

RARETM REVOLUTION

TO BRING ABOUT A DRAMATIC AND WIDE-REACHING CHANGE IN
CONDITIONS AND ATTITUDES FOR THE RARE DISEASE COMMUNITY.
IT'S TIME TO TURN THE TIDE!

Media Kit
TURNING THE TIDE THE RARE REV WAY

LISTEN | CREATE | EDUCATE

RARE reach

RARE Revolution Magazine™ reaches over 3 million people a month globally: 44% USA, 30% Europe, 26% rest of world.



905k

Monthly website traffic



3.4 million

Monthly social media reach



32k readers

Monthly magazine audience



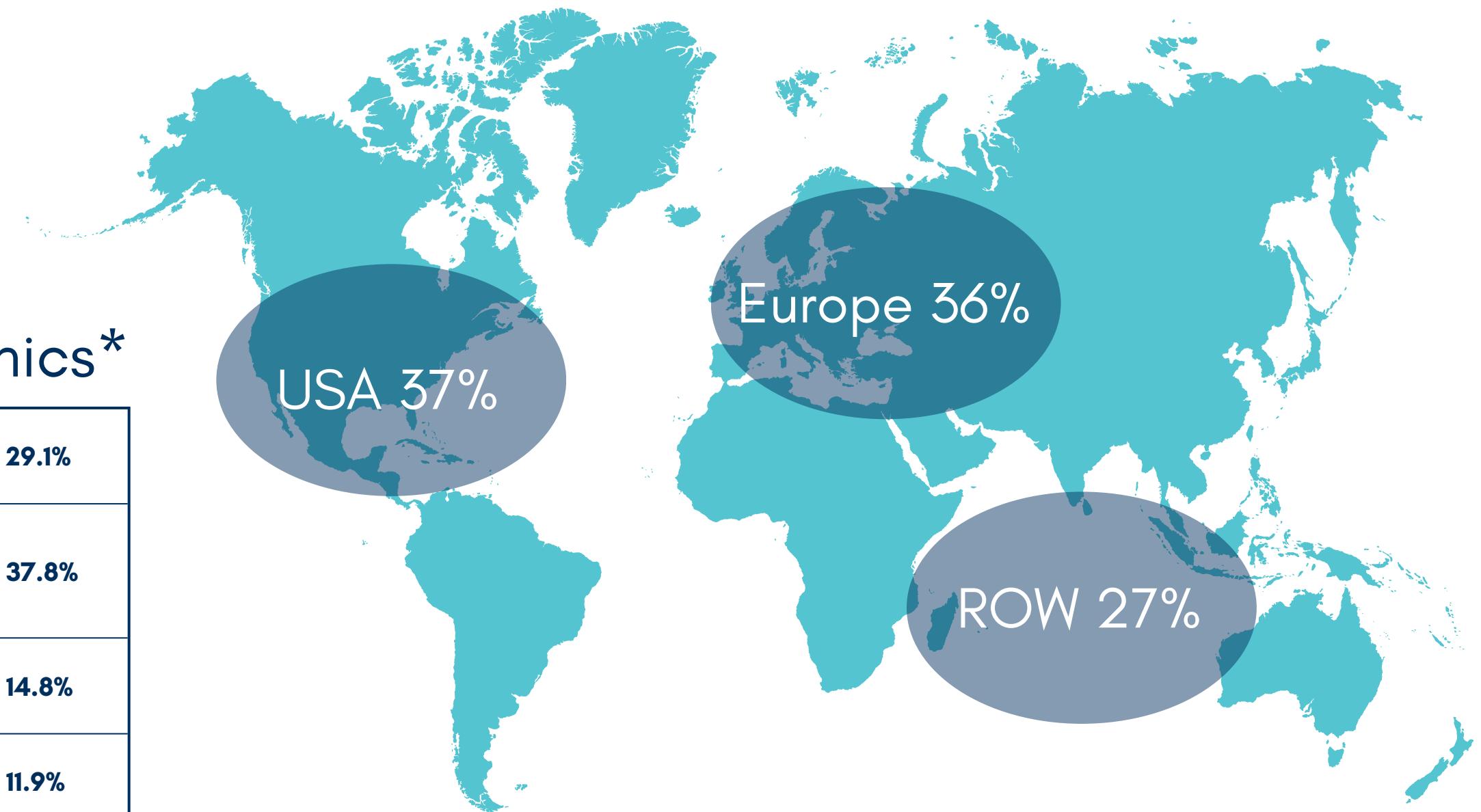
13.9k subscribers

3 weekly newsletters

Demographics*

Industry	29.1%
Individuals and caregivers	37.8%
HCP	14.8%
Charities	11.9%
General interest	6.4%

*Newsletter only



LISTEN

Advisory boards



Landscape/disease mapping

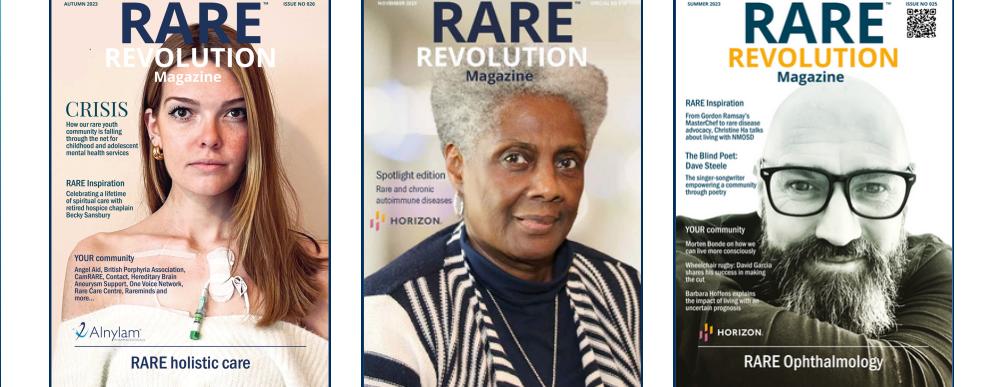


Roundtables



CREATE

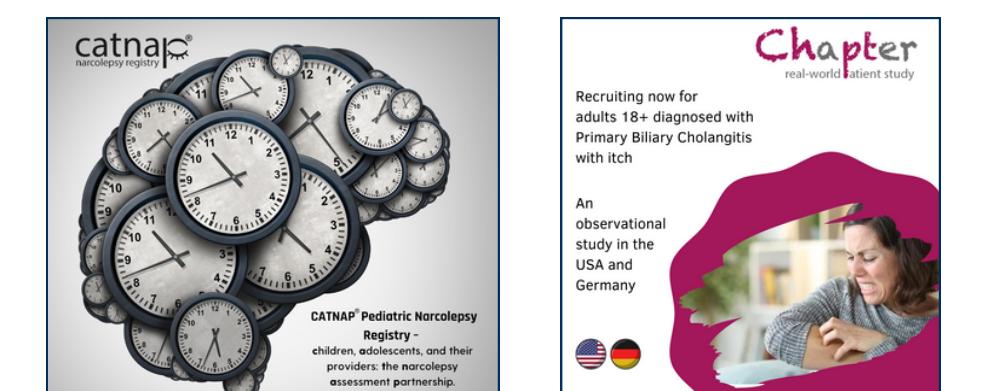
Magazines



Impact reports



RARE recruitment



EDUCATE

RARE REV-inars



Digital spotlights



Thought leadership



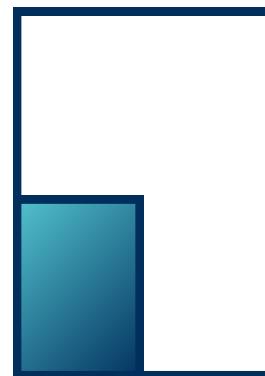
RARETM

REVOLUTION

MAGAZINE

RARE Revolution MagazineTM is a world leading rare disease publication dedicated to elevating the voice of the rare disease ecosystem through our website, digital magazines, patient engagement services, and RARE Youth Revolution. We are renowned for providing trusted insights, thought leadership, personal stories, and advocacy related to rare diseases across industry and patient communities

RATE CARD ADVERTS



Quarter page advert

A6: 105 x 148.5 mm

£350 †



Half page advert

A5: 210 x 148.5 mm

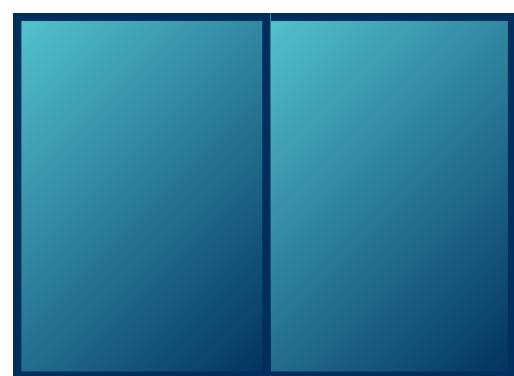
£510 †



Full page advert

A4: 210 x 297 mm

£995 †

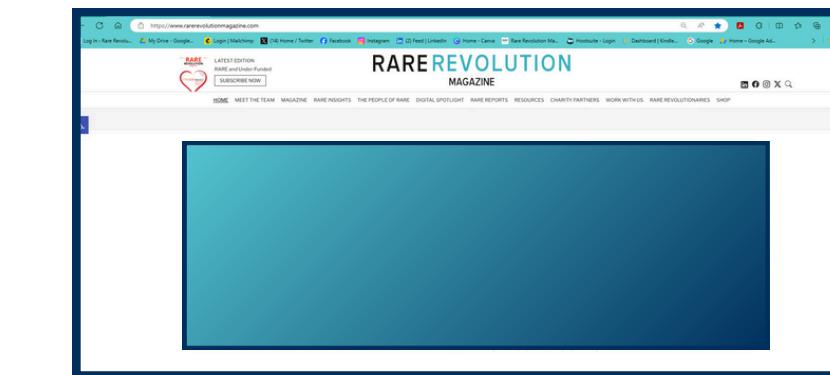


Double page spread

A4 x 2: 210 x 297 mm **or**

A3 landscape: 420 x 594 mm

£1,750 †



Home page web banner

1650 x 620 px

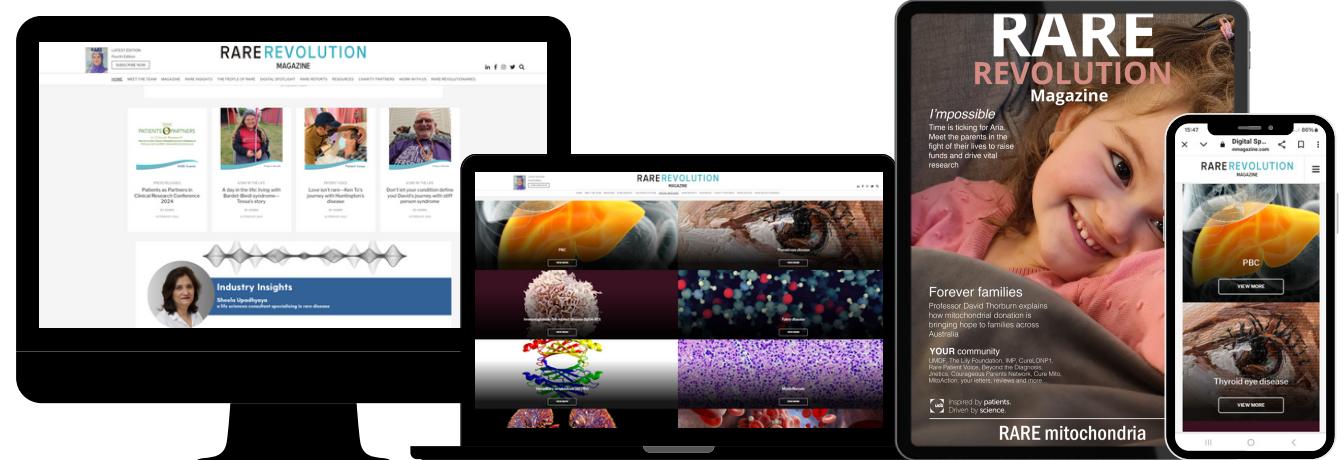
From £995 †

SOCIAL CAMPAIGNS	from £1,495 †
SOLO E-MAIL BLAST	£995 †
EMAIL ADVERT	from £395 †

RATE CARD CONTENT

	MAGAZINE	COMPANY FEATURETTE	DIGITAL	SOCIAL MEDIA	INDUSTRY TAKEOVER
SELF WRITTEN	£1,900 †	£3,500 †	£1,150 †	from £1,495 †	£2,400 †
WRITING SUPPORT	£2,900 †	£5,500 †	£2,150 †	from £2,495 †	£3,500 †

RATE CARD SPONSOR



EDITORIAL – CORE	SPECIAL EDITIONS	RARE REV-INAR	DIGITAL SPOTLIGHTS	RARE DISEASE DAY
<p>Perfect for companies aligned with our editorial theme. Includes;</p> <ul style="list-style-type: none"> • full section branding inc, front page • lead articles in section plus patient insight features • writing support available • printed copies available at extra cost 	<p>Our stand-alone mini issues offer companies a real opportunity to get focused on raising awareness for individual conditions or topic areas. These issues go out to our readership between our regular quarterly issues but are also licensed to clients for distribution through their own channels.</p>	<p>Sponsor one of our webinar series to bring together relevant stakeholders for engaging, roundtable discussions. Available live and on demand. Create evergreen content from your webinar through a video mini-series or article about your webinar ensuring more people reach your content.</p>	<p>Our digital awareness campaigns are designed to bring disease specific awareness, topic education and campaigns to our audience. This opportunity may include;</p> <ul style="list-style-type: none"> • 4 articles • disease infographic • social media promotion • e-newsletter promotion • promotion of campaign in following edition of RARE Revolution Magazine 	<p>Our flagship sponsorship opportunities include</p> <ul style="list-style-type: none"> • Rare Disease Day special magazine edition • digital articles • social media takeovers • RARE REV-inars
£26,500 †	£32,500 †	FROM £6,500 †	FROM £11,800 †	POA



RARE Revolution insiderTM is a life sciences and rare disease publication for professionals, keeping you in the know. Bridging the divide between industry and the community you serve, insider delivers industry insights, key opinion leaders, with exclusive access to our powerful AI librarian, Cognito.

RATE CARD ADVERTS

DIGITAL ADVERTISING RRM AND INSIDER

DYNAMIC ADVERTISING	BILLBOARD ALL PAGES, TWO TITLES	£2,800 PCM †
	BILLBOARD ONE TITLE	£1,800 PCM †
STATIC ADVERTISING	LEADERBOARD	£995 PCM †
	BOTTOM POSITION	£395 PCM †
	LOWER	£995 PCM †

EMAIL ADVERTISING

SOLO BLAST	£995 †
EMAIL ADVERT	£395 †

RATE CARD CONTENT

	THOUGHT LEADERSHIP	COMPANY PROFILE	INDUSTRY TAKEOVER
SELF WRITTEN	£1,150 †	£1,650 †	£2,400 †
WRITING SUPPORT	£2,150 †	£2,650 †	£3,500 †

RATE CARD SPONSOR

VERTICAL SPONSORS	RARE REV-INAR	RARE DISEASE DAY
FROM £3,500 P/A †	FROM £6,500 †	POA

Why RARE Revolution™?

- Award winning team and publication
- Trusted within the rare disease community
- Full, in-house turnkey service from concept to outputs
- Full support from project design to dissemination
- Experiences and up-to-date on global compliance requirements
- Global audience with ability to geo-lock content
- Trusted vendor to some of the biggest pharma/biotech companies



Johnson & Johnson

ALEXION®
AstraZeneca Rare Disease



illumina®

UCB
Inspired by patients.
Driven by science.

ARDEA
OUTCOMES

THE HEALTH NAUTS
Navigate Life Sciences

mendelian

blueprint™
MEDICINES

CSL Behring



MEDISTRAVA
Inizio Medical

IPSEN

iNIZIO
MEDICAL

HORIZON

MERCK

ultragenyx
pharmaceutical

biocryst

PTC
THERAPEUTICS

TREND Community
Turning Anecdotes Into Evidence™

Alnylam®
PHARMACEUTICALS

KIELO
RESEARCH

sanofi

CSL

HEALTH
LUMEN

OPEN HEALTH

AHF
AMERICAS
HEALTH
FOUNDATION

SAVENTIC
Health

RARE
PATIENT
Voice™

CSL Vifor

SIMONS
SEARCHLIGHT

Phormulate

coave
THERAPEUTICS

AscellaHealth.
A RARE AND SPECIAL PERSPECTIVE™

TMC

Pharming®

Chiesi
global rare diseases

VIRIDIAN
PM TRIALS
QUALITY MOBILE RESEARCH

Testimonials

“

I just wanted to extend a heartfelt appreciation for all you've done to bring the special Systemic Mastocytosis (SM) edition to life. You've not only elevated awareness—you've made people living with SM feel seen. That's no small feat. The feedback from the international patient community speaks volumes, and internally, this issue has had a profound impact. It's not often that a publication manages to strike both emotional and strategic chords so clearly, but you've done just that. Thank you for being such thoughtful partners in this work. We are truly grateful—and excited for what's ahead.

CRISTIAN PEREZ, BLUEPRINT MEDICINES

“

I'm excited for the potential impact this issue may have for the undiagnosed, ultra-rare and rare disease communities. Additionally, I hope this issue illuminates the challenges and opportunities for researchers, industry partners and other stakeholders. Thanks again for the opportunity to share my voice. I appreciate all that you and your team does to inform, educate and advocate for the rare disease community.

MARY MORLINO, ADVOCATE

“

If everyone in the world even came close to having the integrity and passion you both have, we would live in Eden! THANK YOU for your partnership, but more importantly, for who you are in this world!! It's been such a pleasure for both Laura and I, and I very much hope to connect and partner again soon.”

MARLENI ARVELO-SAILLANT, SANOFI

“

“It was an honour for me to serve on the panel with such a distinguished group. Thank you to the RARE Revolution team for highlighting this important topic.”

YAEL WEISS, CEO, MAHZI THERAPEUTICS

Meet the Team

GET TO KNOW US



Rebecca Stewart
CEO



Nicola Miller
*Editor-in-chief
and creative
director*



Becky Pender
*Senior
associate -
partnerships and
delivery*



Emma Bishop
*Design and
editorial
associate*



Joe Rumney
*Creative
designer*

rarerevolutionmagazine.com/rare-revolutionaries/meet-the-team

RARE REVOLUTION™



There are [350 million people](#) worldwide living with a [rare disease](#). The zebra is the rare disease mascot.

[It's time to start talking about the zebra in the room.](#)

RARE Revolution Magazine™ is the brainchild of [Nicola Miller](#) and [Rebecca Stewart](#). They came together in 2012 to form Action for XP, a registered charity dedicated to supporting patients and families affected by the rare genetic disorder Xeroderma Pigmentosum (XP), after Nicola's son was diagnosed with the condition aged just 13 months old.

Frustrated with the lack of opportunities available to promote rare disease and source accessible reliable information, they decided to embark on providing a new platform for all rare disease champions to be heard. Thus the #RARERevolution movement was born...

RARE Revolution Magazine™ has gone on to become Scottish Special Interest of the year finalist 2018 and Media NPO of the Year by the Corporate LiveWire Innovation and Excellence Awards 2023 and 2024.

RARE Revolution Publishing Ltd. is a publishing company specialising in rare disease content that creates, produces and distributes RARE Revolution Magazine™, RARE Revolution insider™ the RARE Youth Revolution, RARE Revolutionaries Network Community and TIDE (Together In Driving Excellence – which supports patient engagement initiatives through tailored digital communications strategies and managed advisory boards).

RARE REVOLUTION™

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Are you ready to turn the tide? Discuss your bespoke requirements by booking a discovery call with Rebecca via the QR code or
<https://calendly.com/rstewart-rarerevolutionmagazine/discovery-call>

