of necropsy. (b) Representative images of colonic and cecal sections from control (CD- and CD+) mice and mice receiving FMT with frozen or FD microbiota (day 4 post-infection). Scale bar, 100 μm. (c) Histopathological score measured from CD-infected mice receiving FMT with frozen or FD microbiota. Six mice randomly selected for each point. Statistical significance: * <0.05 , ** <0.01 .

Development of a formulation adapted for the filling of capsules with freeze-dried fecal microbiota for oral administration

To propose a formulation adapted for the filling of hard capsules, we determined the bulk density, particle size and aspect, and the flowability of the FD samples. We observed variable macroscopic and microscopic aspects, depending on the sample

(Figure 7(a,b)). It was not possible to qualify the particle-size distribution of the FD samples due to the extreme intra- and inter-sample diversity of particle nature, shape, and size. Scanning electron microscopy showed a microbiota embedded in its fecal environment (Figure 7(c)). Filling of hard capsules was not possible without excipients. Addition of up to 50% fillers failed to improve

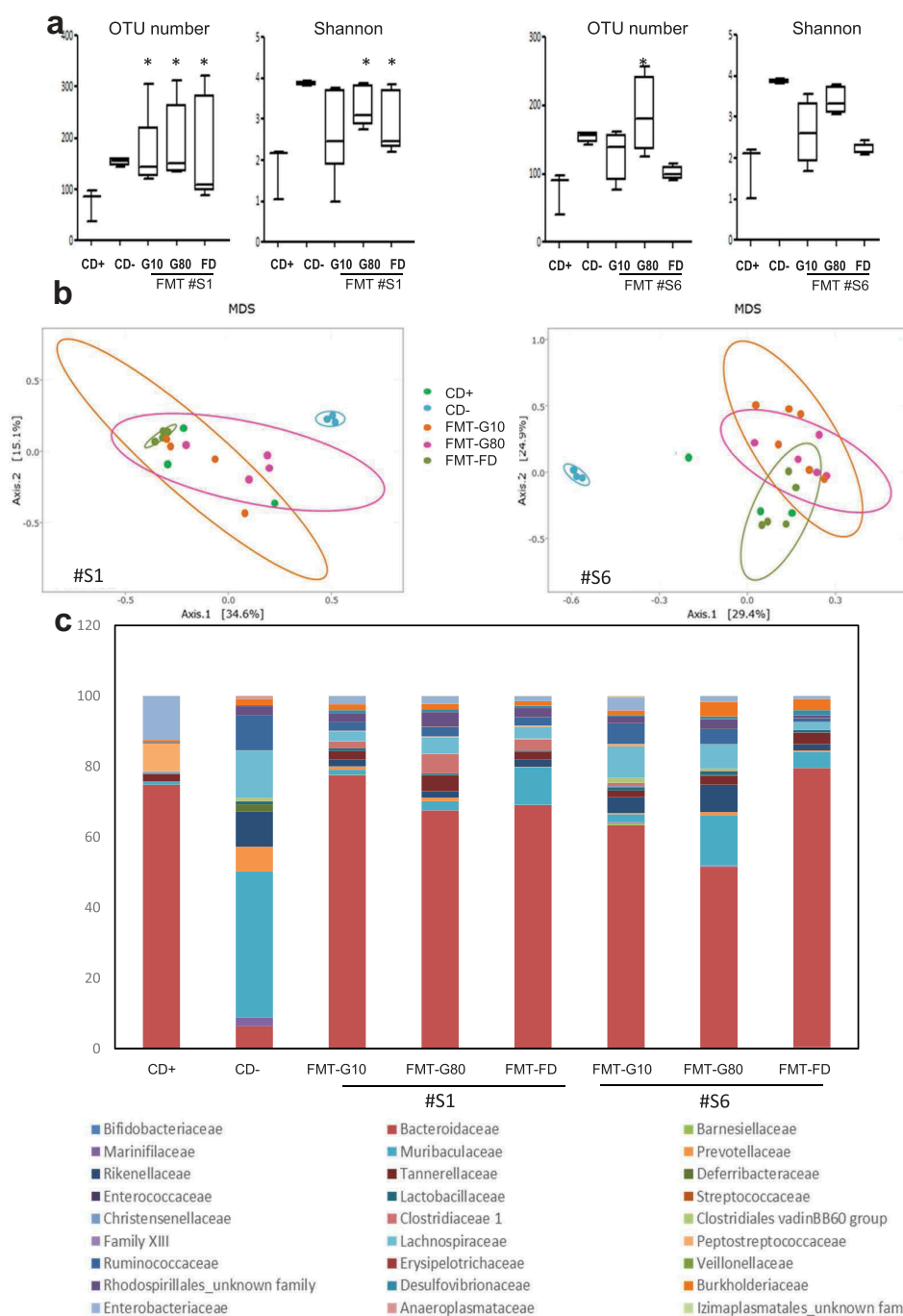


Figure 6. Comparison of the microbiota at day 4 between control mice and *Clostridioides difficile*-infected mice that received, or not, frozen or freeze-dried FMT. (a) α -diversity. (b) β -diversity using Bray–Curtis distance. (c) Composition of the microbiota at the family level. The two stool samples, #S1 and #S6, used for FMT are represented. Statistical significance: $* < 0.05$.

flowability, but 75% allowed the FD samples to flow vertically. Addition of glidants G3 (0.5%) was highly effective in improving the flowability of the FD samples and the bulk density of the free-flowing mixture was 0.30–0.47 g/mL allowing the filling of hard capsules without dilution of the FD samples (Figure 7, lower panel). In such

a condition, 1 g of powder corresponds approximately to 1 g of native stool.

Discussion

In the present study, we showed that bacterial viability and metabolic production were preserved

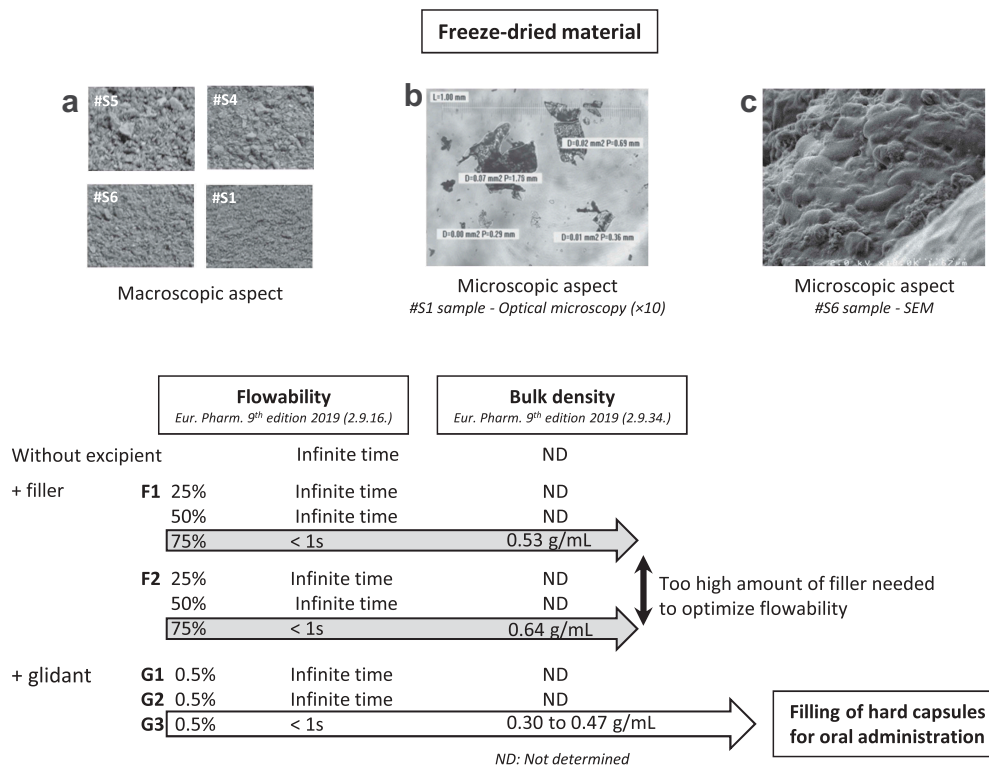


Figure 7. Development of hard capsules for oral administration of freeze-dried fecal microbiota. Upper panel: (a) examples of macroscopic aspects ($n = 4$). (b) microscopic aspects Optical Microscopy ($\times 10$) and (c) SEM. Lower panel: flowability and bulk density of FD samples without excipient ($n = 5$), with 25% to 75% filler ($n = 2$) or with glidant (G1 $n = 1$, G2 $n = 1$, G3 $n = 6$), ND: not determined.

in FD, maltodextrin-trehalose-loaded microbiota stored for at least 12 months at 4°C without significant variation compared to microbiota preserved frozen at -80°C in the presence of glycerol. Using this new FD preparation of fecal microbiota, engraftment was observed to levels usually observed in gnotobiotic rodent models and the therapeutic effectiveness of FMT was confirmed in a preclinical mice model of CD infection with a clear improvement in survival, clinical score, and histopathological lesions in the hindgut. Finally, it was possible to use the FD powder to fill oral hard capsules after the addition of a very small amount of a glidant excipient. Altogether, this work showed that the bacterial viability and clinical efficacy of the fecal microbiota were maintained in this new FD fecal preparation, allowing refinement of the FMT procedure toward oral administration of a dry material that was stable for over 1 year at 4°C.

In recent years, several important advances have been made in an effort to improve the accessibility

of FMT through the administration of frozen^{4,6,16,24} or FD stool.^{19–21} However, these formulations remained to be better standardized and characterized to guarantee the availability and efficacy of fecal transplants.

Here, we assessed the performance of a new freeze-drying process using non-penetrating cryoprotectants, and compared over time results with those obtained with the corresponding samples stored frozen at -80°C after dilution in glycerol, a penetrating cryoprotective agent at both concentrations (G10 and G80) currently used for the preservation of frozen fecal microbiota in clinical assays of FMT.^{14,17,22} We showed the good preservation until 12 months of storage, and even 18 months for two samples regardless the process of preservation, of both viability and diversity of the main targeted bacterial groups, *i.e.* the dominant (EOS and *Bifidobacteria*) or subdominant populations (*Enterobacteria*, *Enterococcus* spp., and *Lactobacillus* spp.) in the inocula. The main variation was observed for *Bacteroides* spp. This

could be due to the aerobic preparation of fecal transplant for samples preserved FD or frozen in G80, leading to damaged cells which would affect the culture and delay colonization which however remain efficient in recipient germ-free mice. As most anaerobic bacteria are not easily cultivable (i.e. *Faecalibacterium* or *Akkermansia*) and even culturability do not imply engraftment, we evaluated the engraftment of those FD microbiota. For *F. prausnitzii*, engraftment was coherent with our previous observations in two gnotobiotic models.^{25,26} Moreover, personal preliminary data suggest that FMT performed with either fresh or FD stools have a similar effect on the epithelium of recipient germ-free mice. Recently, it was suggested that the anaerobic treatment of stool samples lead to a better efficiency of FMT in patients with ulcerative colitis.²⁷ For others, precautions to ensure anaerobiosis during transplant manufacturing was not essential to preserve bacterial viability.²⁸ Here, using germ-free mice, we further showed that the aerobic preparation of transplants did allow a prompt and stable engraftment of the microbiota participating to gut homeostasis. This included major butyrate producers bacteria that are particularly sensitive to growth conditions, such as *F. prausnitzii*²⁹⁻³¹ which is known for its potential anti-CD activity³² and for its importance in human health.³³ Concomitant with bacterial engraftment, we observed deepening of the epithelium, especially for one donor, without hyperproliferation, thus reflecting the establishment of a host-bacteria relationship.³⁴⁻³⁶

We evaluated the protective effect of two fecal FD samples and the same two samples simply frozen in G10 or G80 at -80°C (all samples stored for 3 months) in a preclinical murine model of CD infection. The clinical parameters (low survival rate and major weight loss) observed in the CD-infected mice were consistent with those reported in the literature.³⁷⁻³⁹ FMT performed with rehydrated FD microbiota resulted in a significant decrease in symptomatology, i.e. a significant reduction in body weight loss, clinical score, and mortality relative to untreated CD-infected mice. These results are consistent with those of Jiang et al. who showed that frozen and FD transplants stored up to 7 months were both as effective as a fresh transplant in a mouse model of CD infection.²³ The beneficial effects observed for the

clinical symptoms were consistent with the biological parameters as animals receiving FD-FMT showed a reduced CD load in the cecum, limited colonic inflammation, and limited histological damages. That CD was still detectable in the cecum at the time of sacrifice (i.e. D4 post-infection) coincides with previous results where complete clearance of the pathogen was observed only by D10 after FMT with fresh stool in a murine model of relapsing disease.⁴⁰ Based on our *in vitro* studies showing no significant evolution of microbiota profile between M3 and M12, we can expect that samples stored for up to 12 months should be as efficient for the FMT procedure.

Besides the reconstitution of a diverse microbiota allowing the reinforcement of the barrier effect, various components of the fecal matter can provide health benefits and may work synergistically to restore homeostasis. Among them, SCFA (mainly acetate, propionate, and butyrate) are of major importance in the host-bacteria relationship.⁴¹⁻⁴³ Indeed, a decrease in *Ruminococcaceae*, *Lachnospiraceae*, and butyrate-producing bacteria that belong to *Clostridium* clusters IV and XIVa have been reported for patients with CD infection.⁴⁴ Here, we showed that freeze-drying in maltodextrin-trehalose or freezing in G10 or G80, better preserved SCFA during storage compared to samples frozen without any cryoprotectant. This included butyrate, which has a well-established role as an energetic source for the colonocytes, in promoting the intestinal barrier, and as an anti-inflammatory substrate.⁴⁵ Moreover, high concentrations of acetate, a key element for the development of butyrate-producing microbes through cross-feeding,⁴⁶ was detected in the cryoprotected stool samples.

However, these *in vitro* measurements are only one clue to, not a proof of high SCFA production in the recipient animals.

Overall, our results validate both the freezing and freeze-drying processes for the conservation of an efficient fecal transplant for up to 12 months. Moreover, we showed that the new FD fecal preparation for FMT lead to a significant decrease in symptoms together with an increase in survival rate in mice infected with CD. Such preservation methods would simplify clinical works by facilitating the fecal bio-banking from carefully selected and screened donors. In addition, the FD process allowed producing a powdery material with good stability over 12 months,

offering the possibility to prepare hard capsules for more convenient oral administration to the patient.

Methods

Preparation of stool samples

Fecal samples were collected from 6 healthy volunteers (none of whom suffering from gastrointestinal disorders or taking antibiotics in the previous 3 months), placed immediately in anaerobic conditions using a catalyst (GENbag Anaert; Biomérieux) according to the Human Microbiome Standards⁴⁷ and sent to the laboratory within 2 h of collection. Written informed consent was provided by all volunteers. For the freeze-drying procedure, fresh stool were immediately diluted 1/6 (v/v) in the maltodextrin-trehalose cocktail (patent application WO2017103225A1), homogenized in aerobic condition using a dedicated blender, filtrated with sterile gauze and frozen at -80°C for a minimum of 8 h, and subjected to the freeze-drying process. Vials were sealed under vacuum at the end of the process and stored at $+4^{\circ}\text{C}$ until analysis. At the same time, aliquots of the same stool samples were prepared as described above with dilution 1/6 (v/v) in G10 or G80, used as a cryoprotectant, and stored at -80°C .

Evaluation of microbiota viability in human fecal samples preserved by freeze-drying or freezing

Viability of the fecal microbiota in FD or frozen samples was evaluated after 1 week (W1) and 3, 6, or 12 months (M3, M6, M12) of storage and compared to that of the corresponding fresh stool samples, using culture methods for aerobic and anaerobic bacteria. Two samples were studied until 18 months (M18) of storage. Briefly, samples were serially diluted in sterile peptone water before spreading on several culture media with an automatic spiral system (Chemunex-AES Laboratory, France), allowing the enumeration of total bacteria, *Bifidobacterium* spp., *Bacteroides* spp.; *Enterobacteriaceae*; *Enterococcus* spp., and *Lactobacillus* spp. Plates were incubated at 37°C under aerobic or anaerobic conditions for 2 to 5 days as previously described.⁴⁸ Bacterial counts were expressed as log CFU/g of feces. The threshold of detection was 3 log CFU/g of feces. EOS bacteria were cultured on a rich nonselective culture medium

(YBHI supplemented with cellobiose, maltose, and cysteine) under anaerobic conditions ($\text{N}_2/\text{CO}_2/\text{H}_2$:80/10/10) for enumeration. Each dilution was spread on two plates, one being placed under anaerobic conditions, whereas the other was left out of the anaerobic chamber for 1 h, to eliminate EOS strains, before proceeding to incubation under anaerobic conditions for 48 h. The difference between the number of colonies on the two plates corresponded to the level of EOS strains.²⁹

Evaluation of the fermentative activity of human fecal microbiota

The concentrations of total SCFAs, acetate, propionate, and butyrate were measured by gas-liquid chromatography (Nelson 1020; Perkin-Elmer, France) after a 12-month period of storage in frozen and FD samples in comparison to control frozen samples stored at -80°C without cryoprotectants, as described previously.⁴⁹ Results are presented as $\mu\text{mol/g}$ of feces.

In vitro evaluation of the anti-Clostridioides difficile activity of human fecal microbiota

The agar spot test adapted from Tejero-Sarinena *et al.*⁵⁰ was used for the detection of activity against CD R20291 (027 type strain). Lactic acid (500 mM), *Lactobacillus* sp., and *Bifidobacterium infantis* subsp. *longum* CUETM 89–215 were used as positive chemical and biological controls, whereas *Escherichia coli* CIP 54.8, *Klebsiella pneumoniae* CIP104771, and *Staphylococcus aureus* ATCC 25923 were used as negative controls. An overnight culture of each bacterial suspension (20 μl) was used as control and an equal volume of each fecal suspension of thawed G10 and G80 and rehydrated FD samples were deposited on cellulose disks and placed on WCB agar for overnight incubation at 37°C in an anaerobic atmosphere. Agar plates were then covered with Brucella agar containing a CD R20291 culture (OD = 1). The chemical control, lactic acid, was added to an open well (50 μl). Plates were incubated for another overnight period under anaerobic conditions at 37°C and the zones of inhibition of the CD culture around each spot were measured. Results (from three independent experiments) are expressed as the ratio between inhibition diameters

of pure bacteria or stool samples and chemical control.

Preparation of hard oral capsules containing freeze-dried fecal microbiota

The bulk density (2.9.34. Eur. Pharm. 9th edition 2019),⁵¹ particle size (Optika Vision Pro, Optika SRL, Italy), and aspect (Scanning Electron Microscopy) of the FD samples were assessed. Flow properties were measured (2.9.16. Eur. Pharm. 9th edition 2019) with and without the addition of 25% to 75% directly compressible fillers with good flow properties, designated as F1 and F2, or 0.5% glidant excipients, designated as G1, G2, and G3. Gastro-resistant hydroxypropyl methylcellulose (HPMC) capsules (DRcaps V43-700, size 00, Capsugel, Belgium) were manually filled with the FD samples with optimized flow properties.

Animal experiments

Two experimental models, performed in accordance with the European guidelines for the care and use of laboratory animals, were used for this study.

Evaluation of intestinal colonization by rehydrated freeze-dried microbiota in germ-free mice

Thirteen-week-old female C3 H/HeN germ-free mice (n = 8) (INRAE, Anaxem, Jouy-en-Josas, France), bred under axenic conditions (two/cage), were orally inoculated with 100 μ L (10^7 bacteria) of 2 randomly selected rehydrated FD samples (#S4-FD and #S5-FD, n = 4 for each donor) that had been stored for 12 months at +4°C. Before fecal transfer, the absence of microbes was verified in germ-free mice by microscopic observation of fresh feces and culturing of fecal material on various bacterial culture media. All mice were maintained in Trexler type isolators (one isolator per donor, two cages of two mice/isolator) and received the same standard diet (*ad libitum*, R03-40 SAFE sterilized by gamma irradiation at 45 kGy). Feces were collected per cage at days 1, 4, 11, 18, and 28 post-inoculation. At D28 post-inoculation, mice were euthanized. Total fecal DNA was extracted as previously described.⁵² Total bacterial counts and specific bacterial

profiles were assessed by real-time PCR (qPCR) of the bacterial 16 S rRNA gene using previously described primers and conditions for qPCR amplifications.^{30,53} Histological examination of the transverse colon was performed on 4- μ m sections stained with hematoxylin, eosin, and saffron, or immuno-labeled with rat monoclonal anti-Ki67 antibody (Dako, France). Thirty and sixty crypts per mouse were analyzed to determine the average crypt length and percentage of Ki67-positive cells, respectively. Slides were scanned using Panoramic Scan (3DHistech) and CaseViewer software (3DHistech) was used to count Ki67-positive cells and colon crypt depth. Experimentation was approved by the Regional Council of Ethics for animal experimentation C2EA-45 with authorization APAFIS#3441-2016010614307552 v1.

Evaluation of the inhibitory effect of FMT in a *Clostridioides difficile* infection mouse model

The CD model of infection was developed according to the procedure of Chen *et al.*³⁷ Six- to seven-week-old female C57BL/6 mice (Charles River, France) was housed in groups of 4 to 5 per cage and maintained under a 12-h light-dark cycle. The food (standard chow), water, bedding, and cages were autoclaved. Cage changes and daily assessment of the physical condition and behavior of the animals were performed under a laminar flow hood.

All animals were treated for 3 days with a mixture of antibiotics dissolved in drinking water consisting of kanamycin (0.4 mg/mL), gentamicin (0.035 mg/mL), colistin (850 U/mL), metronidazole (0.215 mg/mL), and vancomycin (0.045 mg/mL) (Sigma Aldrich, France). After 1 day of wash-out, mice received a single intraperitoneal injection of clindamycin (10 mg/kg) (Sigma Aldrich). The day after, mice were orally infected with 8.10^3 CFU of PBS-washed vegetative CD (strain R20291, exponential phase).

At D1 post-infection, the test groups received FMT by gavage (200 μ L) with either frozen and thawed (FMT-G10 and FMT-G80) or rehydrated FD (FMT-FD) microbiota that had been stored for 3 months (n = 10/stool/preservation mode; 2 stools randomly selected, i.e. #S1 and #S6). The positive-control group (CD+) was fed CD alone (n = 10) without any subsequent treatment and the negative control group

(CD-) was fed PBS ($n = 4$). All animals were observed daily from infection until sacrifice for symptoms and mortality, and weight was recorded (clinical scoring, Table 1). At D4 post-infection or before for animals judged to be in a moribund state (scoring > 3), animals were euthanized. Quantification of the CD load was performed on cecal contents on selective agar plates (chromID CDIF, Biomérieux, France). Colonic MPO was measured as a marker of colonic inflammation (Mouse Myeloperoxidase DuoSet kit, R&D Systems, USA). Histological evaluation of the cecum and colon was performed on 4 μm sections stained with hematoxylin-eosin, coded, randomized, and scored by two independent operators in a blind manner.³⁷ Briefly, grading considered the following features: i) edema of the mucosa and hemorrhagic congestion, ii) tissue infiltration by neutrophils, and iii) epithelial damage. A score of 0–3, denoting increasingly severe abnormality was assigned to each parameter and the overall score obtained by summing the component scores. Experimentation was approved by the Regional Council of Ethics for animal experimentation C2EA-34 with authorization APAFIS#7600-2016062016336853 v3.

16 S genomic analysis of microbiota following FMT

Total bacterial DNA was extracted from the fecal content of CD-infected mice taken at D4 post-infection using the DNeasy PowerSoil Kit (Qiagen France). The V3-V4 region of the 16 S rRNA gene was amplified for 30 cycles with an annealing temperature of 65°C using Taq polymerase MTP (Sigma Aldrich) and primers PCR1F_460 (CTTTCCCTACACGACGCTCTTCCGATCTACGGRAGGC-AGCAG) and PCR1R_460 (GGAGTTCAGACG TGTGCTCTTCCGATCTTACCAGGGTATCTAA-TCCT) and sequenced on Illumina MiSeq platform. The raw 16 S rRNA sequences were analyzed using the bioinformatics pipeline FROGS (Find Rapidly OTU with Galaxy Solution).⁵⁴ After sequence processing, affiliations were performed with BLAST using the silva132 16 S database. Data were filtered by retaining only those sequences that were present in at least three samples and contributed 0.005% to the microbial community and multi-affiliation was manually checked with leBiBi (Quick BioInformatic Phylogeny of Prokaryotes). The resulting OTU table

(478 OTUs) was used for subsequent statistical analysis using R software and phyloseq package.⁵⁵ The phylogenetic tree was constructed using Mafft and Fasttree. The samples were standardized to the same depth (7325 sequences). α Diversity was studied by calculating the number of observed OTUs and Shannon indices. β Diversity was studied using Bray–Curtis distance. Differences between groups were evaluated by principal-coordinate analysis (PCoA) and multivariate analysis of permutation variances using distance matrices (PERMANOVA).⁵⁶

Statistical analysis

The stability of microbiota following conservation procedures was analyzed using R.⁵⁷ For all paired data, mixed effect models⁵⁸ were fitted per case, as described below, followed by appropriate contrast calculations, to account for non-independence of the data and avoid increasing the degrees of freedom of error, which could lead to the spurious significance or increased variance, which could mask significant variations. Moreover, we used estimated marginal means⁵⁹ to correct for the unbalanced design due to missing values. The stability over time was assessed by fitting a mixed effect model to the microbiota data over time for each microbiota group. Time and freezing procedure were modeled as fixed effects while the stool sample was modeled as a random effect. Dunnett's contrasts were applied to adjust for multiple comparisons of each time measurement *versus* the first one (W1 or M3), used as a reference.

A mixed-effect model was fitted to the SCFA concentrations, using freezing condition and SCFA as fixed effects and stool sample as a random effect. Dunnett's contrasts were used to adjust for multiple comparisons of each freezing condition *versus* the fresh stool used as a reference.

A Mann Whitney test was used to evaluate the anti-CD growth activity of preserved fecal samples *versus* positive bacterial controls.

For *in vivo* experiment, a mixed effect model was fitted to % baseline weight and clinical score using time and freezing procedure as fixed effects and animal as a random effect. Contrasts were constructed in order to compare the score and body weight evolutions from D0 to D4 of treated *vs* untreated animals, adjusting for multiple

comparisons. Chi-2 test was used for survival analysis and Kruskal–Wallis (1-way ANOVA) followed by Mann Whitney tests were used for a charge of CD in ceecal content, MPO, and histological score.

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Specific author contributions

ChC, FJ, AJW, MT, and NK designed the study; JR was responsible for the study; JR, ChC, JD, CM, CC, AJW, AM, MM, TM, and JA collected data; IN performed statistical analysis; JR and NK drafted the manuscript; NK, AJW, and MT interpreted the data and critically revised the manuscript. All authors discussed the results, data analysis, and preparation of the manuscript. NK and MT should be considered as joint senior authors.


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