

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 5 million people a month globally: 44% USA, 30% Europe, 26% rest of world.



1 million +

Monthly website traffic



4.4 million

Monthly social media reach



35k readers

Monthly magazine audience

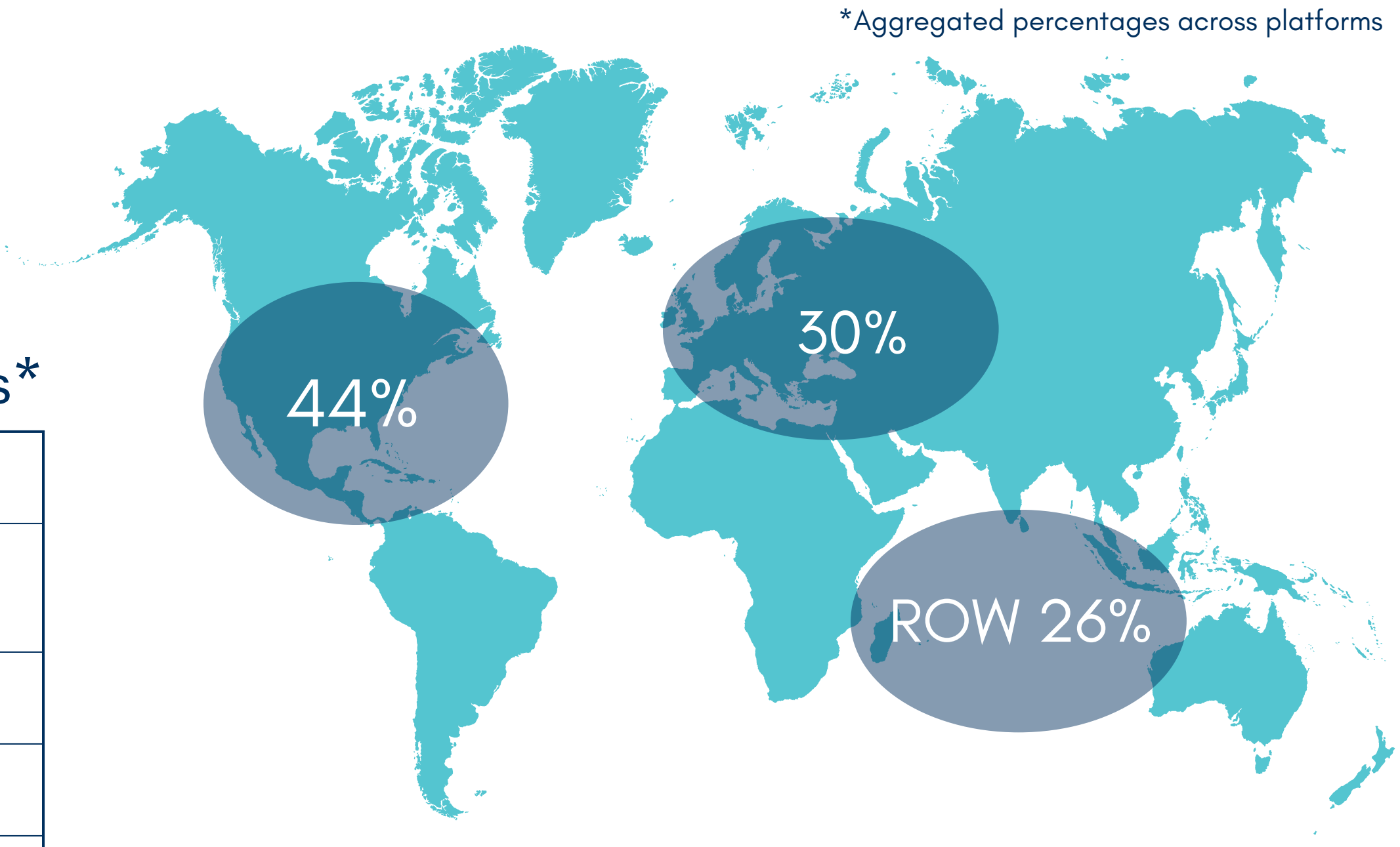


11k subscribers

3 weekly newsletters

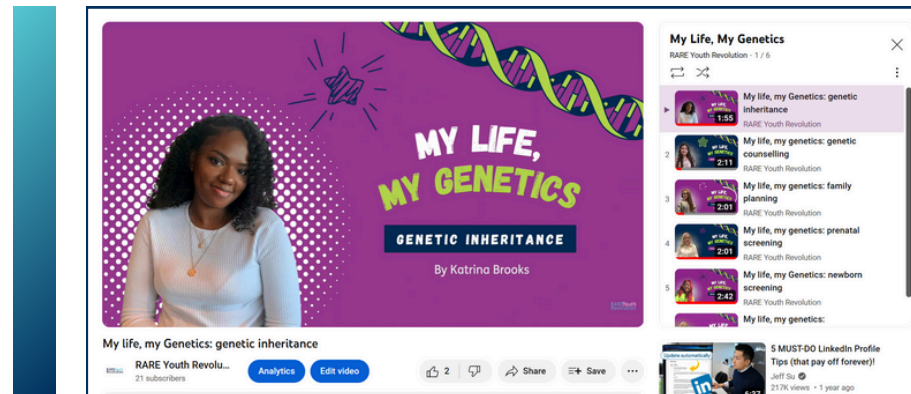
Demographics*

Industry	40.3%
Individuals and caregivers	23.2%
General interest	14.8%
HCP	12%
Charities	9.7%



LISTEN

Advisory boards



Landscape/disease mapping

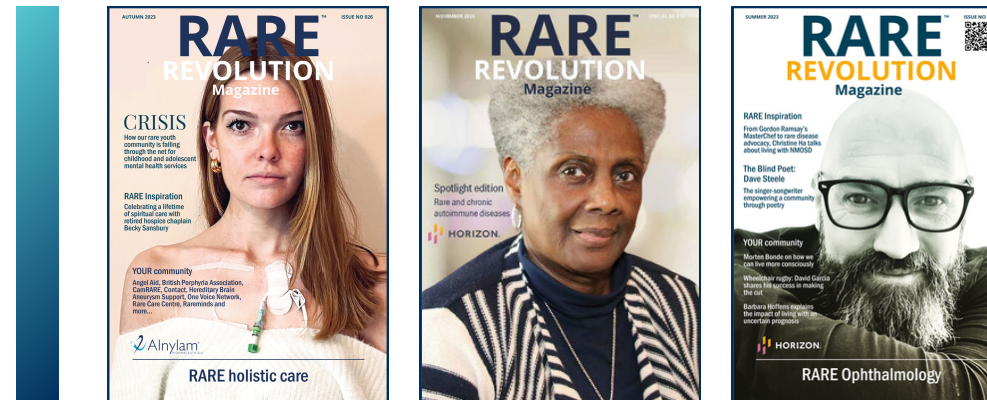


Roundtables



CREATE

Magazines



Impact reports



RARE recruitment



EDUCATE

RARE REV-inars



Digital spotlights



Thought leadership



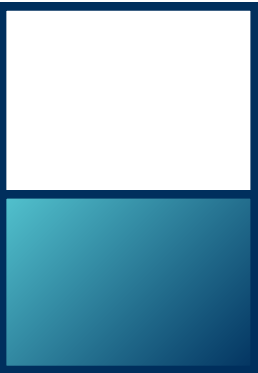
RARETM REVOLUTION MAGAZINE

RARE Revolution MagazineTM is a world leading rare disease publication dedicated to elevating the voice of the rare disease ecosystem through our website, digital magazine, 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 †



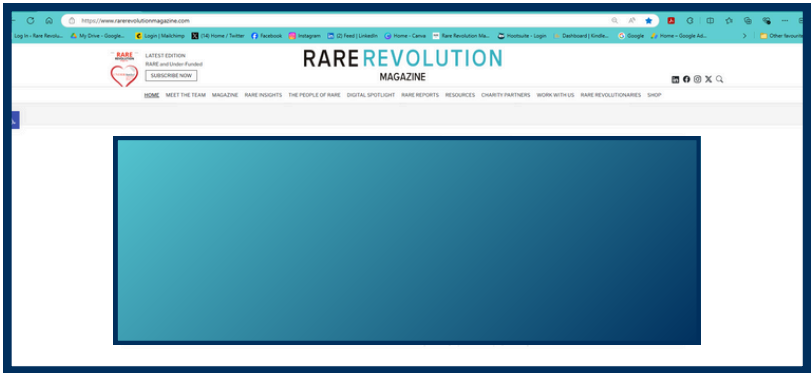
Half page advert
A5: 210 148.5 mm
£510 †



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 †



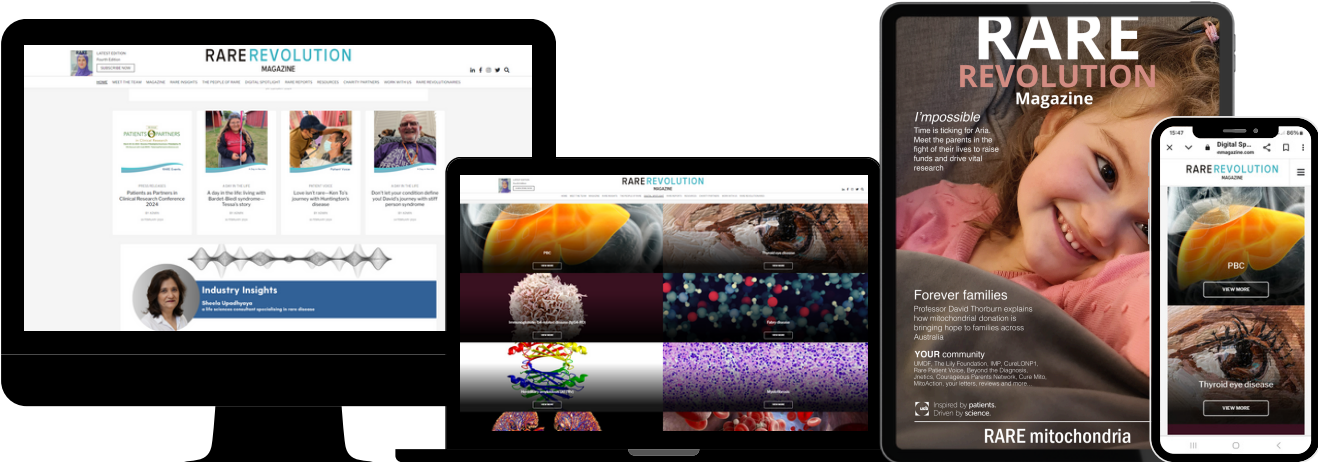
Home page web banner
1650 x 620 px
£650 †

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
£25,500 †	£36,000 †	FROM £6,500 †	FROM £11,500 †	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

BILLBOARD	SIGN UP PAGE	£1,800 PCM †
	TOP OF MAIN MENU	£1,800 PCM †
LEADERBOARD	UPPER	£1,500 PCM †
	MIDDLE	£1,250 PCM †
	LOWER	£995 PCM †

EMAIL ADVERTISING

SOLO BLAST	£1,495 †
EMAIL ADVERT	£995 †

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 ip-to-date on global compliance requirements
- Global audience with ability to geo-lock content
- Trusted vendor to some of the biggest pharma/biotech companies



Testimonials

“

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

“

I have read everything plus a few of the other articles in your Summer edition. So beautiful! You are such a compassionate journalist. Everything so perfectly written and presented. Tears rolling out of my now healed eyes. Thank you, specifically for your incredible work on the two TED articles.”

CHRISTINE GUSTAFSON, PATIENT CONTRIBUTOR

“

So proud to be working with such an amazing team as RARE Revolution. Thank you for your collaboration and support.”

ROBERT MITCHELL-THAIN, CEO, THE PBC FOUNDATION

“

“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
*Community
engagement
senior
associate*



Emma Bishop
*Special editions
editor 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 (formerly Teddington Trust), 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).

RAREREVOLUTION™

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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>

