

Fearless International (FRLE) \$0.19

Fearless International Inc., a luxury performance boat manufacturer, has been the focus of the media for the last several months in magazine such as GQ, Time, Bloomberg Markets, Maxim, and over 20 others.

According to TIME, "When a company bold enough to call itself Fearless Yachts splashed onto the luxury-boat market, it drew considerable attention. Collaborating with Porsche Design Studio/Austria on a series of high-style, high-performance yachts, the brand unveiled its first model, the Fearless 28, in February." Since its release, Fearless Yachts has taken orders for more than 33 Yachts bringing more than \$10 Million in sales and put the production facility at 75% capacity.

Top 5 Reasons To Consider Fearless Yachts:

- 1: Already \$10,000,000 In Sales Since First 7 Months.
- 2: First of a 5 yacht series had huge response from the market.
- 3: Next yacht designs have been released and Debut is in Miami in February
- 4: Company set to begin international marketing.

They're serving industry rather than the public.

The FDA Exposed: An Interview With Dr.

Regarding the determination of drug effectiveness, I think the FDA does a pretty good job.

There is nothing lifesaving there.

MANETTE: Do you think we should have any confidence in the FDA and if so, can you elaborate on what they do that you feel benefits the American people?

There is nothing lifesaving there.

MANETTE: And what about those new drugs?

Clearly Vioxx was the problem.

First, I would separate safety and post-marketing from the pre-marketing.

Rarely will they keep a drug from being marketed or pull a drug off the market.

The people who approve a drug when they see that there is a safety problem with it are very reluctant to do anything about it because it will reflect badly on them.

I have never seen him happier or healthier.

The FDA has never looked at benefit.

"That's all it ever is with me," Wagner said.

To preempt that, he offers me this job, which basically would have been exile to a fancy title with no real ability to have an impact.

If the benefits aren't there we shouldn't be having discussions about labeling the drug.

We see wonderful ads of people demonstrating their health, whether they're skating across the ice or doing their Tai chi.

A very small proportion of drugs represent a new drug that hasn't been marketed before.

What we want in the end are drugs that actually have better benefit.

I would recommend several changes.

The right thing to do would have been to pull the high dose off the market because there is no benefit for short-term relief of acute pain that exceeds this risk.

Americans and Congress should be screaming bloody murder.

This is the first time that my job was actually in jeopardy and where the FDA actually intended to fire me.

MANETTE: How do you cope with that going to work each day?

They need to have larger clinical trials.

They can have some confidence because it turns out that most drugs are remarkably safe.

The assumption is that the drug will be no different than the sugar pill.

You could eat off the ground on that property.

They're hoping that I'll just become very frustrated and disillusioned and leave or that I'll slip up in some way so that they can take some sort of action against me.

Unfortunately, that is the way the FDA is currently structured.

There is nothing new.

The culture also views industry as the client.

Sunday marked the first time he held the lead entering the final round of the tournament.

I knew that was not a sustainable argument.

I've been very fortunate.

Basically, I feel like I'm in the Gulag.

"Attacking the health and welfare of those animals is a direct attack on me, and my credentials are impeccable," said Dinnes.

This guy has told the truth.

Drug safety is about five percent.

Graham, it's truly a pleasure to have the opportunity to interview you.

I've been very fortunate.

This was a conspiracy and it was coordinated and there was collaboration among senior level FDA officials.

Most of them aren't breakthroughs and most of them aren't lifesaving, but they get treated as if they were.

Clearly Vioxx was the problem.

Most often, they'll compare the drug against something called a placebo or a sugar pill.

I've been there with him before, through times when I failed to get a check on time, but they never, ever stuck me, and I was assured they never would again.

It was written, performed, and paid for by industry.

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MANETTE: All of these attacks backfired on them.

"I think I just kind of changed my attitude toward the game," he said.

This is an inherent conflict of interest.

Graham, it's truly a pleasure to have the opportunity to interview you.

It's not a vendetta against any particular drug company.

There is nothing innovative.

The Doctor also responded to an investigation into the welfare of animals at the ranch by the U.

GRAHAM: In a perfect world, I think the FDA would need to be restructured.

On the pre-marketing side, the FDA needs to pay greater attention to safety.

I'm looked up to by the scientific staff because of that expertise.

GRAHAM: As currently configured, the FDA is not able to adequately protect the American public.

It lowers your blood pressure and they have all these glitzy ads, direct-to-consumer advertising.

Haas, who was in second after three rounds, also had his run at the championship end on a double bogey.

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Actually, Senator Grassley and Dodd have recently introduced legislation to create an independent center for post-marketing safety that would serve to protect the American people from unsafe drugs.

They know that they're vulnerable.

He just went directly to the Lancet.

I've had a very different experience.

This intimidation took several forms.

I can talk in my private capacity, but I can't talk about material that would be confidential.

I can't afford to make any mistakes.

All they focus on is efficacy.

MANETTE: How do you cope with that going to work each day?

The culture also views industry as the client.

Tom Devine, as he said publicly, was very interested in doing the right thing.

When a serious safety issue arises at post marketing, the immediate reaction is almost always one of denial, rejection and heat.

These benefit analyses should be done as a matter of routine.

I've been pretty fortunate in that way.

It's a misperception that our lawmakers in Congress have as well and they've been fed this line by industry.

Does the drug work or not?

They're really proud of the fact that I've said it and they're not afraid to be seen with me.

They flunked every test and I passed every test.

"Michael Jackson is pleased to announce that Jay-Z, Mariah Carey, Missy Elliott, R.

The FDA doesn't really care about that.

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This interview is reprinted here with permission from Dr.

Let me begin by asking you how long you've been with the FDA and what your current position is?

The drug might lower your blood pressure by just a few millimeters of mercury, but the FDA will say we can approve it because it does lower your blood pressure.

A very small proportion of drugs represent a new drug that hasn't been marketed before.

What they've done is call me names.

Yes, drugs cause a lot of harm.

He said confidence has helped him turn his game around.

I'm just a scientist doing my job and I have to leave the rest to God to protect me.

GRAHAM: It's a myth that is promulgated not only by industry but also by the FDA itself.

Does the drug work or not?

Industry is our client.

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It's a misperception that our lawmakers in Congress have as well and they've been fed this line by industry.

The quarter-acre site, which holds two villas and a plot of land, is rumoured to be his new home according to contactmusic.

This is no outside critic, either: these are the words from a top FDA employee who has worked at the agency for two decades.

That story is entirely false.

You have to go through the chain of command and if somebody up above you says that they want things done in a particular way well, they want it done in a particular way.

This was a conspiracy and it was coordinated and there was collaboration among senior level FDA officials.

Campbell first took the lead Sunday with back-to-back birdies on Nos.

The blood thinner Coumadin is another example.

"The veterinarian wanted to clarify that the lawsuit was nothing personal against Michael, and that he's glad that it has been settled.

A very small proportion of drugs represent a new drug that hasn't been marketed before.

I've helped to get ten different drugs off the market, and they're from ten different drug companies.

For the American people, it's dropped off the radar screen.

I've had a very different experience.

Regarding the determination of drug effectiveness, I think the FDA does a pretty good job.

A Californian Judge allowed the questions to be asked in London at attorney's requests.

Haas, who was in second after three rounds, also had his run at the championship end on a double bogey.

"Attacking the health and welfare of those animals is a direct attack on me, and my credentials are impeccable," said Dinnes.

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He just went directly to the Lancet.

To preempt that, he offers me this job, which basically would have been exile to a fancy title with no real ability to have an impact.

Did you see the disaster coming?

Today, Michael's publicist, Raymone Bain, has issued a statement on his behalf.

We settled it," said attorney Brent Ayscough, who represented Michael in this case.

Campbell was among six golfers who held or were tied for the lead Sunday at Annandale Golf Club.

This is the first time that my job was actually in jeopardy and where the FDA actually intended to fire me.

For most of these drugs it's more belief.

I've been there with him before, through times when I failed to get a check on time, but they never, ever stuck me, and I was assured they never would again.

But it's the same drug and we already know about the safety of the drug.

I'm just a scientist doing my job and I have to leave the rest to God to protect me.

The incentive is to review and approve the drugs as quickly as possible, and not stand in the way of profit-making.

But I've been a target of retaliation in the past.

The drug might lower your blood pressure by just a few millimeters of mercury, but the FDA will say we can approve it because it does lower your blood pressure.

MANETTE: And what about those new drugs?

We see wonderful ads of people demonstrating their health, whether they're skating across the ice or doing their Tai chi.

If you are the system administrator, please click here.

Lots of patients and lots of doctors will use that medication.

They need to have larger clinical trials.

It was with his business people.

Campbell first took the lead Sunday with back-to-back birdies on Nos.

Sunday marked the first time he held the lead entering the final round of the tournament.

Kelly, Wyclef Jean, Lenny Kravitz, Lauren Hill, Mary J.

Most often, they'll compare the drug against something called a placebo or a sugar pill.

The times when I've done the benefit analysis, I've been chastised, criticized and suppressed by the FDA.

They should be screaming because this can happen again.

Branshaw was done in by a double bogey.

The tournament wasn't decided until the final two holes.

Basically, I feel like I'm in the Gulag.

- Chad Campbell righted his listing game Sunday with a one-stroke victory over Johnson Wagner in the Viking Classic, making two birdies on his last three holes to win.

MANETTE: When did that go into effect?

"I finally started believing that I deserve to be out here and I'm playing like it, so it's nice.

Your Source for cycling tour.

Tell us a little bit about that.

For example, we know that drugs for diabetes can lower your blood sugar.

We see wonderful ads of people demonstrating their health, whether they're skating across the ice or doing their Tai chi.

The FDA cooperates with that mandate.

Ask the FDA to produce its benefit analysis that shows that the benefits exceed the risks.

com AP Photo: Chad Campbell holds the trophy after winning the Viking Classic in Madison, Miss.

Campbell was among six golfers who held or were tied for the lead Sunday at Annandale Golf Club.

Clearly Vioxx was the problem.

I would create a separate center for product safety.

They flunked every test and I passed every test.

GRAHAM: As currently configured, the FDA is not able to adequately protect the American public.

Can you explain what you meant by that?

However Ayscough claimed Michael never paid the full bill as some items were "not what they were represented to be.

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It works on what's called efficacy.

The FDA assumes the drug is safe and now it's up to the company to prove that the drug isn't safe.

"We resolved the lawsuit over the weekend.

These people were posing as whistleblowers themselves ratting on another whistleblower.

The times when I've done the benefit analysis, I've been chastised, criticized and suppressed by the FDA.

Parts of this interview also appear in Dr.

GRAHAM: It's a myth that is promulgated not only by industry but also by the FDA itself.

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I'm just a scientist doing my job and I have to leave the rest to God to protect me.

Ask the FDA to produce its benefit analysis that shows that the benefits exceed the risks.

Industry is saying there are all these lifesaving drugs that the FDA is slow to approve and people are dying in the streets because of it.

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Campbell was among six golfers who held or were tied for the lead Sunday at Annandale Golf Club.

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Regarding the determination of drug effectiveness, I think the FDA does a pretty good job.

It's a misperception that our lawmakers in Congress have as well and they've been fed this line by industry.

I've helped to get ten different drugs off the market, and they're from ten different drug companies.

First, I would separate safety and post-marketing from the pre-marketing.

Obviously he had been tipped off by people in the Senate Finance Committee who are sympathetic to the FDA's status quo that I was going to be called as a witness.

It might also call into question why they approved the drug in the first place.

This high-level FDA official never talked to me about this allegation.

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MANETTE: How do you cope with that going to work each day?

"We resolved the lawsuit over the weekend.

The clinical trials that are done are too small, and as a result it's very unusual to find a serious safety problem in these clinical trials.

This was a conspiracy and it was coordinated and there was collaboration among senior level FDA officials.

There is nothing new.

They're not afraid to work with me.

Whoever these individuals are who are intent on disseminating false information throughout the media regarding Mr.