Applying the Phenotype Approach for Rosacea to Practice and Research

J Tan
M Berg
R L Gallo
James Q. Del Rosso
Touro College, jqdelrosso@yahoo.com

Follow this and additional works at: https://touroscholar.touro.edu/tuncom_pubs

Part of the Dermatology Commons

Recommended Citation

This Article is brought to you for free and open access by the College of Osteopathic Medicine at Touro Scholar. It has been accepted for inclusion in College of Osteopathic Medicine (TUN) Publications and Research by an authorized administrator of Touro Scholar. For more information, please contact touro.scholar@touro.edu.
Applying the phenotype approach for rosacea to practice and research*

J. Tan,1 M. Berg,2 R.L. Gallo3 and J.Q. Del Rosso4

1Department of Medicine, Western University, Windsor, ON, Canada
2Department of Dermatology, Uppsala University, Uppsala, Sweden
3Department of Dermatology, University of California San Diego, La Jolla, CA, U.S.A.
4Department of Dermatology, Touro University Nevada, Henderson, NV, U.S.A.


Correspondence
Jerry Tan.
E-mail: jerrytan@bellnet.ca

Accepted for publication
16 May 2018

Funding sources
Funding for medical writing and editorial assistance was provided by Galderma S.A. (Lausanne, Switzerland). The company was not involved in the review of the manuscript and had no influence over any opinions or recommendations expressed herein.

Conflicts of interest
J.T. has served as an advisor, consultant, clinical investigator and/or speaker for Allergan, Bayer, Galderma and Valeant. M.B. has no conflicts of interest to declare. R.G. has served as a consultant for Sente, as well as for MatriSys, of which he is also co-founder. J.D.R. has served as an advisor, consultant, clinical investigator and/or speaker for Allergan, Almirall/Aqua, Bayer, BioPharmX, Fournix and Galderma.

*Plain language summary available online

DOI 10.1111/bjd.16815

Summary

Background Rosacea diagnosis and classification have evolved since the 2002 National Rosacea Society expert panel subtype approach. Several working groups are now aligned to a more patient-centric phenotype approach, based on an individual’s presenting signs and symptoms. However, subtyping is still commonplace across the field and an integrated strategy is required to ensure widespread progression to the phenotype approach.

Objectives To provide practical recommendations that facilitate adoption of a phenotype approach across the rosacea field.

Methods A review of the literature and consolidation of rosacea expert experience.

Results We identify challenges to implementing a phenotype approach in rosacea and offer practical recommendations to overcome them across clinical practice, interventional research, epidemiological research and basic science.

Conclusions These practical recommendations are intended to indicate the next steps in the progression from subtyping to a phenotype approach in rosacea, with the goals of improving our understanding of the disease, facilitating treatment developments and ultimately improving care for patients with rosacea.

What’s already known about this topic?
- Rosacea diagnosis and classification have evolved from a subtype to a phenotype approach.
- Adoption of the phenotype approach has begun, but more widespread adoption and support across the field are required to ensure a complete transition.

What does this study add?
- We offer practical guidance for clinical practice, interventional and epidemiological research, and basic science, to help overcome challenges and facilitate comprehensive uptake of the phenotype approach in rosacea.

Rosacea diagnosis and classification have evolved. In 2002, the National Rosacea Society (NRS) expert panel proposed diagnostic criteria for rosacea and an associated subtype-based classification that grouped the most common presentations according to disease features.1 These diagnostic criteria and the subtype classification have guided the majority of subsequent publications on rosacea diagnosis and treatment.2–8 However, progress in rosacea research over the last decade has revealed important limitations of this approach.9 As a consequence, three working groups [the American Acne and Rosacea Society (AARS), the global rosacea consensus expert panel (ROSCO) and the NRS] have formalized what many dermatologists already do in clinical practice and updated the methodology for diagnosis based on phenotypes of the disease.9–14
The recommendations for a phenotype approach to rosacea, where the disease is diagnosed and classified according to an individual’s presenting features instead of those grouped by subtypes, acknowledge the limitations of subtyping, and build on previous recommendations to propose an approach based on an individual patient’s presenting signs and symptoms. Table 1 shows a comparison of rosacea according to the NRS 2002 subtype classification and the aligned 2017 ROSCO and revised 2017 NRS phenotype recommendations. The features listed under the 2017 system can be combined into any presentation, provided the diagnostic criteria are met. Conversely, the previous 2002 subtype system combined multiple features into the erythematotelangiectatic, papulopustular, phymatous and ocular subtypes.11,13

Elements of a phenotype approach have been incorporated into recent national treatment recommendations, which advocate targeted treatment of individual rosacea features.15–18 However, subtyping is still commonplace in numerous recent publications.19,20 In order to promote wider implementation of a phenotype approach, it is important first to identify challenges preventing this transition and then to develop strategies to address those challenges.21 Common challenges to implementation include: poor support and organizational structures within the healthcare system; insufficient knowledge, awareness or skill; lack of resources including time; and uncertainty over the legitimacy of new recommendations.22 In the present publication, we identify challenges to implementation of a phenotype approach across clinical practice, interventional research, epidemiological research and basic science in rosacea. We then make practical recommendations to facilitate assessment, management and research progression inclusive of the individual experience of disease.

Table 1 Summary of recommendations for rosacea diagnosis, according to the 2002 subtype and 2017 phenotype approaches

<table>
<thead>
<tr>
<th>NRS 2002</th>
<th>ROSCO 2017/NRS 2017</th>
</tr>
</thead>
<tbody>
<tr>
<td>Primary (diagnostic) features</td>
<td>Flushing/transient erythema</td>
</tr>
<tr>
<td></td>
<td>Nontransient/persistent erythema</td>
</tr>
<tr>
<td></td>
<td>Papules and pustules</td>
</tr>
<tr>
<td></td>
<td>Telangiectasia</td>
</tr>
<tr>
<td>Secondary features</td>
<td>Burning/stinging sensation</td>
</tr>
<tr>
<td></td>
<td>Oedema</td>
</tr>
<tr>
<td></td>
<td>Dry appearance</td>
</tr>
<tr>
<td></td>
<td>Ocular manifestations</td>
</tr>
<tr>
<td></td>
<td>Plaque</td>
</tr>
<tr>
<td>Diagnostic features</td>
<td>Persistent centrofacial erythema with periodic intensification</td>
</tr>
<tr>
<td></td>
<td>by potential trigger factors</td>
</tr>
<tr>
<td></td>
<td>Phymatous changes</td>
</tr>
<tr>
<td>Major features</td>
<td>Transient centrofacial erythema</td>
</tr>
<tr>
<td></td>
<td>Inflammatory papules/pustules</td>
</tr>
<tr>
<td></td>
<td>Telangiectasia</td>
</tr>
<tr>
<td></td>
<td>Ocular manifestations</td>
</tr>
<tr>
<td>Minor/secondary features</td>
<td>Burning/stinging sensation</td>
</tr>
<tr>
<td></td>
<td>Oedema</td>
</tr>
<tr>
<td></td>
<td>Dry sensation/appearance</td>
</tr>
</tbody>
</table>

According to the 2002 subtyping system, multiple features were grouped into defined subtypes. The 2017 phenotyping system allows various combinations of diagnostic, major and minor features into an individual phenotype. NRS, National Rosacea Society; ROSCO, global ROSacea Consensus panel. The presence of one or more of these with a central facial distribution is indicative of rosacea. Individually diagnostic. A combination of at least two is diagnostic, in the absence of a diagnostic feature. Individual publications also discuss the importance of proper history taking, exclusion criteria and trigger factors for specific features.

Challenges

Fifteen years’ usage of the NRS 2002 rosacea diagnostic criteria and subtyping in textbooks, academic papers and podium presentations has engrained these concepts in dermatology. There is a need for clinicians, researchers, authors and scientific reviewers to be apprised of current modifications. Until then, the subtype concept with its shortcomings will persist. Inconsistent nomenclature is a field-wide issue that confounds the reporting of numerous studies. The terms ‘rosacea’ and ‘acne rosacea’ are used variably and often nonspecifically, as are definitions of certain features (e.g. persistent and transient erythema can be confounded with perilesional inflammation). There is a need for better differentiation in research studies of the phenotype of the patients being studied. The challenges in interventional research mostly centre on existing conventions and protocols. As the transition towards a phenotype approach is a recent development, little documentation exists to demonstrate its benefits and convince stakeholders of its value. In addition, older treatments assessed according to subtype in previous literature (e.g. Cochrane systematic reviews) will be disqualified, while new, smaller studies may be required to show an effect of a therapy, either alone or in combination, on specific features.

Industry sponsors of interventional research using subtype-based measures may be reluctant to replace these with outcomes for individual features, unless the value has already
been demonstrated. We believe that the phenotype approach allows investigators to assign outcomes for one or more rosacea features best addressed by investigational treatments, compared with multiple features in some existing scales. Standardizing these outcome measures would also facilitate interstudy comparisons and combining of results. Furthermore, improved severity scales that focus on individual features should be more accurate than those in current usage, thereby offering the potential for a smaller trial population to demonstrate significance. This could translate to lower costs of trials.

Study protocols are a further challenge for adoption of a phenotype approach in interventional and epidemiological research, due to the engrained subtype-based nomenclature. Some researchers accept implicitly that a singular examination predicates the features of an individual patient with rosacea; this fails to account for fluctuation of some features over time (e.g. papules/pustules or erythema with phyma) and may provide an incomplete disease representation for the patient. This is also hindered by the lack of long-term epidemiological studies in the field. Furthermore, darker skin pigment phenotypes are under-represented in epidemiological studies, due to difficulties in diagnosis that often fail to account for the presence of rosacea in these patients.

In basic science, several specific factors also hinder the progression to a phenotype approach to rosacea. These include limited funding opportunities, absence of patient registries or clinical tissue biobanks, and the lack of a satisfactory animal model for rosacea.

Results

Challenges and recommendations for clinical practice

A phenotype approach to rosacea management could improve patient outcomes by targeting the aspects most bothersome to the patient.\textsuperscript{12} It also facilitates use of combination therapy when necessary, to address multiple presenting features and optimize clinical outcomes and quality of life.\textsuperscript{12} Rosacea treatment is moving towards such an approach, which is now recommended by several regional and national bodies in clinical practice.\textsuperscript{15–18} However, further support is needed to facilitate its implementation into daily practice.

Assessment of disease severity and patient impact (including quality of life) would also benefit from a phenotype approach. Several groups have commented that existing scales for assessing these are not sufficient, including the only rosacea-specific tool (Rosacea Quality of Life; RosQoL), which both omits certain features and lacks indication of a clinically relevant difference. As such, there is a need for validated tools that are easy to use in the clinic and address all skin pigment types.\textsuperscript{9,11,13,23}

Recommendations

1. Update the language surrounding rosacea diagnosis and classification to discontinue the concept of subtyping.

2. Initiate an international, multisponsor core meeting of stakeholders to develop a communications plan for the phenotype concept.

3. Educate patients and healthcare providers to raise awareness and improve understanding of the phenotype nomenclature.

4. Demonstrate the proven clinical value of making the change to phenotyping and reinforce the shortcomings of the subtype approach.

5. Develop simple clinical tools for physicians to use with their patients that are applicable to all skin pigment types.

Challenges and recommendations for interventional research

Assessment methodologies used in rosacea clinical trials are variable and could be of higher quality.\textsuperscript{19,23} The subtype approach is likely to hinder treatment progress in two ways. Firstly, a treatment directed at one dimension of a subtype (e.g. inflammatory papules or pustules in papulopustular rosacea) may fail in clinical trials because it does not address the subtype’s other dimensions (i.e. persistent erythema).\textsuperscript{11} Secondly, the multiple dimensions of a subtype can confound commonly used rosacea severity scales, as with clinical practice. In clinical trials, this can interfere with study and assessment of the course of singular disease features.\textsuperscript{9}
Many clinical trials in rosacea still recruit patients using subtype-based inclusion criteria and assess treatment outcomes according to those subtypes (for example, using the NRS Rosacea Clinical Scorecard). However, although some more recent studies have still recruited patients by subtype, they have assessed therapeutic response by change in individual features. This is a positive step towards a phenotype approach.

Recommendations

1. Establish credible, standardized clinical and patient-reported measures for the most common features to make them acceptable to researchers, industry and regulatory authorities.

   → Industry may be especially interested in moving forward new drug entities to address specific signs and symptoms instead of the multiple dimensions required with subtypes.

2. Develop and publish real-life examples of phenotype use in interventional research, to validate the approach and demonstrate best practice throughout the field.

   → Could be based on the five-part AARS series, suggesting that a phenotype evaluation of rosacea correlates with therapy.

   → Enhance with the support of a larger group of interested researchers and clinicians.

Challenges and recommendations for epidemiological research

Epidemiological studies in rosacea provide widely varying disease prevalence estimates. Beyond valid differences relating to location and population differences, this variation may be a result of differences in case determination and study design, and variable assessment of individual features due to differences in skin pigment type.

Characterizing by subtype confounds epidemiological studies because many patients present with varied features, so are inadequately described. For example, there is no subtype for a patient presenting solely with persistent centrofacial erythema without transient erythema and flushing, or vice versa. Similarly, a patient presenting with fixed erythema, inflammatory papules and pustules and telangiectasia could fall into both the erythematotelangiectatic and papulopustular categories, and so might be counted twice. Furthermore, certain features can spontaneously remit and recur, which may result in the patient being classified in multiple subtype categories dependent on the time of observation.

Refining diagnostic criteria based on a phenotype approach could improve the consistency and quality of epidemiological data, which would continue to benefit rosacea diagnosis and overall disease management.

Recommendations

1. Facilitate longitudinal epidemiological studies in rosacea that consider the potential for fluctuation of certain features.

   → There would be great value in extension to a global registry, which would provide longitudinal real-world evidence to bolster epidemiological data and a phenotype treatment approach.

2. Include and support regional groups from all continents in epidemiological studies, to ensure representation of all skin pigment types.

Challenges and recommendations for basic science

Despite ongoing research, the pathophysiology of rosacea remains uncertain. Cutaneous rosacea subtypes demonstrate both differences and overlap in histological, immunohistological and gene expression markers. As such, there is limited information on molecular markers responsible for the initiation and perpetuation of rosacea’s various cutaneous features. Therefore, subtype classification may be hindering biomarker identification for individual features.

Recommendations

1. Work towards highlighting the need for national initiatives to support research in rosacea.

   → For example, the National Institutes for Health Request for Applications.

2. Extend clinical practice updates that emphasize phenotype nomenclature to basic science.

   → Would be of similar benefit in standardizing assessments.

3. Reinforce the concept of phenotype variation over time, as with epidemiological research.

   → Would encourage specimen collections at multiple time points to assess the chronic nature of the disease.

4. Interrogate existing data to unmask key specific factors in rosacea pathophysiology and identify which treatments have the largest or fastest impact on disease.

Discussion

Herein we have discussed strategies to support the phenotype approach in rosacea across clinical practice, interventional and epidemiological research, and basic science. Several strategies span multiple areas, which reflects the need for an integrated approach to ensure consistent and comprehensive uptake.
Audit and feedback lead to small but potentially important improvements in professional practice. It is therefore important to measure the effectiveness of the phenotype approach as it becomes more widely used, which will both enable sharing of successes to promote a virtuous circle of uptake and identify areas for improvement. In studies assessing the effectiveness of clinical guideline dissemination and implementation, the most widely used strategies for measuring change were audit and feedback; however, there is still significant work to be done to ensure a comprehensive uptake. These practical recommendations are intended to indicate the next steps in this process, with the goals of improving our understanding of the disease, facilitating treatment developments, and ultimately improving care for patients with rosacea.

Furthermore, even given an objective tool, the severity of some signs can be subjective; for example, some cultures may consider persistent mild centrofacial erythema presenting as ‘rosy cheeks’ to be a normal or healthy variation. There are also limitations in the use of persistent centrofacial erythema as a definitive diagnostic feature in dark skin phototypes, which will require specific guidance from future work to ensure consistent diagnosis and classification across the patient spectrum.

The transition to a phenotype approach in rosacea is underway; however, there is still significant work to be done to ensure a comprehensive uptake. These practical recommendations are intended to indicate the next steps in this process, with the goals of improving our understanding of the disease, facilitating treatment developments, and ultimately improving care for patients with rosacea.

### Acknowledgments

Medical writing and editorial assistance was provided by Ellie Hughes, PhD; Victoria Smith, BSc (Hons) and Holly Gilbert-Jones, PhD, of Ogilvy Healthworld, London, U.K.
References


Supporting Information

Additional Supporting Information may be found in the online version of this article at the publisher’s website:

Appendix S1 Rosacea phenotype checklist.

Powerpoint S1 Journal Club Slide Set.

Video S1 Author video.