

Winder: Well, good morning everyone. Or afternoon or evening, depending on who and what and where and why you are. This is Winder Lyons, and I have Lindsay Glover on the line with us. And I've got to say, Lindsay, first of all, welcome to the call.

Lindsay: Hi. It's good to be here.

Winder: And I want to get you to make sure you speak up loudly enough so everybody can hear you. And I got to say, I have really, really been looking forward to this. This is going to be one of the most thrilling calls I think I've ever been a part of. Lindsay has this amazingly engaging story about what's happened with the Anovite Colostrum products, and her three-year-old daughter. So first of all what is your daughter's name?

Lindsay: My daughter's name is Ruarie Paula Glover. And she's got kind of a unique spelling, too. She's named after her grandma that recently passed away about a year ago from cancer. That's where the Paula comes from. And the Ruarie means daughter of the Red King. And her father's name is Ray, which means kingly. So it kind of all fits together.

Winder: Well, that's really sweet. Well, tell us a little bit about her story. What happened with your daughter?

Lindsay: Okay, so August of 2015, she started her dance classes at two years old, and she was doing well at first, and then she would come home and she'd be more tired than normal. We assumed it was because she was in dance classes, so she was doing more than she was used to. But it progressively got worse. She had bruising, she had swelling, she had fevers, and this went on for months and months. And in December, we had to finally make the decision to remove her from dance because she couldn't keep up anymore. She was falling down, and none of the doctors could understand what was wrong with her. They said, "Oh, it's cold and flu season. She's probably just sick."

And in January, I had had enough of being pushed around by the doctors locally, so we sought out doctors elsewhere in bigger areas to try to get some answers. She saw a new pediatrician, they ordered up some labs, they looked over the labs initially, and couldn't find anything because something was missed. So we called the head of pediatrics in March, and asked the head to look at her labs again. And the head of pediatrics took one look at them and said, "I don't know how this was missed, but your daughter has arthritis. She has rheumatoid arthritis. All the markers are there. I'm so sorry." Needless to say, we were devastated. To be told that your daughter is going to have, or has arthritis, and it's potentially facing all kinds of challenges that come with it, it was devastating.

By this point, it had progressed. She spent most days sleeping all day, laying around on the couch with her legs propped up, swollen as can be. She had

grapefruits for kneecaps, and she had lost tremendous amounts of weight, her legs were just covered in bruises constantly. On top of that, she developed celiac disease, and they don't know if it was an aftereffect of the rheumatoid arthritis, or if it's what caused the rheumatoid arthritis. But she got to where she was itching all the time. When she'd eat gluten, her neck would swell, so we met with the rheumatology team in May, and they told us, "You know we have to watch things. We have to give her medicine all the time to keep the swelling down, to keep her somewhat normal and keep her fevers under 103," which is easier said than done with somebody with rheumatoid arthritis.

She was facing some pretty horrific medication. So shortly thereafter, we had a plan to go back to rheumatology in October, and I started researching. I wanted to do everything I could to avoid these nasty medications that they were going to start her on in October.

Winder: Let me ask you a question. At that point, did they have her on any sort of meds at all?

Lindsay: The only thing that they had her on, because they needed to get specific blood tests done, was Motrin as needed. And so it was to keep her swelling down, and she was on it daily, multiple times a day. It was every morning, every afternoon, and every evening she had to have a full dose.

Winder: That's not without side effects, right?

Lindsay: Absolutely. Motrin is very, very hard on the stomach. It would make her nauseous. It's definitely not good for the kidneys. And in our family, we have a history of kidney disease. And my son was born with only one kidney, so kidneys are kind of a sacred thing in this family. So things that are filtered out directly through the kidneys, it was scary for me to be medicating this child constantly. So they didn't want her on any of the medications just yet because they wanted to have clear cut, "This is exactly what type of arthritis it is," because she was throwing all kinds of weird markers. They didn't know exactly which form of arthritis it was. She had some rare stuff going on, she had some unexplainable stuff going on.

So they didn't want any tampering with the blood work. And so they wanted to wait the six months to start her on medication, so that we had all those tests done, all the labs, all the scans, everything, so they knew exactly what they were doing. So knowing that we were facing these potential medications, one of which they were saying was Biologics, and if anybody has ever researched them, they are horrendous. I started doing as much research as I could to try to control this and help her heal. Because I know that arthritis can go into remission. It's unlikely, but it can. So I started eliminating foods. We eliminated gluten because the gluten triggered her flairs.

One evening I happened to stumble across a post on Facebook about celiac disease, and one of the independent consultants from Anovite had commented about the amazing healing effects of Colostrum. And as soon as I read it, something in me in my head and in my heart clearly said, “This is what it is. This is what I need to do.” So I messaged her, and I said, “I need some information because this is what I’m facing.” Immediately she set up a conference call with Dr. K, and he shared lots of information with me. He emailed me a lot of information. The next day I read through lots of information, and I don’t know that I’ll be able to read through all of it because there’s so much out there.

We got Ruarie on the sample pack, and within the first week – so we started her on the Colostrum in about May, the second week of May, just two weeks before her third birthday. In the first week, she wasn’t in as much pain, her temperatures had dropped to low fevers, her bruising had seemed to lighten a little bit, and her swelling had seemed to lighten a little bit. By the end of week two, she had no fevers. By the end of the first month, she was herself again. She was running around, she was gaining weight, her swelling was almost completely gone. As of probably, about two weeks ago, so mid-August, beginning of August, she had no swelling, completely normal body temperatures, she has no bruising. It’s phenomenal.

I spoke with her doctors this week, August 25th, was when I spoke with them, and they said, “All of her markers are clear. Not only are they clear, they’re perfect.” And they said, “So is she having symptoms anymore?” And I said, “No. She has no swelling, she has no itching, she has no bruising, she has nothing. She’s running around, she’s gaining weight. She’s herself again. She’s able to focus on school work, she’s able to do the things that she always wanted to do.” She completed dance camp in July, which we never would have been able to do that without this stuff. She never would have been able to do that with just being on Motrin all day every day because she was still so sore, and so debilitated.

And she has, as of August 25th, has been released from care, and is on an as needed basis. And she’s arthritis free. I’m sorry. I’m getting teary eyed just thinking about it because I have my baby back, and she’s herself again. And her arthritis is gone in three months.

Winder: That is absolutely truly one of the most touching stories I’ve ever heard. And you’re not the only one having a hard time. Let me ask. How much were you giving her, and of what?

Lindsay: So we started very small because her father and I both have various sensitivities, so we worried it’s coming from cows, that we were worried that we were going to have issues with it. So we started small. We started

with just a half a capsule twice a day in her almond milk. And then she liked drinking it. She was calling it her special vanilla milk, and still to this day that's what she calls it. But we found that she really enjoys the chews. And we've increased it gradually, and now she takes the chews throughout the day. If we know we're going to be out around people, or there's sick people, or we're traveling, and we know she's going to get overly tired, we ramp it up, so that we know her immune system's got a fighting chance.

So in the morning she'll take one of the pineapple lozenges, and one of the strawberries. And then in the afternoon, we make Colostrum ice cream for the kids, and they love it. And so she'll get ice cream in the afternoon. And it's her special treat. And then in the evening we give her the chews again because she was having problems with her blood sugar dropping in the middle of the night. And she would wake up starving and crying, and from what I understand that's a normal side effect of arthritis and celiac is to have stomach problems. And she is sleeping through the night. She's not having the blood sugar drop like she would have before. She wakes up a happy kid, and she's ready for her moo-chew every morning.

Winder: Wow. So in the ice cream, how much Colostrum were you – using the powder for that?

Lindsay: Yes, we're using the powder. So we make a batch of it, the big batch and use a fourth of a cup in the batch. So she's getting a pretty decent amount in – because it's three cups of cream, and a can of sweetened condensed milk. So then it's a quarter of a cup of the Colostrum powder and it's all mixed in there, so she – and it gives it the most amazing flavor. I'm an ice cream junky, and that is – ice cream is my weakness, and it is amazing, and it doesn't give it a funky taste or anything like that. No, she loves it, and she asks for it all the time.

Winder: That's spectacular. Have you ever used any of the Limu with her?

Lindsay: She's actually tried that one as well. She does like it. She likes it mixed in fruit juice. Her favorite is the V8 Splashes. That's her favorite way to drink it.

Winder: Oh, that's nice. Yeah. That's got to be one of the most spectacular stories I've ever heard anywhere. And I am so happy that you shared this with us because I think this is vitally important that so many people are going to see the truth of how hopeful this stuff really is. And the possibilities that this holds for so many people out there who are in "hopeless situations." It's not hopeless. Our bodies are a miracle. And if you give them the right stuff, things can happen like this. And it's just so thrilling. So I just want to thank you so much for sharing. Anything else you want to add here before we shut this down?

Lindsay: Yeah. I will admit, I was definitely skeptical. I knew the research was there, but I had been trying other natural things and nothing had worked. Everybody that I had talked to, they were like, “There’s no way.” Other RA (rheumatoid arthritis) moms had said that there’s no way. And I went with my gut, and I went with my mommy’s intuition, and I’m so glad that I did because we would not be having this conversation today. It would be a very different conversation today. And I would have a very different story to tell had I not gone with my heart.

Winder: Let me ask something else because that brought something up. When you said that all the other RA (rheumatoid arthritis) moms were saying no way and skeptical. What are they saying now?

Lindsay: I haven’t shared just yet with them because I wanted to take some really good pictures, and show them her knees now, and have it all together. I do know that the moms that I’m closest to, I’ve shared this with, and they can’t believe it. They’re stunned. They’ve seen the improvements in her in just such a short amount of time. Another thing that she’s – she actually ate gluten yesterday, and has had no reaction whatsoever. She had a sandwich, a white bread sandwich, not the best thing for her, but she had it. And she had no itching, she had no stomach upset, she had no swelling, nothing. And this child hasn’t been able to do that, ever.

So it’s definitely incredible, and my whole family is just rallying behind everything, and my phone has been blowing up with people that are the closest to me. Because I wanted to share it with my close family, and friends first. And I wanted to shout out it from the mountaintop, and just share it with everybody because I’m just so excited now.

Winder: Well, this call will be on our website. OhHolyCow.info. We’ll have it up there probably Monday. And along with the transcript and you can send as many people there as want to hear about it. And again, this has been a thrill, Lindsay. Thank you so much, and what a blessing this story is for everyone.

Lindsay: Thank you for having me. I love this product, and it will never not be in my body, and never not be talked about.

Winder: Right. Well, that’s it with all of us. It’s been absolutely life changing for me. And this is another incredible story, so thank you. And so folks if you’re hearing this and you are as inspired as I am, then let me suggest that you get back with whoever referred you to this call, and do yourself the great favor of introducing this miraculous stuff into your life, too. Okay, everyone, thank you.

[End of Recording]