

or because of known analgesic effects. In the small sample studied by Lee *et al*, baseline assessment did not detect significant depression; how effective treatment would be in a larger, more representative, group of patients with CPPS—in whom depression may be a more prominent feature—remains to be determined. Recruitment into adequately powered trials is likely to require a multicentre collaboration.

For many men, time, rather than current treatments, may have the greatest influence on symptoms, with around one third reporting resolution of symptoms after 1 year.³⁰ Overall management of CPPS (investigation and treatment) remains highly variable,³¹ and access to the most appropriate therapists (including pain experts, psychotherapists and, possibly, psychosexual therapists and physiotherapists) is often inadequate. The extent to which genitourinary medicine clinics can develop services for chronic conditions (other than HIV) may be limited by the strengthened focus on sexually acquired infections driven by epidemiological trends and the national sexual health strategy. At this point, the future for improving service provision for men with this often distressing and disabling condition seems very uncertain.

Sex Transm Infect 2005;**81**:96–97.
doi: 10.1136/sti.2004.012310

Correspondence to: G A Luzzi, Department of Genitourinary Medicine, Wycombe Hospital, High Wycombe HP11 2TT, UK; luzg@wycgu.demon.co.uk

REFERENCES

1 NIH Workshop on Chronic Prostatitis. Executive summary. (www.niddk.nih.gov/health/urolog/pubs/cpwork/cpwork.htm). Bethesda, MD: National Institutes of Health-National Institute of

Diabetes and Digestive and Kidney Diseases, 1995;appendix 1:1–5. (www.niddk.nih.gov/health/urolog/pubs/cpwork/cpwork.htm).

2 Krieger JN, Nyberg J, Nickel JC. NIH consensus definition and classification of prostatitis. *JAMA* 1999;**282**:236–7.

3 Litwin MS, McNaughton Collins M, Fowler FJ, *et al*. The National Institutes of Health chronic prostatitis symptom index: development and validation of a new outcome measure. *J Urol* 1999;**162**:369–75.

4 Krieger JN, Riley DE, Cheah PY, *et al*. Epidemiology of prostatitis: new evidence for a worldwide problem. *World J Urol* 2003;**21**:70–4.

5 Schaeffer AJ, Knauss JS, Landis JR, *et al*. Leukocyte and bacterial counts do not correlate with severity of symptoms in men with chronic prostatitis: the National Institutes of Health Chronic Prostatitis Cohort Study. *J Urol* 2002;**168**:1048–53.

6 Nickel JC, Alexander RB, Schaeffer AJ, *et al*. Leukocytes and bacteria in men with chronic prostatitis/chronic pelvic pain syndrome compared to asymptomatic controls. *J Urol* 2003;**170**:818–22.

7 Hellstrom WJG, Schmidt RA, Lue TF, *et al*. Neuromuscular dysfunction in nonbacterial prostatitis. *Urology* 1987;**30**:183–7.

8 Jang TL, Schaeffer AJ. The role of cytokines in prostatitis. *World J Urol* 2003;**21**:95–9.

9 Krieger JN, Riley DE. Bacteria in the chronic prostatitis-chronic pelvic pain syndrome: molecular approaches to critical research questions. *J Urol* 2002;**167**:2574–83.

10 Batstone GRD, Doble A, Gaston JSH. Autoimmune T cell responses to seminal plasma in chronic pelvic pain syndrome (CPPS). *Clin Exp Immunol* 2002;**128**:302–7.

11 Wesselman U. Neurogenic inflammation and chronic pelvic pain. *World J Urol* 2001;**19**:180–5.

12 Yang CC, Lee JC, Kromm BG, *et al*. Pain sensitization in male chronic pelvic pain syndrome: why are symptoms so difficult to treat? *J Urol* 2003;**170**:823–6.

13 Ehler U, Heim C, Hellhammer. Chronic pelvic pain as a somatoform disorder. *Psychother Psychosom* 1999;**68**:87–94.

14 Wessely S, Nimmuan C, Sharpe M. Functional somatic syndromes: one or many? *Lancet* 1999;**354**:936–9.

15 McNaughton Collins M, MacDonald R, Wilt TJ. Diagnosis and treatment of chronic abacterial prostatitis: a systematic review. *Ann Intern Med* 2000;**133**:367–81.

16 Nickel JC, Narayan P, McKay J, *et al*. Treatment of chronic prostatitis/chronic pelvic pain syndrome with tamsulosin: a randomized double blind trial. *J Urol* 2004;**171**:1594–7.

17 Mehik A, Alas P, Nickel JC, *et al*. Alfuzosin treatment for chronic prostatitis/chronic pelvic pain syndrome: a prospective, randomised, double-blind, placebo-controlled pilot study. *Urology* 2003;**62**:425–9.

18 Nickel JC, Pontari M, Moon T, *et al*. A randomized, placebo controlled, multicenter study to evaluate the safety and efficacy of rofecoxib in the treatment of chronic nonbacterial prostatitis. *J Urol* 2003;**169**:1401–5.

19 Dimitrakov JD, Tchitalov J, Dikov D. Efficacy and safety of corticosteroids in the treatment of CP/CPPS: a randomized, double-blind, placebo-controlled trial (abstract). *J Urol* 2004;**171**(Suppl):S61–2.

20 Shoskes DA, Zeitlin SI, Shahed A, *et al*. Quercetin in men with category III chronic prostatitis: a preliminary prospective double-blind, placebo-controlled trial. *Urology* 1999;**54**:960–3.

21 Nickel JC, Downey J, Pontari MA, *et al*. A randomized placebo-controlled multicentre study to evaluate the safety and efficacy of finasteride for male chronic pelvic pain syndrome (category IIIA chronic nonbacterial prostatitis). *BJU Int* 2004;**93**:991–5.

22 De Rose AF, Gallo F, Giglio M, *et al*. Role of mepartricin in category III chronic nonbacterial prostatitis/chronic pelvic pain syndrome: a randomized prospective placebo-controlled trial. *Urology* 2004;**63**:13–16.

23 John H, Ruedi C, Kotting S, *et al*. A new high frequency electrostimulation device to treat chronic prostatitis. *J Urol* 2003;**170**:1275–7.

24 Giannakopoulos X, Entezari K, Schulman C, *et al*. Transurethral needle ablation for chronic nonbacterial prostatitis: a 3-year follow-up study (abstract). *J Urol* 2004;**171**(Suppl):S62.

25 Chen R, Nickel JC. Acupuncture ameliorates symptoms in men with chronic prostatitis/chronic pelvic pain syndrome. *Urology* 2003;**61**:1156–9.

26 Cornel EB, van Haarst EP. Chronic pelvic pain syndrome type 3 successfully treated with biofeedback physical therapy. *J Urol* 2004;**171**(Suppl):S30.

27 De la Rosette JJMCH, Ruijgrok MCM, Jeuken JMG, *et al*. Personality variables involved in chronic prostatitis. *Urology* 1993;**42**:654–62.

28 Berghuis JP, Heiman JR, Rothman I, *et al*. Psychological and physical factors involved in chronic idiopathic prostatitis. *J Psychosom Res* 1996;**41**:313–25.

29 Jick H, Kaye JA, Jick SS. Antidepressants and the risk of suicidal behaviors. *JAMA* 2004;**292**:338–43.

30 Nickel JC, Downey JA, Nickel K, *et al*. Prostatitis-like symptoms: one year later. *BJU Int* 2002;**90**:678–81.

31 Luzzi GA, Bignell C, Mandal D, *et al*. Chronic prostatitis/chronic pelvic pain syndrome: national survey of Genitourinary Medicine clinics. *Int J STD AIDS* 2002;**13**:416–19.

LYMPHOGRANULOMA VENEREUM

Lymphogranuloma venereum in the United Kingdom

P French, C A Ison, N Macdonald

First cases reported from enhanced surveillance

Until 2003 lymphogranuloma venereum (LGV), a disease caused by the more invasive L serovars of *Chlamydia trachomatis*, was considered a rare disease outside resource poor countries. Since then it has emerged as a significant problem among men who

have sex with men (MSM) in Europe. In 2003 an outbreak of LGV was recognised in Rotterdam in the Netherlands.¹ More than 100 men have been reported in this outbreak, most of whom were HIV positive and many had concomitant sexually transmitted infec-

tions including hepatitis C infection. Although many reported unprotected anal sex as a risk factor for acquisition of LGV, fisting and the sharing of sex toys also appeared as possible routes of transmission. Almost all presented with proctitis and symptoms included rectal pain, discharge, tenesmus, and other signs of lower gastrointestinal inflammation including constipation and abdominal pain. Some reported systemic symptoms such as fever and malaise. Genital and inguinal symptoms were rare with only one patient presenting with inguinal lymphadenopathy. Since that report similar outbreaks have been recognised in Antwerp, Hamburg, and Paris.^{2–4} Cases have also been reported from Sweden and more recently from the United States (New York, San Francisco, and Atlanta).⁵ All

the reported cases have been caused by the L2 serovar, although there is some evidence that a number of genetically distinct strains of *C trachomatis* L2 are responsible for these outbreaks.⁶

In October 2004 the Health Protection Agency (HPA) sent out an alert to genitourinary medicine (GUM) clinicians in England and established a case definition, reference service, and reporting system for LGV.⁷ In addition to the information produced by the HPA, the Terence Higgins Trust produced briefings for use in clinics and a leaflet for use in gay venues to increase awareness. The case definition used by the HPA is confirmation of *C trachomatis* and presence of an LGV serovar, L1, L2, or L3, by genotyping. The HPA reference service will test rectal specimens from patients with anorectal symptoms (typically proctitis, rectal discharge) or urethral specimens from patients with inguinal lymphadenopathy that are known to be positive for *C trachomatis*. Serology for *C trachomatis* has been used in Europe and can suggest the possibility of LGV, but does not confirm cases because of a lack of specificity, and has not been used in England as part of the case definition (www.hpa.org.uk/infections/topics_az/hiv_and_sti/LGV/lgv.htm).

In January 2005 the first 24 cases of LGV were reported in the United Kingdom,^{8,9} most from London clinics. Enhanced surveillance data were available for 19 cases and confirmed a picture similar to that reported in the rest of Europe. All were MSM, 17 were HIV positive, four also had hepatitis C infection, and most had symptoms suggesting LGV. Fifteen patients reported a probable country of infection; five in mainland Europe and 10 in the United Kingdom. Up to the middle of February 2005 a total of 34 cases of LGV have been reported in the United Kingdom.

LGV presenting as proctitis in homosexual men is well recognised.¹⁰ The primary (papule/ulcer) of LGV frequently goes unnoticed and patients often present with acute haemorrhagic proctitis and may have pronounced systemic symptoms such as fever and weight loss. Proctoscopy often reveals

marked proctitis, which is usually confined to the distal 10 cm of the anorectal canal. Left untreated, chronic inflammation may lead to stricture and fistula formation as well as local lymphatic obstruction and lymphoedema.¹¹ Patients with acute proctitis related to LGV usually respond well to antibiotic therapy. At present the recommended treatment for LGV in the United Kingdom is either oral doxycycline 100 mg twice daily, or oral erythromycin 500 mg four times a day, both regimens given for 3 weeks.¹² Patients with chronic infection including abscess, fistulas, and strictures often require surgical intervention.

It is likely that LGV has been present for some time in MSM in the United Kingdom, with many cases going undiagnosed. The first UK case identified so far is from a retrospective sample dating from January 2004. The epidemiology and clinical features of LGV in MSM are not fully understood; it is likely that some undiagnosed cases will have progressed to invasive disease, while others may yet prove to be asymptomatic. Clearly, further collaborative research is required.

The first steps in understanding and controlling this outbreak are to increase community and clinician awareness of LGV, to further develop our surveillance system and to monitor clinical manifestations. A national incident team has been established to oversee responses with the aim of developing effective control measures for this outbreak. The key challenge will be to identify and implement appropriate health promotion and prevention measures, particularly addressing the sexual health needs of HIV positive homosexual men, and ensure that potentially severe sequelae of untreated LGV are minimised.

Sex Transm Infect 2005;**81**:97–98.
doi: 10.1136/sti.2005.015263

Authors' affiliations

P French, Mortimer Market Centre, Mortimer Market, and Camden Primary Care Trust and University College London, UK

C A Ison, N Macdonald, Health Protection Agency Centre for Infections, 61 Colindale Avenue, London NW9 5HT, UK

Correspondence to: Patrick French, Mortimer Market Centre, London, UK; PFrench@gum.ucl.ac.uk

A national LGV incident group has been established by the HPA in collaboration with the British Society for Sexual Health and HIV (BASHH), the Terence Higgins Trust (THT), and the Society for Sexual Health Advisers (SHAA) and is chaired by Helen Ward (helen.ward@hpa.org.uk).

Leaflet produced by Terence Higgins Trust. (Single copies can be obtained through THT Direct 0845 12 21 200; multiple copies by emailing james.glavin@tht.org.uk).

REFERENCES

- Götz H**, Nieuwenhuis R, Ossewaarde T, *et al*. Preliminary report of an outbreak of lymphogranuloma venereum in homosexual men in the Netherlands, with implications for other countries in western Europe. *Eurosurveillance Weekly* 2004;**8**(4):22 Jan 2004.
- Vandenbruaene M**. Uitbraak van Lymphogranuloma Venereum in Antwerpen en Rotterdam. *Epidemiologisch Bulletin van de Vlaamse Gemeenschap* 2004;**47**:4–6.
- Robert Koch-Institut**. Zum gehäuftem Auftreten von Lymphogranuloma Venereum in Hamburg im Jahr 2003. *Epidemiologisches Bulletin* 2004;**25**:18 June 2004.
- Institut de Veille Sanitaire**. Emergence de la Lymphogranulomatose vénérienne rectale en France: cas estimés au 31 mars 2004. Synthèse réalisée le 1er Juin, 2004.
- New York City Department of Health and Mental Hygiene**. Two New York City residents diagnosed with rare sexually transmitted infection; same strain found in Europe. Press release, 2 February 2005 (available on www.nyc.gov/html/doh/html/public/press05/pr011-05.html).
- Meyer T**, Arndt R, von Krosigk A, *et al*. Repeated detection of lymphogranuloma venereum caused by Chlamydia trachomatis L2 in homosexual men in Hamburg (letter). *Sex Transm Infect* 2005;**81**:91–4.
- Health Protection Agency**. Enhanced surveillance of lymphogranuloma venereum (LGV) in England. *Commun Dis Rep CDR Wkly* [serial online] 2004;**14**(41): News (available at www.hpa.org.uk/cdr/PDFfiles/2004/cdr4104.pdf).
- Health Protection Agency**. Initial results of enhanced surveillance for lymphogranuloma venereum (LGV) in England. *Commun Dis Rep CDR Wkly* [serial online] 2005;**15**(5): News (available at www.hpa.org.uk/cdr/archives/archive05/News/news0405.htm#botulism).
- Macdonald N**, Ison C, Martin I, *et al*. Initial results of enhanced surveillance for lymphogranuloma venereum (LGV) in England. *Eurosurveillance* 2005;**10**:20. January 2005 (available at www.eurosurveillance.org/ew/2005/050127.asp).
- Quinn TC**, Goodall SE, Mrtichian E, *et al*. Chlamydia trachomatis proctitis. *N Engl J Med* 1981;**305**:195–200.
- Mabey D**. Peeling. Lymphogranuloma venereum. *Sex Transm Infect* 2002;**78**:90–2.
- Clinical Effectiveness Group**. National guideline for the management of lymphogranuloma venereum. *Sex Transm Infect* 1999;**75**(Suppl 1):S40–2.