

Community pharmacists optimising treatment for patients with psoriasis

Submission date 22/05/2014	Recruitment status No longer recruiting	<input checked="" type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 22/05/2014	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
Last Edited 16/03/2017	Condition category Skin and Connective Tissue Diseases	<input type="checkbox"/> Individual participant data

Plain English summary of protocol

Background and study aims

Psoriasis is a long-term inflammatory skin condition that affects up to 2% of the people in the UK. There is no cure for psoriasis and the condition tends to come and go but once it has developed it is normally life-long. The majority of people with psoriasis need to use creams or ointments on a daily basis. This requires a high level of commitment yet research shows that many people don't get enough advice from anyone on how to use them. As a result, psoriasis is often poorly managed by patients. In this study we want to see if advice given by community pharmacists can lead to better management of psoriasis and ultimately improvements in the severity of the condition.

Who can participate?

The study is open to anyone over the age of 18 who is prescribed creams or ointments to manage their psoriasis.

What does the study involve?

Patients who visit participating community pharmacies to collect their prescriptions for topical psoriasis treatments will be invited by the pharmacist or a member of staff to join the study. The study involves two consultations with the pharmacist. At the first appointment the pharmacist will use a specially designed guide to ask the person several questions to assess their understanding of psoriasis and if they know how to use their treatments. The pharmacist will also ask the person to complete two further questionnaires; one of these looks at the impact of psoriasis on their quality of life and the second asks them to rate the severity of their psoriasis. Both of these questionnaires have been used previously in research studies and are very quick and easy to complete. During the initial appointment the pharmacist will provide advice to help the person use their treatments properly. A second appointment will then be arranged for about 4 weeks later to go through the same initial questions to see if the person now has a better understanding of what to do with their treatments. The two questionnaires will also be repeated to see if the person's quality of life is improved and if they think their psoriasis is less severe.

What are the possible benefits and risks to participants?

The benefits for the patients are that they should have a better understanding of their condition and how to manage it. It is also likely that improvements in both quality of life and disease

severity will occur once the person knows more about how to use treatments properly. There are unlikely to be any risks associated with the study.

Where is the study run from?

The study will be run from participating pharmacies and is being co-ordinated by Robert Gordon University, Aberdeen.

When is the study starting and how long it is expected to run for?

We hope to start the study in September 2014 and it will continue until we have recruited and completed two interviews with 30 patients.

Who is funding the study?

The Psoriasis and Psoriatic Arthritis Alliance (UK).

Who is the main contact?

Dr Rod Tucker

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Contact information

Type(s)

Scientific

Contact name

Dr Rod Tucker

Contact details

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

16412

Study information

Scientific Title

Is it feasible for community pharmacists to deliver an educational programme designed to help people with psoriasis self-manage as effectively as possible?

Study objectives

This study attempts to explore whether advice provided by community pharmacists to patients with mild to moderate psoriasis can lead to an improvement in their ability to self-manage their psoriasis.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Office for Research and Ethics Committees Northern Ireland, 16/12/2013, ref. 13/NI/0207

Study design

Non-randomised Interventional Design type: Process of Care

Primary study design

Interventional

Secondary study design

Non randomised controlled trial

Study setting(s)

Other

Study type(s)

Quality of life

Participant information sheet

Not available in web format, please use the contact details below to request a patient information sheet

Health condition(s) or problem(s) studied

Topic: Primary Care; Subtopic: Not Assigned; Disease: All Diseases

Interventions

The pharmacists will opportunistically approach patients who present with prescriptions for themselves for psoriasis. Eligible patients who agree to participate will be asked to complete the SAPASI and DLQI forms and will have a private consultation with the pharmacist to discuss the PEDESI tool. The tool will identify and gaps in patients knowledge and understanding of their condition and its management. A follow-up appointment will be made approximately 4 weeks later to reassess the outcome measures and revisit the PEDESI tool to determine whether patients now feel more confident at self-managing their psoriasis.

Intervention Type

Other

Phase

Not Applicable

Primary outcome measure

Pedesi; Timepoint(s): at follow-up after 6 weeks

Secondary outcome measures

Dermatology quality of life index (DLQI); Timepoint(s): 6 weeks, at the follow-up appointment;
sapasi; Timepoint(s): 6 weeks, at follow-up appointment

Overall study start date

30/07/2014

Completion date

31/12/2014

Eligibility

Key inclusion criteria

1. Patients aged over 18 with a diagnosis of chronic plaque psoriasis currently prescribed topical treatment
2. Male & Female
3. Lower Age Limit: 18 no age limit or unit specified

Participant type(s)

Patient

Age group

Adult

Lower age limit

18 Years

Sex

Both

Target number of participants

Planned Sample Size: 30; UK Sample Size: 30;

Key exclusion criteria

Patients under 18 years, prescribed oral therapy or receiving phototherapy

Date of first enrolment

30/07/2014

Date of final enrolment

31/12/2014

Locations

Countries of recruitment

England

United Kingdom

Study participating centre
LLoyds Pharmacy
4 Market Place
Hornsea
United Kingdom
HU181AW

Sponsor information

Organisation
Robert Gordon University (UK)

Sponsor details
School of Health Sciences
Garthdee Road
Aberdeen
Scotland
United Kingdom
AB10 7QG

Sponsor type
University/education

ROR
<https://ror.org/04f0qj703>

Funder(s)

Funder type
Charity

Funder Name
The Psoriasis and Psoriatic Arthritis Alliance (PAPAA) (UK)

Results and Publications

Publication and dissemination plan
Not provided at time of registration.

Intention to publish date

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	01/04/2017		Yes	No
HRA research summary			28/06/2023	No	No