



Virtual Event Transcript

Medicare Part D in 2026: What Patients Need to Know Featuring Saira Sultan and Kay Scanlan

July 22, 2026

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Hello, and welcome to today's webinar, Medicare Part D in 2026: What CLL Patients Need to Know.

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I am Perla Salazar, Program Coordinator at the CLL Society.

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Prior to beginning, we would like to mention a few pre-event items. All attendees in this webinar are muted and only the speakers are on camera.

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We ask you to please direct all questions to the Q&A section displayed on your screen.

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Questions will be sent directly to our speakers and CLL Society staff and are not visible to the audience. After today's event, you will receive a very brief survey and we greatly appreciate your feedback.

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At this time, I would like to welcome our speaker, Saira Sultan.



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Good morning everybody or good afternoon. Nice to be with you today. We are, I'll introduce both myself and our speaker, Kay Scanlan. I am President and CEO of Connect 4 Strategies and also Director of Government Affairs and Public Policy for the CLL Society...

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and joining me today is Kay Scanlan, Principal at Consilium Strategies.

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I jokingly sometimes say that Kay has forgotten more about Medicare than I ever knew. So, you have a great speaker with you today, and I would like to make sure that you are as interactive as you can be under this context, so please keep your questions coming...

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throughout our webinar, we'll try to get to them as fast as we can. Hold on to your questions, though, because if we don't get to them today on the webinar, we'll ask you to resubmit them at the end of the webinar.

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So, today's discussion on next slide, please.

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We'll focus on what has changed in Medicare in 2026. There are some important changes we previewed last year for you all, and we'll discuss both what's already changed in 2026 and what to look ahead for as well.

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We'll also talk about how Part D plans and sponsors seem to be responding to the changes, both in their financial liability due to the Part D benefit redesign, as well as to formulary changes.

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And with respect to formulary changes, we'll be talking about utilization management and what that means for our community...

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including step therapy for BTK inhibitors, as well as what the National Comprehensive Cancer Network has been able to do with regard to revising CLL guidelines.

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And finally, what to do if you are told that your plan denied coverage for your CLL treatment.

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Next slide, please.

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Of course, why this matters is that Medicare rules and changes in financial responsibilities and also incentives for health plans and for intermediaries has a direct effect on whether our patients are able to receive the treatments that your doctors provide or recommend to you.

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Plans can delay or deny or restrict access through all sorts of administrative hurdles and requirements. And these barriers are often used in high-cost cancer treatments like the ones our patients take.

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Understanding how the system works will help you and us respond quickly and effectively. You directly with your health plan and your doctors, but also the CLL Society in how we approach some of the policy issues that are implicated in these denials or delays you may see.

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Next slide, please.

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So, the next two slides deal with key terms that I understand these slides were provided to you all in advance. So, we laid out some glossary terms, both in this slide, and you can go to the next slide now, and in the next slide,...

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and these, we will try our best to make sure that we are going through these as we explain and as we discuss these topics, but they are here for you as reference. And then, of course, will be provided to you...

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at the end on our website as well.

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So, with that, I'd like to turn it over to my colleague Kay Scanlan to start the discussion on what's changed in 2026. Kay, if you'd like to come off camera...



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and unmute,...

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we're ready to go.

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I am on camera and unmuted.

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Okay, take it away.

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Okay. Well, everything that follows today pretty much starts here. Basically, Congress changed the financial incentives, so plans change their behavior.

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And I think in a lot of ways, this is, this may be the most important slide in the presentation because it explains why patients are facing changes and frustrations and everything else. So, if you felt like your Part D plan suddenly seems more difficult to deal with...

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you're probably not imagining it, but it's not personal. It's that something very important has changed. Congress redesigned the Medicare Part D benefit. One of the biggest goals, and I think we all agree it's a good one, was protecting patients. And that's by putting a cap on annual out-of-pocket prescription drug costs.

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The other thing is negotiated prices. So basically, trying to keep Medicare healthy, trying to save money from some drugs, it'll save patients money. For CLL,...

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probably not. But the trade off in these things that are good...

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is that now Part D plans are responsible for a much bigger share of the cost of expensive medications.

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And one thing I've learned over the years is that whenever...

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Congress or CMS changes financial incentives, behaviors change. That's every single time, and that's exactly what we're seeing now. So,...

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plans aren't responding because they suddenly dislike expensive drugs or actually really prefer one drug over another. They're responding because they're now paying much more of the bill, and how much they pay and how much they get paid can depend on which drug...

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they're providing. So, we're going to be seeing more prior authorization, more step therapy, more formulary management, and generally a bit more or a lot more administrative hurdles before patients can actually get their...

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treatment. And I think understanding why this is happening makes the system much less frustrating. Once you understand the incentives, the behavior starts to make sense.

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So, I think if we go to the next slide,...

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we have,...

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I thought the poll question was up here. Excellent. Okay.

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So, the next what we're seeing...

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is a lot more market consolidation. You may be finding that your...

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Part D...

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premium is higher. You may find that your deductible is higher. You may be finding any of those types of things. You might find that instead of paying \$20 for certain prescriptions, you're paying 20 percent.

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And I think all of these things are emerging this year. I think some plans are going to be taking longer. So, when we do get to the poll question, we might see that some people are experiencing zero problems and maybe because they're not receiving treatment right now. It might be because they started treatment two years ago. It could be any number of things.

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We expect that over time we're going to see more plans...

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catching up with the ones who are really aggressive now...

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and changes over time as well. So, if we want to move to the next slide.

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So, here's our poll question.

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I will read the poll question as well as the answers.

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How did your Part D prescription drug costs - now remember to think about premiums, deductibles, and co-payments -..

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change in 2026...

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compared to last year in 2025? Did you find that they went down?

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They stayed about the same?

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They went up?

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I'm not sure yet...

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Or I don't have a Part D plan in Medicare.



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So, we can go to the next slide and we'll have poll results for you momentarily as Kay continues.

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Thank you, Saira. I'm really curious about this because we're hearing very different experiences around the country. And we're hearing different things from patients and from providers. So, people are experiencing this differently, so it'll be exciting to see what everyone has to say.

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One of the things that...

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we've already spoken about, and something that catches many patients off guard is sometimes you'll hear the word denied, and immediately you think, my insurance won't cover my treatment.

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That's often not what's happening. Very often, the patients, the plan is just saying, we need more information...

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or your doctor needs to request an exemption,...

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or we want to go through a different prior authorizer, another prior authorization process before we'll approve the treatment.

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Those are all frustrating, but they're very different from a final determination that a drug won't be covered.

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So, when you look at the initial claim denials, over 70% of specialty claims face initial denials. If you know that, you know that, okay, 70%, well, I could probably expect that this is what I'm going to face.

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The next question is, how do I deal with it or how do I help my provider deal with it? So, one message that I hope everyone remembers today is this initial denial is often the beginning of the process, not the end of it.



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You don't quietly just, you know, get off the phone with the pharmacy, assuming that nothing can be done. That's when your oncology team needs to know that there's a problem.

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So, if we want to switch to the next slide.

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Actually before we go there, let's see if we can't bring up those poll results now.

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So, it seems like a significant number, a good third, found that their prescription drug costs did indeed go up.

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Another 33% are, don't have Medicare Part D. That's helpful to know for context.

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And it seems like at least less than 20% are seeing that they stayed the same, or they're not sure as yet.

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Yeah.

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So, of the people who do have Part D plans, it looks like a significant number found that they went up Kay, which is, I guess you would expect.

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Yeah, and I think, I mean, the big piece of information here is that we're now two years into...

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Part D redesign...

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where costs, out-of-pocket costs, are supposed to go down. And we're also at the beginning of the price negotiation implementation. So,...

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both of those things were designed to decrease costs. And what we're seeing is three percent. Now that's three percent and you look,...



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a third of people, you know, are not, don't have Medicare, so it's probably closer to eight to nine percent. But still, that's a significant minority when you're looking at implementation of something that was designed to actually reduce costs.

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And Kay, would you say that that is, in part related to the fact that oftentimes we think about there's now a cap on our out-of-pocket spending, and so shouldn't costs have gone down for us, but premiums were not included in the calculation of out-of-pocket costs, yes?

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Right. And I think, yes, premiums were not included in calculation of cost. We included them in the question because we wanted to get to that answer. The other thing is that with these changes in incentives and kind of...

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pushing plans to cover to pay for more of the drugs, that they cover a greater percentage of the costs.

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We've seen a lot of issuers just get out of the Part D business altogether. So, patients have fewer choices, and the choices they have, they have to look very carefully at them, and sometimes those are the more costly ones, maybe...

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you know, a year ago, two years ago, they were in something that was cheaper. They had to switch plans because their plan no longer exists.

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So...

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yep. So, if we go to the next slide, we'll get to a phrase that you'll hear over and over again.

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Aha, what is utilization management?

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It's the name for the insurance rules or processes that control access to treatment.



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So, it sounds like something only health policy people would say. Fortunately, you don't need to remember the phrase. You've probably already experienced it.

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If you've ever heard your doctor needs to send in more information, or we're waiting to hear from your doctor's office, or your insurance wants you to try another drug first...

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or there's a quantity limit or a dosage limit, you're experiencing utilization management.

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From the plan's perspective, these are to manage their spending. From the patient's perspective, it feels like paperwork that stands between you and your treatment. And when you've got cancer, that's the last thing in the world you want.

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Just remember, though, that most utilization management issues are administrative problems or paperwork problems, and they can usually be solved.

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Unfortunately, we're seeing more and more of these tools being used and its impact, it's putting a big burden on your providers.

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So here are some trends, right? We're going to be expecting more utilization management, not less. None of us has a crystal ball...

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but the trends are becoming pretty clear. We're seeing prior authorization use more often, we're seeing greater use of something that we'll talk about later, step therapy, which means basically the plan is saying, hey, you know, I understand you want this patient to take this or this combination or this dosage or this specific brand,...

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great. What we need you to do, though, is have them try this other one that we...

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prefer first, and if they fail or they can't tolerate it or whatever, then fine, we'll let you use the one...



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that you've originally prescribed. So, we're seeing a lot more of that. We're also seeing them pay a lot closer attention to high-cost specialty medications.

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And I'm not saying that to worry anyone. It's because...

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I want you to be prepared,...

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when you understand what's likely to happen when you understand why it's likely to happen, it's easier to respond faster when it does happen.

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And that raises an important question. If we want to move to the next slide.

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Okay, as you get on the previous slide, the last bullet, new negotiated drug controls. And one of the questions we received from a webinar attendee is whether zanubrutinib is going to be affected,...

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Yeah

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affected by the Part D changes. I know you're going to talk a little bit about NCCN later on, but did you have something to say here? Because we, of course, we know that two drugs have been negotiated, two CLL drugs, but not yet zanubrutinib. So does it escape...

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what you're talking about in utilization management, or does it not?

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If I were going to crystal ball, which I said I don't do,...

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what I would say is this,...

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we're going to see a real tightening on zanubrutinib this year, probably last year, this year, the year after, probably right on up to 2029, 2030. And then...

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we're going to see it suddenly looser. So, your plans are going to want you to be taking zanubrutinib.

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And the reason why is acalabrutinib and...

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Imbruvica now, I'm forgetting the technical name for that, so I apologize. I'll say "I".

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Those are both subject to negotiation.

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And that means that Medicare kind of collects a bunch of information from the manufacturer. They look at how much they charge for non-Medicare, non-federal programs. They average it all up. They look at the dosage, they average all that up...

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and they come up with a negotiated price. Now, when I say negotiated, that's kind of a loose term. There's, you know, there are definite ceilings on how much they can...

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put in an offer, start the negotiation with. So, what this does because the negotiated price comes with the theoretical benefit to manufacturers, which is that...

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plans have to put them on their formulary. So that means that the manufacturer isn't going to be going to the plan and saying, hey, you know, we'll give you X amount of a rebate that you can put in your own pocket if you put our drug first, if you prefer it, etc, etc. They can't,...

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they can't do that anymore, you know, that incentive was taken away. So, these negotiated drugs are going to become less attractive because...

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they're less able to get these rebates that they like.

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So, when you look at zanubrutinib, you're going to see something completely different because...

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they will never, they likely will never be subject to negotiation, because the rules changed on orphan drugs so that they're exempted...

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right now, from negotiation ever.

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What they do have is a temporary situation where the plans pay a higher percentage...

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of the cost...

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than they do for the other drugs.

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And that's because, you know, there's this manufacturer discount that really increased how much manufacturers pay, and they, Congress, didn't want small manufacturers to have to bear that burden, because it could crush them. So, they said, okay, tell you what, if you're a small manufacturer,...

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we'll, like, phase this in. So, and instead of Medicare, federal government picking up the slack, they said plans have to. So immediately these drugs became far less attractive, and especially when they're costly.

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I mean, if it's not a very expensive drug, it's not as much of a big deal. If it is expensive, it can be a big deal.

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So, does that answer the question, do you think?

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Okay,...

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I think it does. Thank you, Kay. And I think it answers a related question too about why is it the plans make it so difficult to get these necessary treatments. You know, denying



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and dragging on the process with these extra steps. And yes, it's true that patients are sick and they need timely medicine...

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but I think this webinar and what Kay is talking about will help folks understand some of the financial incentives that make it...

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understandable, but not okay by any means for plans to be doing what they're doing. But it's a financial incentive. And so that's, I think, what we're talking about here today.

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Absolutely, absolutely.

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Thanks, Kay.

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So,...

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here's the big question. Your physician knows you. Your insurance company applies general rules.

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So, this really does get to the heart of today's discussion. Your oncologist knows your CLL/SLL,...

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they know your other medical conditions. They know your treatment history, your medications that you're taking now, medications you've taken in the past and not done well with. They know your goals, they know whether you're a person who travels extensively,...

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foreign travel, do you go across the country. Basically, they know you. Your insurance company doesn't know any of these things.

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What it has are coverage rules. Most of the time, those rules work reasonably well. Sometimes they don't.

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And that's when physicians and patients need to advocate for individualized care.

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So, the goal isn't to fight insurance companies. The goal is to make sure the insurance rules don't override good medical judgment.

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So that is it in a nutshell. They're going to make rules that decide what CLL treatment is quote "best".

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Your doctor looks at you and decides what treatment is best for you.

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Your job is to make sure...

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that you get what your doctor wants rather and what you want, rather than what the insurance company wants. And again, it's a paperwork issue.

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Well, that segues really nicely into the next slide on our next poll question.

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Have you ever had your treatment delayed or denied? We should probably say, due to insurance issues.

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So not because your doctor didn't, you know, is denying you something, but your insurance company denied or delayed you access.

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The answer choices are yes for a BTK inhibitor.

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Yes, for a different medication.

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No, never.

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Or not sure or I don't remember.



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And it is not uncommon...

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for patients to not be sure.

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And say a bit more about that, Kay. Why is that?

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A lot of it's going on behind the scenes.

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So, your doctor may have looked at your insurance before you've even walked into the office, right? So, they know what is the...

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easiest treatment to get through your insurance. They know what's the least expensive, they know what's likely to get approved for today, so that you can get your treatment right away. They know, they may very well know those things before you even walk into the office.

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And all of these prior authorization things, all of these step therapy things, formulary issues aren't...

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affecting things from your perspective. They're affecting things before the decisions are even made. So, I mean, one thing that you can do when you speak with your clinician is ask, ask them whether and how much whether or not,

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if insurance issues didn't matter,..

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what treatment would they prescribe, and just see if you get a different answer, and whether it's worth,..

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you know, and discuss that with your clinician.

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And I know we'll get into that more in a minute. Why don't we go ahead and see the poll results on this one before we keep going?

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Very surprising. No, never. So, I think to Kay's point a moment ago, we really should be perhaps including in our discussions with our physicians, would you have chosen a different medication for me today had it not been but for...

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insurance-related issues. I think that's a really good question, Kay, to arm our community with so that...

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do they really know whether something is being denied to them by insurance and already factored into the doctor's decisions.

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And then, yeah, almost 20% said, yes, they have in fact for a different medication, not a BTK inhibitor, but for a different medication, had their treatment delayed due to insurance issues.

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And that might be related to a combination with venetoclax or any of those things...

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which I know you'll talk about in a little bit.

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Yeah, absolutely.

00:35:49.000 --> 00:35:51.000

So,

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want to keep going?

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Absolutely, let's do that.

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Okay.

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Step therapy.

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It's a problem. And we touched upon it briefly in kind of explaining what it is.

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So, in a way, this is one of my favorite slides because I think it represents a really important development for patients

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Step therapy is built on a really simple assumption. It assumes one drug can be substituted for another.

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Okay, in some diseases, that's a reasonable assumption.

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In CLL, not so much. Your oncologist isn't necessarily choosing a BTK inhibitor because it's their favorite. They're looking at you and your cardiovascular history, whether or not you have headaches, your bleeding risk, drug interactions, all the things we talked about before...

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the patient-specific factors. And those factors really matter.

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And something that I think is pretty exciting is that...

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NCCN, that's the National Comprehensive Cancer Network,...

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and

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NCCN puts out guidelines, and one of the reasons they did it is...

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they wanted to make sure that the care that people receive at comprehensive cancer centers...

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is accessible to patients who aren't close to one of their centers.



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And the way they do that is they put the guidelines out there. They put the guidelines there so that they're accessible. And those guidelines are hugely important. They're important enough that Congress and CMS said, hey, look at NCCN guidelines and make sure that whatever you're doing with your formulary, with your coverage policies...

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aligns with NCCN guidelines.

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And...

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NCCN recently made very explicit what CLL...

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clinicians have understood for a really long time, and that's that BTK inhibitors are not necessarily interchangeable, and they're not interchangeable...

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for some patients, they're very much not interchangeable.

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And that sounds, the change that they made sounds like a small wording change, but it isn't. Since Part D plans are expected to have coverage policies consistent with accepted clinical guidelines,...

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once NCCN said the drugs aren't interchangeable,...

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it becomes or should become much harder to justify forcing patients to fail one BTK inhibitor before receiving another....

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without any clinical rationale. So,...

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basically, NCCN is saying...

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CLL SOCIETY

these aren't interchangeable. NCCN says, look at patient-specific factors and that the clinician makes the treatment. The treating clinician...

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makes the treatment decision.

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So,...

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the right BTK inhibitor depends on the patient, not the price, and not the cost to the plan.

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So,...

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suppose your insurance company still says no.

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If we go to the next slide, we can talk about what to do next.

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As you get into talking about what to do, I want to just make sure we answer a couple of questions here.

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We have a question about whether zanubrutinib and acalabrutinib will be affected the same or differently, and where does Jaypirca fit, I'm never going to say that right, Jaypirca fit into the mix as well.

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Yeah.

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I expect, and I'm going to use initials if I have trouble pronouncing that.

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Well, the questioner used the brand name of the drug, so I'm just repeating it back to you.

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Okay.



CLL SOCIETY

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And I'm going to not repeat it. Because I saw, sometimes you say easy for you to say this time I saw one, so I'm laying off of it.

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So,...

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we do expect to see a lot more...

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prior authorization, tougher prior authorization, and potentially step therapy. And we expect to see that on "J", and we expect to see that on "Z".

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Okay.

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And that's because on both of those it's because they cost the insurance company more...

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and it's simply that. So, they may want you to try a different drug. They may want you to try a drug that they've made a better deal with the manufacturer with.

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But that's, that's what we expect to see is a bit...

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more of a squeeze on "J" and "Z",...

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and less of a squeeze on "I" and "B". And when we get to combination therapy, I think we're going to be, I think we already do see logistics problems on that one. So,...

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let's move from understanding what step therapy denials are...

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to how we might solve them.

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CLL SOCIETY

So, if your insurance company says we want you to try a different BTK inhibitor first,...

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don't assume the conversation is over.

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Like a denial, it's often just the beginning.

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And remember that your insurance company is looking at cost. Your physician is looking at you. Those are two very different questions.

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So NCCN guidelines...

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very much give a strong foundation for an appeal, a reconsideration, a peer to peer,...

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any of those things, instead of simply saying, I prefer this medication, your doctor can now say nationally recognized treatment guidelines that Part D plans are required to a day or two...

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say that these BTK inhibitors are not interchangeable. So, I mean, this isn't about one drug being better. It's about recognizing that different patients need different drugs. And who knows? It's not the insurance company, because they're making one big decision for the entire, their entire cover population.

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It's your doctor, your doctor who knows you.

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And most oncology offices deal with these every day. I will tell you that it is incredibly, incredibly burdensome, but...

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it is less burdensome once you and your clinician know how to frame these appeals.

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So, what does that appeal look like if we want to go to the next slide?

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Okay, the first thing,..

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ask for an expedited clinical exception.

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Time matters most in cancer care. So, this slide is really an action plan for step therapy.

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If step therapy is standing between you and the treatment your clinician prescribed or would have preferred to prescribe,..

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if you don't,..

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if you don't know, then don't, don't settle for that initial answer. Talk to your team, your care team, and ask whether an expedited clinical exception is appropriate. And the key word is expedited.

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Cancer treatment isn't something patients should have to wait weeks to sort out or months or submit documentation multiple, multiple times.

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And going through an appeal isn't saying we don't like your rules. It's saying these rules don't fit this patient. They don't fit this patient and this drug.

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And one thing I've learned is that individualized medicine doesn't stop when your doctor writes a prescription. Sometimes individualized medicine also means individualized insurance review.

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So, if I could give everyone just a piece of advice, stay engaged.

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Stay engaged with your Part D plan and your clinician team, ask questions,..

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follow up. And if you're in the process of trying to push through something like step therapy, don't assume that silence means progress.



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So, let's suppose we're not looking necessarily at a step therapy issue. Let's imagine the problem happens at the pharmacy level. So, if we want to move to the next slide.

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Okay.

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Sometimes...

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you may find that your BTK inhibitor is denied or you get to the pharmacy and it's too expensive.

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Right? And when I say get to the pharmacy, I mean, you know, if this is going to a specialty pharmacy that's delivering it to you, you may hear from the pharmacy. You're interaction is not going to be over the counter, it's going to be by phone.

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So, first off, don't panic, don't get off the phone. Start asking questions.

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And this actually is a fairly practical slide. If you picture this, you get a call from the pharmacy. You're expecting them to say, oh, we've got your prescription. It's coming whenever, or you're looking online to see when it's being delivered. And instead you're told...

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insurance rejected the claim.

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And I can't imagine...

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very many more stressful moments than that. So, what I'd like you to do is first don't panic.

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Second,...

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don't simply walk away or get off the phone and say, okay, or any of that.



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You want to ask the pharmacist what's causing the pharmacist, or whoever you're on the phone with, what's causing the rejection. Is it prior authorization? Is it a need for more documentation? Is it formulary issue?

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Is it step therapy? Is it...

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something to do with, for example, they want you to pay \$2,000...

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at, you know, for your first fill, and you're like, well, wait a minute, I'm on the, the prescription payment plan with Medicare. I'm supposed to pay monthly.

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That's another thing that could happen. The more specific the information you get, the easier it is for you and your oncology team and your care partner to solve the problem.

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So, the most important thing is...

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contact your clinician's office that same day. Don't wait until next week. Don't wait until two days from now. Treatment delays are much easier to solve when everyone knows about them early.

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So again,...

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most insurance problems are paperwork problems, and while paperwork problems are frustrating, they're usually solvable.

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And if the problem can't be solved immediately, there are still additional options.

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So, if we want to get to the next slide.

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Okay, so you have more resources than you may realize.



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Fortunately, patients aren't expected to navigate this stuff alone. You and your clinician...

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can act on your behalf.

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Your clinician can work with you on appeal rights, on reconsiderations, on all of those things.

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A lot of oncology offices have an entire staff behind them who work on these issues every single day. And manufacturers also often have patient support programs. Everyone has the same goal. They want to keep your treatment moving.

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So, for example, bridge programs can sometimes help prevent treatment interruption while insurance issues are being resolved, but they're not appropriate in every situation. They're absolutely worth asking about.

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The important thing is not to struggle with this by yourself.

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You want to let the people whose job it is to help, help.

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So,

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I think we have a poll question coming up. So, I'm curious about how many people have actually experienced step therapy. And we may have,...

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I expect we'll have pretty much a very similar set of answers as we head to the first one.

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Yeah, why don't we go ahead to the next poll question?

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So have you ever been asked to switch drugs or quote "try another one first",...



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before your treatment was covered, before the treatment that was initially prescribed to you by your physician...

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was covered?

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So rather than a delay question, where you delayed in getting it, this time the question has to do with whether you were asked to try another one first.

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And your choices are yes, for my BTK inhibitor.

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Yes, for a different medication.

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No, never.

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Or not sure, I don't remember.

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And as you're thinking about that, and before we pull up the poll results, maybe I'll just take a moment here to...

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ask and maybe help answer a couple of questions along with Kay. Kay, one of the questions we've had is whether the things you're describing on the webinar today apply to Medicare as well as to Medicare Advantage.

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And I think the answer is yes. In fact, maybe more so. But Kay,...

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Yes.

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So, if you have Medicare Advantage, you probably have a Medicare Advantage that has a Part D plan attached to it, right?



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If you have regular Medicare, you have fee-for-service, claims get submitted....

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and your prescription drugs go through a Part D plan.

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So basically, everyone, whether you're Medicare Advantage or Medicare fee-for-service,...

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you're still, if you're looking at BTK inhibitors at oral treatments, you're still dealing primarily with Part D.

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There are some...

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pushes and shoves and things like that that are going to make the Medicare Advantage side for your doctor's office look a little bit...

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more difficult. And that's going to be because...

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a lot of,...

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for example, if you have an oncologist who is...

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giving Medicaid, not giving, where you're getting your medications directly from your clinician's office, or from or if you're at a hospital, from the hospital pharmacy,...

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if it's Medicare Advantage, and if it's one of these drugs that has, you know, been through Medicare negotiation, and that's only "I",...

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ibrutinib first. Next year it'll be acalabrutinib, which is going to be a much bigger volume problem.

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CLL SOCIETY

You're going to see that impacting Medicare Advantage patients a lot more, I believe, when it comes to next year, just because of so many different...

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hands the money has to go through...

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because the pharmacy is going to pay whatever it is the pharmacy is supposed to pay, and then they're supposed to be made whole,...

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meaning get the negotiated price.

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By going through this transition of the transaction facilitator thing,...

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and the Medicare Advantage plan is kind of in the middle on that, so right now we're seeing without Medicare Advantage, it's taking over a month for pharmacies to get their money.

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When you add Medicare Advantage to it, it's going to be much, much longer. So that's something I think we're going to be looking at next year.

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Yeah.

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Is.

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I think we've done, I know we have done previous webinars that should be available on our website on the difference between Medicare and Medicare Advantage plans, and I think it's worth, given how many questions we have,...

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I would refer us to that webinar as well, and we'll certainly make sure there's a link to it in the follow-up that we send out to attendees today. But please don't think that a lower cost, initially lower cost, Medicare Advantage plan...

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CLL SOCIETY

is necessarily beneficial. These same issues will be happening in a Medicare Advantage plan, and as Kay just said, maybe even more exacerbated than what you're seeing in a traditional plan.

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And the lower cost is, while important to help us all manage costs in our healthcare costs, there is an offset in the amount of time and effort and trials and tribulations you'll go through in a Medicare Advantage plan.

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And that will be true whether even though Medicare Advantage plans do have to follow NCCN guidelines as well, and it will be true whether you've been on a treatment already for a year and are being asked again to go through and provide all this documentation and all the different...

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scenarios Kay is laying out. That will be true regardless, unfortunately, whether you've been on your treatment already or not.

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Saira, if it's good with you, we got a couple questions in advance,...

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it's okay if I.

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Yeah, please.

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Answer a couple of those right now. Okay.

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Yeah.

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One was...

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whether there are any changes were coming up on the pricing of acalabrutinib.

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And this is, this individual suggested that they were going to potentially be changing their Part D plan...



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to accommodate potentially needing that, that drug, which is super smart. The pricing changes are not going to impact, the pricing changes themselves. When we see next year,...

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negotiated price will kick in...

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for acalabrutinib. So yes, that will be happening. Will it affect what you pay for it, what your out-of-pocket costs? Will it affect that directly? No. Not over the entirety of the year. You may see slight differences in what,...

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you know, first fill...

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charges, it will definitely be all the way up to the maximum. So...

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you're not going to really see that. But what you will probably be seeing is more prior authorization next year. And when you look at which plans, please be sure, don't just look at the formulary itself,...

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plans have all sorts of documents behind that. Make sure that you're looking at the information on the exact plan that you're thinking about enrolling in.

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Because if you're looking at something like a United Healthcare or whatever, you're going to be seeing...

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a ton of different formularies that they have. Make sure you're looking at the right one. And don't just look at whether or not the particular drug's included. Go deeper on that and look into what types of utilization management...

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they are applying. They should have that information upfront and it should tell you whether it's prior authorization. You'll probably see that. I would expect you'd see that on almost all of them. Some of them are going to say prior authorization with step



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therapy. But make sure that when you enroll, that you know what you're, you know what the coverage rules look like for it. And if you see something different, if you see something, say something. Reach out to the CLL Society, reach out to your doctor,...

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let them know that what this, you switched plans specifically to be able to accommodate this treatment that you absolutely need...

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and what you're seeing from the plan is completely different from what you saw that made you think enrolling with them was a good idea.

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So.

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Yeah, I want to highlight that point that Kay, you just made about reaching out to the CLL Society. It was due to outreach to the CLL Society that in fact the CLL Society spearheaded the effort to get a change made to the NCCN guidelines that Kay referenced earlier.

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And so hearing from you all in response to what you're hearing today, in response, I'll discuss in a minute at the end of our webinar, a survey we'll be sending out to you all about some of these experiences.

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And then also, we'll be putting together a toolkit. So, you don't necessarily have to take notes on every single thing Kay is suggesting you do, because we will be working on a toolkit to distribute to patients and to providers...

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later this year, and so actually at the end of, once we have your survey results, we'll then be able to prepare the toolkit. So, all these things should be responsive to the kinds of experiences you're having. And yes, we hear from some of you, but we certainly would like to hear from more of you.

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Yeah.

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CLL SOCIETY

So, I really appreciate, Kay, the shout out to reach out to the CLL Society as you're experiencing these different things. And while we can't tell you what the right Part D plan is for you or exactly what is being...

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sort of where which drugs will be stepped through or see other utilization management happening in your particular plan, we will provide in the toolkit, as Kay is talking about today, which is where to look...

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for some of these roadblocks, if you will.

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And I had another question from an individual with a Medicare Advantage plan through BCBS of Florida...

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and it's multiple parts. The first part acknowledges something that is very frustrating to patients. You enroll in Medicare Advantage, you find that it is a monster to deal with. Your clinician that you were using before is no longer with them, et cetera, et cetera, et cetera. Lots of changes.

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But you can't go to Medicare fee-for-service and get a supplemental very easily because you can get denied on the supplemental or they can just charge you way too much money. Make it very, very unaffordable. One thing...

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that I think all of this chaos...

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may be giving a bit of a...

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silver lining for patients who are not happy being in Medicare Advantage and feel like they're stuck. Keep an eye on the plan you're enrolled in. There is at least some, if not a lot, of...

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kind of departures from Medicare Advantage, your plan may no longer be offered. The issuer, the under, you know Blue Cross Blue Shield, for example, Cigna, may no longer be working



CLL SOCIETY

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you know, doing plans in your area. If that happens, you do...

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have the ability to go back to Medicare fee-for-service with a Medicare supplement that does not have that medical underwriting, meaning they're not going to ask you for, you know what kind of conditions you have. They're not going to look and see you, see that you have CLL, know you're expensive, and just charge you an arm and a leg. None of that's going to happen.

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The other thing is that...

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always remember that as much as there may be some interplay with this, is the BTK inhibitors are Part D...

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right? So,...

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those are Part D drugs. So, whatever you're,...

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you've got it's not going to necessarily be Medicare Advantage itself. It's going to be the Part D plan associated with it, probably run by the same company, but...

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slightly different.

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One of the other questions was about cost.

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That these medications are expensive and yes, they are. The good news is that there is a cap. There's a cap. It's I believe 2,100 now. Next year it'll go up. It goes up each year. It'll go up each year, maybe 2,200. But then,...

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the other part of it is that you can divide what you pay over time over the entire plan year. So, you don't have to pay the whole amount right up front. Instead, you enroll in the Medicare payment program,...



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you have, you fill your prescription, you stop paying for drugs when they're delivered, you don't have to give a credit card or anything else to the specialty pharmacy. If you go to the pharmacy counter, you don't have to give them any money. Everything is much smoother.

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So, it is huge as far as making it easier for Medicare beneficiaries.

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And I think.

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Why don't we take a look at the poll question results?

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So the question was, have you ever been asked to switch drugs to try another one first before your treatment was covered? And it seems like well over half said, no, that has never happened to them.

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Almost 20% said yes, in fact, that has happened to them, but not for a BTK inhibitor, but for a different medication.

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Okay, any thoughts on that?

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This really does mirror what we saw on the prior authorization question.

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And it's different from what we're hearing from clinicians, which kind of makes me think that a lot of the...

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things that are there that patients may not see them, that the clinician's office, if you're getting the exact prescription, it's because your doctor's office has been fighting...

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for you to be able to get your prescription.

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There's also the possibility that, you know, if your doctor believes that one is as good as the other, they may very well decide to go with the one that's easiest to get through your insurance, which, you know, that's, and again, this is one of the reasons that we suggested earlier...

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that when you're talking to your physician, when you're putting together a treatment plan, you know, as you're asking questions, ask a question about...

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insurance, how much that's kind of influencing. Not that your doctor's going to give you something that doesn't work because it's easier, but...

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is there something they might have done, might do differently if not for insurance?

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Oh, sure.

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Okay, I know Kay is going to switch gears now and start and talk a little bit about fixed duration therapy. We do have a couple of questions about supplemental insurance Kay. So, I might just say very briefly that a Medicare plan allows you to buy...

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a part, a Part D plan and a supplement, whereas a Medicare Advantage plan does not, and a supplement, supplemental plan is just an additional coverage for your out-of-pocket costs....

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but not Part D out-of-pocket costs.

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But not Part D out-of-pocket costs.

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I know when you say for your out-of-pocket costs, it sounds so awesome, right? But no, not your Part D out-of-pocket costs.

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So yeah.

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Great.

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Thank you, Kay.

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Supplemental. You are very welcome.

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Okay.

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All right, so...

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let me find where my...

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and I do apologize, I was flipping around...

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on the slides that I have in front of me.

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You're starting slide 24 at fixed duration.

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You got it.

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As I scroll through and again, I apologize.

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That's all right. Why don't I take a minute to deal with a couple of extra questions that have come in.

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So, we had a question that talks about changing, trying to change from an advantage plan to a supplemental. And I think we've helped answer that question that a supplement is separate from an Advantage plan. Your choices are Medicare versus Medicare Advantage.

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And it's hard to go back to traditional Medicare after being on a Medicare Advantage plan and our other webinar covers that in detail. The exception, which is really important to pay attention to, as Kay said, is if your Medicare Advantage plan goes away for some reason.

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So we can, let me see if I can take care of any others while you're looking for your spot.

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Are you ready, Kay?

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I am just about ready. I think that on the Medicare supplement issue, it is really, really important...

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for people to make sure that they understand that if their plan...

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is no longer available, if their insurer is no longer operating in you know offering plans that that whole thing about not being able to switch is no longer true. And a lot of people don't know that.

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So, make sure you keep that in mind. And if you, you know, if you are unhappy with Medicare Advantage, and you look at, oh, no, my plan's not, I have to find a different plan, instead of thinking I have to find a different plan, it's a decision point.

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Do I want to go to Medicare fee-for-service with a supplement, or do I want to stay in Medicare Advantage? And you're going to want to kind of look at the costs of each of them.

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Okay, so.

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Yeah. And for people in transition and because we had a decent number of people not yet on Medicare, I do want to make note that watching that webinar on Medicare versus Medicare Advantage or listening to what we're saying today may help you understand why the financial...

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behind the scenes stuff that is going on and driving these issues is very different from your employer plan or an ACA plan versus when you transition onto Medicare. So, it's really important to understand that, yes, all these things that you may not have experienced before...

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Perhaps, will be new things you will experience going forward.

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Absolutely. Okay, so this is exciting because it's...

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one of the biggest changes in CLL over the past few years, and that's been the availability of fixed duration combination therapy. And what that means is...

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instead of having one BTK inhibitor...

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that you're going to be taking continuously until it's either no longer working or you'd no longer tolerate it, instead of that...

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you're going to get two...

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cancer drugs, a BTK inhibitor, and another drug, venetoclax, that...

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together will be given to you over a shorter period of time, and it's a fixed time, 12 to 14 months, approximately.

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So as much as it's exciting in that way, it also creates a different insurance experience.

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What we have now is that patients are excited. They like the idea of eventually stopping treatment instead of taking medication indefinitely.

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But the catch is that since there are two specialty drugs instead of one, there are two opportunities for insurance paperwork...



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and you can expect that you will be running into that.

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Insurance companies don't necessarily become simpler just because medications get better.

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What one of the biggest problems that we see is a logistics problem. And the question,..

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even though it's backed by guidelines, NCCN guidelines in particular,..

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you're going to have...

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a logistics problem and we can go through that if we move to the next slide.

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Okay.

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So, you're going to be seeing prior authorization on both drugs. And unfortunately, even though this combination has been in NCCN guidelines for a while, and now acalabrutinib plus venetoclax is on label,..

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a lot of insurers haven't really updated their claims processing systems. They see a claim for one drug, they put it through the rules for that drug. They see a claim for the other drug, they put it through the rules for that drug. So almost always you're going to end up with a prior authorization that is good for one year. Well, what if your treatment is 14 months? You have that...

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little period of time that you've got to make sure you reauthorize.

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There's also a ramp up on venetoclax. It starts low, steps up over a five week schedule..

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so your prescriber is going to be prescribing a certain amount and...



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Quality, quantity limits at the pharmacy can make it, you know, difficult for you to be able to...

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have that combo...

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aligned with the policies on the drug alone.

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One thing that is super important is remember that mid-treatment decisions are incredibly, incredibly...

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difficult to deal with. You don't want to wait until your treatment schedule to begin before asking whether both medications have been approved. Since this is a logistics problem, and you probably are going to need to get some blood work done,...

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various check-ins, etc, etc,...

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talk to your doctor before you're scheduled to start on a combo therapy. Make sure that they have gotten,...

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you know, everything approved through the insurance company before you even start.

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And to me, that's...

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one of the very, very biggest things. Most of these problems can be prevented if you get this paperwork sorted out early.

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And then, if we want to move to the next slide.

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Okay, so...

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when I think about what is the big idea on this, it's to keep the treatment plan intact, right?

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Combination therapy is very carefully designed,...

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mid-course changes, delays, etc. It's not just inconvenience. They can disrupt the treatment strategy your physician selected. And that's one reason that...

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as we talk a little bit later, some of the newer Medicare protections are going to be especially important. Plans can't simply just say, oh, well, we're changing, we're not going to be offering acalabrutinib anymore.

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Right? Where instead we're going to have everyone start on zanu, you know "Z".

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They can't do that without giving you particular rights. And I see you smiling on my use of the initial, Saira. But, and that's really important, and that's going to be important as we because right now, say you're starting a combo therapy, a fixed duration combination therapy...

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next week, right? We're looking at the end of, we can just call it August for practical purposes. You're going to be going into the next plan year. So, what if you change Part D plans? What if your plan is no longer available, et cetera, et cetera. These are things that you need to work out...

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well before. So, if you're going to be starting a fixed duration treatment this year, it's going to be going into the next year, make sure that if you switch plans, that all of those little things are worked out. Also, next year,...

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acalabrutinib is going to be one of the negotiated drugs. Your plan, as much as they like it better than "Z" this year, next year they may not. And they may want you to go switch to "Z". Well,...

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you can't, that's not something that your doctor is going to want you to do. They don't want to work through the insurance denial process, take a month while you're mid-



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treatment. So, make sure that anything, any of those things that happen are thought of in advance and dealt with before,...

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before you even get moving. And if that's done, then things should be pretty smooth.

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Basically, plans can't just make significant changes in the middle of the year either...

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without triggering formal patient protections. So, if something changes unexpectedly, you want to ask questions immediately.

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So, if we want to move to the next slide, we can talk about some of the new appeals protections.

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Okay.

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Lawyers get excited anytime you say appeal rights, right? And I know that that puts me in a fairly small club, but this really does matter.

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And as I said, if your plan makes mid-year coverage changes,...

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you want to alert, alert, alert. It can't just stop covering without giving you formal notice. So, open your mail that you get from your Part D plan.

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I know it can often you get the same thing all the time and you tend not to look at it. Make sure you look at it. That notice that you get will trigger very important appeal rights.

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And it doesn't guarantee you'll always win, but they do guarantee that you have a process. And that process matters.

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So, you're going to get faster dispute resolution, meaning...



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24 to 72 hours. You should be able to leverage that these are oncology drugs and get it on the shorter end of that with 24 hours.

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So, they have to give you that appeal right before they even deny something. If they made a change and you're on something or if it affects you, they have to give you that appeal right before you've run out of your medication, before they've denied anything. So, it's important to remember that.

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They have a kind of a bit of a support for grandfathering, right? It's not an absolute guarantee, but it makes it a lot harder for plans to just abruptly switch you to a different drug while you're already stabilized.

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And the NCCN guidelines saying that they're not interchangeable certainly gives you a lot of ammunition. And if plans are not honoring those NCCN guidelines, we would really like to hear about it.

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Unfortunately, CMS would object to it, but they don't really have a line of sight into it.

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And unless we see what's happening on the ground, we can't fix it.

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So, if we want to move to the next slide.

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Got some takeaways.

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Basically, I just want you to know, you don't need to become a Medicare expert.

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You just need to know what to do. And Saira talked about toolkits and everything and everything else. But with what you've...

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heard today, hopefully what you've learned today,...

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you can remember four things. First,...

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don't panic.

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Insurance patterns are often administrative problems. And administrative problems, paperwork problems are fixable, right? Second, let your oncology team know early.

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Small problems can become big problems when no one knows about them.

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Third, don't assume a denial is the final answer. I mean, we saw early in this presentation...

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the number of denials that are on expensive treatments. It's pretty high. Seventy percent are going to get prior authorization. A lot are going to get denials first time they're submitted.

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Don't think of it as alarming. Think of it as, oh, well, I guess I fall into the 70%, didn't make it into 30% this time. And understand that it's not the final answer.

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And remember that treatment decisions should be individualized. Your physician knows you, and that's why their voice and your voice really, really matters.

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So, you don't need to memorize the Medicare rules or the crazy words that insurance companies use to describe things. You just need to recognize when something doesn't seem right.

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And when you notice that something doesn't seem right, ask for help.

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That's really helpful, Kay. What a nice way to summarize that. And I will, for those of you who did ask for various terminology and what it all means, even though Kay said you don't have to worry about all of it, please do refer back to the initial slides that provided a glossary of terms...



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and that may also help you answer some of those questions.

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I also want to take a moment now to answer a few other questions that are fairly nuts and bolts type questions, and then we'll be able to...

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and...

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I think be able to get to most of your questions at this point, but if we haven't yet answered them,...

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please remember. Oh, sorry, Kay.

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I'm sorry. I have...

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two more advanced questions that I want to make sure that I answer, that we answer those as well.

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Okay.

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Absolutely.

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Okay, you go first then. Why don't you cover those since people were kind enough to submit them in advance, let's deal with those first.

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Yes, early bird gets the something, right? Okay, so a question is on using CHAMPVA...

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as a Medicare eligible person,...

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and, and how that kind of works. So, if you have, you recently got approved for CHAMPVA,...



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that's a veterans program, and when you're Medicare eligible, in order to maintain...

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the CHAMPVA benefits, you're going to need to enroll in Part A and Part B, right?

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Now, when you think about Medicare Advantage, your CHAMPVA is going to be like a supplement, so you have a supplement, okay? You don't have to worry about paying for supplement, so you don't want to say, oh, well, you know, the supplements are really expensive, you already have CHAMPVA.

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It will be a premium free supplement, right? And it covers out-of-pocket costs.

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So that's one thing that's super important. And what you'll end up having is, you know, the doctor's going to submit the bill to Medicare, Medicare is going to pay...

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if it's a drug, then that goes through the Part D. Any of the balances, any of the co-payments, they automatically forward those to CHAMPVA. And it kind of puts you into a very special club that's different from people who are in...

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the regular Medicare with a supplement because the CHAMPVA is going to work with your Part D or what would be your Part D. It's going to work across the board, right? And you can see any doctor that accepts Medicare,...

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any hospital or hospital-based providers that accept Medicare, they all have to accept CHAMPVA.

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So hopefully that answers some of that question.

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The other question was on fixed duration...

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and about what we are expecting. Are these going to go through as two separate drugs? I would love to get that fixed. I would love to see it go through as...



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a combination regimen treatment plan. For the most part, most insurers have not done that yet.

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For right now, it's going through as two separate drugs. You've got the dose escalation issue, you've got the treatment plan lasting beyond 12 months, potentially crossing over, you know, over plan years...

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and also over calendar years where you might be switching plans. So, all of those things make it a logistics problem.

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And the other thing is, it's interesting because if...

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I would think that to the extent that there is stability in a Medicare Advantage plan, for example, the fixed duration ends up saving money over time.

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But since people switch plants so often,...

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plans have, you know, insurers have learned to look at the short term, right? So, they're going to be saying, oh, well, how much is it costing us this year? Because if they're going to be paying all of it this year...

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and then the next insurer ends up getting a break, right? They're not paying for anything. So that's why, unfortunately, even though it's cost effective, plans aren't necessarily seeing it that way.

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So, Saira, if you want to go with your questions.

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Well, a question on the question you just answered. Can you define what CHAMPVA is?

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Yes, it is a,...



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it's out of the Department of Defense and Veterans Administration.

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Okay.

01:21:41.000 --> 01:21:43.000

So it's...

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yeah, it's like it's a benefit for veterans - veterans, spouses, dependents.

01:22:01.000 --> 01:22:02.000

Yeah.

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And do we have any general information on whether VA healthcare pharmacies are better than Medicare Part D pharmacies? I'm not exactly sure that they don't, in fact, overlap quite a bit, but I'm not sure we know that.

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But one thing is that, and this one, you know, CHAMPVA is for it's...

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not veteran. Yes, it's through Veterans Administration, but it's for the spouses and dependents, right? So, with CHAMPVA, you're not necessarily, you know, you're not necessarily a veteran yourself.

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Just to make sure that that's clear. But, you know, from what I understand, they have far fewer prior authorization issues.

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And I would expect that they're going to be also be a lot happier with fixed duration, right? Because they envision covering the entire patient for the patient's entire life.

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Right, so saving money two years from now sounds like a good idea.

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More to them. So, I think that the problem is that a lot of clinicians don't really understand how to work with CHAMPVA.



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But that's kind of been a learning curve. But they are absolutely required if they accept Medicare, then they absolutely must take it.

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Great. We have a really interesting question from someone about the 70% that you've referred to a few times. The 70% denials likely to happen 70% of the time and not to take that as a final answer.

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The question relates to whether all those denials, in fact, end up raising the cost of healthcare in the United States, and is this sort of because of all the cost of the administrative work for the doctor to resubmit the claim, et cetera, et cetera.

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And I'll let you answer that question, but relatedly, the question asked whether this is something we can take to Congress to address. And that gives me a chance to make a plug for the CLL Society's program where we are pulling together...

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and training ambassadors from among the patients in the community or caregivers in the community, and we'll be starting a dialogue with members of Congress in the House and in the Senate at the federal level about a number of these issues.

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So, continuing to hear about your experiences helps inform the kind of conversation we have with members of Congress at the CLL Society, but also there's an opportunity. This is the second year for the program that is kicking off this year.

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The first year was very successful. We had great feedback from the participants, and so if you are at all interested, you'll find information about our virtual fly-in, it's called our ambassadors program on our website

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But with respect to the first half of the question, how does it, you know, how would you say it raises healthcare costs with all this administrative work for the doctors, Kay?

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Yeah, yeah, I...

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I don't think we have enough time to really let me go on a full rant, so I'll abbreviate it a bit. Okay, first off,...

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when you're looking at these denials, this 70% is probably automatic denials because most claims go through,...

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you know, a computer basically, right? And the computer has a specific what they call edits. You know, you see this drug for this diagnosis in this patient deny, deny, deny, deny, right? And a lot of times those edits are really...

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comprehensive...

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and...

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it's frustrating because yes, it does, it absolutely does increase costs. And it also transfers costs, okay? So, let's look at it this way.

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Plans...

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have what they call PBMx, pharmacy benefit managers, and those are the entities that put together the formulary. They say, okay, here's what we are going to do to help you control drug costs, right? They do the formulary, the utilization management, etc, etc.

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There's a whole lot more scrutiny and potentially limits on...

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what kind of money they can get.

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Historically, they, what they've been, what they have been doing is kind of done their own back alley deals with manufacturers to say, hey, look, you know, we're working for Blue Cross Blue Shield. If you want us to put you...

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at the very top of the food chain on all these Blue Cross Blue Shield plans, you've got to grease our palm, right? You've got to give us a rebate, and they get that rebate, and they don't pass the whole thing on...

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to the insurer that they're working for. Now, since those kinds of things have been closed up a bit, and you're going to see them close up further once acalabrutinib has a negotiated price,...

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then, the PBMs don't want to say, oh, well, we don't make as much money. They're going to be saying, so how do we make that much money? And one of the answers is things that they get paid on a, for lack of a better term, per click, right? The per-click things are going to be prior authorizations,...

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medical review, peer-to-peer appeals, all of those things.

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They get paid a flat fee for each of those, each prior authorization that requires them to look over documents, et cetera, et cetera. So, when they put the denial through and the claim is put in for a reconsideration...

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with documentation,...

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that PBM that put that claim edit in there is getting paid.

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And who pays? Well.

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Who pays is your doctor in the form of paperwork burden. You, in the form of delays in care potentially. And the healthcare system to a large degree. So yeah, it's frustrating. There are some things in the works and Saira's talk about Congress is hugely important, and we all were potentially going to be seeing something on these PBMs...

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coming down the pike...

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as far as CMS kind of...



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making it a lot more transparent, and putting some...

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reins on what they can do. Is that helpful, Saira?

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That's very helpful. Thank you.

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So, we have a question about whether patients can self-pay for these medications before all this paperwork and all this utilization management works itself out.

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Is that a possibility?

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If you get a denial,...

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that's interesting. Okay. I'm assuming and I'm going to check on this.

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But I would assume if you get a denial...

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and you go ahead and you pay for it, you get your medication, and you appeal...

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then that's how you would do it.

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Yeah, I think the cautionary note would be that even though, yes, we think of all of these as problems to be solved, all these denials and prior authorizations and utilization management, one can't guarantee that in the end it is a no.

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And so if that happens, your ability to recoup the money you spent, assuming that it was just a paperwork hurdle, is at risk. So, with that cautionary note, I think, I think there's

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you know, value to your suggestion in the quest, to the suggestion in the question.



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And the better play to some extent would be to request..

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emergency...

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dispensing, right? And that's basically saying, hey, look, you know, I've got cancer. I need my treatment. I need my treatment now. I know you guys have things that you want to work out. We're pretty sure we're going to be successful. Look at the NCCN guidelines. Look at, you know, et cetera, et cetera.

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You need to dispense this medication to me for this period of time so that you can get it approved for...

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the full dosage, the full time period. And I think that can be very helpful. And always remember, not just appeal or reconsideration, but expedited...

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so that you get eyes on it quickly and get it resolved quickly.

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That's helpful.

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And also remember, and something that people may not think about is that, you know, say you pay out of pocket for it and you think, oh, well, I've already hit my out-of-pocket cap. You haven't. You haven't even touched it because unless it is covered, quote, covered, meaning that you're getting that drug...

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through your Part D plan and its coverage. It does not count as an out-of-pocket expense.

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And actually, while you're on the topic of out-of-pocket expenses, we have a question about if you start a drug in October, and you choose to divide the payments over a year, do you have to pay the whole \$2,100 during the current year?

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CLL SOCIETY

It's...

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different plans have different plan years. So...

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assuming it's a calendar year, yeah...

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when you say October, I'm going to exactly at the end of your plan year. Okay, let's just call it end of the plan year, because I think that's really where the heart of the question is.

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And one of the things that we've seen is that because...

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as of now, it cannot be spread across...

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years...

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that you're not going to want to do the payment plan if you know if you've, you know, kind of...

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started late because...

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it's pretty much the same as just, you know, the out-of-pocket.

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And that's a shame, and hopefully they will fix that.

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That's.

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Yeah, so essentially just to make sure we understand the amount you'll be asked to pay in October when you start in a year end, in a calendar year plan will be so high in order to recoup the cost during the course of those remaining two months...



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that there really is no point at that, you know, at that juncture to be starting a 12-month cycle plan, because they're going to front-load it all into the two months to make sure that they get their payment in full.

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Right.

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Thanks, Kay.

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Now, I'm not sure I'm going to pronounce the name of this drug correctly, but is there, a question is, is there any way to get MDC coverage for frontline zanubrutinib and sonrotoclax?

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And I'm not sure I know enough to answer that question, and we might have to...

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Yeah.

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do some digging to secure it.

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Thank you.

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Unless you happen to know?

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No, I mean, all of those things I hate to answer on the fly because, you know, it requires looking at what coverage policies are there, what NCCN guidelines say, what the labeled indications are, etc.

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Yeah, I think those are a little too specific for the CLL Society to tackle.

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Yeah, and we can look at it.

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So, we have done so we have done previous...

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webinars last year, and maybe the year before on both. So, we've referenced already Medicare versus Medicare Advantage. We tackled that topic a while back. PBM reform was another one we tackled, and Kay talked a little bit about the...

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charges PBMs have, which is what leads to 70% denials, as the questioner asked before. And then I think there's another one that deals with open enrollment and coming up to open enrollment that may address some of the questions we've had...

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in our Q&A as well

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There's two more questions at the very end here. One really relates to, will there be, the questioner asks, will there be a future update for the community on what is happening with pre-authorization programs being tested and planned to be instituted for traditional Medicare in the future?

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I think I would recommend that to the extent you all have topics that you are interested in learning more about and would like to have more information about, certainly a webinar is one approach. I know we try to,

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we space out the webinars we have throughout a year, but there is always other opportunities to learn about those things as well. So, I'll just say that to the extent we receive questions, your experience, your problems, and we learning, as we learn about those,...

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it really helps to inform the work we are doing throughout the year at the CLL Society's Policy Institute. So, as Kay mentioned, PBM reform will be coming up again.

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There will be a proposed rule later this year on it, we hear. And so it will give us a chance to comment substantively on what our community is facing, and same with any sort of pre-authorization plans, programs being tested and planned for.

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Right now, we're also expecting more on GLOBE and GUARD, which are negotiations for what treatments should cost if they're not one of the IRA negotiated drugs. So, you know, the Federal Government is,...

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keeps us quite busy with proposals and even legislation, proposed legislation, as well as proposed regulations. So, if you hear of things that you wish you knew more about, please do let the CLL Society know, because those definitely come down...

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to myself and to Kay, so we can...

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better understand the questions you all have and try to find ways to be responsive.

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Yeah, and I think Saira also,...

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patients should remember to be their own advocates, right? You and your care partner. When you go to your clinician and you know you talk to them about what the treatment...

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recommendations are. Hopefully you have an excellent relationship with your clinician and you can ask some of the harder questions,...

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like, you know how, how much is...

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my insurance driving what you think of as the treatment that I can receive?

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And we'll be sure to include that information in the toolkits we'll be providing later this year as well. So, let me...

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Yeah. And Saira, I think, oh, I was going to just say that if anybody encounters step therapy...

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If you feel you've, I mean, if your clinician says yes, you can't have this, I think it's something that we want to hear about.

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Right.

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Okay, let's go to the next slide.

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We don't want to forget to thank our donors for this program was made possible through their donations and through the grant support we received from AstraZeneca and BeOne.

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And then finally, I would like to say thank you for attending today. Because you attended today's webinar, you will receive a link to the access to treatment survey. This is the survey that Kay has been talking about throughout this webinar.

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If you're currently being treated for your CLL or have been in the last two years, please take a moment to complete this survey. The results of this survey will help inform the toolkits that we have been talking with you about so that we can help you be better advocates for yourself...

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as you receive, you know, if you were to receive one of those denials that we talked about earlier.

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So please do take a moment to fill out that survey. It really is a very important part of our quite extensive utilization management work we're doing this year on your behalf at the CLL Society.

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Separately, please complete the event survey you'll receive. That's a short one that you do that comes after every webinar. Provide your feedback. That always helps the CLL Society plan for future events.

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This webinar was recorded and will be made available on the website along with the slide deck and the transcript, and we'll be sure to request that links be provided to the



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PBM webinar, as well as the Medicare versus Medicare Advantage webinar, and so on and so forth.

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If you have questions that were not answered today, I had said at the very outset, please remember your questions. You can send them in to support@cllsociety.org.

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Please join the CLL Society for their next webinar: Understanding the Numbers: Patients Guide to CLL Testing and Imaging on August 26th.

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Please remember, the CLL Society is invested in your long life and you can invest in the long life of the CLL Society by supporting their work. Thank you so much.