therapeutic apheresis procedures in our center.

Diseases

Erythropoietic porphyria, liver disease

Hemophagocytic lymphohistiocytosis

Pruritus due to hepatobiliary disease

Vasculitis (Behcet's disease)

peutic plasma exchange.

Total

N-methyl D-aspartate receptor antibody encephalitis

Thrombotic microangiopathy-coagulation mediated

Prevention of RhD alloimmunization after RBC exposure

Hashimoto's encephalopathy

HELLP syndrome

Diseases	
Atopic (neuro-) dermatitis, recalcitrant	
Complex regional pain syndrome	

Progressive multifocal leukoencephalopathy associated with natalizumab

Table 2: The data regarding the rate of diagnosed diseases included in the new ASFA guideline¹ and of the utility of

No: number, TA: therapeutic apheresis, HELLP: hemolysis, elevated liver enzymes, and a low platelet count, RBC: red blood cell, TPE: thera-

Diagnosis No

0

1 (not definite, but

as rule out)

0

6

0

29

117

TA (type/No)

TPE/3

3