trols, are required to better define their pathogenic potential.

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Humans as Source of *Mycobacterium tuberculosis* Infection in Cattle, Spain

To the Editor: *Mycobacterium tuberculosis* is the main causative agent of tuberculosis in humans. However, little attention has been paid to its transmission from humans to animals. We report *M. tuberculosis* infections in 3 cattle farms in Spain. The epidemiologic investigation traced humans as the source of infection, with 1 of the strains showing multidrug resistance. Recent studies have reported isolation of *M. tuberculosis* in cattle with prevalences of 4.7%–30.8% in African and Asian countries (*1–3*). In cattle, this infection occurs in countries with the highest incidence of human tuberculosis in the world. In Europe, only 14 cases of *M. tuberculosis* infection have been described in 3 eastern countries since implementation of eradication programs (*4,5*). The only reported cases of *M. tuberculosis* in cattle in western Europe were described in Great Britain and date back to the 1950s (*6*).

During 2007-2009, three cases of tuberculosis caused by M. tuberculosis were detected in 3 unrelated cattle farms, 2 of them free of tuberculosis (farms 1 and 2). As part of the surveillance system of bovine tuberculosis, a pool of tissue samples from each cow (respiratory lymph nodes and lung) were homogenized with sterile distilled water, and culture was carried out by the BACTEC mycobacteria growth indicator tube 960 system (Beckton Dickinson, Madrid, Spain). Members of the M. tuberculosis complex were identified and genotyped by direct variable repeat spacer olignucleotide typing and mycobacterial interspersed repetitive unit-variable number tandem repeat (MIRU-VNTR) typing (7).

The 3 *M. tuberculosis*–infected animals were ≤ 9 months of age (Table). As described (6), the possibility of infection in young animals could be more probable than infection in older cows.

M. tuberculosis—infected animals from farms 1 and 3 were detected by the intradermal tuberculin test (Table). The animal without immunologic response (farm 2) was detected because an *M. bovis* infection was confirmed in the herd, and all animals were slaughtered. Confirmation of infection by culture without immunologic response is rare, although the high sensitivity of the mycobacteria growth indicator tube system could detect a low bacterial load in the initial stages

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of infection. Recent implementation of liquid systems in animal health laboratories has enabled detection of *M. tuberculosis* when it is compared with results using only conventional methods. Moreover, no tuberculosiscompatible lesions were observed in the 3 animals, similar to previous studies (6). On the basis of these facts, *M. tuberculosis* transmission was not detected among cattle in the following intradermal tuberculin tests.

Co-infection with other mycobacteria (M. avium subsp. hominissuis) was found in the same animal from farm 1 (Table). This co-infection suggested the immunocompromised status of the animal and hence a high susceptibility to M. tuberculosis infection. Moreover, M. bovis was isolated from 52% (16/31) of all animals from farm 2 that showed a positive reaction to the intradermal tuberculin skin test, making remarkable the absence of co-infection with M. bovis in the M. tuberculosis-infected animal. Therefore, the lack of M. tuberculosis transmission within this herd contrasts with the M. bovis dissemination.

The veterinary services reported these findings to the National Public Health System, and an epidemiologic investigation was conducted on the cattle farms to determine the source of infection. In all cases, staff of the farms had active tuberculosis (Table). Three different strains were characterized: SIT2537 (octal 77761777720771), code 253533233433236252211423 (farm 1); SIT1564, 3'5233423245545725121 3423 (farm 2); and SIT58, 254343 243232325262213423 (farm 3) (Table). The MIRU-VNTR pattern and spoligotype are shared by Spanish human and cattle isolates from farm 1; SIT2537 is an uncommon profile that has been detected in Brazil and Spain (according to the SITVIT2 database). The human strain showed multidrug resistance to isoniazid, rifampin, and ethionamide. In cattle and human isolates, genes associated with isoniazid and rifampin resistance were studied (8) and rpoB analysis confirmed rifampin resistance (Ser531Leu). In farm 2, the origin of the farm worker was eastern Europe and the cattle isolate showed an SIT1564 profile, which is found only in 6 human isolates in the SpolDB4 database, all from Poland, Bulgaria, and Russia. On farm 3, human and cattle isolates from Spain shared identical spoligotype and MIRU-VNTR patterns. The profile SIT58 is frequent in Spain (9) and other countries with historical links to Spain, mainly the south American countries (79/114 according to Spol-DB4).

A well-designed program for eradicating bovine tuberculosis helps to detect *M. tuberculosis* infection by immune response or by bacteriologic culture. The use of liquid systems and results of epidemiologic studies (Spanish Database of Animal Mycobacteriosis, mycoDB.es) (S. Rodríguez, unpub. data) are recommended for prompt confirmation of the M. tuberculosis complex infection and for enhancing the sensitivity of culture. In addition, the Spanish Ministry of Environment, Rural and Marine Affairs has reinforced the need to improve cooperation between human and animal health systems to minimize the risk for M. tuberculosis complex transmission from animals to humans or vice versa and to control infection in all susceptible animal species (10).

Variable	Farm 1	Farm 2	Farm 3
Cattle herd			
No. animals	6	54	>200
Production	Beef	Beef	Dairy
Previous status	TB free	TB-free	Non–TB free
No. reactors	1	31	12
M. tuberculosis infection in c	attle		
Year of isolation	2007	2008	2009
Age, mo	9	4	3
IDTB/interferon-γ	Pos/not determined	Neg/neg	Pos/not determined
TB-compatible lesion	No	No	No
Spoligotyping profile	SIT2537	SIT1564	SIT58
MIRU-VNTR profile†	253533233433236252211423	3'52334232455457251213423	254343243232325262213423
Co-infection with	Yes	Yes	No
other mycobacteria	(<i>M. avium</i> subsp. <i>hominissuis</i>)‡	(M. bovis)	
M. tuberculosis infection in h	uman		
Active tuberculosis	Yes	Yes	Yes
Spoligotyping profile	SIT2537	Not available	SIT58
MIRU-VNTR profile†	253533233433236252211423	3'52334232455457251213423	254343243232325262213423
Origin	Spain	Eastern Europe	Spain

*TB, tuberculosis; IDTB, intradermal tuberculin tested according to the European Council Directive 64/432/EEC; pos, positive; neg, negative; MIRU-VNTR, mycobacterial interspersed repetitive unit–variable number tandem repeat.

†MIRU-VNTR profile on the basis of the 24 MIRU-VNTR loci (7).

‡Co-infection in the same animal.

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Porcine and Human Community Reservoirs of *Enterococcus faecalis*, Denmark

To the Editor: Enterococcus faecalis, which exists commensally in the gut in warm-blooded animals and humans, is an opportunistic pathogen that causes a variety of communityacquired and health care–associated infections, such as urinary tract and intraabdominal infections, bacteremia, and endocarditis (1). Only a few studies have assessed the relationships between clinical *E. faecalis* strains; strains endemic to the health care setting; and community strains residing in humans, animals, or animal-origin food (2).

Recently we showed that the emergence of high-level gentamicinresistant (HLGR) *E. faecalis* among patients with infective endocarditis (IE) coincided with an increase in HLGR *E. faecalis* in the pig population in Denmark (3). The majority of isolates belonged to the same clonal group (sequence type [ST] 16), suggesting that pigs constitute a community reservoir of HLGR *E. faecalis*.