Treatment Algorithm for Multiple Sclerosis Disease-modifying Therapies
(Please note the ordering of drugs within each category is alphabetical and not intended to indicate a hierarchy of treatment)

Presentation
- Clinically isolated syndrome without MRI abnormalities allowing the diagnosis of MS
- Clinically isolated syndrome & MRI activity, La McDonald MS
- RRMS: 1 relapse in last 2 years AND radiological activity
- RRMS: 2 significant relapses in last 2 years
- Rapidly evolving severe MS

First-line therapy
- No Treatment
- Interferon beta 1a and 1b (Extavia)
- Alemtuzumab or Ocrelizumab
- Teriflunomide

Alternative First-line therapy due to intolerance
- See note 9
- Interferon beta 1a and 1b (Extavia)
- Glatiramer acetate
- Teriflunomide
- Dimethyl fumarate

Second-line therapy. For disease activity whilst on first-line therapy
- Alemtuzumab

Highly active MS
- Alemtuzumab or Ocrelizumab
- Cladribine
- Fingolimod

Rescue therapy. For continued inflammatory activity whilst on second-line therapy
- No change
- Alemtuzumab or Ocrelizumab
- Cladribine
- Natalizumab
- Autologous hematopoietic stem cell transplant

Switch due to disease activity
Switch due to intolerance
Trials of first-line therapies in people with CIS at high risk of conversion have NOT shown a convincing long-term effect on the accumulation of disability. Therefore it is reasonable to opt for no treatment in many patients in this situation.

Under 2014 NHS England guidance, beta-interferon may be offered for patients within 12 months of a clinically significant clinically isolated syndrome when MRI evidence predicts a high likelihood of recurrent episodes (i.e. development of MS).

In exceptional circumstances, clinical or radiological markers may indicate a poor prognosis for rapidly developing permanent disability even after just one clinical episode, in which case, alemtuzumab (or ocrelizumab if alemtuzumab is contra indicated or otherwise not suitable) may be considered. Physicians and patients should weigh up the considerable risks and burden of monitoring associated with this drug against the potential benefit.

For RRMS (that is not RES), beta-interferon, glatiramer and teriflunomide are effective and safe.

There is some evidence that dimethyl fumarate may be more effective at suppressing relapses than beta-interferon, glatiramer and teriflunomide.

NHS England 2014 policy allowed the use of beta-interferon in “patients with only a single major relapse in the preceding two years, but combined with MRI evidence of continuing disease activity”. The NICE 2018 guidance suggests glatiramer acetate may also be used in this context.

For RRMS (that is not RES), alemtuzumab or ocrelizumab is an option that may be considered, but we note they are more high-risk than the other options. It should be used only when the patient and MS specialists accept the significant risks and burden of monitoring.

Cladribine and alemtuzumab (or ocrelizumab if alemtuzumab is contra indicated or otherwise not suitable) may be considered – by some patients and clinicians – a safer option than natalizumab when JC virus serology is high-index positive.

If a patient satisfies the eligibility criteria for a first-line therapy, and then becomes relapse-free on a drug to which he/she becomes intolerant, they may be switched to another DMT even though their relapses may now fall outside the eligibility window.

The risk of PML on natalizumab is likely to be increased after alemtuzumab, ocrelizumab or cladribine, given the prolonged lymphopenia induced by these drugs. But, where the patient is negative for JC serology, this may be appropriate.

NHS England 2014 policy states that fingolimod can be used as an alternative to natalizumab for those patients receiving natalizumab who are at high risk of developing progressive multifocal leukoencephalopathy (PML) as defined by: (i) JCV exposure indicated by anti-JCV antibody positive status, (ii) Receiving an immunosuppressant prior to receiving natalizumab, or (iii) Natalizumab treatment duration of >2 years. If patients develop a severe adverse effect to natalizumab (e.g. anaphylaxis), and they have not previously received fingolimod, then it may be appropriate to use fingolimod.

Definition of disease activity: treatment failure may be indicated by either clinical or radiological relapse-related changes, after significant exposure to the treating drug, with changes indicating a poor prognosis for future disability. For instance, alemtuzumab and ocrelizumab are specifically licensed for “active disease defined by clinical or imaging features.”

For cladribine, NICE specifically defined treatment failure as “1 relapse in the previous year and MRI evidence of disease activity.”

For fingolimod: under previous guidance, fingolimod may be given if patients have an unchanged or increased relapse rate or ongoing severe relapses compared with the previous year despite treatment with beta interferon or glatiramer acetate. This is now extended to include disease activity on teriflunomide and dimethyl fumarate.

Autologous haematopoietic stem treatment for autoimmunity is commissioned at specialised centres and is currently being offered to some people with MS in some parts of the UK. But there is not yet adequate controlled trial of its efficacy relative to other potent therapies. We recommend that it is made available equitably to all people with MS, but we propose that it should only be considered for people with relapsing disease (not progressive) who have failed high-activity licensed disease modifying therapies, and are prepared to accept the significant risks of the procedure. We recommend that this treatment is offered only by units with expertise both in the management of aggressive multiple sclerosis and the use of autologous haematopoietic stem treatment.

After considering all these options, it may be appropriate to continue the second-line therapy, despite evidence of disease activity.