

Your Ladies Circle Kawasaki Conversation



Let's get going!



societí
The UK Foundation for Kawasaki Disease

Kawasaki Disease Symptoms



Persistent fever



Cracked lips/
'strawberry' tongue



Rash



Swollen fingers/toes



Bloodshot eyes



Swollen glands

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We want everyone in your community to know Kawasaki Disease - and prevent heart damage in children

A word from your President

Each year our President has the privilege of choosing a supported charity and very often there will be a personal connection or reason for choosing their charity. For me, in my year as President this is the case too as I'll be supporting Societi, the UK Foundation for Kawasaki Disease.

Like most parents, I'd hope I know all the serious diseases to look out for as your child (or your grandchild) grows. We're all alert to meningitis and measles for instance, we'd know what to look for and what to do. But did you know Kawasaki Disease is more common than either bacterial meningitis or measles? AND that it's a really serious disease which is the #1 cause of acquired heart disease in UK children. As a mum it's shocking to me that this little-known disease, which is rapidly on the rise (with no known cause) leads to heart damage in 28% of children affected, and it's very serious in babies under one where 39% of infants (yes 39%!) develop lifelong heart damage. This really is a disease that every parent needs to know!

Early recognition of Kawasaki Disease makes all the difference. When treated early, the outlook is generally good. So, I'm on a mission in my year as President to work with you, to raise awareness of Kawasaki Disease and support Societi in their aim to get 'just' 65 million people (everyone in the UK) to know Kawasaki Disease.

*I hope you will join me in raising awareness and getting your brilliant fundraising hats on so that together we can make a massive difference for children during my year as President. **Join me** – let's protect children's hearts in 2019–20.*

Yours in friendship

Lindsay



What is Kawasaki Disease?

It is the leading cause of acquired heart disease in children in the UK. It affects hundreds of children every year and because of low levels of awareness, potentially many more go undiagnosed, with life threatening and life-long consequences. Kawasaki Disease is serious. Awareness is urgent.

What's the plan?

At Societi, we've an ambitious aim to reach everyone in the UK and tell them about Kawasaki Disease but we can't do this alone! Societi works in partnership with organisations, communities and a rapidly growing army of individuals (**#TeamSocieti**) who are just as determined as us to protect children's hearts from Kawasaki Disease. **Together** we can build knowledge and awareness which can protect children's hearts.

What's the aim?

We want every person, every doctor, **every family in the UK to know Kawasaki Disease** - increased awareness will promote early diagnosis and treatment of the disease which is directly linked to better outcomes including reduced heart damage. By being involved in raising Kawasaki Disease awareness, **you can help** us in achieving this aim.

Affected children will only get an early diagnosis if people can recognise Kawasaki Disease. More common than bacterial meningitis and measles, but just as serious, we need everyone to know Kawasaki Disease.

Join us!



Let's work together – and grow our UK Kawasaki Conversation

Your Fundraising

Let's get going!

On the following pages there are ideas for how you can raise awareness and raise funds. But don't be limited by these – your own ideas will be even better! Whatever you'd like to do, please get in touch and let us know! We'll help where we can.

We know you'll have lots of fantastic ideas, but here are a few of our favourite ways our supporters have already raised funds and awareness for Kawasaki Disease:

- Superhero day!
- Cake sale/ coffee morning
- Wear red day at your local school or at work (even red socks count!)
- Non uniform/ dress down day
- Bingo or quiz night!
- Curry night (with the girls!)

...there are lots of other ways to raise funds and our supporters have run marathons, attempted world records, done sponsored walks, held concerts, sports days, dinner parties and evening get-togethers – all in aid of Societí!

There's also lots of support we can give you too:

- Kawasaki Disease Awareness posters
- TEMPERS leaflets/Societí newsletters
- Thank you stickers
- and much more

....all to help you with your awareness raising and fundraising.

Your Ladies Circle Virgin Money Giving fundraising page

So that you can keep an eye your fundraising and that of other Circles, we have created a Virgin Money Giving 'Societí home page' exclusively for Ladies Circle.

Each Circle can create their own fundraising page attached to this 'home page' which will display all of your fundraising activities, allow you to add stories and updates – and will total the amount your Circle and every other Circle has raised.

You can update your page with fundraiser info and activities to inspire others. To add your Circle's fundraising page to the Ladies Circle Virgin Money Giving campaign page, visit virginmoneygiving.com/fund/ladiescirclesocieti, create your page and get going!



Ways to grow awareness and fundraise - A non-uniform approach!

Whether it's a non uniform day at school or a dress down day at work - a "non uniform" day is a great way to raise awareness and raise funds so we can continue our work all year long!

You could promote the day with posters - download the poster below from our website - societi.org.uk If you're holding a non uniform event you might suggest a

donation amount for those taking part - or just have a collection on the day!

A non-uniform day is a great way to raise awareness!

Our TEMPERS information leaflet can also be downloaded - great for sending home to parents or sharing with your work colleagues.

societi Non-Uniform Day!
for Kawasaki Disease Awareness

Write your details here

Kawasaki Disease is the leading cause of acquired heart disease in children in the UK - we want every parent and doctor to be able to recognise the symptoms and help to save tiny hearts



Could you know the symptoms?



Cracked lips/'strawberry' tongue



Rash



Bloodshot eyes



Swollen glands

- ✓ Talk to your local school
- ✓ Choose a date
- ✓ Print out your posters
- ✓ Promote your event
- ✓ Collect a suggested donation of £1 per child





Coffee and cake!

Holding a cake sale is a simply fab way to bring people together, it's a fun... and delicious (!) way to start a Kawasaki Conversation. Cake sales are brilliantly memorable events - small or large! Perhaps at your place of work or children's school, cake sales have happened before - why not make this one a bit different with heart shaped or heart themed cakes! Or red and white as your cake colour theme!

To be honest, chocolate cake works brilliantly (but maybe that's just us!!)

How about a Kawasaki bake off... the options are endless! All of them taste just as good :)

And why not add coffee to your cake sale - there's a coffee morning poster to download if you fancy too, at societi.org.uk

societⁱ Cake Sale!

for Kawasaki Disease Awareness

Write your details here

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Would **you** know the symptoms?

Persistent fever

Cracked lips/'strawberry' tongue

Rash

Swollen fingers/toes

Bloodshot eyes

Swollen glands

@SocietiFoundation
 @Societi_UK_KD

The UK Foundation for Kawasaki Disease

- ✓ Pick a venue
- ✓ Choose a date
- ✓ Print out your posters
- ✓ Have a bake day and make lots of yummy cakes!
- ✓ Have fun selling your delicious cakes



Go red!

'Red' might be the theme for your Kawasaki Conversation – Kawasaki Disease is sometimes known as the 'Red Disease'! Why red....? Well, distinctive features of Kawasaki Disease include red eyes, red lips and mouth, red hands and feet – and a red rash!

pharmacy can be talked into wearing red for a day to raise awareness? Perhaps you could organise this with school, nursery or even at the office? Going 'red' for the day really will help us achieve our aims!

So how about a 'Wear Red Day'? Perhaps staff at your local GP surgery or local

***societⁱ** Wear Red Day !*

for Kawasaki Disease Awareness

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Write your details here

Would **you** know the symptoms?

 Persistent fever	 Cracked lips/'strawberry' tongue	 Rash
 Swollen fingers/toes	 Bloodshot eyes	 Swollen glands

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 @Societi_UK_KD

The UK Foundation for Kawasaki Disease

- ✓ Talk to your local school, nursery, community group or your work place
- ✓ Choose a date
- ✓ Print out your posters
- ✓ Collect a suggested donation for everyone taking part



Do it your own way!

There are lots - and lots - of ways you can start a Kawasaki Conversation in your community. Dropping off TEMPERS leaflets everywhere you go is a great start; our Founder Rachael doesn't go anywhere without a pile of TEMPERS in her handbag...!

Simply changing your profile pic on social media to a Societi heart logo (or choose one of our Societi Foundation frames for

Facebook or Instagram), sharing the link to our website with family and friends, putting a poster up on a community noticeboard in your local supermarket or at work - these are all really effective ways of spreading the word. You can follow us on Facebook, Instagram and Twitter too - a great way to get started. If you'd like to do something unique, or plan a particular event - let us know and we will help where we can!

Follow us!

- Facebook: @SocietiFoundation
- Twitter: @Societi_UK_KD
- Instagram: @societi_uk_kd

THINK Kawasaki Disease

Kawasaki Disease is the leading cause of acquired heart disease in children in the UK. It's time we changed that...
...Together we will

Kawasaki Disease Symptoms:

- Persistent fever
- Bloodshot eyes
- Cracked lips/ 'strawberry' tongue
- Rash

Kawasaki Disease is increasingly common in the UK

If a child has a **persistent high fever** for 5 days or more, with **TWO** or more of these symptoms please **THINK Kawasaki Disease**

Kawasaki Disease can be present with some or all of these symptoms

Please **EXPECT** to see it, be **READY** to treat it!

TREATMENT IS KEY ...PLEASE DON'T DELAY

Children diagnosed and treated in **less than 7 days** from onset of fever have a much reduced risk of life long heart damage

Current UK & Ireland Outcomes for Kawasaki Disease

7+ days	The current average diagnosis time for Kawasaki Disease is 7.8 days This is too slow!	39%	39% of babies with Kawasaki Disease develop coronary artery aneurysms This is too high!
19%	19% of children overall develop coronary artery aneurysms This is too high!	28%	28% of children overall have some heart damage This is too high!

TOGETHER WE CAN CHANGE THIS!

Poster data from Tulloh et al, Kawasaki Disease: a prospective population survey UK & Ireland 2013-15

- ✓ Get in touch with us via the contact form on our website or email info@societi.co.uk to discuss your plan for raising awareness of Kawasaki Disease. We'll help where we can!
- ✓ Download posters and leaflets
- ✓ Talk to local groups, schools and businesses and share Kawasaki Disease information!





Host a curry night!

Curry is the UK's favourite food and a curry night is a great way to get all your friends, family or work colleagues together and unwind after a hard week's work! Combine your curry night with lots of Kawasaki conversations and you'll have a perfect evening!

Speak to a local restaurant or pub to see if they would be willing to help you host your

Kawasaki curry night and help you raise funds for Kawasaki Disease awareness.

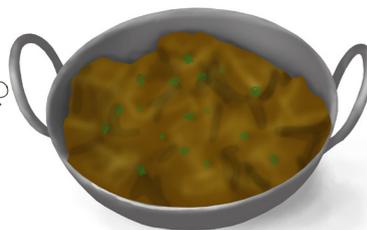
If curry isn't your thing, a restaurant take-over of any 'flavour' would work just as well!

A great approach is to agree a set menu for a price, add a donation amount and you're off! What a fabulous foodie fundraiser!

societⁱ Curry Night! for Kawasaki Disease Awareness

Kawasaki Disease is the leading cause of acquired heart disease in children in the UK - we want every parent and doctor to be able to recognise the symptoms and help to save tiny hearts

Write your details here



Would you know the symptoms?



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Cracked lips/'strawberry' tongue



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Swollen fingers/toes



Bloodshot eyes



Swollen glands



@SocietiFoundation
@Societi_UK_KD



The UK Foundation for Kawasaki Disease



societⁱ.org.uk

- ✓ Choose a restaurant
- ✓ Print out some posters (and leaflets to share on the night)
- ✓ Make a plan with the restaurant
- ✓ Invite everyone!
- ✓ Enjoy a great night out!
- ✓ Send us some pics!

Set Meal - £20
Donation amount - £5

Price per head £25

Kawasaki Disease...
...Let's get it known!





B.I.N.G.O!

Bingo nights are fun and more popular now than ever. All you'll need is some space, some bingo balls and your friends, family and community to take part!

This is a relatively simple to organise but really effective way to raise funds and awareness for Kawasaki Disease. You could even try to work in some heart related 'Kawasaki bingo calls' to raise even more

awareness - two little hearts, 22! - we are sure you could come up with something better!

Maybe there's a local bingo venue close to you - why not ask them to help with a fundraiser? And BINGO... Awareness, fun and funds!

societí Bingo Night!

for Kawasaki Disease Awareness

Write your details here

Kawasaki Disease is the leading cause of acquired heart disease in children in the UK - we want every parent and doctor to be able to recognise the symptoms and help to save tiny hearts

Would you know the symptoms?

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@SocietFoundation

@Societ_UK_KD

societí

The UK Foundation for Kawasaki Disease

societí.org.uk

- ✓ Pick a venue
- ✓ Print out some posters
- ✓ Organise a bingo caller and bingo cards!
- ✓ Enjoy a fun filled evening!
- ✓ Why not think about organising a raffle too!
- ✓ Send us some pics!



Walk a 'Fun Mile'!

Getting out and about in your community to raise awareness of Kawasaki Disease is a great idea, and it's great if everyone can be involved. Organising a 'Fun Mile' is a brilliant way to get out and get moving and also means that everyone in your community can take part.

Everyone from small children to great grandparents will be able to complete this

fun challenge in and around the city, town or village that you live.

It's something you could also fit into a lunchtime at work - or a PE lesson at school - remember to get sponsored and let us know if you'd like leaflets to share - or even medals for super-serious fundraisers!

societ^í Fun Mile!

for Kawasaki Disease Awareness

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 @SocietFoundation

 @Societ_UK_KD

societ^í

The UK Foundation for Kawasaki Disease


societ.org.uk

- ✓ Talk to your school or work place
- ✓ Choose a date
- ✓ Print out your posters (and leaflets to share)
- ✓ Collect sponsorship or choose a donation
- ✓ Enjoy your 'Fun Mile'



Have a Superhero day!

Children love to dress up and what could be more of a boost for your children than being a superhero for a day? Why not talk to your local primary school or nursery about organising a Kawasaki Superhero Day to raise funds and awareness for Kawasaki Disease?

We know that adults enjoy fancy dress too – don't limit your imagination! Just think how your boss dressed as Cat Woman or Super Man would brighten your day!

societí Superhero Day!
for Kawasaki Disease Awareness

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Write your details here

Would you know the symptoms?

Persistent fever	<i>Cracked lips/'strawberry' tongue</i>	<i>Rash</i>
<i>Swollen fingers/toes</i>	<i>Bloodshot eyes</i>	<i>Swollen glands</i>

@SocietFoundation
 @Societ_UK_KD
 The UK Foundation for Kawasaki Disease

- ✓ Talk to your local school, or work place
- ✓ Choose a date
- ✓ Print out your posters (and leaflets for on the day)
- ✓ Collect a suggested donation for everyone taking part
- ✓ Have fun being a Superhero for the day!
- ✓ Send us some pics!



Making Kawasaki Disease memorable in your community

Remember TEMPERS

Through our early work as an organisation, it's really clear that more people simply need to know Kawasaki Disease. As an 'enigmatic' disease which can appear in many guises, there's a reason it's been little known, and often misdiagnosed over time. We want to change that...

So, one Sunday morning, our Founder Rachael created our **TEMPERS** mnemonic to make Kawasaki Disease memorable for both parents and doctors alike. You can download and use this leaflet as a focus for your awareness raising in your own Kawasaki Conversation - just handing out TEMPERS leaflets is a great way to start!

Kawasaki Disease? Remember TEMPERS
Children with **Kawasaki Disease** are characteristically irritable!

- T**emperature - Persistent high fever
- E**rythema - reddened hands and feet with swelling
- M**outh - dry, sore mouth, cracked lips, 'strawberry tongue'
- P**ace - Treat early to reduce potential heart damage
- E**yes - bloodshot, non-sticky conjunctivitis
- R**ash
- S**wollen glands in neck, often just one side

If a child has a **PERSISTENT FEVER** and two or more of these symptoms **THINK KAWASAKI DISEASE!**

5 days of fever? THINK Kawasaki Disease



societi.org.uk
www.justgiving.com/societi

Endorsed by **RCPCH**
Royal College of Paediatrics and Child Health
Leading the way in Children's Health

Kawasaki Disease is the #1 cause of acquired heart disease in children in the UK...



...help us change this.

Kawasaki Disease - who does it affect?
It is mostly a childhood illness with over 75% of those affected being under 5 years old but it affects older children too.

Kawasaki Disease - what's the issue?
In the UK awareness of Kawasaki Disease is low. Currently UK diagnosis and treatment times are too slow. **39%** of babies (under one year) treated for Kawasaki Disease develop serious heart problems. **28%** of diagnosed children experience heart complications. Overall, **19%** of children treated develop serious heart damage. For a few children every year Kawasaki Disease is fatal...help us change this. We need everyone to know Kawasaki Disease as early diagnosis and treatment can prevent heart damage. (Data from BPSU Study, Kawasaki Disease UK & Ireland 2013-2015)

Kawasaki Disease - how common is it?
Hospital admissions in England for Kawasaki Disease have increased fourfold in the last ten years. It's more common than some types of meningitis. About 1 in 10,000 children are currently diagnosed each year and very poor levels of awareness mean many more children may be affected.

Kawasaki Disease - what can I do?
Know the symptoms and remember, symptoms may not appear all at once. Not all children present with all symptoms so - if a child has a **PERSISTENT FEVER** for **5 DAYS** or more with 2 or more of the symptoms overleaf **THINK Kawasaki Disease** and seek **URGENT** medical advice. **You** could save a child's heart.

Kawasaki Disease is serious! Awareness is urgent!
Today, most people haven't heard of Kawasaki Disease. That's one of the biggest challenges we face. Help us get it known because **Kawasaki Disease is increasingly common in the UK**. Too many children and young people today have lifetime heart damage because of Kawasaki Disease...help us change this. For more information visit societi.org.uk

societi.org.uk
www.justgiving.com/societi

Endorsed by **RCPCH**
Royal College of Paediatrics and Child Health
Leading the way in Children's Health





Creating Kawasaki Conversations with care

'Societi Kids'

We're all alert to children's rights and children's privacy. And never-more-so when it comes to their health. So we've taken a conscious decision as an organisation to not use images of child patients in Societi produced material.

But that presents us with a challenge - to convey a serious children's disease to an audience who doesn't know or recognise it. So we created Societi Kids working with our Societi junior artist who has drawn children for us from every walk of life - every part of our society. You'll see them on our Facebook page, in our tweets, GIFs, on our website and on our awareness raising material.

To help develop **your Kawasaki Conversation**, you can use our materials - and with the 'help' of Societi Kids, draw attention to the **serious** nature of Kawasaki Disease - and the **urgent** need for everyone to know the symptoms.

Societi Kids help us to do this in a **fun** way, and a way which we know from experience to date allows people to **engage** well, across all ages and backgrounds.



Kawasaki Disease is the leading cause of acquired heart disease in children in the UK

A bit more about Societi - from Rachael!

Based in Newark, Nottinghamshire, Societi was established to raise awareness of Kawasaki Disease.

Societi works through partnerships to drive transformational change in awareness, about Kawasaki Disease. Our ambition is that everyone: every parent, every doctor and every healthcare professional knows Kawasaki Disease.

As Founder, I established Societi to bring a voice to Kawasaki Disease. It was clear that my expectation for change and action, as a parent with experience of Kawasaki Disease, would not be met without a catalyst for change. And so Societi began!

We work with pace and passion to create high visibility for Kawasaki Disease and bring about the transformation in care our children deserve. We are working to build and share knowledge amongst medical professionals and secure much needed investment. Through our work **we can** stem the now rapidly rising number of children carrying a lifelong legacy of heart damage. They cannot act for themselves.



So Societi must.

Rachael McCormack, Societi Founder

Our Aims

We are an influencing and policy shaping organisation, always working through partnerships to achieve our aims:

Societi has four aims:



1. Awareness raising -
a voice for Kawasaki Disease



2. Clinical research -
sharing knowledge,
influencing funders and
enabling co-ordination



3. Clinical supervision -
shaping new care protocols
and health policy



**4. Support for the Parent
Support Group - KSSG**

Our people

Rachael, our founder, is an energetic and dynamic leadership professional with wide ranging executive experience across public and private sector organisations. She is recognised for her focus on enabling high impact, delivery through partnership, achievement of outcomes and a leadership approach which is collaborative and decisive.



Rob, Professor of Congenital Cardiology and Clinical Lead for Paediatric Cardiology, Bristol, is an internationally recognised expert in Kawasaki Disease. Rob leads the largest Kawasaki Disease clinic in the UK and the only Adult Kawasaki Disease clinic. He is an active researcher and collaborator and supported Societi even before our formal establishment. Rob sits on our Board as a Trustee and leads the work of our Scientific Advisory Board, providing clinical oversight for our organisation.



Claire has 19 years of experience working within the design industry providing comprehensive, technical, graphic and research support across a broad range of projects and disciplines. A valued member of the team, she continues to apply those skills for Societi with her combination of experience and creative flair enabling her to provide proficient and flexible support.



We work together with our Board of Trustees and Scientific Advisory Board to create a voice for Kawasaki Disease across the U.K.



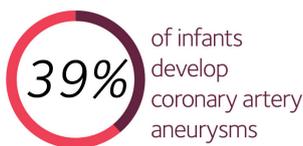
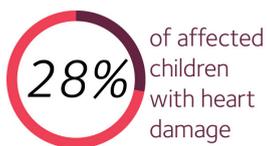
Kawasaki Disease: Facts & outcomes

Kawasaki Disease - Facts:

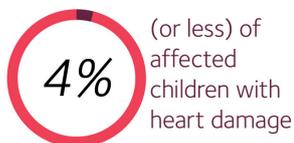
Sometimes called the 'red' disease, Kawasaki Disease is characterised by a **persistent high fever**. You'll see the symptoms in the covers of this document.

- Kawasaki Disease is the UK's leading cause of acquired heart disease in children
- Hundreds of children in the UK are diagnosed with Kawasaki Disease every year
- Across the globe, incidence of Kawasaki Disease has doubled in the last decade. In England hospital admissions have increased FOURFOLD
- Awareness in the UK of Kawasaki Disease is low, research shows delayed treatment = poorer outcomes and increased heart damage
- More UK children today are diagnosed with Kawasaki Disease than some forms of bacterial meningitis or measles
- 20,000 children and young people in the UK have been affected by Kawasaki Disease today
- **39% of infants (babies under 1) will develop coronary artery aneurysms**
- **28% of all affected children will suffer some heart damage**
- **19% will develop coronary artery aneurysms**
- **Kawasaki Disease is serious. Awareness is urgent!**
- **If diagnosed and treated early, children have a much better chance of recovery**

Where we are today



Where we need to be





Societí - creating impact through partnership

Where We Want to Be

Before coming across Societí, it's very likely that you hadn't heard of Kawasaki Disease. We have a vision of a time when Kawasaki Disease is well known, well understood – and as a result, those centres providing expert care, quality treatment and ongoing learning – are well resourced. But there are many hurdles ahead before we achieve our vision.

What is Needed

The image opposite focuses on the main challenges we face. We lag behind our European, American and Asian partners with amongst the poorest levels of awareness across our medical community of Kawasaki Disease in the western world. Amongst parents too, Kawasaki Disease is little known and it is this gap in knowledge which poses such a great threat -- as it is when treatment for Kawasaki Disease is delayed, that the most serious consequences can occur. When diagnosed and treated promptly, children have the greatest chance to make a full recovery.

If we address the knowledge gap with all our audiences, with Governments, with policy makers and funders, with medics and with members of the public, we know we can make real progress.

Key Partnerships in Place

But we are not standing still in the face of these challenges. Each week we take proactive steps, forge new relationships and partnerships, expand our networks, extend our online presence, promote key messages through social

media and raise the profile of Kawasaki Disease in all we do. Today, we are:

- Working actively with **NHS England and Public Health England**
- Working with **RCGP** to develop a clinical E-Learning module for Primary Care
- In dialogue with **professional medical bodies** – working together on a range of projects to raise awareness with the clinical community
- Providing input into the **NHS Improvement Service**, working on patient information
- Working with the UK Support Group, **KSSG** on connected projects
- Actively in partnership **across Europe** pursuing a UK Clinical Trial for Acute Kawasaki Disease
- Working with the **World Heart Federation** through social media to bring global awareness to Kawasaki Disease
- Partnering through our Scientific Advisory Board with **Bristol Children's Hospital, Great Ormond Street Hospital** and **St Mary's Hospital London**.

We have just started building our Societí partnership. We have many more opportunities ahead. We're continually looking for new partners and need to grow! **Your help through Ladies Circle is transformational.**



Ladies... are you ready to build your Kawasaki Conversation? We hope so!

Let us know your plans!

Whatever you do - whether it's taking a handful of leaflets into work or school, or a massive girls night out raising funds, let us know!

We'd love to hear about your projects and initiatives - and can promote your activities on Twitter and Facebook too! A Kawasaki Conversation can start at any time so do keep in touch!

Let's grow our Kawasaki Conversation right across the UK - together, we can prevent lifelong heart damage in our children.

How to send in your funds raised

As you fundraise for Societi, you can keep track of the fundraising you do within your Circle and the activities and targets of other Circles by creating a fundraising page on your exclusive Virgin Money Giving page which has been set up. You'll find it here [virginmoneygiving.com/fund/ladiescircleandsocieti](https://www.virginmoneygiving.com/fund/ladiescircleandsocieti). You'll be able to create a fundraising page for your Circle and post pictures and stories about all your fundraising activities to inspire others! You'll also be able to see what other Circles are doing in their fundraising for Societi!

Ready for posters and leaflets? On the website - go to the **For Families** tab and click the Families Resources and Downloads tab from the drop down menu. Here you'll find lots of printable and downloadable resources.

Check out the **Getting Started!** menu under the Fundraising tab too - where there is even more information to print, which will help you grow your Kawasaki Conversation.

If you're planning a fundraiser or awareness event, please contact us if you need additional support and we will always try to help.

Got Any Questions?

If you've any questions at all, please get in touch and we will help wherever we can!

Contact us by email - info@societi.co.uk

...or through the contact form on our website. We'd love to hear from you!

Your Kawasaki Conversation Toolkit!

societí

Let's make a plan!



My Kawasaki Conversation

A page for your own notes!



Thank you

Over the next year, our partnership with Ladies Circle has potential to reach thousands of people and protect children's hearts across the UK from Kawasaki Disease. We're excited!! We hope you are too!

Sharing knowledge and growing the number of people who can recognise this often pernicious disease is a core aim for Societi.

Thank you for reading and I hope to welcome you and all at Ladies Circle across the UK to #TeamSocieti very soon. Together we can combat Kawasaki Disease and the time you've already invested in getting to know more about Kawasaki Disease means such a lot.

With my very best wishes,

A handwritten signature in blue ink, appearing to read 'Rachael McCormack'.

Rachael McCormack, Founder
for Societi Foundation



Kawasaki Disease Symptoms

societì



Persistent fever



**Cracked lips/
'strawberry' tongue**



Rash



Swollen fingers/toes



Bloodshot eyes



Swollen glands

societì

The UK Foundation for Kawasaki Disease

Victoria Court,
Holme Lane,
Winthorpe, Newark,
Notts NG24 2NU

info@societi.co.uk



societi.org.uk

