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The relationship between *Fusobacterium* species and other flora in mixed infection

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Summary. Mixed infections with three *Fusobacterium* species and seven other bacterial species were studied in a subcutaneous abscess model in mice. Fifteen *Fusobacterium* isolates (eight *F. nucleatum*, four *F. necrophorum*, and three *F. varium*) and one isolate each of *Bacteroides fragilis*, *B. asaccharolyticus*, *Staphylococcus aureus*, Group A β -haemolytic streptococcus, *Escherichia coli*, *Klebsiella pneumoniae* and *Pseudomonas aeruginosa* were studied. Electronmicrographs showed the presence of a thin mucopolysaccharide wall before and after inoculation into mice in 12 isolates which included all of 11 *Fusobacterium* isolates that induced subcutaneous abscesses. After co-inoculation of *Fusobacterium* isolates with other species and selective therapy with antimicrobial agents, *S. aureus* and *K. pneumoniae* were found to be of equal or greater importance in abscess induction than were *Fusobacterium* isolates, while *Fusobacterium* isolates were found to be more important than Group A streptococci and *E. coli*. Mutual enhancement of the numbers of organisms in mixed infections was observed with *Fusobacterium* spp. and *K. pneumoniae*, *P. aeruginosa* or *Bacteriodes* spp. Suppression of *Fusobacterium* spp. was noticed only when they were co-inoculated with Group A streptococci. The additive or synergistic capabilities of *Fusobacterium* species highlighted their potential pathogenicity in infection.

Introduction

Fusobacterium species are involved in various human infections where they are often isolated in mixed cultures with other anaerobic, facultative and aerobic bacteria (Brook, 1983). They are frequently involved in respiratory tract infections such as chronic sinusitis and otitis, peri-tonsillar abscesses, aspiration pneumonia and lung abscesses, and can also induce bacteraemia and intracranial infections. A few studies have investigated the synergistic potentials and importance of *Fusobacterium* spp. relative to other organisms in mixed infections (Altemeier, 1942; Conlon *et al.*, 1977; Hill *et al.*, 1974; Hite *et al.*, 1949). In this study we produced subcutaneous (SC) abscesses in mice to evaluate the synergistic potentials between *Fusobacterium* species and other organisms with which they are commonly found in mixed infections. We determined the relative importance of organisms by selective inhibition with antibiotics and by quantitative cultures of abscess contents.

Materials and methods

Organisms

All organisms were recent clinical isolates from patients admitted to the National Naval Medical Center, Bethesda, Maryland and were kept frozen in skimmed milk at -70°C . They included eight isolates of *Fusobacterium nucleatum*, four *F. necrophorum*, three *F. varium* and one isolate each of *Bacteroides fragilis*, *B. asaccharolyticus*, *Staphylococcus aureus*, Group A β haemolytic streptococcus *Escherichia coli*, *Klebsiella pneumoniae*, and *Pseudomonas aeruginosa*. The isolates were identified by conventional methods (Lennette *et al.*, 1980; Sutter *et al.*, 1980).

Capsular staining

The presence of capsules was established by Hiss's stain before and after inoculation in mice, and was confirmed by electronmicroscopy after staining with ruthenium red (Kasper, 1976).

Animals

Male Swiss Albino mice, 20–25 g, were obtained from the Naval Medical Research Institute mouse colony (NMRI/NIH-CV). The mice were raised in conventional conditions.

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Inoculum

Frozen bacterial suspensions were thawed at room temperature, subcultured on to chocolate or Schaedler-anaerobic blood agar (Difco Laboratoires, Detroit, MI), and incubated for 48 h at 37 °C in an anaerobic glove box for the anaerobic bacteria, or in air with CO₂ 5% for the aerobic bacteria. Twenty-four hours before injection, the bacterial isolates were inoculated on to Schaedler blood-agar plates with brain-heart infusion base (BHI, Difco). From these media, cotton swabs were used to pick colonies from the plates and transfer them to normal saline to give suspensions equivalent to a 10 MacFarland standard. Numbers of colony forming units (cfu) were determined by pour-plate counts in brain-heart infusion agar enriched with vitamin K₁ 10 µg/ml and haemin 5 µg/ml.

Abscess formation

Mice were given subcutaneous (sc) injections in the right medial aspects of their thighs of 0.1 ml of the appropriate bacterial suspensions in saline containing total counts of 10⁸ cfu of each organism. The bacterial counts were estimated by nephelometry with MacFarland standard and the required density obtained by dilution. The doses were chosen to achieve minimal mortality, and maximal abscess formation (Brook *et al.*, 1983). The size of the individual abscesses was determined during necropsy on the fifth day after inoculation. Although the volume of these abscesses could not be determined accurately, their sizes were compared by measuring two perpendicular diameters representing maximum length and width. Assuming that the abscess is an irregular prolate spheroid, the product of the length and width is proportional to the outer surface of the abscess. This product, expressed in mm², was arbitrarily selected as the comparative measure. By this criterion, after 5 days without antibiotic therapy the abscesses caused by a single organism achieved an outer-surface size of 80–200 mm², and abscesses caused by two organisms were 120–300 mm².

Examination of the abscesses

Animals were killed by cervical dislocation and the abscess material was removed aseptically. The site and histology of the abscesses were confirmed, in two mice of each experimental group, by hematoxylin and eosin staining. The number of cfu of each isolate in an abscess was determined individually. The abscesses were homogenised inside a glove box in 1.0 ml of sterile saline in a ground-glass tissue homogeniser. Ten-fold serial dilutions of the homogenates were made with sterile saline and 0.1 ml of each dilution was spread in triplicate on enriched brain-heart infusion (BHI) and blood-agar plates. Colonies were counted after incubation for 48 h in aerobic or anaerobic conditions at 37 °C. Characteristic colonies of all isolates were picked and identified by Gram's stain and biochemical tests (Lennette *et al.*, 1980;

Sutter *et al.*, 1980). The results were analysed by Student's *t* test.

Antimicrobial agents and therapy

Dosage of antimicrobial agents was based upon measurement of levels of drug in the serum of healthy mice after administration of selected doses. In most instances, this was approximately equal to the recommended doses for man (mg/kg).

The daily dosages of antimicrobial agents (administered in divided doses every 8 h) were: penicillin G (Squibb and Sons, Princeton, NJ) 100 mg/kg; metronidazole (Searle, Chicago, IL) 50 mg/kg; gentamicin (Schering, Kenilworth, NJ) 7.5 mg/kg. Treatment was initiated 2 h after inoculation and continued for 5 days. All agents were given by intramuscular injection.

Sensitivity tests

The minimal inhibitory concentrations (MIC) of each agent for all isolates were determined by the agar dilution technique (Thornsberry and Svenson, 1978).

Measurement of antimicrobial agents in serum and abscess fluid

Levels of antimicrobial agents in the sera and in the abscess contents were determined by the following methods: the agar diffusion assay (Lummis *et al.*, 1978) for penicillin with *Micrococcus luteus* ATCC 9341 (American Type Culture Collection, Rockville, MD), and for gentamicin with *Bacillus subtilis* ATCC 6051; high pressure chromatography (Wheeler *et al.*, 1978) for metronidazole. Antibiotic levels were determined in serum and abscess fluid at 0.5 and 8 h after the last injection on day 5 of treatment. These samples were kept frozen at -70 °C until they were assayed.

Induction of abscesses

The ability of an isolate to cause an abscess was determined by sc inoculation of 10⁸ cfu of each isolate alone in 0.1 ml of saline into groups of six mice. Abscesses were also induced by injection of mixtures containing 10⁸ cfu of each isolate in 0.2 ml of saline into groups of six mice. The animals were observed for 28 days, and the abscess sizes were determined by external measurement without killing the animals. This experiment was repeated three times.

Relative importance of isolates

Abscesses were produced by inoculating a single isolate of *Fusobacterium* alone, a single isolate of the aerobic or facultative species alone, or a mixture of both into groups of 40 mice each (table 1). Each experiment was repeated three times. Two hours after inoculation, each group was divided into four sub-groups of 10 mice, receiving therapy by antimicrobial agents directed at (a) *Fusobacterium*

Table 1. Abscess surface size, 5 days after inoculation, in mice infected with *Fusobacterium* spp., alone or with other bacteria, and treated with various antibiotics

<i>Fusobacterium</i> sp. and species co-inoculated	Abscess size* after antimicrobial treatment against				Species of greater significance†
	neither	<i>Fusobacterium</i> sp.	other species	both species	
<i>F. nucleatum</i> (FN)					
None	124 ± 20	35 ± 19	132 ± 37	28 ± 20	
<i>S. aureus</i> (SA)	245 ± 67	134 ± 43	12 ± 8	18 ± 9	SA
Group A streptococcus (GAS)	232 ± 55	65 ± 35	285 ± 52	49 ± 18	FN
<i>E. coli</i> (EC)	246 ± 39	18 ± 10	170 ± 41	24 ± 21	FN
<i>K. pneumoniae</i> (KP)	208 ± 48	112 ± 35	26 ± 21	21 ± 12	KP
<i>P. aeruginosa</i> (PA)	216 ± 72	43 ± 12	27 ± 19	16 ± 8	Equal
<i>F. necrophorum</i> (F.Ne)					
None	156 ± 18	40 ± 16	145 ± 54	25 ± 16	
<i>S. aureus</i> (SA)	192 ± 71	215 ± 34	32 ± 16	46 ± 19	SA
Group A streptococcus (GAS)	278 ± 81	30 ± 24	160 ± 51	38 ± 21	FNe
<i>E. coli</i> (EC)	306 ± 64	78 ± 36	205 ± 37	61 ± 14	FNe
<i>K. pneumoniae</i> (PA)	224 ± 75	58 ± 19	44 ± 27	21 ± 25	Equal
<i>P. aeruginosa</i>	180 ± 50	72 ± 36	58 ± 30	37 ± 14	Equal
<i>F. varium</i> (FV)					
None	192 ± 39	53 ± 28	205 ± 42	41 ± 18	
<i>S. aureus</i>	262 ± 43	188 ± 27	39 ± 18	45 ± 21	SA
Group A streptococcus	257 ± 51	54 ± 24	191 ± 39	59 ± 37	FV
<i>E. coli</i>	285 ± 64	31 ± 26	129 ± 59	50 ± 22	FV
<i>K. pneumoniae</i>	274 ± 51	154 ± 40	24 ± 28	19 ± 25	KP
<i>P. aeruginosa</i>	241 ± 60	163 ± 44	41 ± 36	47 ± 15	PA

* Mean ± standard deviation (mm²)† The organism of greater importance was that which caused abscesses that were significantly smaller ($p < 0.05$). When no statistical difference was noted, the two organisms were considered to be equally important.

alone, (b) other bacteria alone, or (c) combined therapy against both *Fusobacterium* and the other bacteria; (d) control mice received no treatment. For the therapy of mixed infections with fusobacteria and aerobic or facultative Gram-negative aerobic bacilli, penicillin and gentamicin were used, penicillin therapy for the *Fusobacterium* spp. and gentamicin for the aerobic or facultative Gram-negative bacilli. For the therapy of mixed infections between fusobacteria and facultative Gram-positive cocci, gentamicin and metronidazole were used, gentamicin for the cocci and metronidazole for the fusobacteria. The effect of therapy on the combination of *Fusobacterium* spp. and *Bacteroides* spp. could not be tested because of the lack of antimicrobial agents that will selectively inhibit one of them only. Animals infected with two isolates were used to evaluate the relative importance of the isolates, and those infected with single agents were used to ascertain the in-vivo activity of antimicrobial agents.

The relative importance of the organisms causing the abscess was determined by comparing the abscess sizes in the animals treated with one or two antimicrobial agents with those in untreated control mice. For example, an abscess caused by a single organism, which was treated with an antimicrobial agent effective against that organism, was predicted to respond to therapy and be smaller

than that in an untreated control animal. For an abscess caused by two organisms where therapy was directed against only one, the abscess was predicted to be smaller than in untreated control animals when the susceptible bacterial species was the greater contributor to the abscess. Thus, in mixed infections of organisms A and B, therapy directed only against bacterium A could cause (a) no decrease in the abscess size (organism B is more important in mixed infections than organism A), or (b) a larger decrease in abscess size than when organism B was treated (A is more important than B). The statistical significance of the relative importance of the various micro-organisms was estimated by Student's *t* test, comparing the sizes of abscesses caused by both organisms on the fifth day of therapy. The sizes of the abscesses in the group of mice treated with agents effective against the aerobic or facultative bacteria alone, were compared with the sizes of the abscesses in the group where the therapy was directed against the anaerobic bacteria alone. The organisms of greater significance were those with which the reduction in abscess size was significantly greater ($p < 0.05$) with the specific therapy. When no statistical difference was noted, the two organisms were considered to be equally significant. The subgroup infected with mixed flora and treated with two antimicrobial agents was included to observe any synergy

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between the drugs. Synergy was defined by statistically significant reduction ($p < 0.05$) in abscess size associated with addition of a second antimicrobial agent.

Quantitative relationships between isolates

Abscesses were induced by one encapsulated representative of each *Fusobacterium* species alone, or in combination with one isolate each of the other facultative and aerobic species. The abscesses were cultured and bacterial counts were performed on the fifth day after inoculation. Six mice were included in each experimental group and each experiment was repeated three times. The data were analysed by comparing the changes in each of the bacterial isolates present in the abscess with the numbers present in the control abscesses containing the single species (Student's *t* test).

Results

Encapsulation

Electronmicrographs of thin sections of the *Fusobacterium* isolates before inoculation showed a thin dense capsule of mucopolysaccharide (fig. 1) in

seven of the eight *F. nucleatum* isolates, two of the four *F. necrophorum* and all three *F. varium*. The capsule was seen in $> 50\%$ of the organisms seen among 1000 cells on a grid. After animal passage the density and size of the capsule layer were increased in all but two of the isolates used in the study (both *F. nucleatum*) (fig. 2). None of the unencapsulated strains acquired a capsule. Capsules were observed in smears stained by Hiss's method of all of the other isolates except *P. aeruginosa* before and after animal passages.

Abscesses induced by single organisms

Examination of abscesses caused by *Fusobacterium* spp. and other bacteria revealed the presence of a fibrous encapsulated collection of material which contained bacteria and polymorphonuclear leukocytes. Subcutaneous abscesses were found in at least 90% of the mice given 11 of the 15 *Fusobacterium* isolates. Isolates that induced abscesses included seven of the eight *F. nucleatum*, two of the four *F. necrophorum* and two of the three *F. varium*. Electronmicrographs of these isolates showed the



Fig. 1. Electronmicrograph of thin section of 48-h culture of *F. nucleatum* before animal passage; arrow indicates the thin capsule. (Bar = 0.5 μm) ($\times 78\,000$).



Fig. 2. Electronmicrograph of thin section of 48-h culture of *F. nucleatum* (same strain as fig. 1) after animal passage; arrow indicates the thicker capsule (Bar = 0.5 μm) ($\times 78\,000$).

presence of a mucopolysaccharide layer. Three of the four isolates that did not induce an abscess did not possess a capsular layer. Abscesses were formed within 24–48 h after sc inoculation with the fusobacteria. The abscesses reached a maximum diameter of 12–18 mm within 7 days, and most drained spontaneously or were absorbed within 14–21 days. No mice died after inoculation.

Abscesses caused by *Bacteroides* spp. were formed within 24–48 h in 85% of the mice, and the abscesses reached a diameter of 12–18 mm within 4–7 days. Up to 90% of the abscesses drained spontaneously within 9–12 days. No deaths occurred after inoculation.

S. aureus, Group A streptococcus, *K. Pneumoniae*, *P. aeruginosa* and *E. coli* isolates induced abscesses in at least 90% of mice, and the abscesses reached a diameter of 12–16 mm within 5–7 days. A mortality rate of about 10% was observed within the first 48 h after inoculation with all of these organisms except the Group-A streptococcus.

Abscesses induced by mixtures of Fusobacterium isolates and other organisms

After inoculation, abscesses developed in 82% of the animals given any of the combinations of one of the 15 *Fusobacterium* spp. and one of the other bacterial species within 48 h. Abscesses reached a maximum diameter of about 22–26 mm within 5–7 days and, if untreated, drained spontaneously within 14–21 days. When animals died, mortality was similar to that found in animals with abscesses caused by the aerobic or facultative Gram-negative rods or *S. aureus*.

In-vitro sensitivity to antibiotics

The *Fusobacterium* isolates were uniformly susceptible to penicillin (MIC ≤ 0.5 µg/ml) and metronidazole (MIC ≤ 0.1 µg/ml) and resistant to gentamicin (MIC > 128 µg/ml). The aerobic or facultative gram-negative rods and facultative cocci were resistant to penicillin (MIC ≥ 64 µg/ml) and metronidazole (MIC ≥ 128 µg/ml) and susceptible to gentamicin (MIC < 0.5 µg/ml).

Antibiotic levels in serum and abscess fluid

Antibiotic levels were determined on the fifth day of therapy. The mean ($\pm$ SD) serum level of gentamicin determined in a group of 10 mice with abscesses caused by *F. nucleatum* was 6.5 ± 2.2 µg/ml 30 min after last dose, and 1.8 ± 0.6 µg/ml 8 h later. The level in abscesses determined 30 min after

the last dose was 5.6 ± 2.2 µg/ml. The mean levels of penicillin in serum were 27.5 ± 8.2 µg/ml after 30 min, 6.8 ± 2.4 µg/ml after 8 h, and in abscesses 18.2 ± 6.2 µg/ml after 30 min. The mean level of metronidazole in serum was 25.4 ± 5.3 µg/ml at 30 min, 10.8 ± 3.2 µg/ml at 8 h and in abscesses 9.2 ± 4.6 µg/ml at 30 min. It is obvious that sufficient levels were achieved in both locations to inhibit susceptible strains.

Importance of Fusobacterium spp. relative to facultative and aerobic bacteria

The results obtained with combinations of *Fusobacterium* species and aerobic organisms and the assessment of species of greater significance are shown in table I. Abscesses caused by a single species always responded to antibiotic therapy directed at that species. These data were not included in table I. After appropriate therapy, abscesses were smaller but did not disappear within 5 days. Abscesses inappropriately treated increased in size, as did untreated abscesses. No synergy between the antimicrobial agents was noticed. In mixtures of species, *Fusobacterium* spp. were less important than *S. aureus*, equally or less important than *K. pneumoniae* and *P. aeruginosa*, and more important than Group A streptococci and *E. coli* in abscess formation.

Quantitative relationships between Fusobacterium spp. and other bacteria in mixed infections

A summary of the changes that occurred in the numbers of *Fusobacterium* spp. and of the other species in an abscess produced by the mixture is presented in table II. The calculation was done by deducting the average log 10 cfu of an organism when present mixed with another bacteria (see footnote, table II). An example of the calculation illustrates the relationship between *S. aureus* and *F. nucleatum*: the average log 10 cfu in abscesses caused by single isolates were *S. aureus* 9.2 and *F. nucleatum* 7.5; the average log cfu in abscesses caused by two isolates were *S. aureus* 9.6 and *F. nucleatum* 10.9; Ratio of change = $9.6 - 9.2; 10.9 - 7.5 = 0.4/2.4$.

Of the 21 combinations, numbers of fusobacteria were enhanced in 13 and inhibited in three (all with the Group A streptococcus). Numbers of other bacteria were enhanced in 12 of 21 combinations, and were never suppressed (table II). Numbers of *Fusobacterium* spp. were enhanced by *S. aureus* and *K. pneumoniae*. In most instances a concomitant increase occurred in both components of the mixed

Table II. Quantitative relationship between strains in mixed infections

Species co-inoculated	Ratio between change in numbers of <i>Fusobacterium</i> spp. and other bacteria in an abscess*		
	<i>F. nucleatum</i>	<i>F. necrophorum</i>	<i>F. varium</i>
<i>S. aureus</i>	0.4/2.4†	0.8/2.8†	1.2/3.4†
Group A. streptococci	0.6/-2.6†	1.0/-3.5†	1.2/-1.8†
<i>E. coli</i>	0.6/3.8†	1.2/1.6	2.3†/2.8†
<i>K. pneumoniae</i>	3.4†/2.2†	3.8†/2.0†	2.3†/1.9†
<i>P. aeruginosa</i>	2.6†/1.2	2.8†/2.4†	1.4/2.6†
<i>B. fragilis</i>	2.6†/3.2†	3.5†/2.4†	2.6†/1.2
<i>R. asaccharolyticus</i>	3.1†/2.2†	3.4†/1.2	2.0†/1.3
Increase/decrease of <i>Fusobacterium</i>	5/1	4/1	4/1

* Change (vs. control) in average number of other organisms (log 10 cfu)/change (vs. control) in average number of *Fusobacterium* sp. (control = the number of organisms in an abscess caused by the organism alone).

† Significant differences between single and mixed infection ($p < 0.05$).

infection. The combinations of *F. nucleatum* with other flora significantly increased the growth of the fusobacteria in five of the seven combinations and significantly decreased it in one (table II). Other bacteria were enhanced when mixed with *F. nucleatum* in four instances. Numbers of *F. necrophorum* in abscesses were increased in four combinations with other bacteria and decreased in one. Other bacteria were increased when combined with *F. necrophorum* in four instances. The numbers of *F. varium* in abscesses were increased in four combinations and decreased in one. Other bacteria were enhanced when combined with *F. varium* in four instances.

Discussion

Development of suitable animal models of infection are necessary to understand pathogenicity and synergy between species and to evaluate potential therapeutic agents for mixed infection. An important role of bacterial synergy in infection has been suggested by several investigators, and studies of synergistic activity have included the association of anaerobic bacteria with other species in various infections (Altmeier, 1942; Hite *et al.*, 1949; Macdonald *et al.*, 1956; Mergenhagen *et al.*, 1958; Brook and Walker, 1983). Data from clinical specimens indicate that many infections in man are caused by mixed anaerobic or anaerobic and facultative species.

The clinical significance of the Bacteroidaceae, particularly *F. necrophorum*, as opportunistic pathogens in various diseases in animals is also well

documented. Over the years a continuing effort has been made to understand the pathogenicity and importance of this organism in veterinary infections (Conlon *et al.*, 1977).

Several animal models have been used to investigate the pathogenicity of *Fusobacterium* species. The only successful animal models developed with *Fusobacterium* species have been intra-hepatic abscesses in rabbits (Abe *et al.*, 1976) and in mice (Hill *et al.*, 1974) and lymphatic abscesses (Conlon *et al.*, 1977). However, induction, monitoring and culturing such abscesses is complicated and cumbersome. The development, therefore, of subcutaneous abscesses in mice, as described in the present study, establishes a simple animal model that can be used for studies of pathogenicity (Brook and Walker, 1983) and for evaluation of the effect of antimicrobial therapy on single or mixed infections.

Several recent studies have demonstrated the pathogenicity of encapsulated anaerobes and their ability to induce abscesses alone. Onderdonk *et al.* (1977) correlated the virulence of *B. fragilis* strains with the presence of a capsule, and Simon *et al.* (1982) described decreased phagocytosis of the encapsulated *B. fragilis*. Capsular material from *B. melaninogenicus* also inhibits phagocytosis and phagocytic killing of other micro-organisms in an in-vitro system (Okuda and Takazoe, 1973).

We recently studied the relative importance of *Bacteroides* spp. and anaerobic Gram-positive cocci and their ability to cause an abscess in an animal model by virtue of capsule formation (Brook *et al.*, 1983; Brook and Walker, 1983; Brook and Walker, 1984). With few exceptions, possession of a capsule

generally made these organisms more important clinically than their aerobic counterparts. Although unencapsulated organisms did not induce abscesses, many of the strains which had only minimal numbers of encapsulated organisms (<1% of the total population), survived in the abscess after inoculation with other aerobic and anaerobic bacteria and became heavily encapsulated. These heavily encapsulated strains were thereafter able to induce abscesses when injected alone.

As was found with other anaerobic bacteria, the ability of *Fusobacterium* strains to induce subcutaneous abscesses could be correlated with the presence of the thin mucopolysaccharide layer in the cell wall. The production of subcutaneous abscesses could be an indication of the organism's virulence, as has been noticed in other anaerobic strains (Kasper, 1976). In contrast to findings with other anaerobic species such as *Bacteroides* and Gram-positive cocci (Brook and Walker, 1983; Brook *et al.*, 1983; Brook and Walker, 1984), we did not observe minimally encapsulated fusobacterial organisms before animal inoculation. Review of a larger group of fusobacterial isolates may reveal such a phenomenon. However, we did observe an increase in the cell-wall density in fusobacterial cells after passage through mice.

The limitation of our animal model makes it difficult to ascertain the exact role of *Fusobacterium* spp. in mixed infections. We could determine which of the two isolates present in the mixed infection was the greater contributor to the infectious process only in a few isolates and this indirectly by using selective antimicrobial therapy. Furthermore, because many clinical infections involve more than two pathogens, our model may represent a simpli-

fied version of a true infection. The concomitant increase in the numbers of cfu of the two isolates present in the abscess could be due either to a mutual additive effect or is true synergy. However, the increase in abscess size and in numbers of cfu of both isolates suggests a mutual beneficial relationship.

Our study demonstrated the relationship of *Fusobacterium* species with other micro-organisms in mixed infections. The relationship between the bacteria in mixed infections varies with the different combinations. *S. aureus* and *K. pneumoniae* tend to be of greater or equal importance to the *Fusobacterium* species. In contrast, *Fusobacterium* species were of greater importance than the Group A streptococcus and *E. coli*. However, the co-inoculation without suppression by antimicrobial agents produced mutual enhancement of growth when *Fusobacterium* isolates were mixed with *K. pneumoniae*, *P. aeruginosa* or *Bacteroides* spp., and inhibition of growth when mixed with Group A streptococcus.

The additive or synergistic potential between *Fusobacterium* spp. and other bacteria, commonly cultivated in mixed cultures from infected sites, demonstrates the pathogenic potentials of fusobacteria. The synergy between these different bacterial strains may be due to protection from phagocytosis and intracellular killing (Simon *et al.*, 1982), production of essential growth factors (Lev *et al.*, 1971) or lowering of oxidation-reduction potential in host tissues (Mergenhagen *et al.*, 1958).

The opinions and assertions contained herein are those of the authors and are not to be construed as official or reflecting the views of the Navy Department of the naval service at large.

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