Provide the missing starting materials, reagents, or products required to complete the following synthesis. Be sure to provide the compounds in their ionic or neutral form where appropriate.

1) 

2) H₂, Pd/C

3) Adjust pH to 7

1) 

2) 

3) Adjust pH to 7

1) 

2) 

3) Adjust pH to 7
Provide the missing starting materials, reagents, or products required to complete the following synthesis. Be sure to provide the compounds in their ionic or neutral form where appropriate.

1) protect the acid as t-butyl ester

2) adjust the pH

3) Adjust pH to 7

the N-protected amino acid needed to combine with the valine (right) to give the diprotected dipeptide, below right

deprotect the FMOC from amine end

couple the amine with the acid

then deprotect the two benzyl-based sites

adjust pH to 7 (acids deprotonated, amines protonated)

couple the open acid end with the amine

then deprotect the FMOC group (information you can use for an earlier answer)

adjust pH to 7 (acids deprotonated, amines protonated)

Another “road map” of transformations involving peptide synthesis, where identification first is a useful strategy!
Provide the missing starting materials, reagents, or products required to complete the following synthesis. Be sure to provide the compounds in their ionic or neutral form where appropriate.

1) The N-protected amino acid needed to combine with the valine (right) to give the diprotected dipeptide, below right

2) Deprotect the FMOC from amine end

3) The dipeptide, deprotected at the carbon end, opening up that site for a reaction
couple the amine with the acid
then deprotect the two benzyl-based sites
adjust pH to 7 (acids deprotonated, amines protonated)

3) Adjust pH to 7

1) couple the open acid end with the amine
then deprotect the FMOC group (information you can use for an earlier answer)
adjust pH to 7 (acids deprotonated, amines protonated)

3) Adjust pH to 7