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Leprosy. Infliximab has been used¹ in the treatment of recurrent type 2 (erythema nodosum leprosum) lepra reactions (see Leprosy, p.176). However, 2 cases of rapidly progressive leprosy developing in patients given infliximab for rheumatoid arthritis have also been described;2 reversal (type 1) reactions occurred in both when infliximab was stopped.

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Psoriasis. Infliximab is used in the treatment of moderate to sewere plaque psoriasis (p.1583). ¹⁻⁸ In the UK, NICE recommends⁸ that it be reserved for severe cases, unresponsive to standard therapies (including ciclosporin, methotrexate, and PUVA) or where such therapies cannot be used.

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Rheumatoid arthritis. TNF inhibitors play an increasingly important role in the management of rheumatoid arthritis; they tend to be reserved for patients who are unresponsive to more conventional disease-modifying drugs, although some favour use earlier in management.

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Sarcoidosis. For a mention of possible benefit from infliximab in sarcoidosis, see p.1512.

Spondyloarthropathies. References to the use of infliximab in the treatment of ankylosing spondylitis and psoriatic arthritis (p.13). In the UK, NICE considers that TNF inhibitors should be reserved for severe active psoriatic arthritis unresponsive to at least 2 standard disease-modifying drugs; etanercept or adalimumab are preferred to infliximab.

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Uveitis. Infliximab has been tried with some success in the treatment of uveitis (p.1515) including that associated with Behçet's syndrome (p.1499). Uveitis can also develop as a complication of other inflammatory disorders such as rheumatoid arthritis: treatment with infliximab may improve ocular symptoms in addition to its effect on the primary disorder.

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Vasculitic syndromes. For a preliminary report on the use of infliximab in Takayasu's arteritis, see p.1514. Infliximab has also been investigated in the management of Kawasaki disease (p.2228) in patients who are unresponsive to standard treat-

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Preparations

Proprietary Preparations (details are given in Part 3)

Proprietary Preparations (details are given in Part 5)
Arg.: Remicade; Revelled; Austral: Remicade; Belg.: Remicade; Braz.:
Remicade; Canad.: Remicade; Chile: Remicade; Cz.: Remicade; Denn.:
Remicade; Fin.: Remicade; Fr.: Remicade; Ger.: Remicade; Ger.: Remicade; Hong; Remicade; Hung.: Remicade; Inc.: Remicade; Inc.:

Isonixin (HNN)

Isonixine; Isonixino; Isonixinum. 2-Hydroxy-N-(2,6-dimethylphenyl)nicotinamide

 $C_{14}H_{14}N_2O_2 = 242.3.$ CAS — 57021-61-1.

Isonixin is an NSAID (p.96) that has been used in the management of pain and inflammation associated with musculoskeletal and joint disorders. It has been used in doses of 400 mg two to four times daily by mouth or by rectal suppository. It has also been applied topically as a 2.5% cream.

Proprietary Preparations (details are given in Part 3)

Multi-ingredient: Spain: Nixyn

Kebuzone (rINN)

Kebuzona; Kébuzone; Kebuzonum; Ketophenylbutazone. 4-(3-Oxobutyl)-1,2-diphenylpyrazolidine-3,5-dione

 $C_{19}H_{18}N_2O_3 = 322.4.$ CAS — 853-34-9. ATC - MOTAAO6 ATC Vet - QM01AA06.

Kebuzone, a phenylbutazone derivative, is an NSAID (p.96). It has been given for musculoskeletal, joint, and soft-tissue disorders in oral doses of up to 1.5 g daily in divided doses. Kebuzone has also been given as the sodium salt by intramuscular injection in doses equivalent to 1 g of base once or twice daily.

Porphyria. Kebuzone is considered to be unsafe in patients with porphyria because it has been shown to be porphyrinogenic in animals or in-vitro systems.