duration was 40.5±14.8 weeks. The reasons for discontinuation were: lack of effectiveness (n=1), treatment simplification (less-pills regimen) (n=3), abandonment (n=2), drug-drug interactions (n=2), kidney failure (n=1), death (n=1) and follow-up losses (n=2).

Conclusion and relevance A broad spectrum of DAT combinations were used according to patients’ characteristics. Although 14 patients were not at virological suppression at baseline, DAT showed a high durability at 48 weeks and only 2 patients discontinued due to lack of effectiveness or toxicity.

REFERENCES AND/OR ACKNOWLEDGEMENTS
Conflict of interest No conflict of interest

4CPS-177 A QUALITATIVE STUDY ON PHARMACIST PRESCRIBING FOR PATIENTS WITH CHRONIC KIDNEY DISEASE
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Background and importance Chronic kidney disease (CKD) has a high risk of morbidity and mortality. The available evidence worldwide demonstrates that non-medical prescribing by pharmacists in various clinical specialties is a safe and effective approach. There is lack of evidence on the implementation and development of pharmacist prescribing for patients with CKD.

Aim and objectives Aim was to explore the development, implementation and evaluation of pharmacist prescribing for patients with CKD in the UK.

Material and methods This study used a qualitative semi-structured interview. The development of the theory-based semi-structured interview tool followed a rigorous iterative process using findings from the literature, underpinned by the Consolidated Framework for Implementation Research (CFIR) and reviewed independently by an expert panel. A date/time for a telephone interview was arranged following receipt of signed consent. All interviews were transcribed verbatim. Interview data were analysed thematically. The Francis method of checking for data saturation was used. Ethical approval was granted by RGU School of Pharmacy.

Results Data saturation was reached after 14 interviews. Demographic details included: 11 female, 7 had >16 years experience in the profession, all had secondary care as their main practice setting and 8 had >11 years as a prescriber. The interviewees were generally very positive about their prescribing practice and they articulated that they were prescribing in a variety of settings. CFIR helped identify themes related to facilitators and barriers to advancing prescribing practice. There was enthusiasm for the future development of prescribing practice including further establishment of clinics and taking responsibility for groups of patients.

Conclusion and relevance This work provides information relating to the current status of the development of pharmacist prescribing practice in the UK. Further ‘deep dive’ case study work will help explore the practice of leading edge advanced and consultant level practitioners to learn even more about practice development.

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Conflict of interest No conflict of interest

4CPS-181 CLINICAL-PHARMACEUTICAL MEDICATION COUNSELLING FOR PNEUMOLOGICAL PATIENTS IN OUTPATIENT AND INPATIENT AREAS
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Background and importance This project focused on direct interaction between hospital pharmacists and patients, through clinical pharmaceutical counselling. Drug safety and patient well-being are promoted and possible user errors and uncertainties concerning the medication and interactions are communicated directly. During the consultation patients were given the opportunity to ask questions, express uncertainties and receive information regarding their medication. Two different settings for the project (outpatient and inpatient) allowed investigation of which questions were of primary concern in these different situations.

Aim and objectives Most chronic obstructive pulmonary disease (COPD) patients have up to at least three comorbidities in addition to their primary pneumological disease. These can affect the heart, bones, metabolism and/or psyche, among others. Therefore polymedication is almost inevitable. Clinical-pharmaceutical counselling of patients is intended to promote adherence and medication safety. Especially for these patients, adherence to therapy is crucial and close monitoring of the medication is essential. This patient-oriented service is intended to be a tool for optimal drug therapy, since it has already been shown that clinical-pharmaceutical interventions, such as the targeted education of patients, can reduce adverse drug events and readmissions.

Material and methods A guideline with predefined questions about health status, medication scheduling, intake modalities, uncertainties, and a final satisfaction survey was created in order to be able to offer a comparable consultation to all patients. Prior to the consultation, patient records were reviewed and the medication was checked for possible interactions using drug interaction software programs.

Results It could be shown that the consultation for the inpatient area was related to medication changes during the stay and the expected benefits from those changes, thus promoting the patients’ medication knowledge for the time after hospital discharge, whereas most outpatients’ questions were about self-medication and over the counter (OTC) drugs. In this cohort some examples of severe drug interactions could be found.

Conclusion and relevance In summary, this medication consultation benefits the patients by increasing their knowledge regarding their medicine which leads to better adherence and